

The Effects of Hip Musculature on Knee Valgus During a Squat Task With and Without a 2-inch Heel Block

By
Brian J. Vesci

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Approved by

Advisor: Darin A. Padua, PhD, ATC
Reader: Kevin Guskiewicz, PhD, ATC
Reader: Chris Hirth, MS, PT, ATC

ABSTRACT

Brian Vesce: The Effects of Hip Musculature on Knee Valgus During a Squat Task With and Without a 2-inch Heel Block

(Under the direction of Dr. Darin A. Padua)

Objective: To compare hip ROM, strength, muscle activity, and ankle dorsiflexion ROM between groups that demonstrate knee valgus during a squat task and those that do not (control). Also, determine if muscle activity changes after knee valgus is corrected by a two-inch heel block. **Design:** A single-session experimental research design was used to compare the control group and the valgus group. **Participants:** Seventeen (10 Females, 7 Males) control subjects (age[yr] = 23.82 ± 5.76 , height[cm] = 166.12 ± 31.13 , weight[kg] = 69.59 ± 15.37) and fourteen (12 Female, 2 Male) valgus subjects (age[yr] = 22.36 ± 3.08 , height[cm] = 167.21 ± 9.3 , weight[kg] = 65.93 ± 9.9) without lower extremity injury. **Dependent Variables:** Supine hip abduction and external rotation ROM. Ankle dorsiflexion ROM (straight and bent knee). Hip internal rotation, external rotation, extension, and abduction eccentric and concentric peak torque and time to peak torque. EMG mean amplitude of the gluteus maximus, gluteus medius and adductor complex during a squat with and without a two inch heel block. **Data Analysis:** Mixed model analysis of variance tested for difference both between and within groups. **Results:** A significant group by phase interaction effect ($p = 0.02$) existed for mean adductor amplitude between the valgus and the control group. A significant difference ($p < .001$) in straight knee ankle dorsiflexion ROM existed between the control and the valgus group. A significant difference ($p = 0.034$) existed between the internal rotation concentric time to peak torque between the two groups. **Conclusion:** The

adductor complex plays a significant role in hip extension. Increased activity of the adductor during extension could pull the knees into valgus. Gastrocnemius tightness might contribute to knee valgus during a squat. There appears to be three distinct populations when considering this knee valgus squatting position. 1. Normal individuals who do not have valgus. 2. Ankle dysfunction individuals who's valgus position corrects with a heel block. 3. Hip dysfunction individuals who's valgus position does not correct with a heel block.

Keywords: Knee valgus, double leg squat, hip adductor activity, gastrocnemius flexibility

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CHAPTER 1 – INTRODUCTION

Introduction

Baseline screenings are commonly utilized in sports medicine as a method to assess injury status and establish return to play criteria should injury occur. More recently baseline screenings are used to develop corrective exercise programs to prevent injury from occurring. Typically baseline screenings have utilized isolated measures of flexibility, muscle strength, and functional performance; as well as identifying faulty posture. Acquiring this information serves two purposes. Initially this information is required to ensure that the athletes are not injured coming into the season. However, most often this information is important in the ongoing attempt to prevent those injuries from occurring. It is inevitable that certain injuries are going to occur during a season; however, if certain risk factors can be identified and corrected in the preseason, some of these injuries can be prevented.

Posture is a very important and often neglected part of overall health. Ideal posture maintains that structural integrity and optimum alignment of each component of the kinetic chain (Clark, 2001). The kinetic chain consists of the myofascial system, articular system, and the neural system (Clark, 2001). When one component of this system is out of alignment, then the entire system is placed at a disadvantage. Postural malalignment is thought to create predictable patterns of tissue overload and dysfunction, initiating the

cumulative injury cycle (Clark, 2001). This cumulative injury cycle begins with tissue trauma, inflammation, leading to muscle spasm, adhesions, altered neuromuscular control, and muscle imbalance. This cycle is thought to cause decreased athletic performance and eventual injury (Clark, 2001).

Functional movement analyses are becoming increasingly popular as a tool during dynamic postural assessment. Double leg and single leg squatting tasks are common during functional movement analyses. These methods can be easily implemented and are time efficient, thus are attractive measures for the clinician. During functional movement analyses the clinician observes for dysfunctional movement patterns that are believed to represent muscle imbalances caused by lack of flexibility, muscle weakness, and/or muscle activation. For example, individuals who demonstrate excessive knee valgus during a double leg squat are hypothesized to display a tight adductor muscle complex and weak gluteus medius and maximus muscles, as well as a lack of neuromuscular control in the aforementioned muscles. (Clark 2001) However, research has not been performed to validate the idea of functional movement analyses as a method to identify muscle imbalances.

Lack of muscle flexibility is one of the most common thought of risk factors for muscular injury. (van Mechelen, Hlobil et al. 1992; Garrett 1996; Gleim and McHugh 1997) It is theorized that if a muscle is unable to lengthen to a point that is considered normal ROM when stressed during activity, then it has a greater potential to fail. One study has shown that decreased flexibility in certain muscle groups can be considered an important factor in the development of injuries in soccer players. (Witvrouw, Danneels et al. 2003). In contrast, another study has shown that there was no significant difference in preseason adductor flexibility in professional ice hockey players who sustained adductor muscles injuries and

those who did not. (Tyler, Nicholas et al. 2001) The literature is inconclusive as to whether or not flexibility of a muscle is directly related to injury, but most evidence seems to suggest that it is a contributing factor to most muscular injuries. (Worrell 1994; Gleim and McHugh 1997; Thacker, Gilchrist et al. 2004)

Muscular weakness is another potential risk factor to muscular injury. Some studies have examined individual muscle strength and how it relates to injury of that specific muscle. (Worrell 1994) Although, few studies have analyzed whether or not weakness in one muscle has any relationship to injury in another area along the kinetic chain. Muscular weakness may develop secondary to changes in its antagonistic muscle. If a muscle is tight or overactive, then its antagonistic muscle may become weak or inhibited. This concept, called reciprocal inhibition, is considered to be another risk factor that contributes to athletic injury. (Clark M 2001) If a muscle is weak or inhibited, then it cannot fully function to mobilize or stabilize the structure for which is it intended. Muscle weakness can cause certain movement dysfunctions that can put certain muscles or joints in positions considered to be of high risk for injury. (Clark M 2001) This concept has yet to be fully investigated in the literature.

It is logical to assume that athletes, notably lower extremity athletes, may not have lower extremity muscular tightness or weakness. However, the idea of reciprocal inhibition brings to light another aspect of injury risk, which is muscle activation. If athletes do not possess the neuromuscular control to avoid these dysfunctional movements, then strength or flexibility would not matter. Extensive research currently explains the muscle activation patterns of the gluteus medius muscle during functional tasks. (Soderberg and Dostal 1978; Schmitz, Riemann et al. 2002; Earl 2005; Garrison, Hart et al. 2005) Also, research has

looked at the gluteus maximus activation in various functional tasks, although it is less understood. (Zeller, McCrory et al. 2003) However, research has yet to fully investigate hip musculature muscle activation, flexibility, and strength measurements during functional tasks and its relation to knee valgus.

As mentioned before, certain movement dysfunctions have been identified that are believed to be caused by the above mentioned risk factors.(Clark 2001) Knee valgus alignment during dynamic tasks (e.g. squatting) is a common postural dysfunction seen in the lower extremity. Knee valgus alignment during squatting is defined as the mid-patella moving medially and crossing over the ipsilateral great toe as the knee flexes while squatting downward. Knee valgus alignment has been described as a potentially dangerous movement pattern (Ireland 1999). Knee valgus alignment is accompanied by movement of the hip into adduction and internal rotation, which are commonly described mechanisms for anterior cruciate ligament (ACL) injury (Ireland 1999). It has also been shown that patello-femoral compressive forces increase with knee valgus alignment (Escamilla 2001) Thus, knee valgus alignment is also believed to be a contributing factor to patello-femoral pain syndrome. Furthermore, the medial collateral ligament (MCL) is stressed when the knee is exposed to a valgus moment. The MCL can become injured when that valgus stress becomes too great. Individuals who undergo excessive knee valgus motion during functional tasks may place greater stress on the MCL and be at greater risk for injury. These injuries can be detrimental to an individual's physical well-being. Therefore, it is important to understand the factors that influence knee valgus alignment as this may improve our understanding of lower extremity injury risk factor, which may lead to the development of exercise programs to correct knee valgus alignment and reduce injury risk. Currently, knee valgus alignment is

thought to occur due to weakness or inhibition in the hip abductor muscles (e.g. gluteus medius, gluteus maximus) and hip external rotator muscles. Meanwhile, the hip adductor and internal rotator muscles are believed to be tight or overactive (Clark 2001). However, research has not investigated if these hypothesized muscle imbalances actually exist in individuals demonstrating knee valgus alignment during a functional squatting task.

Squat tests, both single and double legged, have been used in the past as functional tests because they put the knee through common motions found in athletics.(Beutler, L.W. et al. 2002; Loudon, Wiesner et al. 2002; Zeller, McCrory et al. 2003) Moreover, studies have been done that examine knee valgus during squat tasks. (Zeller, McCrory et al. 2003) However, currently research has yet to focus on how the abovementioned risk factors associated with injury differ between subjects who experience a position of great knee valgus versus those who do not when performing a squat.]

Statement of Purpose

Closed-chain knee flexion is a very common athletic position. When this position is accompanied by knee valgus stress the risk for a knee injury is increased.(Ireland 1999) A double-legged squatting task (DLST) is a functional test that can be used clinically to mimic this closed-chain knee flexion. When someone's knee adducts during this squatting task, it is logical to assume it would do the same during an athletic activity. As of yet no concrete evidence exists to suggest what exactly causes this knee adduction during a DLST.

Research Questions

1. Do subjects who demonstrate bilateral knee adduction in a DLST have less hip abduction ROM compared to those who do not demonstrate bilateral knee adduction?
2. Do subjects who demonstrate bilateral knee adduction in a DLST have less hip external rotation compared to those who do not demonstrate bilateral knee adduction?
3. Do subjects who demonstrate bilateral knee adduction in a DLST have less straight knee ankle dorsiflexion compared to those who do not demonstrate bilateral knee adduction?
4. Do subjects who demonstrate bilateral knee adduction in a DLST have less bent knee ankle dorsiflexion compared to those who do not demonstrate bilateral knee adduction?
5. Do subjects who demonstrate bilateral knee adduction in a DLST have less gluteus medius muscle strength compared to those who do not demonstrate bilateral knee adduction?
6. Do subjects who demonstrate bilateral knee adduction in a DLST have less gluteus maximus muscle strength compared to those who do not demonstrate bilateral knee adduction?
7. Do subjects who demonstrate bilateral knee adduction in a DLST have less hip external rotator muscle strength compared to those who do not demonstrate bilateral knee adduction?

8. Do subjects who demonstrate bilateral knee adduction in a DLST have more hip internal rotator muscle strength compared to those who do not demonstrate bilateral knee adduction?
9. Do subjects who demonstrate bilateral knee adduction in a DLST have less mean gluteus medius EMG activity compared to those who do not demonstrate bilateral knee adduction?
10. Do subjects who demonstrate bilateral knee adduction in a DLST have less mean gluteus maximus EMG activity compared to those who do not demonstrate bilateral knee adduction?
11. Do subjects who demonstrate bilateral knee adduction in a DLST have more mean adductor complex EMG activity compared to those who do not demonstrate bilateral knee adduction?

Research Hypotheses

1. Subjects who demonstrate bilateral knee adduction in a DLST will have less hip abduction ROM compared to those who do not demonstrate bilateral knee adduction.
2. Subjects who demonstrate bilateral knee adduction in a DLST will have less hip external rotation ROM compared to those who do not demonstrate bilateral knee adduction.
3. Subjects who demonstrate bilateral knee adduction in a DLST will have less straight knee ankle dorsiflexion ROM compared to those who do not demonstrate bilateral knee adduction.

4. Subjects who demonstrate bilateral knee adduction in a DLST will have less bent knee ankle dorsiflexion ROM compared to those who do not demonstrate bilateral knee adduction.
5. Subjects who demonstrate bilateral knee adduction in a DLST will have less gluteus medius muscle strength compared to those who do not demonstrate bilateral knee adduction.
6. Subjects who demonstrate bilateral knee adduction in a DLST will have less gluteus maximus muscle strength compared to those who do not demonstrate bilateral knee adduction.
7. Subjects who demonstrate bilateral knee adduction in a DLST will have less hip external rotator muscle strength compared to those who do not demonstrate bilateral knee adduction.
8. Subjects who demonstrate bilateral knee adduction in a DLST will have more hip internal rotator muscle strength compared to those who do not demonstrate bilateral knee adduction.
9. Subjects who demonstrate bilateral knee adduction in a DLST will have less mean gluteus medius EMG activity compared to those who do not demonstrate bilateral knee adduction.
10. Subjects who demonstrate bilateral knee adduction in a DLST will have less mean gluteus maximus EMG activity compared to those who do not demonstrate bilateral knee adduction.

11. Subjects who demonstrate bilateral knee adduction in a DLST will have more mean adductor complex EMG activity compared to those who do not demonstrate bilateral knee adduction.

Null Hypotheses

1. Subjects who demonstrate bilateral knee adduction in a DLST will have no difference in hip abduction ROM compared to those who do not demonstrate bilateral knee adduction.
2. Subjects who demonstrate bilateral knee adduction in a DLST will have no difference in hip external rotation ROM compared to those who do not demonstrate bilateral knee adduction.
3. Subjects who demonstrate bilateral knee adduction in a DLST will have no difference in straight knee ankle dorsiflexion ROM compared to those who do not demonstrate bilateral knee adduction.
4. Subjects who demonstrate bilateral knee adduction in a DLST will have no difference in bent knee ankle dorsiflexion ROM compared to those who do not demonstrate bilateral knee adduction.
5. Subjects who demonstrate bilateral knee adduction in a DLST will have no difference in gluteus medius muscle strength compared to those who do not demonstrate bilateral knee adduction.
6. Subjects who demonstrate bilateral knee adduction in a DLST will have no difference in gluteus maximus muscle strength compared to those who do not demonstrate bilateral knee adduction.

7. Subjects who demonstrate bilateral knee adduction in a DLST will have no difference in hip external rotator muscle strength compared to those who do not demonstrate bilateral knee adduction.
8. Subjects who demonstrate bilateral knee adduction in a DLST will have no difference in hip internal rotator muscle strength compared to those who do not demonstrate bilateral knee adduction.
9. Subjects who demonstrate bilateral knee adduction in a DLST will have no difference in mean gluteus medius EMG activity compared to those who do not demonstrate bilateral knee adduction.
10. Subjects who demonstrate bilateral knee adduction in a DLST will have no difference in mean gluteus maximus EMG activity compared to those who do not demonstrate bilateral knee adduction.
11. Subjects who demonstrate bilateral knee adduction in a DLST will have no difference in mean adductor complex EMG activity compared to those who do not demonstrate bilateral knee adduction.

Variables

Independent Variable:

1. Group: Subjects will be placed into one of two groups based on their knee motion in an OHST
 - a). Ankle Dysfunction Group: Demonstrates bilateral knee adduction in DLST that is corrected by placing a two-inch heel block under the calcaneus.

- b). Control Group: Does not demonstrate bilateral knee adduction in DLST with or without 2-inch heel block.

Dependent Variables:

- 2. Flexibility as measured in degrees by a manual goniometer will be measured for the following muscles and ROMs:
 - a. Adductor Complex
 - b. Hip IR
 - c. Gastrocnemius
 - d. Soleus
- 3. Concentric and eccentric strength as measured in Newton-meters (N*m) by an isokinetic dynamometer for the following movements:
 - a. Hip Abduction
 - b. Hip Extension
 - c. Hip External Rotation
 - d. Hip Internal Rotation
- 4. Mean amplitude EMG activity during a DLST for the following muscles:
 - a. Gluteus Medius
 - b. Gluteus Maximus
 - c. Adductor Complex

Definition of Terms

- 1. Reciprocal Inhibition – The process whereby a tight or overactive agonist inhibits its functional antagonist.

2. Flexibility – As measured by a goniometer in degrees.
3. Strength – As measured by an isokinetic dynamometer in Newton meters (N*m).
4. Electromyography (EMG) – A measure of the neuromuscular action potential of a muscle; typically measured as peak, average onset time, and mean amplitude.
5. Isokinetic Testing – A measure of muscle contraction in which the length of the muscle is changing while the contraction velocity remains constant.

Operational Definitions

1. Double-Legged Squat Test (DLST) – Subject stands with feet shoulder width apart, feet facing forward, and hands raised above head. Subject then proceeds to squat down as if trying to sit in a chair.
2. Knee Adduction - The subject's knees will be such that the center of their patella is medial to the ipsilateral medial malleolus when they are in a squatted position.
3. Knee Valgus – Adduction and internal rotation of the femur on the tibia.
4. EMG –
 - a. Peak: The highest magnitude of EMG activity during a dynamic maneuver.
 - b. Mean Amplitude: The average magnitude of EMG activity for a given time interval.
 - c. Onset Time: Point in time when EMG amplitude reaches a predetermined magnitude.
5. Torque –
 - a. Peak: The greatest moment produced at one point in a range of motion.

- b. Mean: The average moment produced throughout a range of motion.
- 6. Adductor Complex – Adductor magnus, adductor longus, and adductor brevis

Assumptions

1. Subjects will be truthful in the consent form about previous and current injury status.
2. Subjects will provide maximum effort in performing strength assessments.
3. Tester is reliable in obtaining accurate measurements of strength and flexibility.
4. EMG was collected without noise affecting the signal.
5. Tester will provide each subject with exact same instructions prior to testing.

Limitations

1. EMG may measure different portions of muscles due to skin movement during the DLST.
2. Subjects' previous experience with squatting exercises could affect their form in a squatting activity.

Delimitations

1. Subjects in both groups will be asymptomatic with and without activity and during the DLST
2. Subjects in the hip dysfunction group will demonstrate bilateral knee adduction in the DLST with and without a 2-inch heel block.

3. Subjects in the control group will not demonstrate bilateral knee adduction in the DLST with or without the 2-inch heel block.
4. Subjects in both groups will have no previous history of surgery in either lower extremity.
5. Subjects in control group will be matched to subjects in hip dysfunction group based on height, weight, age, gender, and physical activity level.

Significance of Study

The significance of this study is to ascertain what factors lead to knee valgus during a closed chain flexed knee position. If these factors can be determined, then an intervention protocol can be established to correct them. This correction could then be studied further to examine its impact on decreasing knee injuries thought to be caused by this valgus position

CHAPTER 2 – REVIEW OF THE LITERATURE

Introduction

Preseason screenings are used by sports medicine professionals to gain knowledge about the health of their athletes. These baseline tests are used for documentation, assessment, and prediction of performance.(Crill, Kolba et al. 2004) It is believed that findings such as muscular imbalances and postural faults, both static and dynamic, can help predict not only performance but also risk of injury. Injury has been thought to be due to two different factors, both extrinsic and intrinsic. Extrinsic factors are those thought to be environment related such as weather, playing surfaces, ect., while intrinsic are those that are related to the individual person.(Taimela, Kujala et al. 1990; van Mechelen, Hlobil et al. 1992; Inklaar 1994) Intrinsic factors can be the neuromuscular control and postural imbalances that are sought out during preseason screenings. It is thought that since these factors are related to the athlete, then they can be altered through corrective exercise.

Injuries are inherent in sport. Some injuries, such as contact related traumatic injuries, are difficult, sometimes impossible, to prevent. Other injuries, such as muscular strain injuries are considered to be preventable with proper training. It is these preventable injuries that are of utmost importance to study. If it can be determined why or how muscular strain injuries occur then, theoretically, they can be prevented. The most widely believed cause of these injuries is muscular tightness.

(van Mechelen, Hlobil et al. 1992; Garrett 1996; Gleim and McHugh 1997) It is also believed that muscular tightness can lead to injuries as well, such as sprains and overuse injuries.(Worrell 1994; Krivickas 1997)

Before examining the concept of tightness in a muscle, there are two important terms that must first be defined. These are flexibility and stiffness. Muscular flexibility has most simply been defined as the angle beyond which no further displacement is possible.(Riemann, DeMont et al. 2001) Muscular stiffness represents the amount of deformation proportional to the load applied. (Blackburn, Padua et al. 2004) It is important to keep in mind that although these two concepts are related when assessing muscular length, they are two different characteristics of muscle.

Knee valgus occurs when there is a simultaneous internal rotation and adduction of the femur on the tibia. While these are normal biomechanical motions, this valgus position has been theorized to be a mechanism of injury for the ACL and MCL when the ROM becomes extreme. (Bendjaballah, Shirazi-Adl et al. 1997; Ireland 1999; Boden, Dean et al. 2000) The knee joint ligaments are designed to provide support to the joint; however, knee joint stability is not solely dependent upon these ligaments. As soon as the weight-bearing knee is flexed surrounding musculature is activated to increase joint stability.(Rosse 1997) This surrounding musculature involves muscles that cross the knee joint as well as the hip joint. If these muscles cannot effectively increase the joint stability, then the risk of injury to the knee joint increases. Unfortunately, little research exists to explain the role this musculature has in either leading to, or preventing, injury.

Muscular Tightness – Anatomy

Adductor Complex

When examining the movement of hip adduction, there are five muscles that are typically grouped into the hip adductors. These muscles are the pectineus, adductor longus, adductor brevis, adductor magnus, and the gracilis. However, when referring to the adductor complex, the most commonly thought of muscles are the adductor longus, brevis, and magnus. Although all of these muscles can contribute to hip flexion and internal rotation, together their primary role is hip adduction. Investigation of these muscles and their role in muscular injuries in athletes is limited and inconclusive. A study done by Witvrouw et al showed that there was no significant difference between the adduction flexibility of soccer players who sustained adductor muscle strains as compared to non-injured players.

(Witvrouw, Danneels et al. 2003) Another study on professional ice hockey players found no significant difference in adductor flexibility between players sustaining adductor injuries and players who did not. (Tyler, Nicholas et al. 2001) In contrast, Ekstrand and Gillquist found a correlation between tightness of the adductor muscles and the presence of adductor muscle strains in their prospective study of male soccer players. (Ekstrand and Gillquist 1983) It is important to note that in the study done by Witvrouw et al, players studied as a whole did not suffer many adductor injuries. Thus, their study did not have very much statistical power in regards to adductor muscle involvement in developing muscular injury. Further attention should be devoted towards resolving this conflict, because the current literature has yet to sufficiently explain the hip adductors role in muscular injury.

The pectineus muscle lies just medial to the iliopsoas and forms the medial part of the floor of the femoral triangle. It originates from the pectin of the pubis and the bone anterior

to the pectin, and inserts on the pectineal line, the proximal extension of the linea aspera, of the femur (Rosse 1997). The pectineus receives its vascular supply from the medial femoral circumflex artery and obturator artery, and it is innervated by the femoral nerve (Rosse 1997).

The adductor longus is the most anterior member of the adductor group. It originates from the superior pubic ramus and inserts into the medial lip of the linea aspera on the femur (Rosse 1997). The adductor longus receives its vascular supply from the medial femoral circumflex and obturator arteries, and it is innervated by the anterior division of the obturator nerve (Rosse 1997).

The adductor brevis is deep to the pectineus and adductor longus. It originates from the inferior pubic ramus and inserts into the medial lip of the linea aspera, superior to the adductor longus, on the femur (Rosse 1997). The adductor brevis receives its vascular supply from the medial femoral circumflex and obturator arteries, and it is innervated by the anterior division of the obturator nerve (Rosse 1997).

The adductor magnus is a combination of two muscles with different innervations. Both parts originate from the inferior pubic ramus, the ischiopubic ramus, and the ischial tuberosity. However, the adductor head inserts into the medial lip of the linea aspera on the femur and has horizontal fibers. The ischiocondylar head inserts on the adductor tubercle of the femur (Rosse 1997). The adductor head is innervated by the posterior division of the obturator nerve; while, the ischiocondylar head is innervated by the tibial nerve. Both portions receive their vascular supply from the deep femoral, medial femoral circumflex, and obturator arteries (Rosse 1997).

The gracilis is a long slender muscle lying superficially along the medial side of the thigh. It originates from the inferior pubic ramus and the pubic symphysis, and it inserts on the anteromedial part of the medial tibial condyle, via the pes anserine insertion (Rosse 1997). The gracilis receives its vascular supply from the medial femoral circumflex and obturator arteries, and it is innervated by the anterior division of the obturator nerve (Rosse 1997).

Gastrocnemius/Soleus Complex

The gastrocnemius is the superficial muscle of the triceps surae. It has two heads, medial and lateral, that originate just superior to the medial and lateral femoral condyles respectively. It receives its vascular supply from the posterior tibial and sural arteries and is innervated by the tibial nerve (Rosse 1997). The soleus is the deep muscle of the triceps surae. It originates from the upper part of the fibula and the soleal line on the tibia. The soleus receives its vascular supply from the posterior tibial artery (Rosse 1997). Both the gastrocnemius and soleus insert together into the calcaneus via the Achilles tendon (Rosse 1997).

Muscular Tightness – Assessment

Since it is widely thought that muscular tightness is a factor for muscular injury, it becomes important to have a way to quantify the flexibility of a muscle. Manual goniometers commonly regarded as the easiest and are the most widely used devices to measure range of motion in a joint. (Gajdosik and Bohannon 1987) However, they have their limitations. They have a single hinge and lack the ability to accurately measure movements

made by more than one joint. Furthermore, if proper anatomical landmarks are not established and adhered to for the measurement, then the measurement itself would be useless to a clinician. Despite these limitations though, they seem to be the preferred method of ROM testing. This ROM assessment is used by most studies to make inferences about muscle length and tightness.(Tyler, Nicholas et al. 2001; Witvrouw, Danneels et al. 2003; Decoster, Scanlon et al. 2004)

Hip Abduction

The proper patient positioning for ROM measurement for hip abduction is supine. The fulcrum of the goniometer is placed over the anterior superior iliac spine (ASIS) of the extremity being measured. The stationary arm is placed along an imaginary line from one ASIS to the other. The movement arm is placed along the anterior midline of the femur using the midline of the patella for reference.(Norkin 1995)

Hip Internal Rotation

The proper patient positioning for ROM measurement for hip internal rotation is seated with knees flexed to 90 degrees over the edge of the surface. The hip is in neutral abduction as well as adduction and is flexed to 90 degrees. A towel roll may be placed under the distal end if necessary to maintain the femur in a horizontal plane. The fulcrum of the goniometer is over the anterior center aspect of the patella. The stationary arm is positioned perpendicular to the floor. The movement arm is placed along the anterior midline of the

tibia pointing towards the anterior midpoint of the ankle between the two malleoli.(Norkin 1995)

Ankle Dorsiflexion

The proper patient positioning for ROM measurement of straight knee dorsiflexion was supine with the dominant knee in full extension. The ankle is in neutral inversion/eversion. The axis of the goniometer is the distal lateral malleolus. The stationary arm is positioned with the lateral midline of the fibula, using the fibular head for reference. The movement arm is placed parallel to the lateral aspect of the fifth metatarsal (Norkin 1995).

The proper patient positioning for ROM measurement of bent knee dorsiflexion is the same as straight knee; however, a bolster is placed under the subject's dominant knee so that it is flexed to at least 30 degrees (Norkin 1995).

Muscular Strength – Anatomy

Muscular strength is another intrinsic factor believed to be associated with muscular injury. (Knapik, Bauman et al. 1991; Tyler, Nicholas et al. 2001) If a muscle has inadequate strength then it may break down due to the physical demands placed upon it during functional activities. Unfortunately, the literature fails to fully outline how muscular strength pertains to injury.

Gluteus Medius

The gluteus medius muscle originates on the lateral aspect of the ilium between the anterior and posterior gluteal lines, and it inserts on the greater trochanter of the femur. It receives its vascular supply from the superior gluteal artery and is innervated by the superior gluteal nerve (Rosse 1997). The main action of this muscle is hip abduction and internal rotation of the hip, although it functions effectively as a pelvic stabilizer during standing and as a pelvic rotator during gait. (Gottschalk, Kourosh et al. 1989; Earl 2005) The fibers of the gluteus medius muscle are arranged in such a fashion as to suggest it is capable of performing many functions in many different positions of hip flexion and extension. The muscle itself has been looked at in the past as having three separate sections divided into anterior, middle, and posterior portions.(Gottschalk, Kourosh et al. 1989; Schmitz, Riemann et al. 2002) The anterior and middle portions have been shown to be most active during hip abduction and internal rotation, while the posterior portion is most active in pulling the femoral head into the acetabulum and stabilizing the hip joint.(Earl 2005) Soderberg and Dostal examined the gluteus medius muscle's EMG activity during a variety of function tasks. They found that the three portions each had the highest amount of activity in movements that required maximum control of the femur.(Soderberg and Dostal 1978) This would suggest that the gluteus medius muscle plays an important role in controlling the femur in multiple planes during functional activities.

The common injury of concern when talking about gluteus medius weakness and poor control at the hip is the anterior cruciate ligament (ACL). (Zeller, McCrory et al. 2003) It has been reported that this lack of proper hip control leads to increased hip adduction when the hip is loaded. This increase in adduction causes the femur to internally rotate and provide a

valgus stress to the knee. (Zeller, McCrory et al. 2003) This is considered a deleterious position for the knee to be in and may predispose the ACL to injury.(Beutler, L.W. et al. 2002; Mattacola, Perrin et al. 2002; Zeller, McCrory et al. 2003) While the function of this muscle has been studied quite thoroughly during many functional tasks, such as single leg stance, closed-chain hip rotation, and single leg squat, the step has not yet been taken to fully understand what effects it has on injury. (Schmitz, Riemann et al. 2002; Zeller, McCrory et al. 2003; Earl 2005)

Gluteus Maximus

The gluteus maximus muscle originates from the posterior gluteal line of the ilium, the lateral sacrum and coccyx, and the sacrotuberous ligament, and it inserts into the iliotibial tract and the gluteal tuberosity of the femur. It receives its vascular supply from the inferior gluteal artery and is innervated by the inferior gluteal nerve (Rosse 1997). This muscle is the prime hip extensor and is also involved in hip external rotation.

The external rotation moment arm of the gluteus maximus has been shown to decrease as the hip moves into flexion. Although the external rotation moment arm of the posterior fibers has been shown to remain even with a flexed hip.(Delp, Hess et al. 1999) This suggests that the gluteus maximus can be used to help control femoral internal rotation even when the hip is flexed. Moreover, it has been speculated that increasing strength and neuromuscular control of the gluteus maximus can help correct excessive femoral internal rotation in individuals that are believed to be at a higher risk of knee injury due to increased knee valgus in a position of hip and knee flexion.(Delp, Hess et al. 1999)

As stated before, this muscle is also believed to work in conjunction with the latissimus dorsi to aid in stabilization of the sacroiliac joint. (Clark 2001) Recently though, it has been theorized that its function as an external rotator plays a significant role in controlling functional movements. It is thought that it plays an important role in eccentrically controlling excess femoral internal rotation and avoiding the dangerous knee position previously outlined. (Clark 2001) However, its function in this capacity has currently not been researched enough to fully understand.

Muscular Strength – Assessment

Much like muscular length testing, a clinician must be able to assess and quantify muscular strength. Dynamometers make it possible to evaluate maximal muscular performance clinically. The many measurement methods that can be utilized include static, isokinetic, and, more recently, isotonic, isoinertial, and isoaccelerative. (Nadeau, Gravel et al. 1996) Each method controls one mechanical variable (angle, velocity, load inertia, or acceleration) and measures changes in some of the remaining variables. Static testing is generally considered the easiest way to test muscular strengths at certain positions in the joint's range of motion. While isokinetic testing is commonly considered the more popular method of quantifying dynamic muscular strength. (Osternig 1986; Cabri 1991)

Gluteus Medius

The proper positioning for the gluteus medius to be strength tested is to have the patient side-lying with the test leg being the uppermost. The patient starts with the test limb slightly extended beyond the midline and the pelvis rotated slightly forwards. The lowermost

knee is flexed for stability. The tester stands behind the patient. One hand is used to palpate the gluteus medius just proximal to the greater trochanter of the femur. The hand providing resistance is placed on the lateral surface of the knee joint. Resistance is given in a straight downward direction. The patient abducts the hip through the full range of motion without flexing the hip or rotating it in either direction. (Kendall 1993; Hislop 2002)

Gluteus Maximus

The proper positioning for the gluteus maximus to be strength tested is to have the patient supine with knee flexed to ninety degrees. The tester stands at the side to be tested at the level of the pelvis. One hand is placed over the sacrum to stabilize the alignment of the pelvis. The hand providing resistance is placed over the posterior thigh just above the knee. Resistance is provided in a straight downward manner. The patient extends the hip through the full available range of motion while maintaining knee flexion. (Kendall 1993; Hislop 2002)

Proper Double-Legged Squat Technique

The National Strength and Conditioning Association (NSCA) has outlined the proper body mechanics for a squat. The subject is to hold the chest up and out while pulling the scapulae towards each other. The head is tilted slightly upward. The feet are positioned should-width apart, even with each other, with the toes pointed straight ahead or just slightly outward (externally rotated). The subject allows the hips and knees to slowly flex while keeping the torso-to-floor angle relatively constant during the downward movement phase. The heels of the feet are to remain on the floor and the knees stay aligned over the feet. The

torso is not to be flexed forward or the low back rounded. The subjects continues the simultaneous hip and knee flexion until the thighs are parallel to the floor.(Baechle 2000)

Isokinetic Testing

Isokinetic testing is a form of strength testing that offers an objective measure of muscular strength. Isokinetic machines allow the subject to offer a maximal contraction force throughout a range of motion at a predetermined angular velocity. The objective measures that can be obtained from an isokinetic machine are muscular strength isometrically, concentrically, and eccentrically.(Arnheim 2000)

Strength measurement on isokinetic devices are measured in torque or moment. Torque is measured in foot-pounds or Newton-meters. Peak torque is the maximum moment recorded in the range of motion. Average torque is the mean torque produced by the muscle throughout the entire range of motion. Subjects who differ in body size make it difficult to compare isokinetic data in Newton-meters. This difficulty is avoided by dividing the torque measurements by the subject's body weight.

Different hip motion isokinetic testing positions currently exist. Some studies have suggested that hip abductor/adductor strength testing should be done in the side-lying position with the hip extended, and hip external rotation should be tested with the subject supine.(Burnett, Betts et al. 1990) Kincom (Chattanooga Group Inc.) recommends in their clinical desk reference that hip flexion/extension be tested in a standing position, hip abduction/adduction be tested in a supine position, and hip external/internal rotation be tested in a seated position.

Electromyography (EMG)

In their resting state, all body cells exhibit a resting membrane potential. This potential typically ranges from -20 to -200 millivolts (mV).(Marieb 1999) This potential can be characterized as the relative charge between the inside and outside of the cell and is negative because the inside of the cell is negative as compared to the outside.(Marieb 1999)

When a muscle fiber is activated by a motor neuron, an electrical impulse is conducted along the length of its axon. This response is called an action potential. This potential is always the same, regardless of the source or type of stimulus.(Marieb 1999) Muscles respond to these action potentials by forming stable bonds between contractile myofibrils and creating force.(Marieb 1999)

Surface EMG uses electrodes placed on the skin to detect these action potentials. Typically, two electrodes are placed in a parallel over the muscle 2-4 cm apart. The extracellular charge is positive before muscular contraction. As the muscle fibers depolarize under the first electrode, the action potential under that electrode becomes negative with respect to the second electrode. The EMG system can measure this difference between the electrodes.

One problem with EMG measurements is that they can vary greatly between subjects. This makes it difficult to compare the raw data between different subjects. A reference standard must first be defined before data can be interpreted across different subjects. This process is known as normalization. A common way to normalize EMG data across different subjects is through a measurement of maximal voluntary isometric contraction (MVIC). The EMG data is expressed as a percentage of the MVIC.

CHAPTER 3 - METHODS

Subjects

Approximately 70 subjects were screened by the principal investigator by watching them perform a double leg squat. Those subjects that were observed to exhibit a valgus motion during the squat were then observed with a two-inch heel block underneath their calcaneus. These subjects were recruited through volunteers from emails, fliers, and personal recruiting in classes. Subjects were divided into two groups based upon their results on the double leg squat task (DLST). The valgus group consisted of subjects who demonstrated bilateral knee adduction during the DLST that was corrected by placing a two-inch heel block under their calcaneus. The control group consisted of subjects who did not demonstrate knee adduction on either leg during the DLST. Subjects in the control group were matched to subjects in the hip dysfunction group by age, height, and weight. The control group consisted of 17 subjects while the valgus group consisted of 14 subjects whose ages ranged from 18-26 years. Subject demographics are located in Table 1. Subjects were otherwise healthy individuals that had no current musculoskeletal injuries or had not sustained an injury to either lower extremity in the past six months. Subjects were also excluded if they had surgery to either lower extremity in the past year. Furthermore, subjects in both groups were asymptomatic with and without activity, and subjects were asymptomatic during the DLST. Prior to all testing all subjects read and signed an informed

consent form approved by a University of North Carolina at Chapel Hill Biomedical Institutional Review Board (IRB).

Measurement and Instrumentation

A manual goniometer was used to measure joint angles in degrees for ROM assessment of the muscles of interest. Flexibility measurements were taken for the hip adductors ($ICC_{(2,1)} .89$ SEM = 1.743), the hip internal rotators ($ICC_{(2,1)} .97$ SEM 2.625), gastrocnemius, and soleus. ICCs were not calculated for these final two range of motion measures.

Concentric and eccentric muscle strength was evaluated using a Biodex System 3 Pro isokinetic dynamometer (Biodex Medical Systems, Shirley, NY), measured in Foot-pounds (Ft*lbs) of torque. The data were then analyzed with a customized Matlab 7.0 program (The Mathworks, Inc., Natick, MA) to determine the average peak torque, as well as time to peak torque, for the hip external rotators, internal rotators, hip abductors, and hip extensors. Each average was measured over the middle three of five total trials.

The mean amplitude electromyographic (EMG) activity of the gluteus medius, gluteus maximus, and adductor group was measured using an eight channel DelSys Bagnoli EMG System (Boston, MA). DE-2.1 single differential surface electrodes (DelSys Boston, MA) with a contact dimension of 1.0cm x 0.1cm and a contact spacing of 1.0cm over the muscle bellies and parallel to the fibers of the gluteus medius, gluteus maximus, and adductor muscles. Electrodes were plugged into a belt mounted I/O unit that plugged into the DelSys Bagnoli EMG system. This system amplified (x10,000 Hz) the EMG signal as it passed into the computer and was stored for analysis. The EMG data were processed with passive

demeaning (0.0 ms begin/ 10.0 ms end), a Butterworth notch filter at 60.0 Hz, a band pass Butterworth filter from 10.0 Hz – 350 Hz, and finally RMS smoothing was used at a time constant of 25 msec. All EMG data were normalized to the percentage of maximum voluntary isometric contraction to allow for comparison between subjects. All EMG data collection and processing was done using Datapac2K2 (Run Technologies, Mission Viejo, CA).

An electro-goniometer was placed over the knee joint so that knee joint angle could be recorded and related to EMG activity during the DLST. Electro-goniometer data were filtered with a low pass Butterworth at 15.0 Hz.

A tripod was used to standardize squat depth; as well as a metronome to standardize squat speed at 66 beats per minute. Subjects used two beats to descend, two beats to ascend and one beat to rest between squats.

Procedures

Students and faculty of the University of North Carolina at Chapel Hill were asked through email and fliers to volunteer for screening for this study. Screening consisted of a questionnaire pertaining to previous history of injury or surgery to the lower extremity. Each subject completed their individual testing in one session. Prior to testing, subjects had the testing procedures explained to them and were asked to read and sign an informed consent form. Each testing session began with the recording of the subject's gender, age (years), height (cm), and weight (kg). Subjects were then video taped performing the DLST both with and without a two-inch heel block to ensure that they met the inclusion criteria for their assigned group. The tester then assessed the subject's dominant leg, defined as the leg they

would use to kick a soccer ball for maximum distance, for the following variables: the subject's passive hip abduction ROM; passive external rotation ROM; passive straight knee ankle dorsiflexion ROM; passive bent knee dorsiflexion ROM; Concentric/eccentric isokinetic strength for hip external and internal rotation, hip abduction, and hip extension; and EMG activity during a DLST in the gluteus medius, gluteus maximus, and adductor complex. The valgus group also had EMG recorded for a DLST while standing on a two-inch heel block. Prior to testing subjects were allowed to warm up for five minutes on a stationary bike at a self-determined pace.

ROM measurements

Subject positioning for ROM measurement of hip abduction was supine. The axis of the goniometer was placed over the ASIS of the extremity being measured. The stationary arm was placed along an imaginary line from one ASIS to the other. The movement arm was placed along the anterior midline of the femur using the midline of the patella for reference. The subject was then passively abducted until the tester felt the contralateral ASIS begin to move (Norkin 1995). Three separate measurements were taken and the mean was recorded.

Subject positioning for ROM measurement for hip internal rotation was supine with the dominant hip and knee flexed to 90 degrees. The hip was in neutral abduction/adduction. The axis of the goniometer was the anterior center aspect of the patella. The stationary arm was positioned parallel to the table. The movement arm was placed along the anterior midline of the tibia pointing towards the anterior midpoint of the ankle between the two malleoli. (Norkin 1995) The subject's hip was passively externally rotated until the tester felt the end range. Three separate measurements were taken and the mean was recorded.

Subject positioning for ROM measurement of straight knee dorsiflexion was supine with the dominant knee in full extension. The ankle was in neutral inversion/eversion. The axis of the goniometer was distal lateral malleolus. The stationary arm was positioned with the lateral midline of the fibula, using the fibular head for reference. The movement arm was placed parallel to the lateral aspect of the fifth metatarsal (Norkin 1995). The subject's ankle was then passively dorsiflexed until the tester felt the end range. Three separate measurements were taken and the mean was recorded.

Subject positioning for ROM measurement of bent knee dorsiflexion was the same as straight knee; however, a bolster was placed under the subject's dominant knee so that it was flexed to at least 30 degrees (Norkin 1995). The subject's ankle was then passively dorsiflexed until the tester felt the end range. Three separate measurements were taken and the mean was recorded.

Electromyography

The sites for electrode placement were shaven and cleansed with alcohol to improve signal transmission from the muscles. Electrodes were placed over the muscle bellies of the gluteus medius, gluteus maximus, and adductor complex. A reference electrode was placed on the tibial tuberosity. Electrodes were placed in a parallel to muscle fibers.

The electrodes for the gluteus medius were placed halfway between the iliac crest and the greater trochanter of the femur (Kleissen 1990). The electrodes for the gluteus maximus were placed 20% of the distance between the spinous process of S2 and a point 10 cm distal to the greater trochanter. The electrodes for the adductor complex were placed on the muscle belly at the mid point of the femur (Leveau 1992). All electrode placements were confirmed

with an isometric manual muscle test and checked for cross talk. Maximal voluntary isometric contraction (MVIC) was used to normalize muscle activity between subjects (% MVIC).

Strength Measurements

Reliability for all strength tests was demonstrated by Halverson and Hawkey, abstract published in Journal of Athletic Training supplement 2004.

The hip external rotators were tested in a seated position. The subject's hip and knee was flexed to 90 degrees. The dynamometer was aligned with the long axis of the femur. Pressure was applied to the medial aspect of the distal tibia. The thigh was stabilized to the chair using straps. A towel was also placed between the subjects' knees to act as a fulcrum for external rotation and prevent adduction. The external rotators were tested through twenty degrees of motion that began at five degrees of external rotation and ended at fifteen degrees of internal rotation.

The hip internal rotators were tested in a seated position. The subject's hip and knee were flexed to 90 degrees. The dynamometer was aligned with the long axis of the femur. Pressure was applied to the lateral aspect of the distal tibia. The thigh was stabilized to the chair using straps. The internal rotators were tested through twenty degrees of motion that began at five degrees of internal rotation and ended at fifteen degrees of external rotation.

The hip abductors were tested with the subject in a side-lying position to isolate the gluteus medius. The joint axis was located at 0.5 inches medial to the ASIS at the level of the greater trochanter (Lyons, Perry et al. 1983). The hip was abducted and externally rotated, making sure not to let the trunk and pelvis rotate backward. Pressure was applied

against the thigh in the direction of adduction. Pressure was not applied against the external rotation component of the start position (Kendall 1993). The trunk was stabilized by strapping the subject to the chair. The muscle was tested from 0-20 degrees of hip abduction.

The hip extensors were tested in a supported, standing position in an attempt to isolate the gluteus maximus and maximize stabilization. The dynamometer axis of rotation was aligned with the anterior superior tip of the greater trochanter (Lyons, Perry et al. 1983). The subjects stood in front of the Biodex chair, and the seat was raised to the level of the subjects' ASIS. If the chair was unable to reach the subjects' ASIS, the subjects were then asked to flex the contralateral knee until the chair was even with the level of the ASIS. The subjects flexed their trunk to ninety degrees and laid their chest on the chair. The trunk was stabilized to the Biodex chair prior to testing using straps. This stabilization was aimed at preventing accessory trunk motions that might influence strength testing. The knee of the test leg was flexed to ninety degrees. The stance leg (non-test leg) was flexed at the knee such that the subjects' chest was comfortably resting on the chair. The subjects were then asked to actively extend the hip through a range of motion that began at ninety degrees of hip flexion (femur perpendicular to the ground) and ended at fifty degrees of hip flexion. Pressure was applied against the distal portion of the posterior thigh in the direction of hip flexion (Kendall 1993).

All muscles were tested concentrically and eccentrically at $60 \text{ deg} \cdot \text{sec}^{-1}$. The testing procedure accounted for gravity corrections during hip abduction and hip extension testing since the test limb was sufficiently close to the horizontal plane. The test limb did not come close enough to the horizontal plane to warrant a gravity correction during the testing of the hip external or internal rotators.

Following the isokinetic testing of each muscle group, the subject's strength was tested isometrically. The positioning and alignment of the subjects was exactly the same for the isometric testing as it was used during the isokinetic testing. The test limb was positioned in the middle of the isokinetic ROM or neutral position for isometric testing. The test limb was placed at seventy degrees of hip flexion while testing the hip extensors. The test limb was placed at ten degrees of hip abduction while testing the hip abductors. The test limb was placed in a neutral position (zero degrees of abduction/adduction) while testing the hip adductors. Each subject performed three maximal isometric contractions. The contractions were held for five seconds each, and there was 10-12 seconds rest between each repetition. The mean amplitude of the three trials during the middle three seconds of each trial was recorded.

The EMG data for the MVIC of the gluteus maximus was recorded while testing hip extension. The EMG data for the MVIC of the gluteus medius was recorded while testing hip abduction. The EMG data for the MVIC of the adductor complex was recorded while still in the testing position for hip abduction. The subjects were asked to maximally adduct the hip against the Biodex dynamometer for five seconds. EMG from these MVIC's was recorded so that the EMG data recorded during the DLST could be normalized and used to compare between subjects.

Double-Legged Squat Task

Subjects were asked to stand with hips, feet, and knees facing forward and arms fully extended overhead parallel to their ears. The subjects then were instructed to go down into a squat position as if they were trying to sit in a chair. Squat depth was standardized to 70

degrees of knee flexion to ensure that each subject went through at least 60 degrees of motion. Subjects were instructed to squat while their knee joint angle was measured with a manual goniometer. When the desired amount of 70 degrees of knee flexion was achieved a tripod was set up to assist the subject in knowing the proper squat depth. An electrogoniometer was then placed on the subject's knee so that EMG data could be examined in relation to knee position. The subjects then performed the squat task five times. EMG activity was recorded for the gluteus medius, gluteus maximus, and the adductor complex from 0-60 degrees of knee flexion during each of the squat tasks. Subjects in the control group only performed the DLST without the heel block. Subjects in the valgus group performed the DLSQ with and without the heel block while EMG data were recorded.

Data Analysis

One way analysis of variance tests were used to analyze differences between groups in the ROM measurements. A 2 x 2 analysis of variance was used to analyze the straight and bent knee ankle dorsiflexion ROMs. 2 x 2 mixed model analysis of variance tests were used to analyze differences between groups in concentric and eccentric peak torque as well as time to peak torque. EMG data differences were analyzed both between groups and between phases within groups with separate 2 x 2 mixed model analyses of variances. Tukey Post Hoc testing was used to identify where significance was found for the ankle dorsiflexion and EMG data. All data were analyzed using SPSS 13.0 (Chicago, IL). All null hypotheses were tested for significance at $p < .05$.

CHAPTER 4 – RESULTS

ROM Measurements

Means, standard deviations, and effect size for hip abduction ROM are presented in Table 2. There was no significant difference [$F_{(1,29)} = 0.01$, $p = .920$] for hip abduction range of motion between the control and the valgus groups. Means, standard deviations, and effect size are presented in Table 2. There was no significant difference [$F_{(1,29)} = 0.859$, $p = .362$] for hip external rotation ROM between the control and the valgus groups. There was a significant group by knee position interaction effect for ankle dorsiflexion [$F_{(1,26)} = 17.4$, $p < .001$]. Tukey Post Hoc tests revealed that the difference between groups with straight knee dorsiflexion was significant. This difference is illustrated in Figure 1.

Hip External Rotation Concentric/Eccentric Peak Torque

Means, standard deviations, and effect sizes are presented in Table 3. There was no significant difference for hip external rotation concentric [$F_{(1,29)} = 1.309$, $p = .262$] or eccentric [$F_{(1,29)} = .514$, $p = .479$] peak torque between the control and valgus groups.

Hip Internal Rotation Concentric/Eccentric Peak Torque

No significant differences were observed for hip internal rotation concentric [$F_{(1,29)} = 0.00$, $p = .985$] or eccentric [$F_{(1,29)} = 0.06$, $p = .808$] peak torque between the control and valgus groups.

Hip Extension Concentric/Eccentric Peak Torque

There was no significant difference [$F_{(1,29)} = 0.046$, $p = .833$] for hip extension concentric peak torque between the control and valgus groups. There was no significant difference [$F_{(1,29)} = 0.305$, $p = .585$] for hip extension eccentric peak torque between the control and valgus groups.

Hip Abduction Concentric/Eccentric Peak Torque

There was no significant difference [$F_{(1,29)} = 3.682$, $p = .065$] for hip abduction concentric peak torque between the control and valgus groups. However, the p value suggests a trend and the finding might reach significance if the statistical power was greater. This trend is illustrated in Figure 2. There was no significant difference [$F_{(1,29)} = 0.006$, $p = .939$] for hip abduction eccentric peak torque between the normal and valgus groups.

Hip External Rotation Concentric/Eccentric Time to Peak Torque

Means, standard deviations, and effect size are presented in Table 4. There was no significant difference [$F_{(1,29)} = 0.376$, $p = .544$] for hip external rotation concentric time to peak torque between the control and valgus groups. There was no significant difference [$F_{(1,29)} = 0.038$, $p = .847$] for hip external rotation eccentric time to peak torque between the control and valgus groups.

Hip Internal Rotation Concentric/Eccentric Time to Peak Torque

There was a significant difference [$F_{(1,29)} = 4.976$, $p = .034$] for hip internal rotation concentric time to peak torque between the control and valgus groups. The normal group

achieved peak torque almost a full second faster (2 sec vs. 3 sec) than the valgus group (fig 3). There was no significant difference [$F_{(1,29)} = 0.023$, $p = .880$] for hip internal rotation eccentric time to peak torque between the normal and valgus groups.

Hip Extension Concentric/Eccentric Time to Peak Torque

There was no significant difference [$F_{(1,29)} = 0.456$, $p = .501$] for hip extension concentric time to peak torque between the control and valgus groups. There was no significant difference [$F_{(1,29)} = 1.235$, $p = .276$] for hip extension eccentric time to peak torque between the control and valgus groups.

Hip Abduction Concentric/Eccentric Time to Peak Torque

There was no significant difference [$F_{(1,29)} = 3.218$, $p = .083$] for abduction concentric time to peak torque between the control and valgus groups. However, the p value suggests a trend that the valgus group required a half second longer (1.1 sec vs. 1.6 sec) to reach peak torque. This trend is illustrated in Figure 4. There was no significant difference [$F_{(1,29)} = 1.725$, $p = .199$] for hip abduction rotation eccentric time to peak torque between the normal and valgus groups.

Gluteus Maximus EMG Mean Amplitude During a Squatting Task

Means, standard deviations, and effect size are presented in Table 5. There was no main effect for group [$F_{(1,27)} = 0.002$, $p = .962$] for gluteus maximus activity during the ascending and descending phases of the squat. No main effect for phase [$F_{(1,27)} = 0.024$, $p = .877$] was observed for gluteus maximus activity between the ascending and descending

phases of the squat for the control or valgus groups. There was no group by phase interaction effect [$F_{(1,27)} = 0.020$, $p = .888$] for gluteus maximus activity during the squat task.

Gluteus Medius EMG Mean Amplitude During a Squatting Task

There was no main effect for group [$F_{(1,27)} = 1.037$, $p = .318$] during the ascending or descending phases of the squat between the control or valgus groups. There was no main effect for phase [$F_{(1,27)} = 0.101$, $p = .753$] of gluteus medius activity between ascending and descending phases of the squat for either group. No group by phase interaction effect [$F_{(1,27)} = 0.098$, $p = .757$] was observed for gluteus medius activity during the squat task.

Adductor Mean EMG Amplitude During Squatting

There was a group by phase interaction [$F_{(1,27)} = 6.095$, $p = .020$] for adductor activity during the squat task. The control group had higher mean amplitude during the descending portion of the squat and decreased during the ascending phase. In contrast, the valgus group had higher adductor activity during ascending than descending. Tukey post hoc testing revealed that the difference between adductor activity of the control group and the valgus group during the ascent phase was significant. This significance is illustrated in Figure 5. No main effect for group [$F_{(1,27)} = 0.163$, $p = .690$] was observed for adductor activity during the ascending or descending phases of the squat. There was no main effect for phase [$F_{(1,27)} = 0.196$, $p = .661$] for adductor activity between the ascending and descending phases of the squat for either the normal or the valgus group.

Gluteus Maximus EMG Mean Amplitude During Squatting with Heel Block

Means, standard deviations, and effect size for mean gluteus maximus amplitude during the squatting task on the heel block are presented in Table 6. There was no main effect for the heel block [$F_{(1,11)} = 0.134$, $p = .721$] for gluteus maximus activity during the ascending and descending phases of the squat between the valgus group with and without the heel block. There was no main effect for phase [$F_{(1,11)} = 1.525$, $p = .241$] for gluteus maximus activity between the ascending and descending phases of the squat for the valgus group either with or without the heel block. There was no heel block x phase interaction effect [$F_{(1,27)} = 1.430$, $p = .255$] for gluteus maximus activity during the squat task.

Gluteus Medius Mean EMG Amplitude During Squatting with a Heel Block

Means, standard deviations, and effect size for gluteus medius EMG mean amplitude during the squatting task on the heel block are presented in Table 6. No main effect was observed for gluteus medius activity [$F_{(1,11)} = 0.022$, $p = .885$] during ascending and descending phases of a squat between the valgus group with and without the heel block. There was no main effect for phase [$F_{(1,11)} = 2.034$, $p = .179$] of gluteus medius activity between the ascending and descending phases of the squat for the valgus group either with or without the heel block. There was no heel block by phase interaction effect [$F_{(1,11)} = 1.955$, $p = .187$] for gluteus medius activity during the squat task.

Adductor EMG Mean Amplitude During Squatting with and without a Heel Block

Means, standard deviations, and effect size for mean adductor amplitude during the squatting task on the heel block are presented in Table 6. There was no main effect for heel

block [$F_{(1,11)} = 2039$, $p = .148$] for adductor activity during the ascending and descending phases of the squat between the valgus group with and without the heel block. There was a main effect for phase [$F_{(1,11)} = 10.951$, $p = .006$] of adductor activity between the ascending and descending phases of the squat for the valgus group while on the heel block. Adductor activity during the descending phase of the squat was less than the ascending phase. This significance is illustrated in Figure 6. There was no heel block by phase interaction effect [$F_{(1,11)} = 1.7$, $p = .217$] for adductor activity during the squat task.

CHAPTER 5 – DISCUSSION

The most important findings from this study are that individuals who demonstrated knee valgus during a squat and have the valgus position corrected with a two-inch heel block have different adductor muscle activity during a squat. Our results also show these valgus individuals have less passive ankle dorsiflexion ROM (with a straight knee) when compared to a control group. Moreover, the passive bent knee ankle dorsiflexion ROM of the valgus group was significantly increased over their straight knee ankle dorsiflexion. The control group reached concentric peak torque of the internal rotators quicker than the valgus group. Trends in the data suggest that the control group has larger concentric peak torque in the abductors and achieve peak torque during concentric abduction faster when compared to the valgus group.

Muscle Activity During Squatting

An analysis of the control group shows that muscle activity during different phases of a squat (descending and ascending) is not different. This means the control group uses hip musculature equally. This is not the case for those who display a valgus motion during a squat and the difference seems to be in the adductor complex. The other two muscles examined (gluteus maximus and gluteus medius) showed similar activity patterns when compared to the control group.

A group by phase interaction was demonstrated by the adductor muscle complex during the squat task. When comparing maximal muscle activity, the control group's adductor muscle had the greatest activity during the decent phase while the valgus group was most active during the ascent phase (Figure 5). Thus, subjects performing proper squat posture utilized all three muscles simultaneously to eccentrically lower themselves, while the valgus group adductor complex was not as active in the eccentric lowering during the descent phase. However, during the ascent phase the control group's adductors were not as active and relied on the gluteus maximus to assist with hip extension. It is important to note that neither the decrease in the adductor activity nor the increase in the gluteus maximus activity from the descent phase to the ascent phase in the normal group were deemed significant. The opposite situation was found to exist with the adductor complex of the valgus group. Adductor activity during the decent phase was significantly less than adductor activity during the ascent phase. Furthermore, their adductor activity during the ascent phase was significantly greater than that of the normal group during the same phase.

Subjects in the valgus group in this study had their valgus corrected when squatting on a two-inch heel block. Current theory (Clark 2001) suggests valgus may result from tight musculature of the ankle rather than weak hip musculature. The heel block allows for increased length of the gastrocnemius and soleus complex, allowing normal motion and correcting valgus. Further research needs to identify individuals that present with valgus that is not corrected by a heel block and validate that hip musculature is the issue and what role it may play in knee valgus.

Increased adductor activity in the valgus group during the ascent phase of a squat could be explained an improved length-tension relationship of the muscle group due to

increased hip flexion. One study examined moment arms of hip extensor muscles (gluteus maximus and adductor magnus) and found that the gluteus maximus moment arm decreased while the adductor magnus moment arm increased while the hip was flexed. The adductor magnus extension moment arm was shown to continue to increase until 75 degrees of hip flexion (Nemeth and Ohlsen 1985). Furthermore, it has been shown that women demonstrate significantly more hip flexion and adduction during single-legged squats (Zeller, McCrory et al. 2003). While our study examines double-legged squats, this is still relevant because valgus was part of our inclusion criteria and a majority of our valgus subjects were women. Females are being put into a position of hip flexion and adduction that may increase their adductor muscle activity during the ascent phase. While in the control group the gluteus maximus increases during the ascent phase, it actually slightly decreases in the ascent phase for the valgus group. This may be explained by the increase in hip adduction decreasing the gluteus maximus' ability to function as an extensor. This position of knee adduction is associated with femoral internal rotation (Ireland 1999). During the ascent phase the gluteus maximus may be functioning more in its role as a hip external rotator to compensate for femoral internal rotation. If this is the case, then it makes sense that the adductor activity increases as they function more in extension. However, the changes in gluteus maximus muscle activity in our study were not found to be significant. Moreover, we did not measure hip flexion angles in our subjects. Therefore this theory needs to be explained further through future research.

A final possible explanation of the difference seen between our groups in adductor muscle activity arises from stance width. Research suggests stance width has a significant effect on adductor longus muscle activity during a squat task (McCaw and Melrose 1999).

Stance width was not measured in this study but we used a self-selected stance width set at shoulder width apart and researcher verified. We think this minimizes any effect stance width may have on our data.

There were no significant differences found in any of the muscles tested in the valgus group between the regular squat and the squat on the heel block, except for the significant difference in adductor activity between phases that had already been discussed in comparison to the control group. This is the first study to examine how hip muscle activity changes when individuals with knee valgus squat on a two-inch heel block. More research needs to be done to fully understand how plantar flexing the ankle by placing an individual on a two-inch heel block affects squatting.

The lack of significance in the muscle activity between the valgus group with and without the heel block begins to outline three populations. These populations are a normal or control group, an ankle dysfunction group and a hip dysfunction group. The control group demonstrates a proper squat technique and keeps their knees centered over their toes (Clark 2001). The ankle dysfunction group is the group examined in our study and demonstrates knee valgus during a squat that is corrected by placing a two inch block under their heel. The hip dysfunction group consists of individuals who demonstrate knee valgus during a squat that is not corrected by the heel block. Currently no study has focused on the hip dysfunction group to verify current theory as to what is causing their knee valgus.

Straight and Bent Knee Ankle Dorsiflexion ROM

Another significant finding of this study was the difference of the straight knee ankle dorsiflexion ROM between the valgus group and control group and the difference of the

valgus group between straight knee and bent knee. The valgus group had significantly less straight knee dorsiflexion. The gastrocnemius is a bi-articulate muscle crossing the ankle and knee. When the knee is straight, ROM of the ankle decreases because the gastrocnemius reaches end range, limiting motion. Tightness in the gastrocnemius muscle may have played a role in valgus during the DLST. This point is illustrated further when the knee was bent. The valgus group's dorsiflexion range of motion was significantly increased and approximately equal to the control group. This idea is the foundation of current theory! When the valgus group squatted with the heel block, the gastrocnemius shorts and the resulting tension is removed from the squat, correcting knee valgus. This suggests that the tightness in the gastrocnemius muscle played a role in the knee valgus. It is important to note that when the valgus group was on the heel block, there were no significant changes in hip musculature activity.

Ankle dorsiflexion has been shown to be around 10 degrees during the stance phase of gait (Murray 1967). During more function tasks, such as sit-to-stand, stair climbing, and sport specific activities, the requirement for ankle dorsiflexion ROM can increase to about 25 degrees (Andriacchi, Andersson et al. 1980; Lindsjo, Danckwardt-Lilliestrom et al. 1985; Livingston, Stevenson et al. 1991). The double-legged squat used in our study can be related to the function activities that can require the greater dorsiflexion. Thus, the lack of motion seen in the valgus group can explain their inability to perform the task properly. However, it becomes hard to explain why gastrocnemius tightness in and of itself would contribute to knee valgus. It may be a unilateral issue. If the lateral gastroc is tight or short, it may be responsible for pulling the knee into a valgus position. Currently it is believed that during a squat decreased ankle dorsiflexion would lead to the heels of the individual rising up off of

the floor (Clark 2001). This idea makes more sense when examining the gastrocnemius muscle by itself; however, when considering the entire kinetic chain there appears to be more to the picture. It makes more sense to think that this tightness in the gastrocnemius would pull, from its insertion, the calcaneus into eversion, thus causing the foot to pronate. Our study did not examine foot and ankle pronation but if this is the case, then our results seem to suggest that when the arch flattens and pronation ceases the gastrocnemius begins to influence an increase in knee valgus. Therefore it seems that more research needs to be done to determine whether or not the gastrocnemius muscle itself can cause knee valgus or if it only contributes to increased valgus when it is already present.

Internal Rotation Time to Peak Torque

Our study found that the control group reached their internal rotation concentric peak torque approximately a full second (2 sec vs. 3 sec) before the valgus group. This difference was deemed statistically significant and further strengthens the idea that there is a difference in adductor activity between groups. The current literature suggests that the gluteus maximus and gluteus medius are important hip stabilizers (Gottschalk, Kourosh et al. 1989; Clark 2001; Schmitz, Riemann et al. 2002) Our study showed that the gluteus maximus and gluteus medius activity between the control and the valgus groups were similar even though the valgus group demonstrated poor proximal hip control. This suggests that it may not be the gluteal muscles that are responsible for stabilization but rather other hip internal and external rotators. This can be further explained by the differences discussed above about the adductor activity between the two groups, as the adductors also function as hip internal rotators.

Abduction Concentric Peak Torque and Time to Peak Torque

There was a trend discovered in our study that suggests the control group produced greater concentric peak torque in the abductors than the valgus group and the control group achieved peak torque a half second faster (1.1 sec vs. 1.6 sec). The lack of significance of this may be due to lack of statistical power; an increase in power may draw out the significance of this finding. Regardless, this finding is clinically significant as it suggests that one potential cause of the knee valgus position is that the valgus group lacks abduction strength as well as the ability to achieve maximum strength in time to prevent hip adduction. This could be related to the previous stated findings of ankle dorsiflexion range of motion. It may be that the adduction begins due to a lack of abduction stabilization strength and is increased by the lack of gastrocnemius flexibility and decreased ankle dorsiflexion

Ireland has described a “position of no return” as when an athlete has poor hip control and the hip moves into adduction, leading to femoral internal rotation and a position of knee valgus (Ireland 1999). These trends discovered in our study seem to support this and further explain it by demonstrated ankle dorsiflexion ROM’s role in this knee adduction. However, further research is required to further explore these trends and determine if they are significant.

Hip Abduction and Internal Rotation ROM

No difference was observed in hip abduction or internal rotation ROM between the control and valgus groups. Initially, we hypothesized that tight adductors or tight internal

rotators may pull individuals into knee valgus. However, our study failed to identify this as a contributing factor. In fact the means of each ROM between groups was almost identical.

Hip Extension and External Rotation Concentric and Eccentric Peak Torque

No statistical difference was observed between group concentric or eccentric peak torques. This finding is clinically significant because it suggests that the gluteus maximus muscle does not play a significant role in preventing knee valgus during a squat. It is currently thought that individuals who demonstrate knee valgus have weak gluteus maximus control during squatting (Clark 2001). Again it becomes important to note that it appears that the subjects tested had ankle dysfunction and not necessarily hip dysfunction, and that individuals not corrected by the heel block may have differences in gluteus maximus strength.

Limitations

The greatest limitation of this study was the lack of statistical power for some of the analyses. Ideally, each group would have around 20 subjects to have enough power to draw out significant differences that may exist. This was unable to be achieved in this study due to the difficulty in identifying subjects who met the valgus group criteria. However, most of the analyses that lacked power also had small effect sizes. Therefore, if power was increased such that significance was found, it still may not be clinically significant.

Another limitation for this study is that the order of testing was not randomized. Every subject followed the same protocol in the same order. However, since the subjects were observed to exhibit the bilateral knee valgus both before and after ROM and strength

testing it is not believed that either altered the way in which the subjects performed the squat task.

The inability to identify many males who fit the inclusion criteria for the valgus group limits the generalizability this study. The control group is more evenly divided between males and females; however, the valgus group consists of primarily females. Although, the demographics between each group suggest that subjects were closely matched by age, height, and weight.

The lack of lower leg EMG activity is also a limitation of this study. While the ROM of ankle dorsiflexion was measured, no information about the muscle activity around the ankle of the ankle dysfunction group was available to help further understand what may be contributing to their valgus knee position.

Future Research

Current theory places subjects into categories if they are identified as having knee valgus (Clark 2001). There is a control population that demonstrates correct squat technique. There is a hip dysfunction population that demonstrates bilateral knee valgus both with and without a two-inch heel block. Finally, there is an ankle dysfunction population that demonstrates bilateral knee valgus during a squat that is corrected by a two-inch heel block. This study contrasted the control population against the ankle population and how hip musculature and ankle ROM contributes to knee valgus. Future research needs to verify that this classification system is correct and document differences between groups. Areas of focus should be the hip musculature (gluteus maximus, gluteus medius and adductor complex), ankle musculature (medial and lateral gastrocnemius, tibialis anterior, and soleus),

and these variables relation to this position of knee valgus during a squat task. Future research should also focus on the kinematics of the squat to verify that valgus is corrected or not with the heel block. Finally, once the theory has been established, research should attempt to implement training programs to correct valgus.

APPENDIX A: TABLES

Table 1: Subject Demographics; Mean and Standard Deviation

Group	Gender		Age(yr)		Height(cm)		Weight(kg)	
	Male	Female	Mean	SD	Mean	SD	Mean	SD
Normal	7	10	23.82	5.76	166.12	31.13	69.59	15.37
Valgus	2	12	22.36	3.08	167.21	9.3	65.93	9.9

Table 2: Ranges of Motion; Means and Standard Deviations

ROM	Normal			Valgus			Effect Size
	Mean	±	SD	Mean	±	SD	
Hip Abd	36.88	±	5.34	35.29	±	3.97	0.35
Hip ER	73.92	±	17.51	74.23	±	7.02	0.04
AnkleSKD	17.88	±	7.93	13.43	±	7.85*	0.59
AnkleBKD	19.86	±	8.37	19.88	±	8.37**	0.003

p < .001 for group by ROM interaction

* Denotes Significant Difference from Normal Group AnkleSKD

** Denotes Significant Difference from Valgus Group AnkleSKD

Abd - Abduction

ER - External Rotation

SKD - Straight Knee Dorsiflexion

BKD - Bent Knee Dorsiflexion

Table 3: Isokinetic Peak Torque Normalized to Body Weight (ft * lbs/kg); Means and Standard Deviations

ROM	Contraction	Normal			Valgus			Effect Size
		Mean	±	SD	Mean	±	SD	
Hip ER	Concentric	0.315	±	0.075	0.273	±	0.134	0.43
	Eccentric	0.605	±	0.143	0.557	±	0.229	0.27
Hip IR	Concentric	0.806	±	0.293	0.804	±	0.201	<.000
	Eccentric	0.444	±	0.18	0.428	±	0.188	0.09
Hip Ext	Concentric	1.281	±	0.521	1.244	±	0.424	0.08
	Eccentric	0.806	±	0.415	0.73	±	0.335	0.21
Hip Abd	Concentric	0.438	±	0.175	0.327	±	0.142*	0.76
	Eccentric	0.508	±	0.277	0.501	±	0.217	0.03

* Denotes trend that Valgus group shows less concentric hip abduction peak torque (p = .065)

ER - External Rotation

IR - Internal Rotation

Ext - Extension

Abd - Abduction

Table 4: Isokinetic Time to Peak Torque (seconds); Means and Standard Deviations

ROM	Contraction	Normal			Valgus			Effect Size
		Mean	±	SD	Mean	±	SD	
Hip ER	Concentric	3.35	±	1.63	2.98	±	1.75	0.23
	Eccentric	3.23	±	1.24	3.12	±	1.83	0.07
Hip IR	Concentric	2.29	±	1.11	3.21	±	1.18*	0.83
	Eccentric	2.76	±	1.35	2.86	±	2.16	0.06
Hip Ext	Concentric	4.49	±	1.76	4.9	±	1.58	0.25
	Eccentric	2.51	±	1.98	3.31	±	1.92	0.41
Hip Abd	Concentric	1.18	±	0.58	1.62	±	0.79**	0.7
	Eccentric	2.73	±	1.42	3.36	±	1.22	0.49

* Denotes that Valgus group took significantly longer to reach IR concentric peak torque ($p = .034$)

** Denotes trend that Valgus group shows longer time to concentric peak torque ($p = .083$)

ER - External Rotation

IR - Internal Rotation

Ext - Extension

Abd - Abduction

**Table 5: Mean EMG Amplitude (Normalized to MVIC) During a Squat Task;
Means and Standard Deviations**

		Normal		Valgus		Effect Size
		Mean	SD	Mean	SD	
Gluteus Maximus	Descending	10.98 ± 12.53		11.37 ± 5.65		0.02
	Ascending	11.52 ± 8.03		11.4 ± 5.72		0.02
Gluteus Medius	Descending	14.99 ± 10.04		19.02 ± 9.72		0.39
	Ascending	15.5 ± 11.02		19.03 ± 9.75		0.39
Adductors	Descending	29.68 ± 39.02		18.85 ± 14.26		0.16
	Ascending	18.02 ± 7.62		35.6 ± 33.78*†		0.16

p = .020 for group x phase interaction

* Denotes significant difference from valgus group descending activity

† Denotes significant difference from normal group ascending activity

Table 6: Mean EMG Amplitude (Normalized to MVIC) During a Squat Task with Heel Block; Means and Standard Deviations

		Valgus (no heel block)			Valgus (heel block)			Effect Size
		Mean	±	SD	Mean	±	SD	
Gluteus Maximus	Descending	11.37	±	5.65	10.58	±	5.06	0.5
	Ascending	11.4	±	5.72	13.07	±	8.6	0.5
Gluteus Medius	Descending	19.02	±	9.72	18.14	±	9.43	0.58
	Ascending	19.03	±	9.75	20.26	±	11.86	0.58
Adductors	Descending	18.85	±	14.26	18.09	±	14.08	0.73
	Ascending	35.6	±	33.78	28.49	±	20.95	0.73

APPENDIX B: FIGURES

Figure1: Ankle Dorsiflexion ROM

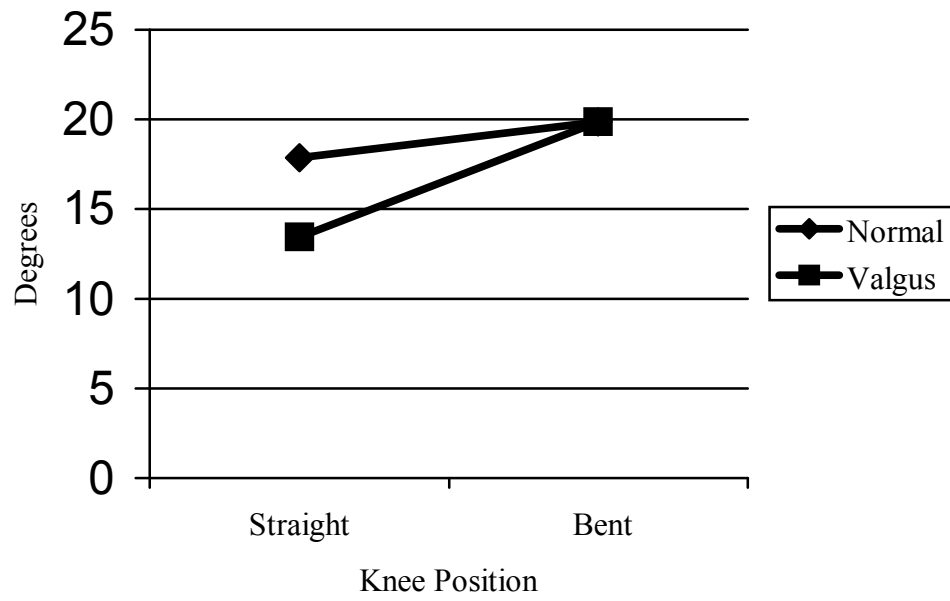


Figure 2: Hip Abduction Isokinetic Strength (Normalized to Body Weight)

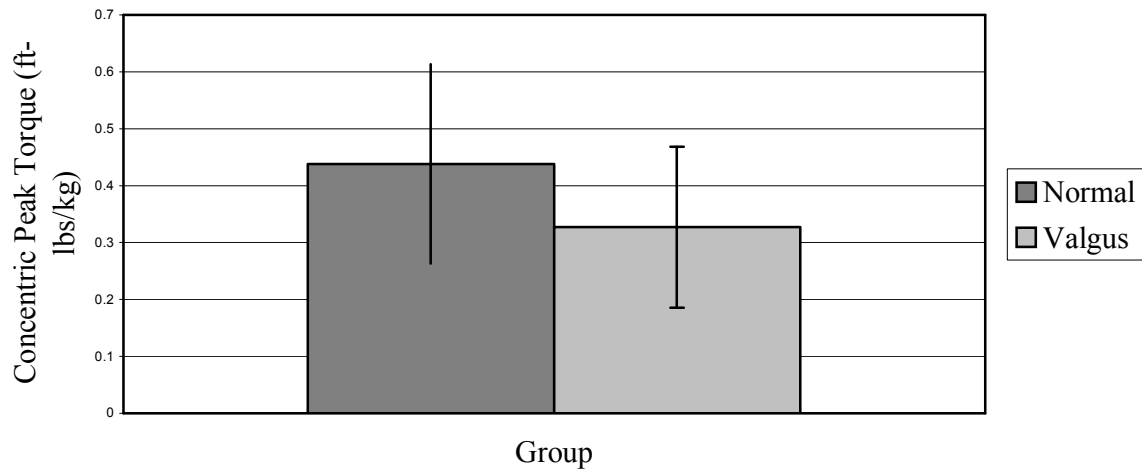


Figure 3: Internal Rotation Time to Peak Torque

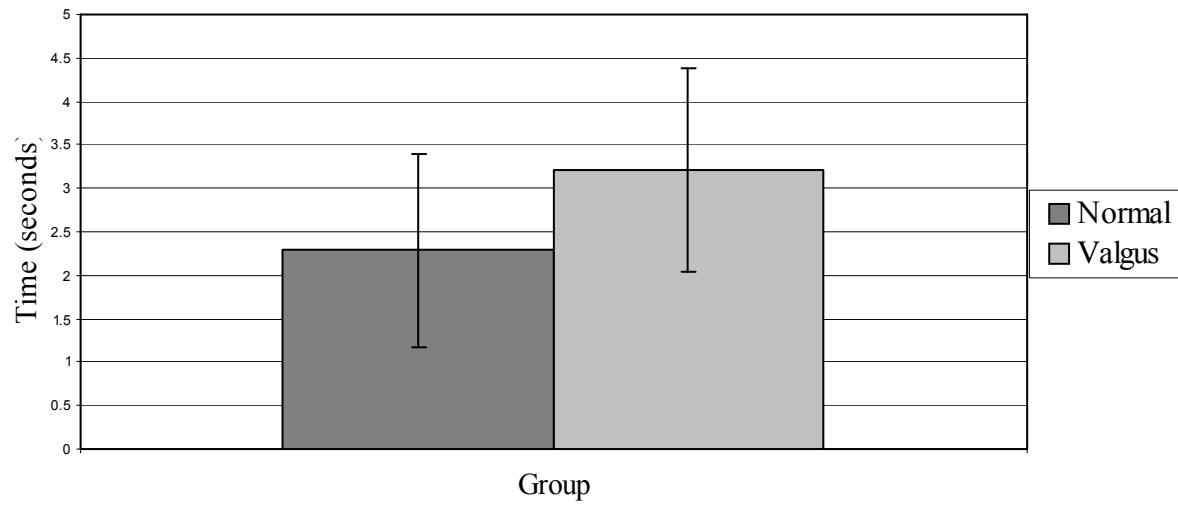


Figure 4: Abduction Concentric Time to Peak Torque

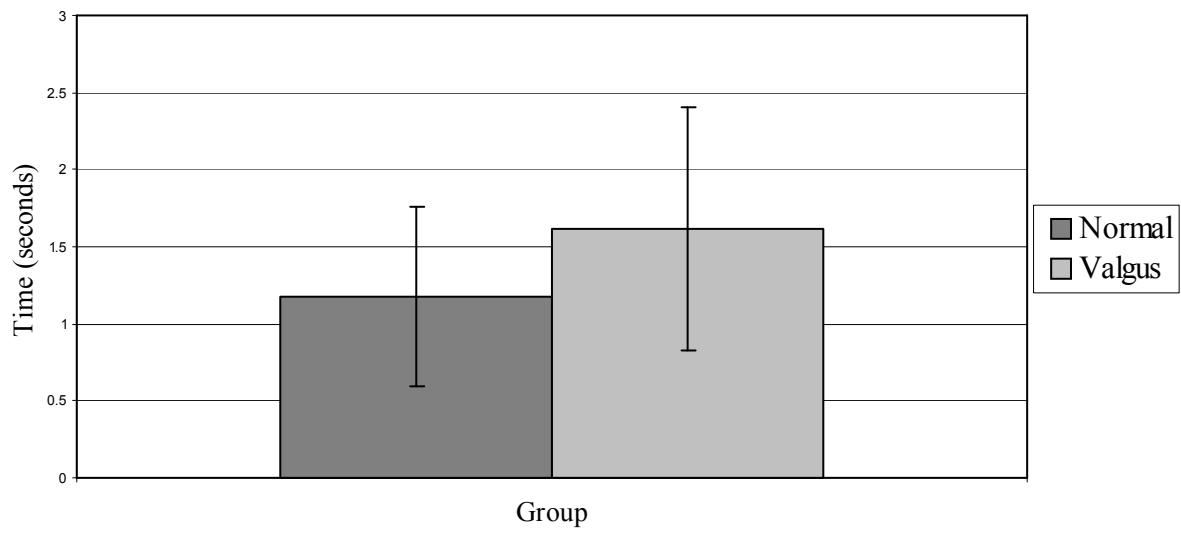


Figure 5: Adductor Mean EMG Amplitudes

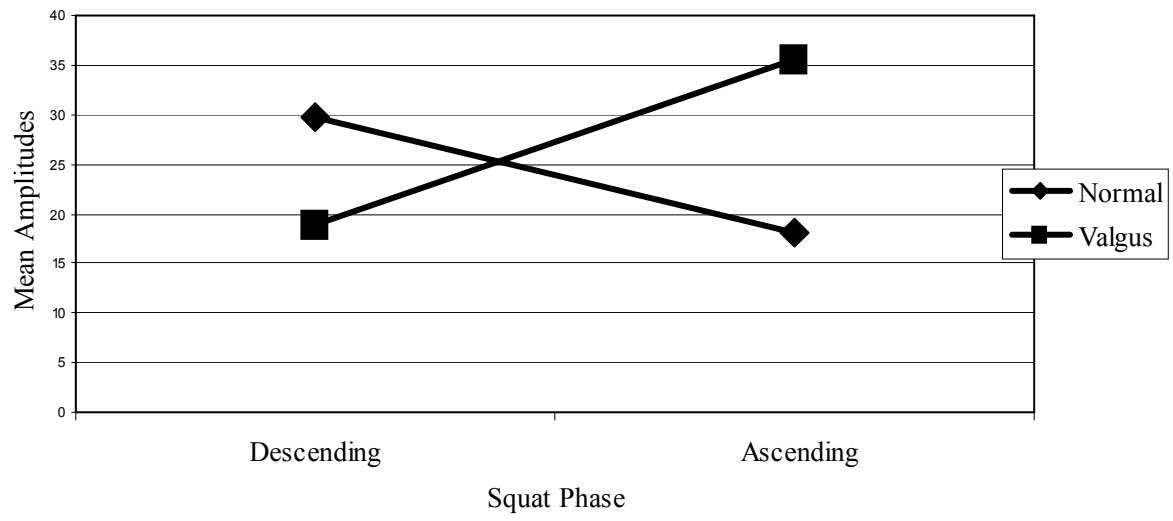


Figure 6: Hip Abduction ROM Measurement Positioning

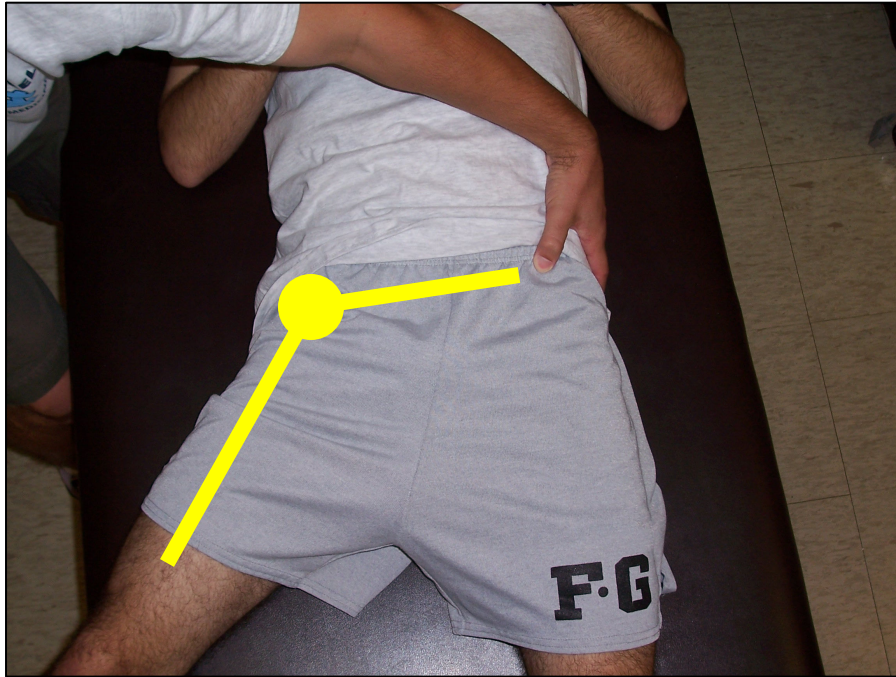


Figure 7: Hip External Rotation Measurement Positioning



Figure 8: Straight Knee Ankle Dorsiflexion ROM Measurement Positioning

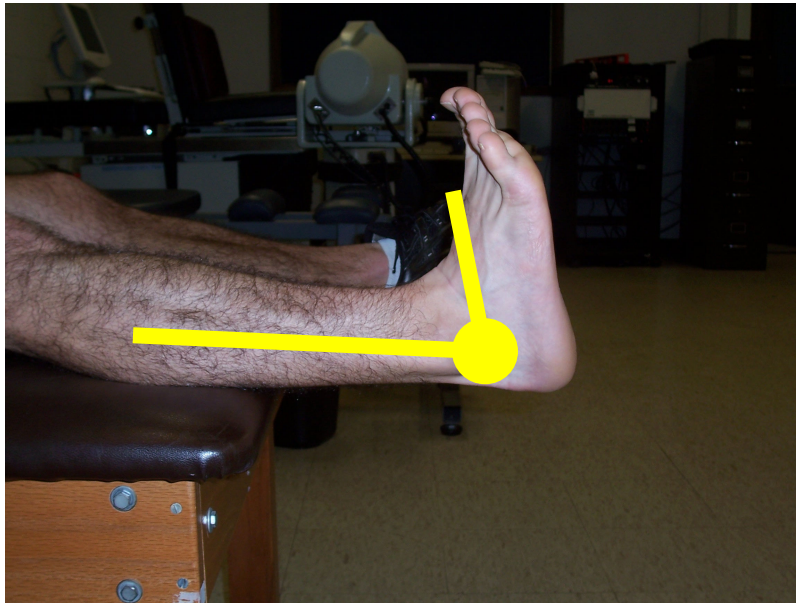


Figure 9: Bent Knee Ankle Dorsiflexion ROM Measurement Positioning



Figure 10: Gluteus Medius Electrode Placement

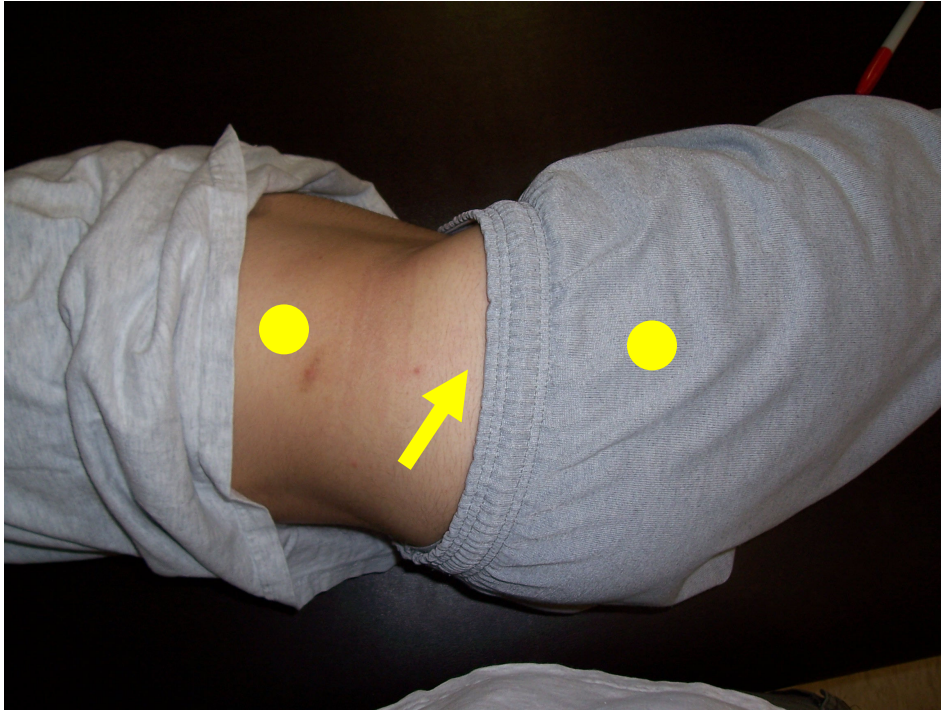


Figure 11: Gluteus Maximus Electrode Placement

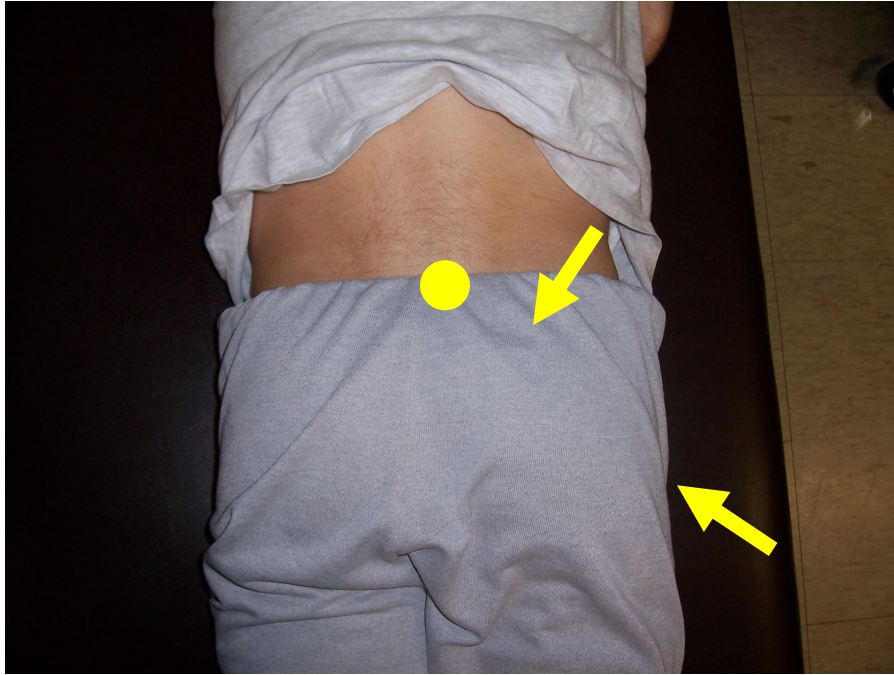


Figure 12: Adductor Electrode Placement

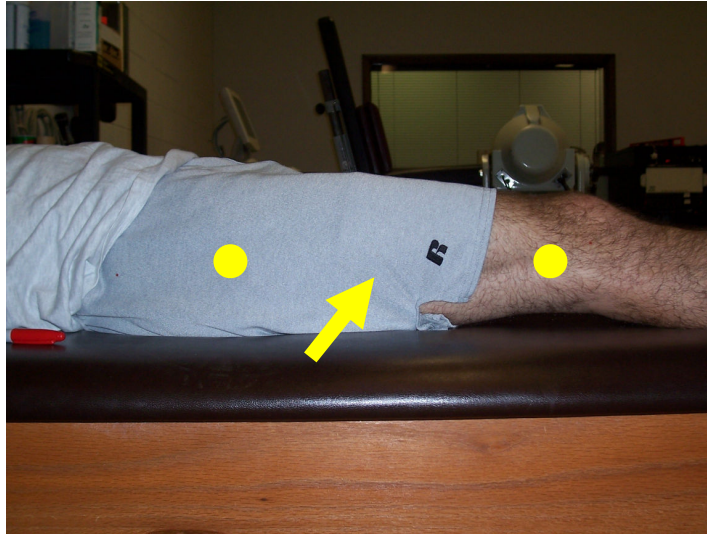


Figure 13: Hip External Rotation Strength Measurement Positioning



Figure 14: Hip Internal Rotation Strength Measurement Positioning

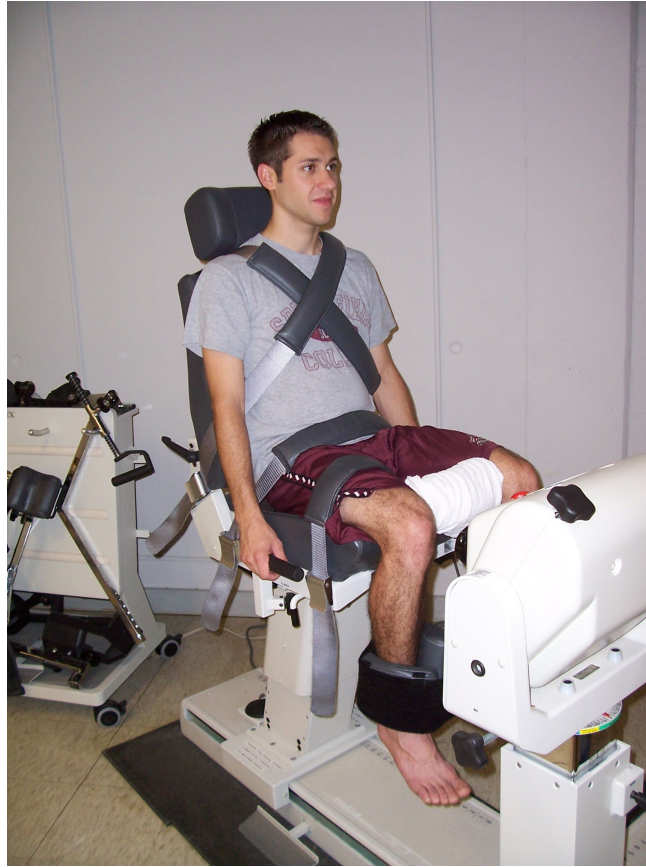


Figure 15: Hip Extension Strength Measurement Positioning

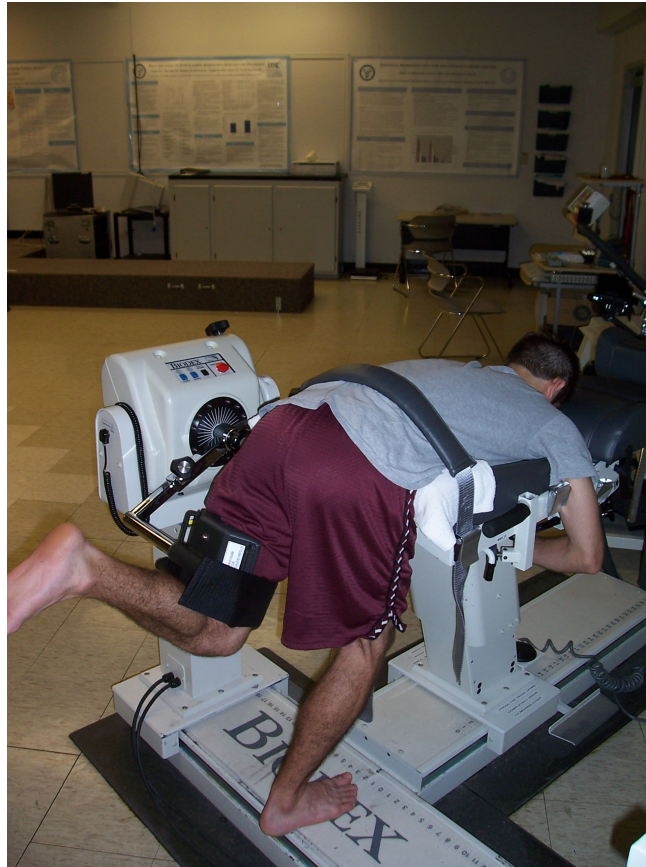


Figure 16: Hip Abduction Strength Measurement Positioning



Figure 17: Squat Task Positioning



Figure 18: Squat Task Positioning – Heel Block



APPENDIX C: MANUSCRIPT

The Effects of Hip Musculature on Knee Valgus During a Squat Task; and How Muscle Activity Changes When Standing on a Two-Inch Heel Block

Objective: To compare hip ROM, strength, muscle activity, and ankle dorsiflexion ROM between groups that demonstrate knee valgus during a squat task and those that do not (control). Also, determine if muscle activity changes after knee valgus is corrected by a two-inch heel block.

Design: A single-session experimental research design was used to compare the control group and the valgus group

Participants: Seventeen (10 Females, 7 Males) control subjects ($\text{age}[\text{yr}] = 23.82 \pm 5.76$, $\text{height}[\text{cm}] = 166.12 \pm 31.13$, $\text{weight}[\text{kg}] = 69.59 \pm 15.37$) and fourteen (12 Female, 2 Male) valgus subjects ($\text{age}[\text{yr}] = 22.36 \pm 3.08$, $\text{height}[\text{cm}] = 167.21 \pm 9.3$, $\text{weight}[\text{kg}] = 65.93 \pm 9.9$) without lower extremity injury.

Dependent Variables: Supine hip abduction and external rotation ROM. Ankle dorsiflexion ROM (straight and bent knee). Hip internal rotation, external rotation, extension, and abduction eccentric and concentric peak torque and time to peak torque. EMG mean amplitude of the gluteus maximus, gluteus medius and adductor complex during a squat with and without a two inch heel block.

Data Analysis: Mixed model analysis of variance tested for difference both between and within groups.

Results: A significant group by phase interaction effect ($p = 0.02$) existed for mean adductor amplitude between the valgus and the control group. A significant difference ($p < .001$) in straight knee ankle dorsiflexion ROM existed between the control and the valgus group. A

significant difference ($p = 0.034$) existed between the internal rotation concentric time to peak torque between the two groups.

Conclusion: The adductor complex plays a significant role in hip extension. Increased activity of the adductor during extension could pull the knees into valgus. Gastrocnemius tightness might contribute to knee valgus during a squat. There appears to be three distinct populations when considering this knee valgus squatting position. 1. Normal individuals who do not have valgus. 2. Ankle dysfunction individuals who's valgus position corrects with a heel block. 3. Hip dysfunction individuals who's valgus position does not correct with a heel block.

Keywords: Knee valgus, double leg squat, hip adductor activity, gastrocnemius flexibility

INTRODUCTION

Baseline screenings are commonly utilized in sports medicine as a method to assess injury status and establish return to play criteria should injury occur (McKeag 1985; Adirim and Cheng 2003). More recently baseline screenings have been used to develop corrective exercise programs to prevent injury from occurring (Nicholas and Tyler 2002). Typically baseline screenings have utilized isolated measures of flexibility, muscle strength, and functional performance; as well as identifying faulty posture (Wang, Chen et al. 2006). Acquiring this information serves two purposes. Initially this information is required to ensure that the athletes are not injured at the beginning of a competitive season. However, most often this information is used to identify possible problematic alignments and movements that might predispose an athlete to injury and address the problem before injury occurs. If risk factors can be identified and corrected in the preseason, some of these injuries may be prevented.

Posture is a very important and often neglected part of overall health. Ideal posture maintains that structural integrity and optimum alignment of each component of the kinetic chain (Clark, 2001). The kinetic chain consists of the myofascial system, articular system, and the neural system (Clark, 2001). When one component of this system is out of alignment, then the entire system is placed at a disadvantage. Postural malalignment is thought to create predictable patterns of tissue overload and dysfunction, initiating the cumulative injury cycle (Clark, 2001). This cumulative injury cycle begins with tissue trauma, inflammation, leading to muscle spasm, adhesions, altered neuromuscular control, and muscle imbalance. This cycle is thought to cause decreased athletic performance and eventual injury (Clark, 2001).

Functional movement analyses are becoming increasingly popular as a tool during dynamic postural assessment. Double leg and single leg squatting tasks are common during

functional movement analyses. These methods can be easily implemented and are time efficient, thus are attractive measures for the clinician. During functional movement analyses, the clinician observes for dysfunctional movement patterns that are believed to represent muscle imbalances caused by lack of flexibility, muscle weakness, and/or muscle activation. For example, individuals who demonstrate excessive knee valgus during a double leg squat are hypothesized to display a tight adductor muscle complex, tight gastrocnemius/soleus complex, and weak gluteus medius and maximus muscles, as well as a lack of neuromuscular control in the aforementioned muscles. (Clark 2001) However, research has not been performed to validate the idea of functional movement analyses as a method to identify muscle imbalances.

The adductor muscle complex functions both as a hip adductor and internal rotator. (Rosse 1997) Tightness in this complex would not allow full motion and cause the femur to pull into adduction and internal rotation. During a squat task, this may result in a valgus motion at the knee. The gluteus medius muscle has various functions. The anterior fibers function to internally rotate the hip. (Earl 2005) However, the most common thought of function of the gluteus medius is its role in hip abduction and stabilizing the hip and limiting adduction. (Schmitz, Riemann et al. 2002; Earl 2005) A weakness in this muscle that affected motion would allow the hip to excessively adduct that may cause the knee to exhibit a valgus motion. The gastrocnemius muscle is the prime mover in ankle plantar flexion. (Rosse 1997) A tightness in this muscle could affect both the ankle or the knee joint because it is bi-articular. During functional movements, it may cause either the heel to rise or the knee to flex excessively. Also due to its insertion into the calcaneus it may have an impact on the foot flattening. These muscle imbalances can lead to movement dysfunctions that cause an increase in stress to certain structures and contribute to the cumulative injury cycle.

Knee valgus alignment during dynamic tasks (e.g. squatting) is a common postural dysfunction seen in the lower extremity. Knee valgus alignment during squatting is defined as the mid-patella moving medially and crossing over the ipsilateral great toe as the knee flexes while squatting downward. Knee valgus alignment has been described as a potentially dangerous movement pattern (Ireland 1999). Knee valgus alignment is accompanied by movement of the hip into adduction and internal rotation, which are commonly described mechanisms for anterior cruciate ligament (ACL) injury (Ireland 1999). It has also been shown that patello-femoral compressive forces increase with knee valgus alignment (Escamilla 2001). Thus, knee valgus alignment is also believed to be a contributing factor to patello-femoral pain syndrome. Furthermore, the medial collateral ligament (MCL) is stressed when the knee is exposed to a valgus moment. The MCL can become injured when that valgus stress becomes too great. Individuals who undergo excessive knee valgus motion during functional tasks may place greater stress on the MCL and be at greater risk for injury. These injuries can be detrimental to an individual's physical well-being. Therefore, it is important to understand the factors that influence knee valgus alignment as this may improve our understanding of lower extremity injury risk factor, which may lead to the development of exercise programs to correct knee valgus alignment and reduce injury risk. Currently, knee valgus alignment is thought to occur due to weakness or inhibition in the hip abductor muscles (e.g. gluteus medius, gluteus maximus) and hip external rotator muscles. Meanwhile, the hip adductor and internal rotator muscles are believed to be tight or overactive (Clark 2001). However, research has not investigated if these hypothesized muscle imbalances actually exist in individuals demonstrating knee valgus alignment during a functional squatting task.

Squat tests, both single and double legged, have been used in the past as functional tests because they put the knee through common motions found in athletics.(Beutler, L.W. et al. 2002;

Loudon, Wiesner et al. 2002; Zeller, McCrory et al. 2003) Moreover, studies have been done that examine knee valgus during squat tasks. (Zeller, McCrory et al. 2003) However, currently research has yet to focus on how the abovementioned risk factors associated with injury differ between subjects who experience a position of knee valgus versus those who do not when performing a squat.

METHODS

Subjects

Approximately 70 subjects were screened by the principal investigator by watching them perform a double leg squat. Those subjects that were observed to exhibit a valgus motion during the squat were then observed with a two-inch heel block underneath their calcaneus. These subjects were recruited through volunteers from emails, fliers, and personal recruiting in classes. Subjects were divided into two groups based upon their results on the double leg squat task (DLST). The valgus group consisted of subjects who demonstrated bilateral knee adduction during the DLST that was corrected by placing a two-inch heel block under their calcaneus. The control group consisted of subjects who did not demonstrate knee adduction on either leg during the DLST. Subjects in the control group were matched to subjects in the hip dysfunction group by age, height, and weight. The control group consisted of 17 subjects while the valgus group consisted of 14 subjects whose ages ranged from 18-26 years. Subject demographics are located in Table 1. Subjects were otherwise healthy individuals that had no current musculoskeletal injuries or had not sustained an injury to either lower extremity in the past six months. Subjects were also excluded if they had surgery to either lower extremity in the past year. Furthermore, subjects in both groups were asymptomatic with and without activity, and subjects were asymptomatic during the DLST. Prior to all testing all subjects read and signed an informed

consent form approved by a University of North Carolina at Chapel Hill Biomedical Institutional Review Board (IRB).

Measurement and Instrumentation

A manual goniometer was used to measure joint angles in degrees for ROM assessment of the muscles of interest. Flexibility measurements were taken for the hip adductors ($ICC_{(2,1)} = 0.89$ SEM = 1.743), the hip internal rotators ($ICC_{(2,1)} = 0.97$ SEM = 2.625), gastrocnemius, and soleus. ICCs were not calculated for these final two range of motion measures.

Concentric and eccentric muscle strength was evaluated using a Biodex System 3 Pro isokinetic dynamometer (Biodex Medical Systems, Shirley, NY), measured in Foot-pounds (Ft*lbs) of torque. The data were then analyzed with a customized Matlab 7.0 program (The Mathworks, Inc., Natick, MA) to determine the average peak torque, as well as time to peak torque, for the hip external rotators, internal rotators, hip abductors, and hip extensors. Each average was measured over the middle three of five total trials.

The mean amplitude electromyographic (EMG) activity of the gluteus medius, gluteus maximus, and adductor group was measured using an eight channel DelSys Bagnoli EMG System (Boston, MA). DE-2.1 single differential surface electrodes (DelSys Boston, MA) with a contact dimension of 1.0cm x 0.1cm and a contact spacing of 1.0cm over the muscle bellies and parallel to the fibers of the gluteus medius, gluteus maximus, and adductor muscles. Electrodes were plugged into a belt mounted I/O unit that plugged into the DelSys Bagnoli EMG system. This system amplified ($\times 10,000$ Hz) the EMG signal as it passed into the computer and was stored for analysis. The EMG data were processed with passive demeaning (0.0 ms begin/ 10.0 ms end), a Butterworth notch filter at 60.0 Hz, a band pass Butterworth filter from 10.0 Hz – 350 Hz, and finally RMS smoothing was used at a time constant of 25 msec. All EMG data were normalized to the percentage of maximum voluntary isometric contraction to allow for

comparison between subjects. All EMG data collection and processing was done using Datapac2K2 (Run Technologies, Mission Viejo, CA).

An electro-goniometer was placed over the knee joint so that knee joint angle could be recorded and related to EMG activity during the DLST. Electro-goniometer data were filtered with a low pass Butterworth at 15.0 Hz.

A tripod was used to standardize squat depth; as well as a metronome to standardize squat speed at 66 beats per minute. Subjects used two beats to descend, two beats to ascend and one beat to rest between squats.

Procedures

Students and faculty of the University of North Carolina at Chapel Hill were asked through email and fliers to volunteer for screening for this study. Screening consisted of a questionnaire pertaining to previous history of injury or surgery to the lower extremity. Each subject completed their individual testing in one session. Prior to testing, subjects had the testing procedures explained to them and were asked to read and sign an informed consent form. Each testing session began with the recording of the subject's gender, age (years), height (cm), and weight (kg). Subjects were then video taped performing the DLST both with and without a two-inch heel block to ensure that they met the inclusion criteria for their assigned group. The tester then assessed the subject's dominant leg, defined as the leg they would use to kick a soccer ball for maximum distance, for the following variables: the subject's passive hip abduction ROM; passive external rotation ROM; passive straight knee ankle dorsiflexion ROM; passive bent knee dorsiflexion ROM; Concentric/eccentric isokinetic strength for hip external and internal rotation, hip abduction, and hip extension; and EMG activity during a DLST in the gluteus medius, gluteus maximus, and adductor complex. The valgus group also had EMG recorded for a DLST

while standing on a two-inch heel block. Prior to testing subjects were allowed to warm up for five minutes on a stationary bike at a self-determined pace.

ROM Measurements

Subject positioning for ROM measurement of hip abduction was supine. The axis of the goniometer was placed over the ASIS of the extremity being measured. The stationary arm was placed along an imaginary line from one ASIS to the other. The movement arm was placed along the anterior midline of the femur using the midline of the patella for reference. The subject was then passively abducted until the tester felt the contralateral ASIS begin to move (Norkin 1995). Three separate measurements were taken and the mean was recorded.

Subject positioning for ROM measurement for hip internal rotation was supine with the dominant hip and knee flexed to 90 degrees. The hip was in neutral abduction/adduction. The axis of the goniometer was the anterior center aspect of the patella. The stationary arm was positioned parallel to the table. The movement arm was placed along the anterior midline of the tibia pointing towards the anterior midpoint of the ankle between the two malleoli. (Norkin 1995) The subject's hip was passively externally rotated until the tester felt the end range. Three separate measurements were taken and the mean was recorded.

Subject positioning for ROM measurement of straight knee dorsiflexion was supine with the dominant knee in full extension. The ankle was in neutral inversion/eversion. The axis of the goniometer was distal lateral malleolus. The stationary arm was positioned with the lateral midline of the fibula, using the fibular head for reference. The movement arm was placed parallel to the lateral aspect of the fifth metatarsal (Norkin 1995). The subject's ankle was then passively dorsiflexed until the tester felt the end range. Three separate measurements were taken and the mean was recorded.

Subject positioning for ROM measurement of bent knee dorsiflexion was the same as straight knee; however, a bolster was placed under the subject's dominant knee so that it was flexed to at least 30 degrees(Norkin 1995). The subject's ankle was then passively dorsiflexed until the tester felt the end range. Three separate measurements were taken and the mean was recorded.

Electromyography

The sites for electrode placement were shaven and cleansed with alcohol to improve signal transmission from the muscles. Electrodes were placed over the muscle bellies of the gluteus medius, gluteus maximus, and adductor complex. A reference electrode was placed on the tibial tuberosity. Electrodes were placed in a parallel to muscle fibers.

The electrodes for the gluteus medius were placed halfway between the iliac crest and the greater trochanter of the femur (Kleissen 1990). The electrodes for the gluteus maximus were placed 20% of the distance between the spinous process of S2 and a point 10 cm distal to the greater trochanter. The electrodes for the adductor complex were placed on the muscle belly at the mid point of the femur (Leveau 1992). All electrode placements were confirmed with an isometric manual muscle test and checked for cross talk. Maximal voluntary isometric contraction (MVIC) was used to normalize muscle activity between subjects (% MVIC).

Strength Measurements

Reliability for all strength tests was demonstrated by Halverson and Hawkey, abstract published in Journal of Athletic Training supplement 2004.

The hip external rotators were tested in a seated position. The subject's hip and knee was flexed to 90 degrees. The dynamometer was aligned with the long axis of the femur. Pressure was applied to the medial aspect of the distal tibia. The thigh was stabilized to the chair using

straps. A towel was also placed between the subjects' knees to act as a fulcrum for external rotation and prevent adduction. The external rotators were tested through twenty degrees of motion that began at five degrees of external rotation and ended at fifteen degrees of internal rotation.

The hip internal rotators were tested in a seated position. The subject's hip and knee were flexed to 90 degrees. The dynamometer was aligned with the long axis of the femur. Pressure was applied to the lateral aspect of the distal tibia. The thigh was stabilized to the chair using straps. The internal rotators were tested through twenty degrees of motion that began at five degrees of internal rotation and ended at fifteen degrees of external rotation.

The hip abductors were tested with the subject in a side-lying position to isolate the gluteus medius. The joint axis was located at 0.5 inches medial to the ASIS at the level of the greater trochanter (Lyons, Perry et al. 1983). The hip was abducted and externally rotated, making sure not to let the trunk and pelvis rotate backward. Pressure was applied against the thigh in the direction of adduction. Pressure was not applied against the external rotation component of the start position (Kendall 1993). The trunk was stabilized by strapping the subject to the chair. The muscle was tested from 0-20 degrees of hip abduction.

The hip extensors were tested in a supported, standing position in an attempt to isolate the gluteus maximus and maximize stabilization. The dynamometer axis of rotation was aligned with the anterior superior tip of the greater trochanter (Lyons, Perry et al. 1983). The subjects stood in front of the Biodex chair, and the seat was raised to the level of the subjects' ASIS. If the chair was unable to reach the subjects' ASIS, the subjects were then asked to flex the contralateral knee until the chair was even with the level of the ASIS. The subjects flexed their trunk to ninety degrees and laid their chest on the chair. The trunk was stabilized to the Biodex chair prior to testing using straps. This stabilization was aimed at preventing accessory trunk

motions that might influence strength testing. The knee of the test leg was flexed to ninety degrees. The stance leg (non-test leg) was flexed at the knee such that the subjects' chest was comfortably resting on the chair. The subjects were then asked to actively extend the hip through a range of motion that began at ninety degrees of hip flexion (femur perpendicular to the ground) and ended at fifty degrees of hip flexion. Pressure was applied against the distal portion of the posterior thigh in the direction of hip flexion (Kendall 1993).

All muscles were tested concentrically and eccentrically at $60 \text{ deg} \cdot \text{sec}^{-1}$. The testing procedure accounted for gravity corrections during hip abduction and hip extension testing since the test limb was sufficiently close to the horizontal plane. The test limb did not come close enough to the horizontal plane to warrant a gravity correction during the testing of the hip external or internal rotators.

Following the isokinetic testing of each muscle group, the subject's strength was tested isometrically. The positioning and alignment of the subjects was exactly the same for the isometric testing as it was used during the isokinetic testing. The test limb was positioned in the middle of the isokinetic ROM or neutral position for isometric testing. The test limb was placed at seventy degrees of hip flexion while testing the hip extensors. The test limb was placed at ten degrees of hip abduction while testing the hip abductors. The test limb was placed in a neutral position (zero degrees of abduction/adduction) while testing the hip adductors. Each subject performed three maximal isometric contractions. The contractions were held for five seconds each, and there was 10-12 seconds rest between each repetition. The mean amplitude of the three trials during the middle three seconds of each trial was recorded.

The EMG data for the MVIC of the gluteus maximus was recorded while testing hip extension. The EMG data for the MVIC of the gluteus medius was recorded while testing hip abduction. The EMG data for the MVIC of the adductor complex was recorded while still in the

testing position for hip abduction. The subjects were asked to maximally adduct the hip against the Biodex dynamometer for five seconds. EMG from these MVIC's was recorded so that the EMG data recorded during the DLST could be normalized and used to compare between subjects.

Double-Legged Squat Task

Subjects were asked to stand with hips, feet, and knees facing forward and arms fully extended overhead parallel to their ears. The subjects then were instructed to go down into a squat position as if they were trying to sit in a chair. Squat depth was standardized to 70 degrees of knee flexion to ensure that each subject went through at least 60 degrees of motion. Subjects were instructed to squat while their knee joint angle was measured with a manual goniometer. When the desired amount of 70 degrees of knee flexion was achieved a tripod was set up to assist the subject in knowing the proper squat depth. An electrogoniometer was then placed on the subject's knee so that EMG data could be examined in relation to knee position. The subjects then performed the squat task five times. EMG activity was recorded for the gluteus medius, gluteus maximus, and the adductor complex from 0-60 degrees of knee flexion during each of the squat tasks. Subjects in the control group only performed the DLST without the heel block. Subjects in the valgus group performed the DLSQ with and without the heel block while EMG data were recorded.

Data Analysis

One way analysis of variance tests were used to analyze differences between groups in the ROM measurements. A 2 x 2 analysis of variance was used to analyze the straight and bent knee ankle dorsiflexion ROMs. 2 x 2 mixed model analysis of variance tests were used to analyze differences between groups in concentric and eccentric peak torque as well as time to

peak torque. EMG data differences were analyzed both between groups and between phases within groups with separate 2 x 2 mixed model analyses of variances. Tukey Post Hoc testing was used to identify where significance was found for the ankle dorsiflexion and EMG data. All data were analyzed using SPSS 13.0 (Chicago, IL). All null hypotheses were tested for significance at $p < .05$.

RESULTS

ROM Measurements

Means, standard deviations, and effect size for hip abduction ROM are presented in Table 2. There was no significant difference [$F_{(1,29)} = 0.01$, $p = .920$] for hip abduction range of motion between the control and the valgus groups. Means, standard deviations, and effect size are presented in Table 2. There was no significant difference [$F_{(1,29)} = 0.859$, $p = .362$] for hip external rotation ROM between the control and the valgus groups. There was a significant group by knee position interaction effect for ankle dorsiflexion [$F_{(1,26)} = 17.4$, $p < .001$]. Tukey Post Hoc tests revealed that the difference between groups with straight knee dorsiflexion was significant. This difference is illustrated in Figure 1.

Hip External Rotation Concentric/Eccentric Peak Torque

Means, standard deviations, and effect sizes are presented in Table 3. There was no significant difference for hip external rotation concentric [$F_{(1,29)} = 1.309$, $p = .262$] or eccentric [$F_{(1,29)} = .514$, $p = .479$] peak torque between the control and valgus groups.

Hip Internal Rotation Concentric/Eccentric Peak Torque

No significant differences were observed for hip internal rotation concentric [$F_{(1,29)} = 0.00$, $p = .985$] or eccentric [$F_{(1,29)} = 0.06$, $p = .808$] peak torque between the control and valgus groups.

Hip Extension Concentric/Eccentric Peak Torque

There was no significant difference [$F_{(1,29)} = 0.046$, $p = .833$] for hip extension concentric peak torque between the control and valgus groups. There was no significant difference [$F_{(1,29)} = 0.305$, $p = .585$] for hip extension eccentric peak torque between the control and valgus groups.

Hip Abduction Concentric/Eccentric Peak Torque

There was no significant difference [$F_{(1,29)} = 3.682$, $p = .065$] for hip abduction concentric peak torque between the control and valgus groups. However, the p value suggests a trend and the finding might reach significance if the statistical power was greater. This trend is illustrated in Figure 2. There was no significant difference [$F_{(1,29)} = 0.006$, $p = .939$] for hip abduction eccentric peak torque between the normal and valgus groups.

Hip External Rotation Concentric/Eccentric Time to Peak Torque

Means, standard deviations, and effect size are presented in Table 4. There was no significant difference [$F_{(1,29)} = 0.376$, $p = .544$] for hip external rotation concentric time to peak torque between the control and valgus groups. There was no significant difference [$F_{(1,29)} = 0.038$, $p = .847$] for hip external rotation eccentric time to peak torque between the control and valgus groups.

Hip Internal Rotation Concentric/Eccentric Time to Peak Torque

There was a significant difference [$F_{(1,29)} = 4.976$, $p = .034$] for hip internal rotation concentric time to peak torque between the control and valgus groups. The normal group achieved peak torque almost a full second faster (2 sec vs. 3 sec) than the valgus group (fig 3). There was no significant difference [$F_{(1,29)} = 0.023$, $p = .880$] for hip internal rotation eccentric time to peak torque between the normal and valgus groups.

Hip Extension Concentric/Eccentric Time to Peak Torque

There was no significant difference [$F_{(1,29)} = 0.456$, $p = .501$] for hip extension concentric time to peak torque between the control and valgus groups. There was no significant difference

[$F_{(1,29)} = 1.235$, $p = .276$] for hip extension eccentric time to peak torque between the control and valgus groups.

Hip Abduction Concentric/Eccentric Time to Peak Torque

There was no significant difference [$F_{(1,29)} = 3.218$, $p = .083$] for abduction concentric time to peak torque between the control and valgus groups. However, the p value suggests a trend that the valgus group required a half second longer (1.1 sec vs. 1.6 sec) to reach peak torque. This trend is illustrated in Figure 4. There was no significant difference [$F_{(1,29)} = 1.725$, $p = .199$] for hip abduction rotation eccentric time to peak torque between the normal and valgus groups.

Gluteus Maximus EMG Mean Amplitude During a Squatting Task

Means, standard deviations, and effect size are presented in Table 5. There was no main effect for group [$F_{(1,27)} = 0.002$, $p = .962$] for gluteus maximus activity during the ascending and descending phases of the squat. No main effect for phase [$F_{(1,27)} = 0.024$, $p = .877$] was observed for gluteus maximus activity between the ascending and descending phases of the squat for the control or valgus groups. There was no group by phase interaction effect [$F_{(1,27)} = 0.020$, $p = .888$] for gluteus maximus activity during the squat task.

Gluteus Medius EMG Mean Amplitude During a Squatting Task

There was no main effect for group [$F_{(1,27)} = 1.037$, $p = .318$] during the ascending or descending phases of the squat between the control or valgus groups. There was no main effect for phase [$F_{(1,27)} = 0.101$, $p = .753$] of gluteus medius activity between ascending and descending phases of the squat for either group. No group by phase interaction effect [$F_{(1,27)} = 0.098$, $p = .757$] was observed for gluteus medius activity during the squat task.

Adductor Mean EMG Amplitude During Squatting

There was a group by phase interaction [$F_{(1,27)} = 6.095$, $p = .020$] for adductor activity during the squat task. The control group had higher mean amplitude during the descending portion of the squat and decreased during the ascending phase. In contrast, the valgus group had higher adductor activity during ascending than descending. Tukey post hoc testing revealed that the difference between adductor activity of the control group and the valgus group during the ascent phase was significant. This significance is illustrated in Figure 5. No main effect for group [$F_{(1,27)} = 0.163$, $p = .690$] was observed for adductor activity during the ascending or descending phases of the squat. There was no main effect for phase [$F_{(1,27)} = 0.196$, $p = .661$] for adductor activity between the ascending and descending phases of the squat for either the normal or the valgus group.

Gluteus Maximus EMG Mean Amplitude During Squatting with Heel Block

Means, standard deviations, and effect size for mean gluteus maximus amplitude during the squatting task on the heel block are presented in Table 6. There was no main effect for the heel block [$F_{(1,11)} = 0.134$, $p = .721$] for gluteus maximus activity during the ascending and descending phases of the squat between the valgus group with and without the heel block. There was no main effect for phase [$F_{(1,11)} = 1.525$, $p = .241$] for gluteus maximus activity between the ascending and descending phases of the squat for the valgus group either with or without the heel block. There was no heel block x phase interaction effect [$F_{(1,27)} = 1.430$, $p = .255$] for gluteus maximus activity during the squat task.

Gluteus Medius Mean EMG Amplitude During Squatting with a Heel Block

Means, standard deviations, and effect size for gluteus medius EMG mean amplitude during the squatting task on the heel block are presented in Table 6. No main effect was observed for gluteus medius activity [$F_{(1,11)} = 0.022$, $p = .885$] during ascending and descending phases of a squat between the valgus group with and without the heel block. There was no main

effect for phase [$F_{(1,11)} = 2.034$, $p = .179$] of gluteus medius activity between the ascending and descending phases of the squat for the valgus group either with or without the heel block. There was no heel block by phase interaction effect [$F_{(1,11)} = 1.955$, $p = .187$] for gluteus medius activity during the squat task.

Adductor EMG Mean Amplitude During Squatting with and without a Heel Block

Means, standard deviations, and effect size for mean adductor amplitude during the squatting task on the heel block are presented in Table 6. There was no main effect for heel block [$F_{(1,11)} = 2039$, $p = .148$] for adductor activity during the ascending and descending phases of the squat between the valgus group with and without the heel block. There was a main effect for phase [$F_{(1,11)} = 10.951$, $p = .006$] of adductor activity between the ascending and descending phases of the squat for the valgus group while on the heel block. Adductor activity during the descending phase of the squat was less than the ascending phase. This significance is illustrated in Figure 6. There was no heel block by phase interaction effect [$F_{(1,11)} = 1.7$, $p = .217$] for adductor activity during the squat task.

DISCUSSION

The most important findings from this study are that individuals who demonstrated knee valgus during a squat and have the valgus position corrected with a two-inch heel block have different adductor muscle activity during a squat. Our results also show these valgus individuals have less passive ankle dorsiflexion ROM (with a straight knee) when compared to a control group. Moreover, the passive bent knee ankle dorsiflexion ROM of the valgus group was significantly increased over their straight knee ankle dorsiflexion. The control group reached concentric peak torque of the internal rotators quicker than the valgus group. Trends in the data suggest that the control group has larger concentric peak torque in the abductors and achieve peak torque during concentric abduction faster when compared to the valgus group.

Muscle Activity During Squatting

An analysis of the control group shows that muscle activity during different phases of a squat (descending and ascending) is not different. This means the control group uses hip musculature equally. This is not the case for those who display a valgus motion during a squat and the difference seems to be in the adductor complex. The other two muscles examined (gluteus maximus and gluteus medius) showed similar activity patterns when compared to the control group.

A group by phase interaction was demonstrated by the adductor muscle complex during the squat task. When comparing maximal muscle activity, the control group's adductor muscle had the greatest activity during the decent phase while the valgus group was most active during the ascent phase (Figure 5). Thus, subjects performing proper squat posture utilized all three muscles simultaneously to eccentrically lower themselves, while the valgus group adductor complex was not as active in the eccentric lowering during the descent phase. However, during the ascent phase the control group's adductors were not as active and relied on the gluteus maximus to assist with hip extension. It is important to note that neither the decrease in the adductor activity nor the increase in the gluteus maximus activity from the descent phase to the ascent phase in the normal group were deemed significant. The opposite situation was found to exist with the adductor complex of the valgus group. Adductor activity during the decent phase was significantly less than adductor activity during the ascent phase. Furthermore, their adductor activity during the ascent phase was significantly greater than that of the normal group during the same phase.

Subjects in the valgus group in this study had their valgus corrected when squatting on a two-inch heel block. Current theory (Clark 2001) suggests valgus may result from tight musculature of the ankle rather than weak hip musculature. The heel block allows for increased

length of the gastrocnemius and soleus complex, allowing normal motion and correcting valgus. Further research needs to identify individuals that present with valgus that is not corrected by a heel block and validate that hip musculature is the issue and what role it may play in knee valgus.

Increased adductor activity in the valgus group during the ascent phase of a squat could be explained an improved length-tension relationship of the muscle group due to increased hip flexion. One study examined moment arms of hip extensor muscles (gluteus maximus and adductor magnus) and found that the gluteus maximus moment arm decreased while the adductor magnus moment arm increased while the hip was flexed. The adductor magnus extension moment arm was shown to continue to increase until 75 degrees of hip flexion (Nemeth and Ohlsen 1985). Furthermore, it has been shown that women demonstrate significantly more hip flexion and adduction during single-legged squats (Zeller, McCrory et al. 2003). While our study examines double-legged squats, this is still relevant because valgus was part of our inclusion criteria and a majority of our valgus subjects were women. Females are being put into a position of hip flexion and adduction that may increase their adductor muscle activity during the ascent phase. While in the control group the gluteus maximus increases during the ascent phase, it actually slightly decreases in the ascent phase for the valgus group. This may be explained by the increase in hip adduction decreasing the gluteus maximus' ability to function as an extensor. This position of knee adduction is associated with femoral internal rotation (Ireland 1999). During the ascent phase the gluteus maximus may be functioning more in its role as a hip external rotator to compensate for femoral internal rotation. If this is the case, then it makes sense that the adductor activity increases as they function more in extension. However, the changes in gluteus maximus muscle activity in our study were not found to be significant.

Moreover, we did not measure hip flexion angles in our subjects. Therefore this theory needs to be explained further through future research.

A final possible explanation of the difference seen between our groups in adductor muscle activity arises from stance width. Research suggests stance width has a significant effect on adductor longus muscle activity during a squat task (McCaw and Melrose 1999). Stance width was not measured in this study but we used a self-selected stance width set at shoulder width apart and researcher verified. We think this minimizes any effect stance width may have on our data.

There were no significant differences found in any of the muscles tested in the valgus group between the regular squat and the squat on the heel block, except for the significant difference in adductor activity between phases that had already been discussed in comparison to the control group. This is the first study to examine how hip muscle activity changes when individuals with knee valgus squat on a two-inch heel block. More research needs to be done to fully understand how plantar flexing the ankle by placing an individual on a two-inch heel block affects squatting.

The lack of significance in the muscle activity between the valgus group with and without the heel block begins to outline three populations. These populations are a normal or control group, an ankle dysfunction group and a hip dysfunction group. The control group demonstrates a proper squat technique and keeps their knees centered over their toes (Clark 2001). The ankle dysfunction group is the group examined in our study and demonstrates knee valgus during a squat that is corrected by placing a two inch block under their heel. The hip dysfunction group consists of individuals who demonstrate knee valgus during a squat that is not corrected by the heel block. Currently no study has focused on the hip dysfunction group to verify current theory as to what is causing their knee valgus.

Straight and Bent Knee Ankle Dorsiflexion ROM

Another significant finding of this study was the difference of the straight knee ankle dorsiflexion ROM between the valgus group and control group and the difference of the valgus group between straight knee and bent knee. The valgus group had significantly less straight knee dorsiflexion. The gastrocnemius is a bi-articulate muscle crossing the ankle and knee. When the knee is straight, ROM of the ankle decreases because the gastrocnemius reaches end range, limiting motion. Tightness in the gastrocnemius muscle may have played a role in valgus during the DLST. This point is illustrated further when the knee was bent. The valgus group's dorsiflexion range of motion was significantly increased and approximately equal to the control group. This idea is the foundation of current theory! When the valgus group squatted with the heel block, the gastrocnemius shorts and the resulting tension is removed from the squat, correcting knee valgus. This suggests that the tightness in the gastrocnemius muscle played a role in the knee valgus. It is important to note that when the valgus group was on the heel block, there were no significant changes in hip musculature activity.

Ankle dorsiflexion has been shown to be around 10 degrees during the stance phase of gait (Murray 1967). During more function tasks, such as sit-to-stand, stair climbing, and sport specific activities, the requirement for ankle dorsiflexion ROM can increase to about 25 degrees (Andriacchi, Andersson et al. 1980; Lindsjo, Danckwardt-Lilliestrom et al. 1985; Livingston, Stevenson et al. 1991). The double-legged squat used in our study can be related to the function activities that can require the greater dorsiflexion. Thus, the lack of motion seen in the valgus group can explain their inability to perform the task properly. However, it becomes hard to explain why gastrocnemius tightness in and of itself would contribute to knee valgus. It may be a unilateral issue. If the lateral gastroc is tight or short, it may be responsible for pulling the knee into a valgus position. Currently it is believed that during a squat decreased ankle dorsiflexion

would lead to the heels of the individual rising up off of the floor (Clark 2001). This idea makes more sense when examining the gastrocnemius muscle by itself; however, when considering the entire kinetic chain there appears to be more to the picture. It makes more sense to think that this tightness in the gastrocnemius would pull, from its insertion, the calcaneus into eversion, thus causing the foot to pronate. Our study did not examine foot and ankle pronation but if this is the case, then our results seem to suggest that when the arch flattens and pronation ceases the gastrocnemius begins to influence an increase in knee valgus. Therefore it seems that more research needs to be done to determine whether or not the gastrocnemius muscle itself can cause knee valgus or if it only contributes to increased valgus when it is already present.

Internal Rotation Time to Peak Torque

Our study found that the control group reached their internal rotation concentric peak torque approximately a full second (2 sec vs. 3 sec) before the valgus group. This difference was deemed statistically significant and further strengthens the idea that there is a difference in adductor activity between groups. The current literature suggests that the gluteus maximus and gluteus medius are important hip stabilizers (Gottschalk, Kourosh et al. 1989; Clark 2001; Schmitz, Riemann et al. 2002) Our study showed that the gluteus maximus and gluteus medius activity between the control and the valgus groups were similar even though the valgus group demonstrated poor proximal hip control. This suggests that it may not be the gluteal muscles that are responsible for stabilization but rather other hip internal and external rotators. This can be further explained by the differences discussed above about the adductor activity between the two groups, as the adductors also function as hip internal rotators.

Abduction Concentric Peak Torque and Time to Peak Torque

There was a trend discovered in our study that suggests the control group produced greater concentric peak torque in the abductors than the valgus group and the control group

achieved peak torque a half second faster (1.1 sec vs. 1.6 sec). The lack of significance of this may be due to lack of statistical power; an increase in power may draw out the significance of this finding. Regardless, this finding is clinically significant as it suggests that one potential cause of the knee valgus position is that the valgus group lacks abduction strength as well as the ability to achieve maximum strength in time to prevent hip adduction. This could be related to the previous stated findings of ankle dorsiflexion range of motion. It may be that the adduction begins due to a lack of abduction stabilization strength and is increased by the lack of gastrocnemius flexibility and decreased ankle dorsiflexion

Ireland has described a “position of no return” as when an athlete has poor hip control and the hip moves into adduction, leading to femoral internal rotation and a position of knee valgus (Ireland 1999). These trends discovered in our study seem to support this and further explain it by demonstrated ankle dorsiflexion ROM’s role in this knee adduction. However, further research is required to further explore these trends and determine if they are significant.

Hip Abduction and Internal Rotation ROM

No difference was observed in hip abduction or internal rotation ROM between the control and valgus groups. Initially, we hypothesized that tight adductors or tight internal rotators may pull individuals into knee valgus. However, our study failed to identify this as a contributing factor. In fact the means of each ROM between groups was almost identical.

Hip Extension and External Rotation Concentric and Eccentric Peak Torque

No statistical difference was observed between group concentric or eccentric peak torques. This finding is clinically significant because it suggests that the gluteus maximus muscle does not play a significant role in preventing knee valgus during a squat. It is currently thought that individuals who demonstrate knee valgus have weak gluteus maximus control during squatting (Clark 2001). Again it becomes important to note that it appears that the

subjects tested had ankle dysfunction and not necessarily hip dysfunction, and that individuals not corrected by the heel block may have differences in gluteus maximus strength.

Limitations

The greatest limitation of this study was the lack of statistical power for some of the analyses. Ideally, each group would have around 20 subjects to have enough power to draw out significant differences that may exist. This was unable to be achieved in this study due to the difficulty in identifying subjects who met the valgus group criteria. However, most of the analyses that lacked power also had small effect sizes. Therefore, if power was increased such that significance was found, it still may not be clinically significant.

Another limitation for this study is that the order of testing was not randomized. Every subject followed the same protocol in the same order. However, since the subjects were observed to exhibit the bilateral knee valgus both before and after ROM and strength testing it is not believed that either altered the way in which the subjects performed the squat task.

The inability to identify many males who fit the inclusion criteria for the valgus group limits the generalizability this study. The control group is more evenly divided between males and females; however, the valgus group consists of primarily females. Although, the demographics between each group suggest that subjects were closely matched by age, height, and weight.

The lack of lower leg EMG activity is also a limitation of this study. While the ROM of ankle dorsiflexion was measured, no information about the muscle activity around the ankle of the ankle dysfunction group was available to help further understand what may be contributing to their valgus knee position.

Future Research

Current theory places subjects into categories if they are identified as having knee valgus (Clark 2001). There is a control population that demonstrates correct squat technique. There is a hip dysfunction population that demonstrates bilateral knee valgus both with and without a two-inch heel block. Finally, there is an ankle dysfunction population that demonstrates bilateral knee valgus during a squat that is corrected by a two-inch heel block. This study contrasted the control population against the ankle population and how hip musculature and ankle ROM contributes to knee valgus. Future research needs to verify that this classification system is correct and document differences between groups. Areas of focus should be the hip musculature (gluteus maximus, gluteus medius and adductor complex), ankle musculature (medial and lateral gastrocnemius, tibialis anterior, and soleus), and these variables relation to this position of knee valgus during a squat task. Future research should also focus on the kinematics of the squat to verify that valgus is corrected or not with the heel block. Finally, once the theory has been established, research should attempt to implement training programs to correct valgus.

APPENDIX D: IRB MATERIALS

OFFICE OF HUMAN RESEARCH ETHICS -- Institutional Review Board
INSTRUCTIONS FOR APPLICATION FOR IRB APPROVAL
OF HUMAN SUBJECTS RESEARCH
Version 28-Sep-2005

What is the purpose of this form?

This application is to seek *initial* IRB approval for a research study.

What parts of this application should you submit?

- For ***all studies***, submit Part A, which consists of these sections:
 - Part A.1. Contact Information, Agreements, and Signatures
 - Part A.2. Summary Checklist
 - Part A.3. Conflict of Interest Questions and Certification
 - Part A.4. Questions Common to All Studies
 - Part A.5. The Consent Process and Consent Documentation (including Waivers)
- For ***studies that involve direct interaction*** with human subjects (any contact with subjects including questionnaires, interviews, focus groups, observation, treatment interventions, etc), submit:
 - Part B. Questions for Studies that Involve Direct Interaction with Human Subjects
- For ***studies*** that use data, records or human biological specimens ***without direct subject contact***, submit:
 - Part C. Questions for Studies using Data, Records or Human Biological Specimens without Direct Contact with Subjects

Note: You should submit Parts B or C only as applicable. If the study involves *both* direct interaction *and* data collection without contact, use both Parts B and C in addition to Part A.

Who can serve as principal investigator (PI)?

The PI is the person who will personally conduct or supervise this research study. Under most circumstances, this will be a faculty member. For IRB communication purposes, a trainee/student may be listed as PI. However, a faculty advisor must be identified, who holds ultimate responsibility for ensuring that this project complies with all University, regulatory, and fiscal requirements.

→ *See next page for additional instructions*

---- **Instructions – Do not submit this page with your application** ----

Unless otherwise instructed, submit to the IRB that typically serves the home department of the principal investigator (PI). How many copies to submit depends on the IRB and type of review required for a given study (table). Complete submission instructions can be found at http://ohre.unc.edu/submit_instructions.php. All application and consent materials should be copied or printed on one side only.

Special submission instructions apply for studies that require additional review. Examples include the General Clinical Research Center (GCRC; <http://gcr.med.unc.edu/investigators/admin/gcraapp.htm>) or the Oncology Protocol Review Committee (PRC; <http://cancer.med.unc.edu/research/prc/default.asp>). See their web sites for details.

Number of Copies to be Submitted by Type of Review and IRB (number below must include one original)			
IRB	Exempt or Expedited	Full Board	Address for mailing complete application
Behavioral	1	16	CB# 3378, Bank of America Center Chapel Hill, NC 27599-3378
Biomedical (includes Dental)	3	3	CB# 7097, Medical Building 52 Chapel Hill, NC 27599-7097
Nursing	1	14	CB# 7460, Carrington Hall Chapel Hill, NC 27599-7460
Public Health	2	14	CB# 7400, Rosenau Hall Chapel Hill, NC 27599-7400

Types of Review

There are three levels of IRB Review (full board, expedited, and exempt), determined by the nature of the project, level of potential risk to human subjects, and the subject population. *The final determination of type of review applicable to a particular study is made by the IRB.* Regardless of the kind of review, all applications use the same submission form.

Exempt and expedited review can be given to studies that constitute no more than minimal risk to the human subjects, i.e., the risk one experiences in daily living. These reviews are done in the IRB office on a continual basis.

Full board review is required for studies that involve greater than minimal risk or vulnerable populations that require special protection by the IRB. These require review by the convened IRB at the next scheduled meeting. See http://ohre.unc.edu/guide_to_irb.php for additional guidance.

---- Instructions – Do not submit this page with your application ----

OFFICE OF HUMAN RESEARCH ETHICS
Institutional Review Board

APPLICATION FOR IRB APPROVAL OF
HUMAN SUBJECTS RESEARCH
Version 28-Sep-2005

<i>For IRB Use</i>				
Behav	Bio	Dent	Nurs	PH
IRB Study # _____				
Rec'd _____				
Full	Expedited	Exempt		

Part A.1. Contact Information, Agreements, and Signatures

Title of Study: The Effects of Hip Strength, Range of Motion, and Muscle Activity on a Knee Valgus Position During Functional Movement Tasks
Date: 2-1-05

Name and degrees of Principal Investigator: Brian Vesce BS, LAT, ATC, PES
Department: Exercise and Sports Science Mailing address/CB #: Sports Medicine Research Lab

Fetzer Gymnasium CB 8700
UNC-CH
Chapel Hill NC, 27514

UNC-CH PID: 711073070 Pager: N/A
Phone #: 412-855-3080 Fax #: 919-962-0489 Email Address: vesce@email.unc.edu

For trainee-led projects: ___ undergraduate X graduate ___ postdoc ___ resident ___ other

Name of faculty advisor: Dr. Darin Padua
Department: Exercise and Sports Science Mailing address/CB #: Sports Medicine Research Lab

Fetzer Gymnasium CB 8700
UNC-CH
Chapel Hill NC, 27514

Phone #: 919-843-5117 Fax #: 919-962-0489 Email Address: dpadua@email.unc.edu

Name, phone number, email address of project manager or coordinator, if any:

List **all other project personnel** including co-investigators, and anyone else who has contact with subjects or identifiable data from subjects: Lindsay Strickland, David Bell

Name of funding source or sponsor:

X not funded ___ Federal ___ State ___ industry ___ foundation ___ UNC-CH
___ other (specify):

Sponsor or award number:

Include following items with your submission, where applicable. Check the items below and **include in order listed**.

- ☒ This application. One copy must have original PI signatures.
- ☒ Consent and assent forms, fact or information sheets; include phone and verbal consent scripts
- ☐ HIPAA authorization addendum to consent form
- ☒ All recruitment materials including scripts, flyers and advertising, letters, emails
- ☐ Questionnaires, scripts used to guide phone or in-person interviews, etc.
- ☐ Focus group guides
- ☐ Data use agreements (may be required for use of existing data from third parties)
- ☐ Addendum for Multi-Site Studies where UNC-CH is the Lead Coordinating Center

- Documentation of reviews from any other committees (e.g., GCRC, Oncology)
- Documentation of training in human research ethics for all study personnel
- Investigator Brochure if a drug study
- Protocol, grant application or proposal supporting this submission; (e.g., extramural grant application to NIH or foundation, industry protocol, student proposal)

Principal Investigator: I will personally conduct or supervise this research study. I will ensure that this study is performed in compliance with all applicable laws, regulations and University policies regarding human subjects research. I will obtain IRB approval before making any changes or additions to the project. I will notify the IRB of any other changes in the information provided in this application. I will provide progress reports to the IRB at least annually, or as requested. I will report promptly to the IRB all unanticipated problems or serious adverse events involving risk to human subjects. I will follow the IRB approved consent process for all subjects. I will ensure that all collaborators, students and employees assisting in this research study are informed about these obligations. All information given in this form is accurate and complete.

Signature of Principal Investigator

Date

Faculty Advisor if PI is a Student or Trainee Investigator: I accept ultimate responsibility for ensuring that this study complies with all the obligations listed above for the PI.

Signature of Faculty Advisor

Date

Department or Division Chair, Center Director (or counterpart) of PI: (or Vice-Chair or Chair's designee if Chair is investigator or otherwise unable to review): I certify that this research is appropriate for this Principal Investigator, that the investigators are qualified to conduct the research, and that there are adequate resources (including financial, support and facilities) available. I support this application, and hereby submit it for further review.

Signature of Department Chair or designee

Date

Print Name of Department Chair or designee

Department

Part A.2. Summary Checklist

Are the following involved?

	Yes	No
A.2.1. Existing data, research records, patient records, and/or human biological specimens?	<u> </u>	<u>X</u>
A.2.2. Surveys, questionnaires, interviews, or focus groups with subjects?	<u>X</u>	<u> </u>
A.2.3. Videotaping, audiotaping, filming of subjects?	<u>X</u>	<u> </u>
A.2.4. Do you plan to enroll subjects from these vulnerable or select populations:		
a. UNC-CH students or UNC-CH staff?	<u>X</u>	<u> </u>
b. Non-English-speaking?	<u> </u>	<u>X</u>
c. Decisionally impaired?	<u> </u>	<u>X</u>
d. Patients?	<u> </u>	<u>X</u>
e. Prisoners, parolees and other convicted offenders?	<u> </u>	<u>X</u>
f. Pregnant women?	<u> </u>	<u>X</u>
g. Minors (less than 18 years)? If yes , give age range: to years	<u> </u>	<u>X</u>
A.2.5. a. Is this a multi-site study (i.e., involves organization(s) outside UNC-CH)?	<u> </u>	<u>X</u>
b. Will any of these sites be outside the United States?	<u> </u>	<u>X</u>
If yes , provide contact information for the foreign IRB.		
c. Is UNC-CH the sponsor or lead coordinating center?	<u> </u>	<u>X</u>
If yes , include the <u>Addendum for Multi-site Studies where UNC-CH is the Lead Coordinating Center.</u>		
A.2.6. Will there be a data and safety monitoring committee (DSMB or DSMC)?	<u> </u>	<u>X</u>
A.2.7. a. Are you collecting sensitive information such as sexual behavior, HIV status, recreational drug use, illegal behaviors, child/physical abuse, immigration status, etc?	<u> </u>	<u>X</u>
b. Do you plan to obtain a federal Certificate of Confidentiality for this study?	<u> </u>	<u>X</u>
A.2.8. a. Investigational drugs? (provide IND #)	<u> </u>	<u>X</u>
b. Approved drugs for “non-FDA-approved” conditions?	<u> </u>	<u>X</u>
<i>All studies testing substances in humans must provide a letter of acknowledgement from the <u>UNC Health Care Investigational Drug Service</u> (IDS).</i>		
A.2.9. Placebo(s)?	<u> </u>	<u>X</u>
A.2.10. Investigational devices, instruments, machines, software? (provide IDE #)	<u> </u>	<u>X</u>
A.2.11. Fetal tissue?	<u> </u>	<u>X</u>
A.2.12. Genetic studies on subjects’ specimens?	<u> </u>	<u>X</u>
A.2.13. Storage of subjects’ specimens for future research?	<u> </u>	<u>X</u>
If yes , see instructions within the form <u>Consent for Stored Samples.</u>		
A.2.14. Diagnostic or therapeutic ionizing radiation, or radioactive isotopes, which subjects would not receive otherwise?	<u> </u>	<u>X</u>
If yes , approval by the <u>UNC-CH Radiation Safety Committee</u> is required.		
A.2.15. Recombinant DNA or gene transfer to human subjects?	<u> </u>	<u>X</u>
If yes , approval by the <u>UNC-CH Institutional Biosafety Committee</u> is required.		
A.2.16. Does this study involve UNC-CH cancer patients?	<u> </u>	<u>X</u>
If yes , submit this application directly to the <u>Oncology Protocol Review Committee.</u>		
A.2.17. Will subjects be studied in the General Clinical Research Center (GCRC)?	<u> </u>	<u>X</u>
If yes , obtain the <u>GCRC Addendum</u> from the GCRC and submit complete application (IRB application and Addendum) to the GCRC.		

Part A.3. Conflict of Interest Questions and Certification

The following questions apply to **all investigators and study staff** engaged in the design, conduct, or reporting results of this project **and/or their immediate family members**. For these purposes, "family" includes the individual's spouse and dependent children. "Spouse" includes a person with whom one lives together in the same residence and with whom one shares responsibility for each other's welfare and shares financial obligations.

<p>A.3.1. Currently or during the term of this research study, does any member of the research team or his/her family member have or expect to have:</p> <p>(a) A personal financial interest in or personal financial relationship (including gifts of cash or in-kind) with the sponsor of this study?</p> <p>(b) A personal financial interest in or personal financial relationship (including gifts of cash or in-kind) with an entity that owns or has the right to commercialize a product, process or technology studied in this project?</p> <p>(c) A board membership of any kind or an executive position (paid or unpaid) with the sponsor of this study or with an entity that owns or has the right to commercialize a product, process or technology studied in this project?</p>	<p>___ yes</p> <p>___ yes</p> <p>___ yes</p>	<p><u>X</u> no</p> <p><u>X</u> no</p> <p><u>X</u> no</p>
<p>A.3.2. Has the University or has a University-related foundation received a cash or in-kind gift from the Sponsor of this study for the use or benefit of any member of the research team?</p>	<p>___ yes</p>	<p><u>X</u> no</p>
<p>A.3.3. Has the University or has a University-related foundation received a cash or in-kind gift for the use or benefit of any member of the research team from an entity that owns or has the right to commercialize a product, process or technology studied in this project?</p>	<p>___ yes</p>	<p><u>X</u> no</p>

If the answer to ANY of the questions above is yes, the affected research team member(s) must complete and submit to the Office of the University Counsel the form accessible at <http://coi.unc.edu>. List name(s) of all research team members for whom any answer to the questions above is yes:

Certification by Principal Investigator: By submitting this IRB application, I (the PI) certify that the information provided above is true and accurate regarding my own circumstances, that I have inquired of every UNC-Chapel Hill employee or trainee who will be engaged in the design, conduct or reporting of results of this project as to the questions set out above, and that I have instructed any such person who has answered “yes” to any of these questions to complete and submit for approval a Conflict of Interest Evaluation Form. I understand that as Principal Investigator I am obligated to ensure that any potential conflicts of interest that exist in relation to my study are reported as required by University policy.

Signature of Principal Investigator

Date _____

Faculty Advisor if PI is a Student or Trainee Investigator: I accept ultimate responsibility for ensuring that the PI complies with the University's conflict of interest policies and procedures.

Signature of Faculty Advisor

Date

Part A.4. Questions Common to All Studies

For all questions, if the study involves only secondary data analysis, focus on your proposed design, methods and procedures, and not those of the original study that produced the data you plan to use.

A.4.1. Brief Summary. Provide a *brief* non-technical description of the study, which will be used for internal and external communications regarding this research. Include purpose, methods, and participants. Typical summaries are 50-100 words.

Lower extremity postural alignment, such as knee valgus (e.g. knock knees), is commonly described as a potential risk factor for various lower extremity injuries (e.g. anterior cruciate ligament sprain, patello-femoral pain). Thus, an understanding of factors that influence knee valgus is believed to be an important aspect of understanding potential causes of lower extremity injuries. Muscle strength, activation, and flexibility are all believed to be contributing factors to knee valgus alignment. However, research has not investigated the influence of muscle strength, activation, and flexibility on knee valgus alignment. Therefore, the purpose of this study is to determine if there are significant differences in hip muscle strength, activation, and flexibility in individuals with knee valgus alignment compared to those with normal knee alignment. A secondary purpose of this study is to investigate the effects of a 6-week exercise program on knee valgus alignment, muscle strength, muscle activation, and flexibility. Sixty participants will be recruited from the general population at the University of North Carolina at Chapel Hill as part of this study.

A.4.2. Purpose and Rationale. Provide a summary of the background information, state the research question(s), and tell why the study is needed. If a complete rationale and literature review are in an accompanying grant application or other type of proposal, only provide a brief summary here. If there is no proposal, provide a more extensive rationale and literature review.

Posture is a very important and often neglected part of overall health. Ideal posture maintains that structural integrity and optimum alignment of each component of the kinetic chain (Clark, 2001). The kinetic chain consists of the myofascial system, articular system, and the neural system (Clark, 2001). When one component of this system is out of alignment, then the entire system is placed at a disadvantage. Postural malalignment is thought to create predictable patterns of tissue overload and dysfunction, initiating the cumulative injury cycle (Clark, 2001). This cumulative injury cycle begins with tissue trauma, inflammation, leading to muscle spasm, adhesions, altered neuromuscular control, and muscle imbalance. This cycle is thought to cause decreased athletic performance and eventual injury (Clark, 2001).

Knee valgus alignment during dynamic tasks (e.g. squatting) is a common postural dysfunction seen in the lower extremity. Knee valgus alignment during squatting is defined as the mid-patella moving medially and crossing over the ipsilateral great toe as the knee flexes while squatting downward. Knee valgus alignment has been described as a potentially dangerous movement pattern (Ireland, 1999). Knee valgus alignment is accompanied by movement of the hip into adduction and internal rotation, which are commonly described mechanisms for anterior cruciate ligament (ACL) injury (Ireland, 1999). It has also been shown that patello-femoral compressive forces increase with knee valgus alignment (Escamilla, 2001). Thus, knee valgus alignment is also believed to be a contributing factor to patello-femoral pain syndrome. Furthermore, the medial collateral ligament (MCL) is stressed when the knee is exposed to a valgus moment. The MCL can become injured when that valgus stress becomes too great. Individuals who undergo excessive knee valgus motion during functional tasks may place greater stress on the MCL and be at greater risk for injury. These injuries can be detrimental to an individual's physical well-being. Therefore, it is important to

understand the factors that influence knee valgus alignment as this may improve our understanding of lower extremity injury risk factor, which may lead to the development of exercise programs to correct knee valgus alignment and reduce injury risk. Currently, knee valgus alignment is thought to occur due to weakness or inhibition in the hip abductor muscles (e.g. gluteus medius, gluteus maximus) and hip external rotator muscles. Meanwhile, the hip adductor and internal rotator muscles are believed to be tight or overactive (Clark, 2001). However, research has not investigated if these hypothesized muscle imbalances actually exist in individuals demonstrating knee valgus alignment during a functional squatting task.

The purpose of this study is to ascertain what factors lead to knee valgus during squatting tasks. Once these factors are determined, a one-time intervention protocol will be administered to test whether or not these factors can be altered with an immediate intervention. This intervention will focus on lengthening tight hip musculature, strengthening weakened hip musculature, and performing functional lower extremity exercises intending to improve muscular activation. Furthermore, a six-week long intervention protocol will also be implemented to study how these factors can be changed over time.

Specific Aim 1: To compare hip muscle strength, hip muscle flexibility, and hip muscle activation between individuals with knee valgus alignment and those without knee valgus alignment (normal alignment)

Specific Aim 2: To compare knee valgus alignment, hip muscle strength, hip muscle flexibility, and hip muscle activation before and after a one-time supervised exercise program.

Specific Aim 3: To compare knee valgus alignment, hip muscle strength, hip muscle flexibility, and hip muscle activation before and after a six-week long exercise program.

A.4.3. Full description of the study design, methods and procedures. Describe the research study. Discuss the study design; study procedures; sequential description of what subjects will be asked to do; assignment of subjects to various arms of the study if applicable; doses; frequency and route of administration of medication and other medical treatment if applicable; how data are to be collected (questionnaire, interview, focus group or specific procedure such as physical examination, venipuncture, etc.). Include information on who will collect data, who will conduct procedures or measurements. Indicate the number and duration of contacts with each subject; outcome measurements; and follow-up procedures. If the study involves medical treatment, distinguish standard care procedures from those that are research. If the study is a clinical trial involving patients as subjects and use of placebo control is involved, provide justification for the use of placebo controls.

This study's design incorporates a single testing session for the normal knee alignment (NKA) control group and a repeated measures design for the dysfunction groups, the knee valgus intervention and knee valgus control (KVI and KVC). Volunteers from the university community will attend a general screening session in the Sports Medicine Research Laboratory. During this screening session, volunteers will be asked to perform a double-leg squat test and assessed by the researchers for the knee valgus alignment in real time. Forty volunteers who demonstrate a noticeable knee valgus position, as defined by demonstrating medial mid-patella movement crossing over the ipsilateral great toe bilaterally, during the final descent phase of the squat, with and without a 2-inch heel block under their heel, will be asked to volunteer for the study. From this group of subjects, twenty subjects will be randomly assigned to one of the dysfunction groups, either the knee valgus intervention (KVI)

group or knee valgus control (KVC) group. Twenty volunteers who do not demonstrate the knee valgus position will also be asked to volunteer for the study and be placed in the NKA control group.

Following the general screening session, all subjects will report to the Sports Medicine Research Laboratory for a single testing session. Upon arrival, all subjects will complete an informed consent form approved by the Institutional Review Board at the University of North Carolina at Chapel Hill along with a general health and activity level questionnaire. All subjects will be dressed in a t-shirt, athletic shorts and shoes. Subjects will be video taped performing the double leg squat task used in the initial screening. Subjects will then perform a five-minute warm-up on a stationary bicycle at a pace deemed comfortable to them. All testing will be performed on the subject's dominant leg, which will be defined as the leg used to kick a ball for maximal distance.

Range of Motion Assessment

After the warm-up, range of motion of the hip internal rotators and adductor complex will be recorded using a manual goniometer. For both range of motion measurements, the subjects will be in a supine position. In order to measure the hip internal rotator's range of motion, the subject's leg will be placed into ninety degrees of hip flexion and knee flexion. The axis of the goniometer will be placed over the inferior pole of the patella while the stationary arm will be aligned with the subject's trunk and the movement arm will be placed along the anterior midline of the tibia. The tester will passively move the hip into external rotation until resistance is felt by the tester. At this position, the hip internal rotators' range of motion will be recorded from the goniometer. In order to measure the hip adductor complex's range of motion, the leg will be in full knee extension and zero degrees of hip flexion/extension with the toe perpendicular to the floor. The axis of the goniometer will be placed over the dominant leg's ASIS with the stationary arm aligned with the opposite ASIS while the movement arm is placed along the anterior midline of the femur using the anterior midline of the patella for reference. The tester will passively abduct the leg until resistance is felt by the tester. At this position, hip adductor range of motion will be recorded from the goniometer.

Muscle Activation Assessment

Following muscle range of motion measurements, surface electromyography (EMG) will be used to measure the electrical activity of several hip muscles. Two surface electrodes will be placed over the muscle bellies of the gluteus maximus, gluteus medius, and the adductor complex. A single reference electrode will be placed over the tibial tuberosity to serve as an electrical ground reference. To reduce impedance to the EMG signal and allow for proper electrode fixation, electrode sites will be prepared by shaving any hair from the immediate vicinity of the muscle belly, lightly abrading the skin with an abrasive pad, and cleansing the skin with isopropyl alcohol. To prevent movement of the electrodes and subsequent alteration of the EMG signal, electrodes will be secured to the leg using prewrap and athletic tape. Manual muscle test will be performed to serve as a normalization of muscle activation. Three five-second maximal voluntary contraction trials will be performed during manual muscle testing on the isokinetic dynamometer. Signals from the electrodes will be passed to a battery operated FM transmitter worn by the subject, thus allowing for uninhibited movement. A receiver and analog-to-digital converted will convert the analog signal into digital data whereby it may be further analyzed by a computer utilizing custom software. These procedures for electromyography are similar to a study conducted by Michael DiStefano (IRB # 03-EXSS-648).

EMG data will be collected while subjects perform the identical squat task as was performed during the initial screening procedures. Following the double leg squat, instructions will then be given to each subject regarding the two jump landings and the researcher will demonstrate the maneuvers. Before any trials are recorded, each participant will have three to five trials to practice the jump landings. The practice trials will conclude when the subjects can demonstrate consistent movements with the jump landing. During the first jump landing, subjects will be required to jump forward off of a 30-centimeter box onto a force plate located on the ground. The distance between

the front of the box and the center of the force plate will be standardized for each subject, half their body height. Subjects will be instructed to land with their dominant foot in the center of the force plate and then jump as high as they can. This jump landing maneuver has been previously described in a study conducted by Lindsay Strickland (IRB # 04-EXSS-450). The second jump landing task is a transverse jump, instead of a forward jump involving similar procedures as the first jump landing task described. Instead of jumping forward onto the force plate, subjects will rotate ninety degrees in the air before landing. After landing, the subjects will then jump again in the air as high as they can as in the first jump described. The ninety-degree turn will occur in the air during the first part of the jump landing. A total of 5 trials will be performed with a 30 second rest period in between each trial. Trials will be repeated if there is a problem with the recording, if the subject does not land properly on the force plate, or if the subject does not perform the jump landing correctly. During all jump landing trials, EMG and force plate data will be recorded.

Strength Assessment

Strength measurements for the hip internal rotators, external rotators, abductors, and extensors will be recorded by an isokinetic dynamometer at 60 degrees per second (Strength testing procedures are adopted from previous study IRB# 03-EXSS-538). For each strength test, subjects will perform one trial with five repetitions.

For hip internal rotation and external rotation strength, subjects will be seated with the hip and knee flexed to ninety degrees and the thigh stabilized with the chair straps. For hip internal rotation strength, the dynamometer will be aligned along the long axis of the femur, pressure will be applied to the distal lateral fibula, and will be tested through twenty degrees of motion beginning at five degrees of internal rotation through fifteen degrees of external rotation. The hip external rotation strength will be tested in the same position however the pressure will be applied to the distal medial tibia. Motion will start at five degrees of external rotation and end at fifteen degrees of internal rotation. Also a towel will be placed between the subjects knees to reduce the risk of hip adduction.

Hip extension strength will be tested in a supported standing position with the chair at the subject's ASIS level. The subject's trunk will be stabilized to the chair by straps, and the dynamometer axis of rotation will be aligned with the anterior superior tip of the greater trochanter. Pressure will be applied to the distal posterior thigh and motion will start at ninety degrees of hip flexion and end at fifty degrees of hip flexion.

Hip abduction strength will be tested in a side lying position with the subject strapped in for stability. The dynamometer axis of rotation will be aligned .5 inches medial to the ASIS at the level of the greater trochanter. Pressure will be applied at the distal femur in the direction of adduction. Motion will be tested from zero to twenty degrees of hip abduction. MVIC's for the muscles having EMG activity recorded will be taken while the subject is in position after each individual isokinetic test. MVIC data will be collected for three seconds.

Intervention

The KVI group will perform 2 interventions. The first intervention will occur during the first testing session (acute intervention) and the second intervention will take place over a six-week period (long term intervention) both consisting of the same exercises. The intervention consists of and subjects will be asked to perform a series of two myofascial techniques and three static stretching exercises, which will take approximately 10 minutes to complete. Following completion of the myofascial release and stretching exercises the subjects will perform two strengthening exercises and two integration exercises, which will take approximately 10 minutes to complete. We will ask the participants to perform all exercises four times a week over the six-week long term intervention period. These specific exercises include: 1) myofascial technique to inhibit the hip adductors (1minute), 2) myofascial technique to inhibit the tensor fascia latae (1minute), 3) stretching exercise

for the hip adductors (2 repetitions for 30 seconds), 4) stretching exercise for the tensor fascia latae (2 repetitions for 30 seconds), 5) stretching exercise for the biceps femoris (2 repetitions for 30 seconds) 6) strengthening exercise for the gluteus medius (3 sets of 10 repetitions) 7) strengthening exercise for the gluteus maximus (3 sets of 10 repetitions) 8) lateral tube walking (3 sets of 10 repetitions) and 9) single leg reach for balance (3 sets of 3 repetitions). (Please see the included sheet entitled “Exercise Program Quick Reference Guide” for pictures of these exercises). The KVC group will be asked to sit and relax for 20 minutes before repeating the testing again (double leg squat, jump landing, transverse jump landing. Each dysfunction group, (KVI, KVC), will be asked to return 6 weeks after the initial testing session in order to be video taped during a squat task, -re-assess measures of strength, flexibility, and muscle activation during the three tasks. The NKA group will be finished after their initial testing session.

A.4.4. Benefits to subjects and/or society. Describe any potential for direct benefit to individual subjects, as well as the benefit to society based on scientific knowledge to be gained; these should be clearly distinguished. Consider the nature, magnitude, and likelihood of any direct benefit to subjects. If there is no direct benefit to the individual subject, say so here and in the consent form (if there is a consent form). Do not list monetary payment or other compensation as a benefit.

All participants will receive an intervention protocol aimed at correcting the movement dysfunction through lengthening, strengthening, and improving muscle activity in the hip musculature; however, this protocol is one of the aspects of the study being researched, therefore the potential exists that it will not benefit the subject whatsoever. In addition, all participants will also receive rehabilitation equipment necessary for the intervention program. The subjects in the KVI and the KNA group will receive these materials at the time of their first testing session. The KVC will receive the package when they have completed all of their testing. The sports medicine community will gain insight into what role flexibility, strength, and muscle activity play in contributing to knee valgus during a closed-chain knee flexion position. Also, if these causative factors can be corrected in a one-time intervention or through a six week intervention protocol, then correction of these factors could potentially lead to a reduction in knee injuries from athletics.

A.4.5. Full description of risks and measures to minimize risks. Include risk of psychosocial harm (e.g., emotional distress, embarrassment, breach of confidentiality), economic harm (e.g., loss of employment or insurability, loss of professional standing or reputation, loss of standing within the community) and legal jeopardy (e.g., disclosure of illegal activity or negligence), as well as known side effects of study medication, if applicable, and risk of pain and physical injury. Describe what will be done to minimize these risks. Describe procedures for follow-up, when necessary, such as when subjects are found to be in need of medical or psychological referral. If there is no direct interaction with subjects, and risk is limited to breach of confidentiality (e.g., for existing data), state this.

As with any physical activity, participation in this study carries a risk of bodily injury. The motions that subjects will be asked to perform are ones that repeatedly occur during physical activity. Therefore, subjects will be familiar and able to perform the tasks with minimal injury risk. To further minimize injury risk, participants will be allowed a warm up on a stationary bike to prepare themselves for testing. In case of injury, medical personnel (certified athletic trainers) will be located in the same building as testing, and ice will be available if needed. The subjects will be free to cease participation at any time.

There exists a risk of embarrassment or discomfort to the subject due to the location of the EMG electrode placement. This will be minimized by the fact that at no time will any subject be asked to

remove any article of clothing. All electrode placements can be easily reached without removing clothing and with exposing only a small amount of skin. To further minimize this risk, a researcher of the same sex as the subject will do the electrode placement. Also, a towel will be used to drape and further minimize the amount skin that is exposed.

There exists a small risk of muscle imbalances developing in subjects who are in the intervention group. Through stretching and strengthening certain muscle groups and not others there always exists a small chance of injury due to these imbalances. However, the exercises that will be given to subjects are aimed at correcting imbalances that may already exist, which should minimize any risk of injury associated with the intervention. Also, the potential for muscle strain injuries exist when stretching or strengthening, or bruising can occur from myofascial techniques. To minimize these risks, all subjects will receive the same instruction from certified athletic trainers who are trained in how to properly execute these exercises so that there is little risk for injury. Subjects will be given a handout describing how to properly perform exercises and at any time have the right to contact the researchers to answer any questions they may have about how to properly perform the exercises.

There exists a risk of minor pain being experienced by the subject from the shaving and abrading of the areas prior to electrode placement. Subjects will be informed of this risk in the consent form they read and sign; however, due to the necessity of these steps to ensure a clear EMG signal there are no steps to take to avoid this possibility of pain.

A.4.6. Data analysis. Tell how the qualitative and/or quantitative data will be analyzed. Explain how the sample size is sufficient to achieve the study aims. This might include a formal power calculation or explanation of why a small sample is sufficient (e.g., qualitative research, pilot studies).

The data of each dependent variable will be analyzed using two-way repeated measures analyses of variance comparing groups and testing sessions. Prior studies using similar techniques indicate for an estimated power of .80, 20 subjects will be needed for each group.

A.4.7. Will you collect or receive any of the following identifiers as part of the study data? Does not apply to consent forms.

☐ No ☒ Yes *If yes, check all that apply:*

- | | |
|--|---|
| a. <input checked="" type="checkbox"/> Names | d. <input type="checkbox"/> Any geographic subdivisions smaller than a State, including street address, city, county, precinct, zip code and their equivalent geocodes, except for the initial three digits of a zip code |
| b. <input checked="" type="checkbox"/> Telephone numbers | e. <input type="checkbox"/> Fax numbers |
| c. <input type="checkbox"/> Any elements of dates (other than year) for dates directly related to an individual, including birth date, admission date, discharge date, date of death. For ages over 89: all elements of dates (including year) indicative of such age, except that such ages and elements may be aggregated into a single category of age 90 and older | f. <input checked="" type="checkbox"/> Electronic mail addresses |
| | g. <input type="checkbox"/> Social security numbers |
| | h. <input type="checkbox"/> Medical record numbers |

- i. ___ Health plan beneficiary numbers
- j. ___ Account numbers
- k. ___ Certificate/license numbers
- l. ___ Vehicle identifiers and serial numbers (VIN), including license plate numbers
- m. ___ Device identifiers and serial numbers (e.g., implanted medical device)
- n. ___ Web universal resource locators (URLs)
- o. ___ Internet protocol (IP) address numbers
- p. ___ Biometric identifiers, including finger and voice prints
- q. ___ Full face photographic images and any comparable images
- r. ___ Any other unique identifying number, characteristic or code, other than dummy identifiers that are not derived from actual identifiers and for which the re-identification key is maintained by the health care provider and not disclosed to the researcher

A.4.8. Data sharing. With whom will *identifiable* (contains any of the 18 identifiers listed in question 7 above) data be shared outside the immediate research team? For each, explain confidentiality measures. Include data use agreements, if any.

- ☒ No one
- ☐ Coordinating Center:
- ☐ Statisticians:
- ☐ Consultants:
- ☐ Other researchers:
- ☐ Registries:
- ☐ Sponsors:
- ☐ External labs for additional testing:
- ☐ Journals:
- ☐ Publicly available dataset:
- ☐ Other:

A.4.9. Confidentiality of the data. Describe procedures for maintaining confidentiality of the data you will collect or will receive. Describe how you will protect the data from access by those not authorized. How will data be transmitted among research personnel? Where relevant, discuss the potential for deductive disclosure (i.e., directly identifying subjects from a combination of indirect IDs). Describe your plan to destroy identifiers. When will identifiers be destroyed?

No subjects will be identified in any report or publication about this study. All subjects will be assigned an identification number (ID) for data collection. This ID number will be matched to the identifiers listed above in an excel document. This will be the only place in which a subjects identifiers and ID number will co-exist. This document will be stored on a separate cd apart from all other data that will be collected. These identifiers will be collected during the screening session for the sole purpose of contacting subjects to schedule subsequent testing sessions. Once a subject has completed all of his/her testing sessions, then their identifiers will be deleted from the excel document until all subjects have been tested and the document is destroyed. All information on the data collection form used for testing will be referenced with the subject ID number. At no time will the identifiers above be listed on the same document as data collected during testing. All data will be stored on cds which will be kept in the Sports Medicine Research Lab. All videotapes will be stored in a secure area of the Sports Medicine Research Lab. All data analysis will be performed on computers in the Sports Medicine Research Lab where a password is necessary for access to the computers. Only members performing research have access to these computers, therefore identification of any subjects or data is very unlikely. If disclosure is ever required, UNC-CH will take all steps allowable by law to protect the privacy of personal information.

Personal privacy during testing sessions will be maintained through limiting the people within the research lab to current employees of the lab and the testers themselves. The only door to enter the lab is locked with key card access to ensure privacy. Patients will be properly draped with a towel during electrode placement to ensure privacy.

A.4.10. Data security for storage and transmission. Please check all that apply.

For electronic data:

- ☐ Secure network ☐ Password access ☐ Encryption
☐ Other (describe):
☐ Portable storage (e.g., laptop computer, flash drive)
Describe how data will be protected for any portable device:

For hardcopy data (including human biological specimens, CDs, tapes, etc.):

- ☐ Data de-identified by research team (stripped of the 18 identifiers listed in question 7 above)
☒ Locked suite or office
☒ Locked cabinet
☒ Data coded by research team with a master list secured and kept separately
☐ Other (describe):

Part A.5. The Consent Process and Consent Documentation (including Waivers)

The standard consent process is for all subjects to sign a document containing all the elements of informed consent, as specified in the federal regulations. Some or all of the elements of consent, including signatures, may be altered or waived under certain circumstances.

- If you will obtain consent in any manner, complete **section A.5.1**.
- If you are obtaining consent, but requesting a waiver of the requirement for a signed consent document, complete **section A.5.2**.
- If you are requesting a waiver of any or all of the elements of consent, complete **section A.5.3**.

You may need to complete more than one section. For example, if you are conducting a phone survey with verbal consent, complete sections A.5.1, A.5.2, and possibly A.5.3.

A.5.1. Describe the process of obtaining informed consent from subjects. If children will be enrolled as subjects, describe the provisions for obtaining parental permission and assent of the child. If decisionally impaired adults are to be enrolled, describe the provision for obtaining surrogate consent from a legally authorized representative (LAR). If non-English speaking people will be enrolled, explain how consent in the native language will be obtained. Address both written translation of the consent and the availability of oral interpretation. *After you have completed this part A.5.1, if you are not requesting a waiver of any type, you are done with Part A.5.; proceed to Part B.*

Informed consent will be obtained from all subjects prior to any testing. Subjects will report to the University of North Carolina-Chapel Hill, Sports Medicine Research Lab in Fetzer Gym. Before beginning the testing session the subject will be asked to read an informed consent agreement outlining the procedures, protocols and potential risks of the study. This informed consent agreement form will be in accordance with the standards set forth by the Medical IRB at the University of North Carolina at Chapel Hill. After the subjects sign the consent form, a copy will be given to them, and testing will commence.

A.5.2. Justification for a waiver of *written* (i.e., signed) consent. *The default is for subjects to sign a written document that contains all the elements of informed consent.* Under limited circumstances, the requirement for a signed consent form may be waived by the IRB if either of the following is true:

- a. The only record linking the subject and the research would be the consent document and the principal risk would be potential harm resulting from a breach of confidentiality (e.g., study involves sensitive data that could be damaging if disclosed). ___ yes ___ no

Explain.

- b. The research presents no more than minimal risk of harm to subjects and involves no procedures for which written consent is normally required outside of the research context (e.g., phone survey). ___ yes ___ no

Explain.

If you checked “yes” to either, will consent be oral? Will you give out a fact sheet? Use an online consent form, or include information as part of the survey itself, etc?

A.5.3. Justification for a full or partial waiver of consent. *The default is for subjects to sign a written document that contains all the elements of informed consent.* A waiver might be requested for research involving only existing data or human biological specimens (see also Part C). More rarely, it might be requested when the research design requires withholding some study details at the outset (e.g., behavioral research involving deception). In limited circumstances, parental permission may be waived. This section should also be completed for a waiver of HIPAA authorization if research involves Protected Health Information (PHI) subject to HIPAA regulation, such as patient records.

___ Requesting **waiver of some elements** (specify; see SOP 28 on the IRB web site):

___ Requesting **waiver of consent entirely**

If you check either of the boxes above, answer items a-f.. To justify a full waiver of the requirement for informed consent, you must be able to answer “yes” (or “not applicable” for question c) to items a-f. **Insert brief explanations that support your answers.**

- a. Will the research involve no greater than minimal risk to subjects or to their privacy? ___ yes ___ no

Explain.

- b. Is it true that the waiver will *not* adversely affect the rights and welfare of subjects? *(Consider the right of privacy and possible risk of breach of confidentiality in light of the information you wish to gather.)* ___ yes ___ no

Explain.

- c. When applicable to your study, do you have plans to provide subjects with pertinent information after their participation is over? *(e.g., Will you provide* ___ yes ___ not applicable

details withheld during consent, or tell subjects if you found information with direct clinical relevance? This may be an uncommon scenario.)

Explain.

d. Would the research be impracticable without the waiver? *(If you checked “yes,” explain how the requirement to obtain consent would make the research impracticable, e.g., are most of the subjects lost to follow-up or deceased?).* ☐ yes ☐ no

Explain.

e. Is the risk to privacy reasonable in relation to benefits to be gained or the importance of the knowledge to be gained? ☐ yes ☐ no

Explain.

If you are accessing patient records for this research, you must also be able to answer “yes” to item f to justify a waiver of HIPAA authorization from the subjects.

f. Would the research be impracticable if you could not record (or use) Protected Health Information (PHI)? *(If you checked “yes,” explain how not recording or using PHI would make the research impracticable).* ☐ yes ☐ no

Explain.

Part B. Questions for Studies that Involve Direct Interaction with Human Subjects

→ *If this does not apply to your study, do not submit this section.*

B.1. Subjects. Specify number, gender, ethnicity, race, and age. Specify whether subjects are healthy volunteers or patients. If patients, specify any relevant disease or condition and indicate how potential subjects will be identified.

Subjects will be divided into two groups based upon their results on the double leg squat task (DLST). The dysfunction group will consist of subjects who demonstrate bilateral knee adduction during the DLST both with and without a two-inch heel block place under the calcaneus. The heel block will be used to ensure that tightness in the posterior lower leg is not influencing a knee valgus alignment. The dysfunction group will be further divided, randomly, into an intervention group (KVI) and a control group (KVC). The normal alignment control group (NKA) will consist of subjects who do not demonstrate knee adduction on either leg during the DLST with and without a two-inch heel block. Subjects in the control group will be matched to subjects in the hip dysfunction groups by age, height, weight, gender and physical activity level. Each group will consist of 20 subjects whose ages range from 18-35 years. Subjects will be otherwise healthy volunteers that have no current musculoskeletal injuries or have not sustained an injury to either lower extremity in the past six months. No potential subject will be excluded due to gender, ethnicity, or race.

B.2. Inclusion/exclusion criteria. List required characteristics of potential subjects, and those that preclude enrollment. Justify exclusion of any group, especially by criteria based on gender, ethnicity, race, or age. If pregnant women are excluded, or if women who become pregnant are withdrawn, specific justification must be provided.

Inclusion Criteria for Dysfunction Groups: Aysmptomatic with & without activity and during the double leg squat; Demonstrates bilateral knee adduction during double leg squat (no heel block); Demonstrates bilateral knee adduction during double leg squat with 2-inch heel block placed underneath calcaneus.

Inclusion Criteria for Normal Alignment Group: Aysmptomatic with & without activity and during the double leg squat; Does not demonstrate bilateral knee adduction during double leg squat with and without a two-inch heel block.

No potential subject will be excluded due to gender, ethnicity, or race. Pregnant women will be excluded and women who become pregnant will be withdrawn due to the complexity and activity necessary for the functional tasks.

B.3. Methods of recruiting. Describe how and where subjects will be identified and recruited. Indicate who will do the recruiting, and tell how subjects will be contacted. Describe efforts to ensure equal access to participation among women and minorities. Describe how you will protect the privacy of potential subjects during recruitment. *For prospective subjects whose status (e.g., as patient or client), condition, or contact information is not publicly available (e.g., from a phone book or public web site), the initial contact should be made with legitimate knowledge of the subjects' circumstances. Ideally, the individual with such knowledge should seek prospective subjects' permission to release names to the PI for recruitment. Alternatively, the knowledgeable individual could provide information about the study, including contact information for the investigator, so that interested prospective subjects can contact the investigator.* Provide the IRB with a copy of any document or script that will be used to obtain the patients' permission for release of names or to introduce the study. Check with your IRB for further guidance.

All potential subjects will be recruited through informational emails and flyers written by the principal investigator. The contact number in the email and the flyer will be that of the investigator, Brian Vesci. The potential subjects will have the freedom to either respond or ignore the emails; therefore, they will not feel obligated to participate in the study.

Emails will be sent to the current student and faculty population of the University of North Carolina at Chapel Hill via listservs. Efforts to ensure equal access to participation among women and minorities will be limited only by the current population of women and minorities in the student body and faculty population.

B.4. Protected Health Information (PHI). If you need to access Protected Health Information (PHI) to identify potential subjects who will then be contacted, you will need a *limited waiver of HIPAA authorization*. If this applies to your study, please provide the following information.

- a. Will the information collected be limited only to that necessary to contact the subjects to ask if they are interested in participating in the study?
- b. How will confidentiality/privacy be protected prior to ascertaining desire to participate?
- c. When and how will you destroy the contact information if an individual declines participation?

B.5. Duration of entire study and duration of an individual subject's participation, including follow-up evaluation if applicable. Include the number of required contacts and approximate duration of each contact.

All subjects will have an initial contact during the screening process. This will last approximately five minutes per subject.

Subjects in the normal alignment group (NKA) will only require one more contact for the testing session. This will last approximately one hour.

Subjects in the dysfunction groups (KVI, KVC) will require two further contacts beyond the initial screening. Both of these testing sessions will last approximately one and a half hours. The first of these testing sessions will occur following the general screening session and the final testing session will occur six weeks later.

B.6. Where will the subjects be studied? Describe locations where subjects will be studied, both on and off the UNC-CH campus.

All subjects will be studied in the Sports Medicine Research Lab in the Fetzer Gymnasium building on the campus of the University of North Carolina at Chapel Hill

B.7. Privacy. Describe procedures that will ensure privacy of the subjects in this study. Examples include the setting for interviews, phone conversations, or physical examinations; communication methods or mailed materials (e.g., mailings should not indicate disease status or focus of study on the envelope).

No subjects will be identified in any report or publication about this study. All subjects will be assigned an identification number (ID) for data collection. All data will be stored on cds which will be kept in the Sports Medicine Research Lab. All videotapes will be stored in a secure area of the Sports Medicine Research Lab. All data analysis will be performed on computers in the Sports Medicine Research Lab where a password is necessary for access to the computers. Only members performing research have access to these computers, therefore identification of any subjects or data is very unlikely. If disclosure is ever required, UNC-CH will take all steps allowable by law to protect the privacy of personal information.

Personal privacy during testing sessions will be maintained through limiting the people within the research lab to current employees of the lab and the testers themselves. The only door to enter the lab is locked with key card access to ensure privacy. Patients will be properly draped with a towel during electrode placement to ensure privacy.

B.8. Inducements for participation. Describe all inducements to participate, monetary or non-monetary. If monetary, specify the amount and schedule for payments and how this will be prorated if the subject withdraws (or is withdrawn) from the study prior to completing it. For compensation in foreign currency, provide a US\$ equivalent. Provide evidence that the amount is not coercive (e.g., describe purchasing power for foreign countries). Include food or refreshments that may be provided.

All participants in this study will receive an exercise package valued at approximately thirty to forty dollars per participant following completion of the study. This package will include exercise tubing, foam rollers, and the exercise protocol.

B.9. Costs to be borne by subjects. Include child care, travel, parking, clinic fees, diagnostic and laboratory studies, drugs, devices, all professional fees, etc. If there are no costs to subjects other than their time to participate, indicate this.

There will be no cost borne by subjects other than their time.

Part C. Questions for Studies using Data, Records or Human Biological Specimens without Direct Contact with Subjects

→ *If this does not apply to your study, do not submit this section.*

C.1. What records, data or human biological specimens will you be using? (check all that apply):

- ☐ Data already collected for another research study
- ☐ Data already collected for administrative purposes (e.g., Medicare data, hospital discharge data)
- ☐ Medical records (custodian may also require form, e.g., HD-974 if UNC-Health Care System)
- ☐ Electronic information from clinical database (custodian may also require form)
- ☐ Patient specimens (tissues, blood, serum, surgical discards, etc.)
- ☐ Other (specify):

C.2. For each of the boxes checked in 1, how were the original data, records, or human biological specimens collected? Describe the process of data collection including consent, if applicable.

C.3. For each of the boxes checked in 1, where do these data, records or human biological specimens currently reside?

C.4. For each of the boxes checked in 1, from whom do you have permission to use the data, records or human biological specimens? Include data use agreements, if required by the custodian of data that are not publicly available.

C.5. If the research involves human biological specimens, has the purpose for which they were collected been met before removal of any excess? For example, has the pathologist in charge or the

clinical laboratory director certified that the original clinical purpose has been satisfied? Explain if necessary.

☐ yes ☐ no ☐ not applicable (explain)

C.6. Do all of these data records or specimens exist at the time of this application? If not, explain how prospective data collection will occur.

☐ yes ☐ no If no, explain

University of North Carolina-Chapel Hill
Consent to Participate in a Research Study
Adult Subjects
Biomedical Form

IRB Study # 05-EXSS-839

Consent Form Version Date: 3/2/06

Title of Study: The Effects of Hip Strength, Range of Motion, and Muscle Activity on a Knee Valgus Position During Functional Movement Tasks.

Principal Investigator: Brian Vesci, BS, ATC, LAT, PES

UNC-Chapel Hill Department: Department of Exercise and Sport Science

UNC-Chapel Hill Phone number: (919) 962-7187

Email Address: vesci@email.unc.edu

Co-Investigators: David R. Bell, MEd, ATC, LAT; Lindsay Strickland, MS, ATC, LAT

Faculty Advisor: Darin Padua, PhD, ATC, LAT

Funding Source:

Study Contact telephone number: 919-962-7187

Study Contact email: vesci@email.unc.edu

What are some general things you should know about research studies?

You are being asked to take part in a research study. To join the study is voluntary.

You may refuse to join, or you may withdraw your consent to be in the study, for any reason.

Research studies are designed to obtain new knowledge that may help other people in the future. You may not receive any direct benefit from being in the research study. There also may be risks to being in research studies.

Deciding not to be in the study or leaving the study before it is done will not affect your relationship with the researcher, your health care provider, or the University of North Carolina-Chapel Hill.

Details about this study are discussed below. It is important that you understand this information so that you can make an informed choice about being in this research study. You will be given a copy of this consent form. You should ask the researchers named above, or staff members who may assist them, any questions you have about this study at any time.

What is the purpose of this study?

The position of knee valgus (a.k.a. knock knee), which occurs as the knees move inward towards the great toe while moving into a squatted position, has been theorized as a position that may predispose an individual for various knee and lower extremity injuries. Limited research has been performed to investigate potential causes of knee valgus. Also, there is no research investigating exercises to correct knee valgus alignment. Therefore, the primary purpose of this study is to investigate the differences in muscle strength, muscle flexibility, and muscle activation between subjects with and without knee valgus alignment during squatting. A secondary purpose of this study is to examine the effectiveness of an exercise intervention on changing knee valgus alignment, muscle strength, muscle flexibility, and muscle activation.

Are there any reasons you should not be in this study?

You should not participate in this study if any of the following apply to you:

- You are not interested in doing strength or flexibility exercises
- You have any current symptoms of injury (redness, swelling, pain)
- You have used an external ankle brace/support on a regular basis within the past three months
- You have had leg surgery in the past year
- You have had a lower extremity injury in the past three months that required you to miss three or more consecutive days of physical activity

How many people will take part in this study?

If you decide to be in this study, you will be one of approximately 60 participants in this research study.

How long will your part in this study last?

Your participation in this study will depend upon your group assignment. If you are in the group identified with normal knee alignment (NKA), you will only take part in one testing session that will last approximately one hour. If you are in the groups identified with a knee valgus alignment your involvement will consist of two testing sessions. The first testing session will last approximately one and a half hours. The second testing session will take place six weeks after the first session and will last approximately one and a half hours.

What will happen if you take part in the study?

Subjects in the knee valgus intervention group (KVI) and the normal knee alignment group (NKA) will receive a foam roller, exercise band, and exercise program aimed at improving their knee valgus alignment at the time of their first testing session. The other group, knee valgus control (KVC), will receive the same exercise package at the completion of their testing. It is important to understand that as a subject you will receive the exercise package however you may receive no benefit from the proposed exercises as they are part of what is being analyzed in the study and their ability to improve knee valgus alignment has not yet been demonstrated.

Subjects will be asked to attend the testing sessions wearing a t-shirt, shorts and athletic shoes. Subjects will be divided into three groups. One group will consist of those who have been identified with normal knee alignment (NKA). The two additional groups will be randomly assigned and are those participants that have been identified as having knee valgus during a squat. One of these groups will be another control group, the knee valgus control (KVC), and the other will receive the exercise program, the knee valgus intervention (KVI).

Testing Session 1 (NKA, KVC, KVI):

The session will begin with you being video taped during a squat task to confirm and document whether or not you demonstrate knee valgus. You will then ride a stationary bicycle for five minutes to warm-up. After you are sufficiently warmed up, we will measure the range of motion of your hip. Adhesive electrodes designed to measure electrical activity of your muscles will be placed on the outside and back (high buttocks/low back) of your hip as well as approximately half way between your hip and your knee on the inside of your thigh. A small area will be shaven and cleansed with alcohol to ensure that electrodes will be secured in place. An examiner of the same sex as yourself will perform all electrode placements. At no time will you be asked to remove any articles of clothing. The electrodes can be placed properly with minimal skin exposure and a towel will be used for draping in an attempt to remove any feelings of discomfort. Examiners will then measure your hip muscle strength on an isokinetic-strength testing machine, meaning that your maximal strength will be tested at a predetermined and constant speed.

You will then perform three different activities during which the investigators will measure your muscle activity. They consist of 5 squats, 5 straight ahead jump landings, and 5 jump landings in which the subject will rotate 90 degrees in the air, from an approximately 12-inch high box to a distance half of your body height away.

At this time the NKA group will be finished with testing and will receive the exercises package. The KVI group will be asked to perform a series of exercises that will take approximately 20 minutes to complete. The exercises include five stretching exercises, two strengthening exercises, and two functional exercises. These specific exercises include:

- 1) Foam roller stretch to the inside of the thigh (1 minute)
- 2) Foam roller stretch to the outside of the thigh (1 minute)
- 3) Stretching exercise for the inside of the thigh (2 repetitions for 30 seconds)
- 4) Stretching exercise for the outside of the thigh (2 repetitions for 30 seconds)
- 5) Stretching exercise for the back of the thigh (2 repetitions for 30 seconds)

- 6) Strengthening exercise for the outside of the thigh (3 sets of 10 repetitions)
- 7) Strengthening exercise for the back of the thigh (3 sets of 10 repetitions)
- 8) Side stepping with resistance band around ankles (3 sets of 10 repetitions)
- 9) Single leg squat reach for balance (3 sets for 3 repetitions)

The KVC group will rest for 20 minutes instead of performing exercises. After the rest or exercise period each subject will perform the squats, jump landing, and transverse jump landing for a second time. Subjects will not have to repeat the strength testing after the rest or exercises. Finally, the exercise group (KVI) will be asked to perform the exercise program at home, four times a week over a six-week period. Each subject in the exercise group will be given a handout with pictures depicting each exercise, as well as a daily exercise log to chart when and which exercises were performed.

Testing Session 2 (KVC, KVI):

Both groups identified with a knee valgus alignment will return to the Sports Medicine Research Laboratory after 6-weeks. The second testing session will be very similar to the first. All groups will: 1) warm-up on the stationary bike, 2) have the electrodes placed again, and 3) have their strength test again exactly as the first session. Finally, each group will perform 5 squats, 5 jump landings (half the distance of their height), and 5 transverse landings (half the distance of their height). No exercise intervention will take place in the second session. At this point after the second testing session the KVC group will be given the exercise package and will have the experimental exercise protocol explained to them.

What are the possible benefits from being in this study?

Research is designed to benefit society by gaining new knowledge about knee valgus. The benefits to you from being in this study may include a prescriptive exercise program designed to attempt to correct hip muscle weakness and tightness that may exist. The researcher will discuss your performance on the jumping and squatting maneuvers with you if you wish. This information may help educate you on improving your jumping and squatting technique. The results of this study may benefit society by providing data on what factors may influence a knee valgus position during functional movements that may predispose someone to various knee injuries as well as if any intervention exercise programs can be implemented to correct these factors.

What are the possible risks or discomforts involved with being in this study?

This study involves a jumping and squatting maneuver that may involve the following risks and/or discomforts to you:

- Possibility of a ligament injury to the joints of your lower extremities
- Possibility of muscle strains/pulls/soreness in your lower extremities
- Possibility of pain due to shaving and abrading sites of electrode placement
- Possible embarrassment due to electrode placement
- There may be uncommon or previously unrecognized risks that might occur

While the possibility of the above mentioned risks exist, all appropriate precautions will be taken to minimize these risks.

No penalty will be incurred if you decide to not participate or if you withdraw yourself from testing during the study. Please do not feel pressured to participate, or continue with the study if at any point you feel uncomfortable.

What if we learn about new findings or information during the study?

You will be given any new information gained during the course of the study that might affect your willingness to continue your participation.

How will your privacy be protected?

No subjects will be identified in any report or publication about this study. All subjects will be identified as a number throughout data collection. All data storage and analysis will be on computers the sports medicine research lab where a password is necessary for access to the computers. Only members performing research have access to the lab and use of its computers. Although every effort will be made to keep research records private, there may be times when federal or state law requires the disclosure of such records, including personal information. This is very unlikely, but if disclosure is ever required, The University of North Carolina at Chapel Hill will take all steps allowable by law to protect the privacy of personal information.

What will happen if you are injured by this research?

All research involves a chance that something bad might happen to you. This may include the risk of personal injury. In spite of all safety measures, you might develop a reaction or injury from being in this study. If such problems occur, the researchers will help you get medical care, but any costs for the medical care will be billed to you and/or your insurance company. The University of North Carolina at Chapel Hill has not set aside funds to pay you for any such reactions or injuries, or for the related medical care. However, by signing this form, you do not give up any of your legal rights.

What if you want to stop before your part in the study is complete?

You can withdraw from this study at any time, without penalty. The investigators also have the right to stop your participation at any time. This could be because you have had an unexpected reaction, or have failed to follow instructions, or because the entire study has been stopped.

Will you receive anything for being in this study?

You will be receiving an exercise package valued at approximately thirty to forty dollars consisting of exercise and rehabilitation equipment (foam rollers, exercise tubing), as well as the exercise protocol, following completion of the study. If you are in exercise group (KVI) or the normal knee alignment group (NKA) you will receive the exercise equipment and the exercise protocol at the time of your first testing session. If you are not in this group, you will receive the exercise package, as well as the exercise intervention, when you have completed all testing associated with the study.

Will it cost you anything to be in this study?

The only cost to you will be in your time and transportation to the University of North Carolina at Chapel Hill Sports Medicine Research Laboratory for your testing session.

What if you are a UNC student?

You may choose not to be in the study or to stop being in the study before it is over at any time. This will not affect your class standing or grades at UNC-Chapel Hill. You will not be offered or receive any special consideration if you take part in this research.

What if you are a UNC employee?

Taking part in this research is not a part of your University duties, and refusing will not affect your job. You will not be offered or receive any special job-related consideration if you take part in this research.

Who is sponsoring this study?

The National Academy of Sports Medicine is donating the equipment provided to the subjects for this study. However they are not sponsoring the study. Researchers for this study are not receiving any type of compensation for their work.

What if you have questions about this study?

You have the right to ask, and have answered, any questions you may have about this research. If you have questions, or if a research-related injury occurs, you should contact the researchers listed on the first page of this form.

What if you have questions about your rights as a research subject?

All research on human volunteers is reviewed by a committee that works to protect your rights and welfare. If you have questions or concerns about your rights as a research subject you may contact, anonymously if you wish, the Institutional Review Board at 919-966-3113 or by email to IRB_subjects@unc.edu.

IRB Study # 05-EXSS-839

Subject's Agreement:

I have read the information provided above. I have asked all the questions I have at this time.
I voluntarily agree to participate in this research study.

Signature of Research Subject

Date

Printed Name of Research Subject

Signature of Person Obtaining Consent

Date

Printed Name of Person Obtaining Consent

University of North Carolina – Chapel Hill
Research Study Questionnaire
Adult Subjects

Medical IRB Study # 05-EXSS-839

Title of Study: The Effects of Hip Strength, Range of Motion, and Muscle Activity on a Knee Valgus Position During Functional Movement Tasks.

Principal Investigator: Brian Vesci, BS, ATC, LAT, PES

UNC-CH Department: EXSS

Phone Number: 919-962-7187

Co-Investigators: David Bell, Med, ATC, LAT; Lindsay Strickland, MS, ATC, LAT

Sponsor: None

1. Are you currently in good general health?

YES / NO

2. Have you had a lower extremity injury in the past three months that required three consecutive days missed from physical activity?

YES / NO

3. Do you have any current symptoms of injury?

YES / NO

4. How often do you exercise per week? _____ Days

5. Approximately how many minutes do you exercise per day? _____ Minutes

6. What type of exercise activity do you most often participate in (walking, running, aerobics, basketball, etc.)?

Email to potential subjects

Subject:

Are you interested in contributing to research aimed at preventing lower extremity knee injuries?

Script:

You may be eligible to participate in a research study investigating lower body posture during squatting and jumping. This study is sponsored by the UNC Department of Exercise and Sport Science.

If chosen to participate in this study you will be asked to attend either one (control subjects) or two (non-control subjects) testing sessions. The control subjects session will last approximately one hour. Sessions for the non control subjects will last approximately one and a half hours. During testing sessions you will be video taped while performing a squatting task, have strength measurements taken of various hip muscles, EMG muscle activity recorded for various hip muscles (including electrode placement on the buttocks and halfway down the inside of your thigh), and may asked to participate in an exercise program for six weeks at which point you will be tested again. If you qualify for the control group you will not have to participate in any exercise program.

Criteria for inclusion in the control group consists of having no pain while performing a squat and having what is judged by the tester as normal knee alignment while squatting. Inclusion criteria for the intervention group consists of having no pain while squatting and having what the tester judges to be knee valgus (knock knees) alignment while squatting.

Volunteers will be asked to attend a general screening session to determine if they are eligible. To volunteer you must be healthy, free from lower extremity injury, and between the ages of 18-35.

If selected to participate as a subject in this study you will receive an exercise package including equipment and exercises aimed at improving your lower body posture

All interested individuals should contact the principal investigator, Brian Vesci, at vesci@email.unc.edu

ARE YOU INTERESTED IN CONTRIBUTING TO RESEARCH AIMED AT PREVENTING KNEE INJURIES?

You may be eligible to participate in a research study investigating lower body posture during squatting and jumping. This study is being conducted by the UNC Department of Exercise and Sport Science

If chosen to participate in this study you will be video taped while performing a squatting task, have strength measurements taken of various hip muscles, EMG muscle activity recorded for various hip muscles (including electrode placement on the buttocks and halfway down the inside of your thigh), Control subjects, being judged to have normal knee alignment, will have one testing session lasting approximately one hour. Intervention subjects will be asked to participate in a six week exercise program and will be tested twice, both sessions will last approximately one and a half hours.

Volunteers will be asked to attend a general screening session to determine if they are eligible. To volunteer you must be healthy, free from lower extremity injury, have no pain when performing a squat, and between the ages of 18-35.

If selected to participate as a subject in this study you will receive an exercise package including equipment and exercises aimed at improving your lower body posture

To receive more information, contact:

Brian Vesci LAT, ATC, PES
vesci@email.unc.edu
412-855-3080

UNC Sports Medicine Research Lab
919-962-7187

APPENDIX E: DATA COLLECTION MATERIALS

Data Collection Form

Subject ID _____

Group _____
Height _____ cm

Leg L R
Weight _____ kg

Sex M F
Age _____

Range of Motion

Passive Hip Abduction 1) _____ 2) _____ 3) _____ Avg(SPSS) _____

Passive Hip ER 1) _____ 2) _____ 3) _____ Avg(SPSS) _____

Electrode Placement

Iliac crest to greater trochanter _____ / 2 = _____

S2 – 10 cm distal to greater trochanter _____ x .2 = _____

Greater trochanter to lateral epicondyle _____ / 2 = _____

Isokinetic Strength (Average Torque)

Hip ER Arm Level _____

Hip IR Arm level _____

Hip Extension Arm level _____

Hip Abduction Arm level _____

MVIC Testing Order: Gmax _____ Gmed _____ Add _____

Activity Testing Order: DLSQ: _____ JL: _____ TJP: _____

File information: Sub#: _____ Session: 1 or 2 _____ Pre or Post _____

Please write in all final concatenate files saved for this subject:

1) MVIC File:

2) DLSQ File:

3) Jump Landing File (JL):

4) Transverse Landing File (TJP):

APPENDIX F: RAW DATA

Subject	Group	Height	Weight	Age	Leg	Sex	HipAB1	HipAB2	HipAB3	HipABAVG
1	1	175	98	28	2	1	45	37	35	39
2	1	173	99	25	1	1	40	35	41	38.666667
3	1	173	66	23	2	1	39	32	36	35.666667
4	1	193	79	23	1	1	32	34	35	33.666667
5	1	175	59	24	1	2	45	42	42	43
6	1	163	54	22	1	2	42	37	38	39
7	1	168	61	18	2	2	27	24	22	24.333333
8	1	170	64	18	1	2	38	37	35	36.666667
9	1	173	54	23	1	2	42	36	41	39.666667
10	1	170	77	23	2	2	28	32	35	31.666667
11	1	175	79	24	1	1	34	37	36	35.666667
12	1	170	64	22	1	2	41	45	38	41.333333
13	1	48	43	44	1	2	48	43	44	45
14	1	183	86	25	1	1	36	34	30	33.333333
15	1	170	59	21	1	2	45	47	43	45
16	1	170	66	22	1	2	31	33	36	33.333333
17	1	175	75	20	1	1	32	29	35	32
18	2	163	61	21	2	2	33	31	32	32
19	2	168	61	20	1	2	26	40	35	33.666667
20	2	163	61	20	1	2	34	43	41	39.333333
21	2	165	54	25	1	2	45	45	44	44.666667
22	2	165	64	26	1	2	41	39	38	39.333333
23	2	157	61	22	1	2	35	30	32	32.333333
24	2	157	61	25	1	2	32	30	34	32
25	2	183	73	20	1	1	33	34	27	31.333333
26	2	157	57	21	1	2	35	33	32	33.333333
27	2	170	84	23	1	2	38	42	40	40
28	2	163	59	20	1	2	35	36	35	35.333333
29	2	175	79	20	1	2	37	31	34	34
30	2	188	84	20	1	1	28	31	40	33
31	2	167	64	30	1	2	35	29	37	33.666667

HipER1	HipER2	HipER3	HipERAVG	Gast1	Gast2	Gast3	GastAVG	Sol1	Sol2
40	45	43	42.666667	1	1	3	1.666667	5	6
64	66	65	65	14	17	16	15.66667	16	12
76	74	76	75.333333	24	24	23	23.66667	24	23
49	47	48	48	8	10	11	9.666667	13	14
105	102	103	103.33333	18	18	22	19.33333	19	26
83	84	85	84	17	16	15	16	15	16
62	64	65	63.666667	16	18	16	16.66667	20	22
72	75	72	73	22	24	26	24	24	25
105	108	108	107				0		
72	77	73	74	18	17	15	16.66667	21	18
60	59	58	59				0		
93	93	96	94	18	21	20	19.66667	22	24
78	83	77	79.333333	23	18	19	20	13	17
68	71	66	68.333333				0		
86	89	89	88	33	36	35	34.66667	40	42
68	65	67	66.666667	25	22	24	23.66667	24	24
64	64	68	65.333333	11	8	8	9	12	11
78	79	79	78.666667	3	3	4	3.333333	6	8
63	68	71	67.333333	8	8	9	8.333333	14	16
78	79	79	78.666667	6	8	7	7	9	12
63	62	65	63.333333	10	13	13	12	21	20
68	67	68	67.666667	26	25	24	25	28	27
78	79	78	78.333333	18	15	17	16.66667	26	27
86	85	87	86	16	14	17	15.66667	24	27
65	66	63	64.666667	23	25	25	24.33333	29	35
68	72	74	71.333333	14	17	13	14.66667	21	23
75	72	74	73.666667	3	3	3	3	10	12
65	75	80	73.333333	28	26	30	28	34	29
84	85	84	84.333333	14	8	12	11.33333	16	15
78	68	78	74.666667	9	8	11	9.333333	22	20
76	81	83	80	9	9	10	9.333333	12	13

GlutMax1	T1	T2	GlutMax2	T1	T2	GlutMax3	T1	GlutMaxAVG
0.637	1.99	4.99	0.752	11.41	14.41	0.796	19.91	0.728333333
0.539	1.99	4.99	0.475	10.99	13.99	0.48	18.5	0.498
0.93	2.5	5.5	1.498	10	13	1.268	16	1.232
1.002	3	6	1.272	9.5	12.5	1.459	17.5	1.244333333
0.198	1.99	4.99	0.188	8.5	11.5	0.237	15	0.207666667
0.687	1.85	4.85	0.593	10.15	13.15	0.601	17.65	0.627
1.24	1.75	4.75	0.75	10.25	13.25	0.63	17.75	0.873333333
0.788	3.93	6.93	0.804	12.25	15.25	0.837	19.75	0.809666667
0.925	1.99	4.99	0.933	10.5	13.5	1.069	18.99	0.975666667
0.24	1.99	4.99	0.259	10.3	13.3	0.25	19.8	0.249666667
0.689	2.79	5.79	0.913	11.3	14.3	1	19.3	0.867333333
0.546	2.58	5.58	0.719	10.46	13.46	0.652	20.16	0.639
0.558	2.18	5.18	0.455	10.18	13.18	0.348	18.68	0.453666667
0.513	3.381	6.381	0.622	13.58	16.58	0.803	21.58	0.646
1.031	2.38	5.38	0.996	10.38	13.38	0.936	18.88	0.987666667
0.71	2.58	5.58	0.558	9.88	12.88	0.486	18.38	0.584666667
0.38	2.18	5.18	0.329	11.18	14.18	0.371	19.18	0.36
0.591	1.267	4.267	0.517	7.766	10.766	0.488	14.27	0.532
1.296	1.77	4.77	1.423	8.27	11.27	1.321	15.77	1.346666667
0.495	1.99	4.99	0.686	9.77	12.77	0.464	16.77	0.548333333
0.562	1.59	4.59	0.612	9.59	12.59	0.475	17.09	0.549666667
0.576	1.79	4.79	0.584	8.99	11.99	0.69	16.99	0.616666667
0.76	1.69	4.69	0.984	9.19	12.19	0.801	16.09	0.848333333
0.424	7.24	10.24	0.384	16.74	19.74	0.278	26.24	0.362
1.612	1.99	4.99	1.467	9.5	12.5	1.472	16.99	1.517
0.31	2.354	5.354	0.22	9.15	12.15	0.22	15.65	0.25
0.23	1.45	4.45	0.21	9.95	12.95	0.21	16.95	0.216666667
0.39	1.78	4.78	0.4	8.95	11.95	0.36	16.95	0.383333333
0.59	1.38	4.38	0.53	9.38	12.38	0.59	16.38	0.57
0.2	2.89	5.89	0.28	9.28	12.28	0.32	17.78	0.266666667
0.44	1.89	4.89	0.3	8.89	11.89	0.53	15.89	0.423333333

GlutMed1	T1	T2	GlutMed2	T1	T2	GlutMed3	T1	GlutMedAVG
0.409	26.91	29.91	0.638	34.91	37.91	0.571	42.91	0.539333333
0.548	25.99	28.99	0.567	36.5	39.5	0.532	44.5	0.549
0.872	22.5	25.5	0.811	30	33	0.683	36.5	0.788666667
0.94	23.5	26.5	0.762	31	34	0.602	37.5	0.768
0.311	20.1	23.1	0.263	28.6	31.6	0.268	34.09	0.280666667
0.439	26.15	29.15	0.336	35.65	38.65	0.319	44.15	0.364666667
0.555	25.25	28.25	0.557	34.25	37.25	0.523	41.25	0.545
0.676	26.25	29.25	0.831	34.75	37.75	0.923	43.25	0.81
0.341	26.5	29.5	0.317	33.99	36.99	0.38	41.5	0.346
0.206	26.8	29.8	0.236	39.3	42.3	0.32	48.8	0.254
0.944	28.79	31.79	1.121	36.73	39.73	0.934	45.09	0.999666667
0.642	29.16	32.16	0.6	37.16	40.16	0.437	45.36	0.559666667
0.552	26.18	29.18	0.471	34.18	37.18	0.46	43.18	0.494333333
0.138	29.48	32.48	0.205	36.88	39.88	0.239	43.18	0.194
0.494	26.38	29.38	0.51	34.38	37.38	0.501	42.88	0.501666667
0.737	26.38	29.38	0.684	34.38	37.38	0.671	41.38	0.697333333
0.687	26.18	29.18	0.588	34.18	37.18	0.614	42.68	0.629666667
0.39	22.77	25.77	0.441	29.77	32.77	0.409	36.77	0.413333333
0.649	21.77	24.77	0.732	28.27	31.27	0.563	35.27	0.648
0.196	21.77	24.77	0.253	30.27	33.27	0.373	37.77	0.274
0.456	22.09	25.09	0.372	29.59	32.59	0.33	36.59	0.386
0.681	22.99	25.99	0.642	28.99	31.99	0.571	36.99	0.631333333
0.293	116.42	119.42	0.201	123.92	126.92	0.093	131.42	0.195666667
0.152	34.74	37.74	0.164	43.24	46.24	0.214	50.72	0.176666667
1.879	22.99	25.99	2.088	30.49	33.49	1.928	37.5	1.965
0.17	22.15	25.15	0.16	31.15	34.15	0.18	38.15	0.17
0.37	23.45	26.45	0.29	30.45	33.45	0.37	37.45	0.343333333
0.32	22.95	25.95	0.33	30.45	33.45	0.22	37.95	0.29
0.29	23.38	26.38	0.33	31.38	34.38	0.33	38.38	0.316666667
0.43	25.28	28.28	0.34	32.28	35.28	0.28	38.78	0.35
0.38	22.89	25.89	0.35	30.39	33.39	0.29	37.89	0.34

Add1	T1	T2	Add2	T1	T2	Add3	T1	AddAVG	GMaxDecAmp1
0.332	50.4	53.4	0.373	56.9	59.9	0.533	65.4	0.41267	0.1
0.427	52.5	55.5	0.515	60.5	63.5	0.606	68.5	0.516	0.04
0.263	42.5	45.5	0.256	49	52	0.25	55.5	0.25633	0.03
0.501	43	46	0.453	49.5	52.5	0.448	56.5	0.46733	0.07
0.314	40.59	43.59	0.33	48.59	51.59			0.322	0.04
0.283	53.15	56.15	0.232	61.65	64.65	0.272	69.65	0.26233	0.06
0.356	48.25	51.25	0.426	57.25	60.25	0.314	64.75	0.36533	0.14
0.311	50.75	53.75	0.349	58.75	61.75	0.348	66.75	0.336	0.03
0.378	49.5	52.5	0.703	56.5	59.5	0.547	64.5	0.54267	0.05
0.175	58.83	61.83	0.152	68.33	71.33	0.168	75.83	0.165	0.04
0.324	51.59	54.59	0.477	58.59	61.59	0.484	66.09	0.42833	0.03
0.163	53.36	56.36	0.354	62.86	65.86	0.563	72.86	0.36	0.04
0.309	50.68	53.68	0.208	58.68	61.68	0.231	66.48	0.24933	0.04
0.362	51.68	54.68	0.42	59.18	62.18	0.381	66.48	0.38767	0.04
0.612	50.38	53.38	0.634	58.38	61.38	0.549	65.88	0.59833	0.06
0.478	49.38	52.38	0.588	56.88	59.88	0.587	63.88	0.551	0.06
0.331	49.68	52.68	0.255	57.18	60.18	0.371	64.18	0.319	
0.644	47.77	50.77	0.592	55.27	58.27	0.601	62.76	0.61233	
0.471	41.27	44.27	0.443	48.27	51.27	0.276	54.77	0.39667	0.03
0.429	43.77	46.77	0.5	50.27	53.27	0.593	56.77	0.50733	0.05
0.829	43.09	46.09	0.953	50.09	53.09	0.996	57.59	0.926	0.05
1.044	43.99	46.99	0.987	50.99	53.99	1.004	57.99	1.01167	0.08
0.693	137.9	140.9	0.567	144.9	147.9	0.583	152.5	0.61433	0.03
0.156	58.25	61.25	0.137	65.24	68.24	0.133	73.74	0.142	0.04
0.248	45.99	48.99	0.3	52.99	55.99	0.282	60.5	0.27667	0.06
0.21	45.15	48.15	0.2	52.65	55.65	0.18	60.15	0.19667	0.04378
0.25	44.45	47.45	0.21	51.45	54.45	0.31	58.95	0.25667	0.04598
0.54	44.95	47.95	0.66	52.45	55.45	0.64	59.95	0.61333	0.05065
0.29	45.38	48.38	0.48	52.88	55.88	0.47	59.88	0.41333	0.06411
0.35	47.28	50.28	0.43	54.28	57.28	0.5	61.78	0.42667	0.02558
0.23	45.93	48.93	0.21	53.48	56.48	0.15	62.41	0.19667	0.05139

GmDecAmp1	AddDecAmp1	GMAscAmp1	GmAscAmp1	AddAscAmp1	GMDecAmp2
0.06	0.06	0.07	0.07	0.05	0.06
0.04	0.04	0.04	0.03	0.04	0.06
0.47	0.47	0.03	0.13	0.04	0.04
0.09	0.09	0.18	0.09	0.05	0.06
0.03	0.03	0.09	0.04	0.05	0.04
0.05	0.05	0.07	0.09	0.06	0.04
0.12	0.12	0.08	0.07	0.05	0.12
0.04	0.04	0.05	0.07	0.1	0.04
0.04	0.04	0.08	0.09	0.06	0.07
0.07	0.07	0.02	0.03	0.05	0.05
0.03	0.03	0.04	0.04	0.04	0.03
0.04	0.04	0.07	0.05	0.07	0.04
0.03	0.03	0.05	0.04	0.07	0.05
0.03	0.03	0.08	0.03	0.11	0.04
0.03	0.03	0.06	0.03	0.06	0.06
0.04	0.04	0.05	0.05	0.05	0.05
0.05	0.05	0.03	0.05	0.07	0.03
0.05	0.05	0.05	0.05	0.5	0.07
0.03	0.03	0.05	0.03	0.06	0.05
0.05	0.05	0.08	0.05	0.015	0.07
0.06	0.06	0.03	0.06	0.17	0.04
0.05	0.05	0.04	0.05	0.12	0.04
0.04	0.04	0.06	0.04	0.06	0.06
0.03126	0.03126	0.04298	0.03171	0.09713	0.04857
0.04852	0.04852	0.04588	0.04856	0.08147	0.04327
0.05612	0.05612	0.05058	0.05623	0.08286	0.04936
0.08532	0.08532	0.06391	0.08647	0.10334	0.06213
0.05875	0.05875	0.02462	0.05577	0.06336	0.02808
0.11185	0.11185	0.05249	0.11192	0.009565	0.04807

GmDecAmp2	AddDecAmp2	GMAscAmp2	GmAscAmp2	AddAscAmp2	GMDecAmp3
0.07	0.05	0.05	0.08	0.06	0.05
0.04	0.04	0.06	0.04	0.07	0.05
0.69	0.69	0.04	1.11	0.07	0.04
0.07	0.07	0.15	0.07	0.05	0.07
0.03	0.03	0.08	0.03	0.06	0.03
0.08	0.08	0.08	0.1	0.07	0.04
0.11	0.11	0.09	0.09	0.05	0.12
0.05	0.05	0.05	0.08	0.09	0.04
0.04	0.04	0.08	0.09	0.06	0.06
0.06	0.06	0.03	0.05	0.05	0.04
0.05	0.05	0.04	0.04	0.04	0.03
0.05	0.05	0.08	0.06	0.07	0.04
0.04	0.04	0.06	0.04	0.07	0.05
0.03	0.03	0.09	0.03	0.09	0.04
0.4	0.04	0.06	0.04	0.06	0.06
0.05	0.05	0.05	0.06	0.05	0.03
0.05	0.05	0.03	0.05	0.06	0.04
0.06	0.06	0.07	0.06	0.05	0.06
0.04	0.04	0.05	0.04	0.05	0.05
0.06	0.06	0.07	0.06	0.018	0.08
0.06	0.06	0.04	0.06	0.19	0.04
0.04	0.04	0.04	0.04	0.11	0.05
0.06	0.06	0.06	0.07	0.07	0.07
0.04258	0.04258	0.04829	0.04214	0.11685	0.04992
0.04948	0.04948	0.04342	0.04923	0.08965	0.04115
0.05217	0.05217	0.04961	0.05216	0.08979	0.05329
0.09892	0.09892	0.06211	0.09839	0.11926	0.06444
0.05995	0.05995	0.02785	0.05898	0.06287	0.02653
0.11058	0.11058	0.04679	0.11034	0.08518	0.05407

GmDecAmp3	AddDecAmp3	GMAscAmp3	GmAscAmp3	AddAscAmp3	GMDecAmp4
0.07	0.07	0.04	0.07	0.05	0.07
0.05	0.05	0.05	0.06	0.07	0.05
0.14	0.14	0.05	0.17	0.05	0.03
0.08	0.08	0.2	0.09	0.06	0.06
0.06	0.06	0.09	0.07	0.08	0.04
0.08	0.08	0.09	0.1	0.07	0.06
0.1	0.1	0.08	0.07	0.05	0.15
0.06	0.06	0.04	0.08	0.1	0.04
0.04	0.04	0.09	0.08	0.06	0.06
0.07	0.7	0.03	0.04	0.05	0.06
0.04	0.04	0.05	0.05	0.04	0.04
0.06	0.06	0.08	0.06	0.07	0.04
0.04	0.04	0.05	0.04	0.06	0.04
0.03	0.03	0.1	0.03	0.09	0.04
0.04	0.04	0.06	0.03	0.07	0.05
0.05	0.05	0.06	0.08	0.05	0.03
0.05	0.05	0.04	0.05	0.07	0.04
0.05	0.05	0.06	0.05	0.06	0.07
0.05	0.05	0.05	0.05	0.06	0.05
0.05	0.05	0.08	0.05	0.16	0.07
0.05	0.05	0.04	0.05	0.17	0.04
0.05	0.05	0.05	0.05	0.17	0.05
0.07	0.07	0.07	0.07	0.06	0.1
0.03516	0.03516	0.05011	0.0353	0.08845	0.07649
0.0478	0.0478	0.04102	0.04759	0.07969	0.04368
0.06444	0.06444	0.06265	0.06459	0.08658	0.05538
0.09866	0.09866	0.06403	0.09936	0.12706	0.06453
0.05682	0.05682	0.0263	0.05709	0.06567	0.02872
0.11078	0.11078	0.05418	0.11088	0.09993	0.05769

GmDecAmp4	AddDecAmp4	GMAscAmp4	GmAscAmp4	AddAscAmp4	GMDecAmp5
0.07	0.07	0.05	0.06	0.06	0.09
0.05	0.05	0.06	0.05	0.08	0.07
0.3	0.3	0.04	0.43	0.06	0.04
0.09	0.09	0.17	0.08	0.05	0.06
0.04	0.04	0.08	0.05	0.06	0.05
0.08	0.08	0.08	0.12	0.06	0.05
0.12	0.12	0.08	0.07	0.04	0.1
0.05	0.05	0.05	0.08	0.11	0.04
0.05	0.05	0.07	0.08	0.06	0.06
0.08	0.08	0.03	0.04	0.05	0.5
0.05	0.05	0.05	0.05	0.04	0.04
0.05	0.05	0.09	0.07	0.09	0.05
0.04	0.04	0.06	0.04	0.06	0.04
0.02	0.02	0.1	0.03	0.11	0.04
0.03	0.03	0.05	0.08	0.07	0.05
0.04	0.04	0.06	0.07	0.05	0.03
0.05	0.05	0.03	0.04	0.07	0.04
0.05	0.05	0.07	0.05	0.06	0.08
0.05	0.05	0.05	0.05	0.07	0.05
0.06	0.06	0.07	0.05	0.17	0.06
0.05	0.05	0.04	0.05	0.18	0.04
0.06	0.06	0.05	0.06	0.15	0.05
0.09	0.09	0.1	0.09	0.07	0.07
0.05763	0.05736	0.08034	0.05718	0.09661	0.0598
0.04582	0.04582	0.04309	0.04507	0.08115	0.04533
0.06387	0.06387	0.05597	0.06564	0.083	0.06581
0.10838	0.10832	0.06465	0.10746	0.011638	0.06866
0.05742	0.05742	0.02901	0.05846	0.07349	0.02581
0.11272	0.11272	0.05533	0.11293	0.09435	0.04425

GmDAmp5	AddDAmp5	GMAAmp5	GmAmp5	AddAAmp5	GMDAmpAVG
0.06	0.06	0.08	0.06	0.05	0.074
0.04	0.04	0.07	0.05	0.07	0.054
0.14	0.14	0.04	0.16	0.05	0.036
0.09	0.09	0.19	0.09	0.05	0.064
0.04	0.4	0.06	0.04	0.05	0.04
0.09	0.09	0.09	0.09	0.06	0.05
0.12	0.12	0.08	0.06	0.04	0.126
0.07	0.07	0.04	0.11	0.12	0.038
0.04	0.04	0.07	0.06	0.06	0.06
0.06	0.06	0.04	0.05	0.05	0.138
0.05	0.05	0.04	0.04	0.04	0.034
0.07	0.07	0.1	0.07	0.08	0.042
0.04	0.04	0.08	0.04	0.06	0.044
0.02	0.02	0.1	0.03	0.1	0.04
0.03	0.03	0.05	0.08	0.07	0.056
0.05	0.05	0.05	0.05	0.06	0.04
					0
					0
0.05	0.05	0.04	0.05	0.06	0.036
0.05	0.05	0.08	0.05	0.05	0.066
0.05	0.05	0.05	0.06	0.06	0.05
0.05	0.05	0.06	0.05	0.16	0.072
0.06	0.06	0.04	0.06	0.2	0.038
0.05	0.05	0.05	0.05	0.13	0.046
0.06	0.06	0.07	0.06	0.07	0.072
0.0408	0.0408	0.05934	0.04108	0.08328	0.055712
0.05675	0.05675	0.04539	0.05607	0.08619	0.043882
0.05994	0.05994	0.0667	0.0619	0.08267	0.054898
0.11466	0.11466	0.06867	0.11413	0.1045	0.064774
0.05647	0.05647	0.02589	0.05688	0.07069	0.026944
0.10953	0.10953	0.04372	0.10998	0.8704	0.051094

GMDesAmpNorm	GmDesAmpAVG	GmDesAmpNorm	ADesAmpAVG	ADesAmpNorm
10.16018307	0.066	12.23733004	0.062	15.02423263
10.84337349	0.044	8.014571949	0.044	8.527131783
2.922077922	0.348	44.12510566	0.348	135.7607282
5.143316368	0.084	10.9375	0.084	17.9743224
19.26163724	0.04	14.25178147	0.112	34.7826087
7.974481659	0.076	20.84095064	0.076	28.9707751
14.42748092	0.114	20.91743119	0.114	31.20437956
4.69328942	0.054	6.666666667	0.054	16.07142857
6.149641271	0.042	12.13872832	0.042	7.73955774
55.27369826	0.068	26.77165354	0.194	117.5757576
3.920061491	0.044	4.401467156	0.044	10.27237354
6.572769953	0.054	9.648600357	0.054	15
9.698750918	0.038	7.687120701	0.038	15.24064171
6.191950464	0.026	13.40206186	0.026	6.706792777
5.669929126	0.106	21.12956811	0.034	5.682451253
6.841505131	0.046	6.596558317	0.046	8.34845735
0	0	0	0	0
0	0	0	0	0
2.673267327	0.05	7.716049383	0.05	12.60504202
12.03647416	0.052	18.97810219	0.052	10.24967148
9.096422074	0.044	11.39896373	0.044	4.75161987
11.67567568	0.054	8.553326294	0.054	5.337726524
4.479371316	0.056	28.62010221	0.056	9.115572436
12.70718232	0.05	28.30188679	0.05	35.21126761
4.746209624	0.064	3.256997455	0.064	23.13253012
22.2848	0.041486	24.40352941	0.041432	21.06711864
20.25323077	0.049674	14.46815534	0.049674	19.35350649
14.32121739	0.059308	20.45103448	0.059308	9.669782609
11.36385965	0.101188	31.95410526	0.101176	24.47806452
10.104	0.057882	16.53771429	0.057882	13.56609375
12.06944882	0.111092	32.67411765	0.111092	56.48745763

GMAAscAmpAVG	GMAAscAmpNorm	GmAscAmpAVG	GmAscAmpNorm	AddAAmpAVG
0.058	7.963386728	0.068	12.60815822	0.054
0.056	11.24497992	0.046	8.378870674	0.066
0.04	3.246753247	0.4	50.71851226	0.054
0.178	14.30484865	0.084	10.9375	0.052
0.08	38.52327448	0.046	16.38954869	0.06
0.082	13.07814992	0.1	27.42230347	0.064
0.082	9.389312977	0.072	13.21100917	0.046
0.046	5.68135035	0.084	10.37037037	0.104
0.078	7.994533652	0.08	23.12138728	0.06
0.03	12.01602136	0.042	16.53543307	0.05
0.044	5.073020753	0.044	4.401467156	0.04
0.084	13.14553991	0.062	11.07802263	0.076
0.06	13.22556943	0.04	8.091706001	0.064
0.094	14.55108359	0.03	15.46391753	0.1
0.056	5.669929126	0.052	10.3654485	0.066
0.054	9.236031927	0.062	8.891013384	0.052
0	0	0	0	0
0	0	0	0	0
0.034	2.524752475	0.048	7.407407407	0.066
0.066	12.03647416	0.052	18.97810219	0.144
0.05	9.096422074	0.046	11.91709845	0.06
0.072	11.67567568	0.052	8.236536431	0.1046
0.038	4.479371316	0.056	28.62010221	0.182
0.046	12.70718232	0.05	28.30188679	0.136
0.072	4.746209624	0.066	3.358778626	0.066
0.056212	22.4848	0.041482	24.40117647	0.096464
0.04376	20.19692308	0.049304	14.36038835	0.08363
0.057102	14.89617391	0.060104	20.72551724	0.08498
0.064674	11.34631579	0.101162	31.94589474	0.0931596
0.026734	10.02525	0.057436	16.41028571	0.067216
0.050502	11.9296063	0.11121	32.70882353	0.231885

AddAscAmpNorm	GMaxDecAmp1board	GMedDecAmp1board	AddDecAmp1board
13.08562197			
12.79069767			
21.0663199			
11.12696148			
18.63354037			
24.39644219			
12.59124088			
30.95238095			
11.05651106			
30.3030303			
9.338521401			
21.11111111			
25.6684492			
25.79535684			
11.03064067			
9.43738657			
0			
0	0.06	0.07	0.07
16.63865546	0.03	0.05	0.05
28.38370565	0.06	0.05	0.05
6.479481641	0.05	0.03	0.03
10.33937397	0.06	0.06	0.06
29.62561042	0.04	0.04	0.04
95.77464789	0.04	0.04	0.04
23.85542169	0.09	0.09	0.09
49.04949153	0.04033	0.03527	0.03527
32.58311688	0.04026	0.05816	0.05816
13.85543478	0.05102	0.05213	0.05213
22.5386129	0.08254	0.07655	0.07655
15.75375	0.02713	0.05337	0.05337
117.9076271	0.03356	0.1032	0.1032

GMaxAscAmp1board	GmedAscAmp1board	AddAscAmp1board	GMaxDecAmp2board
0.06	0.06	0.09	0.07
0.03	0.05	0.06	0.03
0.06	0.05	0.05	0.09
0.05	0.03	0.05	0.04
0.05	0.06	0.17	0.05
0.04	0.04	0.11	0.04
0.05	0.07	0.11	0.05
0.09	0.09	0.07	0.05
0.04059	0.03548	0.08776	0.04304
0.03981	0.05612	0.06958	0.0385
0.05158	0.05162	0.08114	0.05037
0.08306	0.07663	0.09325	0.08231
0.10029	0.11744	0.11024	0.02447
0.05893	0.10929	0.11087	0.03253

GMaxDesAmpboardNorm	GMedDesAmpboardAVG	GMedDesAmpboardNorm
0	0	0
0	0	0
0	0	0
0	0	0
0	0	0
0	0	0
0	0	0
0	0	0
0	0	0
0	0	0
0	0	0
0	0	0
0	0	0
0	0	0
0	0	0
0	0	0
0	0	0
0	0	0
0	0	0
0	0	0
13.90977444	0.076	18.38709677
2.524752475	0.048	7.407407407
13.49544073	0.05	18.24817518
8.004851425	0.038	9.844559585
8.756756757	0.046	7.286166843
4.950884086	0.05	25.55366269
14.91712707	0.056	31.69811321
3.823335531	0.056	2.849872774
16.5536	0.036406	21.41529412
18.61476923	0.04791	13.95436893
13.42852174	0.058426	20.14689655
14.24421053	0.08901	28.10842105
9.9945	0.064644	18.46971429
8.203937008	0.104936	30.86352941

AddDesAmpboardAVG	AddDesAmpboardNorm	GMaxAscAmpboardAVG
0	0	0
0	0	0
0	0	0
0	0	0
0	0	0
0	0	0
0	0	0
0	0	0
0	0	0
0	0	0
0	0	0
0	0	0
0	0	0
0	0	0
0	0	0
0	0	0
0	0	0
0	0	0
0	0	0
0	0	0
0.076	12.41154056	0.072
0.048	12.10084034	0.034
0.05	9.855453351	0.074
0.038	4.103671706	0.044
0.046	4.546952224	0.052
0.05	8.138903961	0.042
0.056	39.43661972	0.05
0.056	20.24096386	0.06
0.036406	18.51152542	0.041492
0.04791	18.66623377	0.040042
0.058426	9.525978261	0.05226
0.08901	21.53467742	0.08132
0.064644	15.1509375	0.096226
0.104936	53.35728814	0.066548

GMaxAscAmpboardNorm	GMedAscAmpboardAVG	GMedAscAmpboardNorm
0	0	0
0	0	0
0	0	0
0	0	0
0	0	0
0	0	0
0	0	0
0	0	0
0	0	0
0	0	0
0	0	0
0	0	0
0	0	0
0	0	0
0	0	0
0	0	0
0	0	0
0	0	0
0	0	0
0	0	0
13.53383459	0.07	16.93548387
2.524752475	0.044	6.790123457
13.49544073	0.05	18.24817518
8.004851425	0.036	9.32642487
8.432432432	0.05	7.919746568
4.950884086	0.05	25.55366269
13.8121547	0.068	38.49056604
3.955174687	0.056	2.849872774
16.5968	0.036716	21.59764706
18.48092308	0.04731	13.77961165
13.63304348	0.058368	20.12689655
14.26666667	0.089464	28.25178947
36.08475	0.130684	37.33828571
15.72	0.112514	33.09235294

AddAscAmpboardAVG	AddAscAmpboardNorm	ERConPkTq	ERConPkTqNorm
0	0	31.7667	0.32415
0	0	32.1	0.324242424
0	0	30.067	0.455560606
0	0	24.733	0.313075949
0	0	16.267	0.275711864
0	0	13.167	0.243833333
0	0	18.467	0.302737705
0	0	20.633	0.322390625
0	0	19.933	0.36912963
0	0	17.467	0.226844156
0	0	21.467	0.271734177
0	0	12.067	0.188546875
0	0	20.933	0.486813953
0	0	31	0.360465116
0	0	16.633	0.281915254
0	0	17.733	0.268681818
0	0	25.767	0.34356
0.102	16.6575939	17.533	0.28742623
0.062	15.6302521	19.567	0.320770492
0.05	9.855453351	20.7	0.339344262
0.058	6.26349892	13.933	0.258018519
0.172	17.00164745	32.767	0.511984375
0.112	18.23114487	2.0333	0.033332787
0.112	78.87323944	12.567	0.206016393
0.066	23.85542169	31.367	0.429684932
0.089528	45.52271186	6.8	0.119298246
0.076198	29.68753247	14.5	0.172619048
0.083722	13.65032609	8.0667	0.136723729
0.099162	23.99080645	21.633	0.273835443
0.123262	28.88953125	37.2	0.442857143
0.115976	58.97084746	17.4	0.271875

ERConAngPkTq	ERConTmPkTq	EREccPkTq	EREccPkTqNorm	EREccAngPkTq
14	3	56.9	0.580612245	13.6667
15	1.3333	60.733	0.613464646	15
14.667	2.6667	53.6	0.812121212	14.667
15	2.6667	51.9	0.656962025	15
15	1.6667	18.767	0.318084746	14.667
15	1.6667	26.7	0.494444444	14.667
12	2.6667	40.333	0.661196721	12
14	3.6667	48.267	0.754171875	14
15	3	31	0.574074074	15
15	5.3333	32.767	0.425545455	15
12.667	6.3333	58.7	0.743037975	14
13.667	4	35	0.546875	14.333
15	1.3333	34.4	0.8	14
12	6.3333	33.167	0.385662791	13.667
14.667	2.3333	39.733	0.673440678	14.333
13	5.3333	35.6	0.539393939	13.333
14	3.6667	53.167	0.708893333	14.333
15	1	39.8	0.652459016	15
14.333	2	42.167	0.691262295	12.667
13.667	3.6667	30.333	0.497262295	13
14.667	3.3333	34.567	0.64012963	13
13	3	13.067	0.204171875	13.333
14.667	3.6667	24.933	0.408737705	15
15	1.6667	29.233	0.479229508	15
14.333	3.6667	55.2	0.756164384	15
13.667	4	27.633	0.484789474	14.667
9	1.3333	42.467	0.505559524	9
10.667	8	12.067	0.204525424	12.667
13.667	2.6667	45.733	0.578898734	14
6	1	94.333	1.123011905	4.6667
14.333	2.6667	36.467	0.569796875	14

EREccTmPkTq	IRConPkTq	IRConPkTqNorm	IRConAngPkTq	IRConTmPkTq
4.3333	30.067	0.306806122	1	2
2.6667	89.3	0.902020202	6	1.6667
2.3333	85.1	1.289393939	5	1.6667
2.3333	55.3	0.7	5	3.3333
6	24.167	0.409610169	5	1.6667
4.3333	52.367	0.969759259	6	2.3333
2.6667	31.433	0.515295082	2	4
2	74.867	1.169796875	5	1.3333
1	49.733	0.920981481	3	3.3333
3.3333	37.333	0.484844156	5	1
4	49.233	0.623202532	5	1
2.6667	57.667	0.901046875	4	1.3333
	54.3	1.262790698	5	3
4	51.067	0.593802326	5	1
2.6667	47.533	0.805644068	5	2.3333
4.6667	53.033	0.803530303	5	3.6667
2.6667	78	1.04	5	4.3333
1.3333	52.7	0.863934426	5	3.3333
4.6667	62.9	1.031147541	5	2
5.6667	40.8	0.668852459	5	1
6.6667	33.567	0.621611111	6	3.6667
2.6667	65.133	1.017703125	5	2
2	57.1	0.936065574	4	4.6667
1.6667	26.4	0.432786885	5	2
2	84.933	1.163465753	5	3.3333
1.6667	50.767	0.890649123	5	4
1.3333	70.1	0.83452381	5	4.3333
5.3333	39.4	0.66779661	5	3.3333
1.3333	44.4	0.562025316	4	3
3.3333	66.133	0.787297619	5	5.3333
4	49.8	0.778125	5	3

IREcPkTq	IREcPkTqNorm	IREcAngPkTq	IREcTmPkTq	ExtCoPkTq	ExtCoPkTqNorm
26.367	0.26905102	1	3.3333	81.867	0.835377551
63.333	0.639727273	6	1	204.7	2.067676768
50.067	0.758590909	5	1.3333	79.433	1.203530303
29	0.367088608	5		88.067	1.114772152
17.2	0.291525424	5	1	69.867	1.184186441
29.567	0.547537037	6	3	78.833	1.45987037
14.867	0.243721311	2	4.6667	38.967	0.638803279
35.567	0.555734375	5	4	135.97	2.12453125
31.6	0.585185185	3	2.6667	113.07	2.093888889
23.733	0.308220779	5	4.6667	54.767	0.71125974
25.7	0.325316456	5	2	120.33	1.523164557
23.8	0.371875	4	1.3333	95.867	1.497921875
31.6	0.734883721	4.6667	3.6667	37.867	0.880627907
10.167	0.11822093	5	4.3333	129.2	1.502325581
26.2	0.444067797	5	3	26.433	0.448016949
28.733	0.435348485	5		55.333	0.838378788
41.533	0.553773333	5	1.3333	124.4	1.658666667
25.267	0.414213115	5	4	85.867	1.407655738
38.3	0.627868852	5	1.6667	40.167	0.65847541
27.4	0.449180328	5	5	75.967	1.245360656
18.833	0.348759259	6	1.3333	54.933	1.017277778
43.133	0.673953125	5	2	100.43	1.56921875
9.7	0.159016393	3	8.6667	51.767	0.848639344
6.3	0.103278689	5	2	110.3	1.808196721
56.9	0.779452055	5	1	134.9	1.847945205
22.533	0.395315789	5	2	49.633	0.870754386
32.7	0.389285714	5	1	114.47	1.362738095
23.667	0.401135593	5	3	60.767	1.029949153
19.5	0.246835443	4	5	47.533	0.601683544
45	0.535714286	5	2.3333	154.33	1.837261905
29.8	0.465625	5	1	84.233	1.316140625

ExCoAngPkTq	ExCoTmPkTq	ExEcPkTq	ExEcPkTqNorm	ExEcAngPkTq	ExEcTmPkTq
39	2.6667	62.7	0.639795918	39	2
0	4.3333	126	1.272727273	0	1
0.66667	3.3333	49.033	0.742924242	0	1
40	4	38.967	0.493253165	40	3.6667
1.6667	3.6667	54.467	0.923169492	1.3333	3.6667
-1	4.6667	39.833	0.737648148	1	1.6667
1.6667	4	32.2	0.527868852	0	1
0	2.3333	18.6	0.290625	0.33333	4
0	3.3333	101.77	1.88462963	0	4.3333
0	3	21.767	0.282688312	0	1
0	4.6667	92.8	1.174683544	0	1
40	6.6667	43.067	0.672921875	40	1
1	3	56	1.302325581	0.33333	2
39	6	48.867	0.56822093	39	1
1.6667	9.3333	53.567	0.907915254	0.66667	6.3333
2.6667	5.3333	28.967	0.438893939	3.6667	7
3.3333	6	63.433	0.845773333	0	1
2	5.6667	44.9	0.736065574	0.66667	2.3333
40	6.6667	81.1	1.329508197	40	4.3333
31	3	47.967	0.786344262	31	1
39	4.6667	41.6	0.77037037	39	3
0.33333	6.3333	56.067	0.876046875	0.33333	4.3333
1	2	61.9	1.014754098	1	1.6667
40	3.3333	33.967	0.556836066	40	3.3333
2	4	88.4	1.210958904	2	3
40	6.6667	17.733	0.311105263	40	8.3333
1	6	42.267	0.503178571	1	3
2	6	9.8	0.166101695	0	
4	6.6667	31.867	0.403379747	1	1
40.333	3.3333	81.3	0.967857143	39.667	4.6667
0	4.3333	37.867	0.591671875	0	3

AbCoPkTq	AbCoPkTqNorm	AbCoAngPkTq	AbCoTmPkTq	AbEcPkTq	AbEcPkTqNorm
54.3667	0.554762245	0	3.3333	45.1667	0.460884694
38.9	0.392929293	0	1	41.933	0.423565657
35.833	0.542924242	0	1	56.5	0.856060606
37.8	0.478481013	0	1	41.767	0.528696203
21.1	0.357627119	0	1	7.1667	0.121469492
32.1	0.594444444	0	1.6667	39.067	0.723462963
20.267	0.332245902	0	1	8.1667	0.133880328
39.733	0.620828125	0	1	64.2	1.003125
23.967	0.443833333	0	1	30.6	0.566666667
18.033	0.234194805	0	1	27.3	0.354545455
52.667	0.666670886	0	1	56.5	0.715189873
11.567	0.180734375	0	1	36.6	0.571875
28.033	0.651930233	0	1	24.233	0.56355814
4.6667	0.054263953	0	1	38.633	0.44922093
25.933	0.439542373	0	1	2.9333	0.049716949
21.533	0.326257576	0	1	15	0.227272727
43.467	0.57956	0	1	66.733	0.889773333
32.967	0.540442623	0	1	47.667	0.78142623
32.467	0.532245902	0	1	42.033	0.689065574
30.667	0.502737705	-1	1	37.7	0.618032787
16.533	0.306166667	-1	1	18.933	0.350611111
25.167	0.393234375	0	2.6667	37.1	0.5796875
9.8667	0.16174918	0	1	6.9667	0.114208197
14.767	0.242081967	-1	1	50.833	0.833327869
27.467	0.376260274	-1.6667	3	36.033	0.49360274
7.2333	0.1269	-0.66667	2.6667	28.4	0.498245614
26.3	0.313095238	-1	1	53.8	0.64047619
14.333	0.242932203	0	1.3333	24.667	0.418084746
13	0.164556962	-1	1.3333	9.2667	0.1173
38.233	0.455154762	-0.33333	2	43.7	0.520238095
13.933	0.217703125	-1	2.6667	23.2	0.3625

AbEccAngPkTq	AbEccTmPkTq
0	2.3333
0	1
-1.6667	4
0	2
0	3.6667
0	1.3333
0	3.6667
0	2.3333
0	2.6667
-0.33333	2
0	2.6667
0	1
-1.3333	4.3333
0	1
-0.66667	6
-0.33333	2
-0.66667	4.3333
-0.66667	4
-1.6667	4
-2	3.6667
-1.6667	4
0	2.6667
-0.66667	4.3333
-2	4
-1	1
-1.6667	5
-2.6667	4.6667
0	1.6667
-1	2.6667
0	1.6667
-1.3333	3.6667

APPENDIX G: SPSS OUTPUTS

General Linear Model

Within-Subjects Factors

Measure: MEASURE_1

muscle	Dependent Variable
1	GastAVG
2	SolAVG

Between-Subjects Factors

Group	Value Label	N
1.0000	Normal	14
2.0000	Valgus	14

Descriptive Statistics

	Group	Mean	Std. Deviation	N
GastAVG	Normal	17.880952	7.9255156	14
	Valgus	13.428571	7.8451123	14
	Total	15.654762	8.0632694	28
SolAVG	Normal	19.857143	8.3724357	14
	Valgus	19.880952	8.3692632	14
	Total	19.869048	8.2143803	28

Multivariate Tests

Effect		Value	F	Hypothesis df	Error df	Sig.	Partial Eta Squared	Noncent. Parameter	Observed Power ^a
muscle	Pillai's Trace	.704	61.695 ^b	1.000	26.000	.000	.704	61.695	1.000
	Wilks' Lambda	.296	61.695 ^b	1.000	26.000	.000	.704	61.695	1.000
	Hotelling's Trace	2.373	61.695 ^b	1.000	26.000	.000	.704	61.695	1.000
	Roy's Largest Root	2.373	61.695 ^b	1.000	26.000	.000	.704	61.695	1.000
muscle * Group	Pillai's Trace	.401	17.400 ^b	1.000	26.000	.000	.401	17.400	.980
	Wilks' Lambda	.599	17.400 ^b	1.000	26.000	.000	.401	17.400	.980
	Hotelling's Trace	.669	17.400 ^b	1.000	26.000	.000	.401	17.400	.980
	Roy's Largest Root	.669	17.400 ^b	1.000	26.000	.000	.401	17.400	.980

a. Computed using alpha = .05

b. Exact statistic

c.

Design: Intercept+Group

Within Subjects Design: muscle

Mauchly's Test of Sphericity

Measure: MEASURE_1

		Approx. Chi-Square	df	Sig.	Epsilon ^a		
					Greenhouse-Geisser	Huynh-Feldt	Lower-bound
Within Subjects Effects	Mauchly's W						
muscle	1.000	.000	0	.	1.000	1.000	1.000

Tests the null hypothesis that the error covariance matrix of the orthonormalized transformed dependent variables is proportional to an identity matrix.

a. May be used to adjust the degrees of freedom for the averaged tests of significance. Corrected tests are displayed in the Tests of Within-Subjects Effects table.

b.

Design: Intercept+Group

Within Subjects Design: muscle

Tests of Within-Subjects Effects

Measure: MEASURE_1

Source		Type III Sum of Squares	df	Mean Square	F	Sig.	Partial Eta Squared	Noncent. Parameter	Observed Power ^a
muscle	Sphericity Assumed	248.643	1	248.643	61.695	.000	.704	61.695	1.000
	Greenhouse-Geisser	248.643	1.000	248.643	61.695	.000	.704	61.695	1.000
	Huynh-Feldt	248.643	1.000	248.643	61.695	.000	.704	61.695	1.000
	Lower-bound	248.643	1.000	248.643	61.695	.000	.704	61.695	1.000
muscle * Group	Sphericity Assumed	70.127	1	70.127	17.400	.000	.401	17.400	.980
	Greenhouse-Geisser	70.127	1.000	70.127	17.400	.000	.401	17.400	.980
	Huynh-Feldt	70.127	1.000	70.127	17.400	.000	.401	17.400	.980
	Lower-bound	70.127	1.000	70.127	17.400	.000	.401	17.400	.980
Error(muscle)	Sphericity Assumed	104.786	26	4.030					
	Greenhouse-Geisser	104.786	26.000	4.030					
	Huynh-Feldt	104.786	26.000	4.030					
	Lower-bound	104.786	26.000	4.030					

a. Computed using alpha = .05

Tests of Within-Subjects Contrasts

Measure: MEASURE_1

Source	muscle	Type III Sum of Squares	df	Mean Square	F	Sig.	Partial Eta Squared	Noncent. Parameter	Observed Power ^a
muscle	Linear	248.643	1	248.643	61.695	.000	.704	61.695	1.000
muscle * Group	Linear	70.127	1	70.127	17.400	.000	.401	17.400	.980
Error(muscle)	Linear	104.786	26	4.030					

a. Computed using alpha = .05

Tests of Between-Subjects Effects

Measure: MEASURE_1

Transformed Variable: Average

Source	Type III Sum of Squares	df	Mean Square	F	Sig.	Partial Eta Squared	Noncent. Parameter	Observed Power ^a
Intercept	17667.175	1	17667.175	137.787	.000	.841	137.787	1.000
Group	68.643	1	68.643	.535	.471	.020	.535	.109
Error	3333.738	26	128.221					

a. Computed using alpha = .05

Estimated Marginal Means

1. Grand Mean

Measure: MEASURE_1

Mean	Std. Error	95% Confidence Interval	
		Lower Bound	Upper Bound
17.762	1.513	14.652	20.872

2. Group

Measure: MEASURE_1

Group	Mean	Std. Error	95% Confidence Interval	
			Lower Bound	Upper Bound
Normal	18.869	2.140	14.470	23.268
Valgus	16.655	2.140	12.256	21.053

3. muscle

Measure: MEASURE_1

muscle	Mean	Std. Error	95% Confidence Interval	
			Lower Bound	Upper Bound
1	15.655	1.490	12.592	18.718
2	19.869	1.582	16.617	23.121

4. Group * muscle

Measure: MEASURE_1

Group	muscle	Mean	Std. Error	95% Confidence Interval	
				Lower Bound	Upper Bound
Normal	1	17.881	2.107	13.549	22.213
	2	19.857	2.237	15.259	24.456
Valgus	1	13.429	2.107	9.097	17.761
	2	19.881	2.237	15.282	24.480

Oneway

Descriptives

	N	Mean	Std. Deviation	Std. Error	95% Confidence Interval for Mean		Minimum	Maximum
					Lower Bound	Upper Bound		
HipERAVC Normal	17	3.921569	17.5144898	.2478877	64.916449	82.926688	42.6667	107.0000
Valgus	14	4.428571	7.0169862	.8753684	70.377084	78.480059	63.3333	86.0000
Total	31	4.150538	13.6016989	.4429372	69.161394	79.139681	42.6667	107.0000
HipABAVC Normal	17	6.882353	5.3384626	.2947674	34.137569	39.627137	24.3333	45.0000
Valgus	14	5.285714	3.9696652	.0609376	32.993698	37.577731	31.3333	44.6667
Total	31	6.161290	4.7624076	.8553536	34.414425	37.908156	24.3333	45.0000

ANOVA

		Sum of Squares	df	Mean Square	F	Sig.
HipERAVG	Between Groups	1.973	1	1.973	.010	.920
	Within Groups	5548.213	29	191.318		
	Total	5550.186	30			
HipABAVG	Between Groups	19.572	1	19.572	.859	.362
	Within Groups	660.844	29	22.788		
	Total	680.416	30			

T-Test

Group Statistics

	Group	N	Mean	Std. Deviation	Std. Error Mean
HipABAVG	Normal	17	36.882353	5.3384626	1.2947674
	Valgus	14	35.285714	3.9696652	1.0609376
HipERAVG	Normal	17	73.921569	17.5144898	4.2478877
	Valgus	14	74.428571	7.0169862	1.8753684
GastAVG	Normal	14	17.880952	7.9255156	2.1181831
	Valgus	14	13.428571	7.8451123	2.0966945
SolAVG	Normal	14	19.857143	8.3724357	2.2376276
	Valgus	14	19.880952	8.3692632	2.2367797

Independent Samples Test

		Levene's Test for Equality of Variances		t-test for Equality of Means						
		F	Sig.	t	df	Sig. (2-tailed)	Mean Difference	Std. Error Difference	95% Confidence Interval of the Difference	
									Lower	Upper
HipABAVG	Equal variance assumed	1.031	.318	.927	29	.362	.5966387	.7228318	-1.92695	1202252
	Equal variance not assumed			.954	28.748	.348	.5966387	.6739209	-1.82822	0214957
HipERAVG	Equal variance assumed	6.408	.017	-.102	29	.920	-.5070028	.9919466	-10.7167	7026743
	Equal variance not assumed			-.109	21.824	.914	-.5070028	.6434424	-10.1414	1274008
GastAVG	Equal variance assumed	.093	.763	1.494	26	.147	-.4523810	.9804073	-1.67393	10.57870
	Equal variance not assumed			1.494	25.997	.147	-.4523810	.9804073	-1.67396	10.57873
SolAVG	Equal variance assumed	.399	.533	-.008	26	.994	-.0238095	.1638838	-6.52727	4796467
	Equal variance not assumed			-.008	26.000	.994	-.0238095	.1638838	-6.52727	4796467

T-Test

Paired Samples Statistics

		Mean	N	Std. Deviation	Std. Error Mean
Pair 1	GastAVG	13.428571	14	7.8451123	2.0966945
	SolAVG	19.880952	14	8.3692632	2.2367797

Paired Samples Correlations

		N	Correlation	Sig.
Pair 1	GastAVG & SolAVG	14	.930	.000

Paired Samples Test

	Paired Differences					t	df	Sig. (2-tailed)
	Mean	Std. Deviation	Std. Error	95% Confidence Interval of the Difference				
				Lower	Upper			
Pair 1 GastAVG - SolA	-6.45238	3.0734795	.8214219	-8.22696	-4.67781	-7.855	13	.000

Oneway

Descriptives

		N	Mean	Std. Deviation	Std. Error	95% Confidence Interval for Mean		Minimum	Maximum
						Lower Bound	Upper Bound		
ERConPkTqNorm	Normal	17	.315258	.0753943	.0182858	.276494	.354023	.1885	.4868
	Valgus	14	.271699	.1335344	.0356886	.194599	.348799	.0333	.5120
	Total	31	.295586	.1060385	.0190451	.256691	.334482	.0333	.5120
EREccPkTqNorm	Normal	17	.605175	.1431187	.0347114	.531590	.678760	.3181	.8121
	Valgus	14	.556857	.2291982	.0612558	.424522	.689192	.2042	1.1230
	Total	31	.583354	.1851635	.0332563	.515436	.651273	.2042	1.1230
IRConPkTqNorm	Normal	17	.805796	.2927646	.0710058	.655270	.956321	.3068	1.2894
	Valgus	14	.803999	.2006404	.0536234	.688153	.919845	.4328	1.1635
	Total	31	.804984	.2513124	.0451370	.712802	.897166	.3068	1.2894
IREccPkTqNorm	Normal	17	.444110	.1804319	.0437612	.351340	.536879	.1182	.7586
	Valgus	14	.427831	.1876576	.0501536	.319481	.536181	.1033	.7795
	Total	31	.436758	.1808061	.0324737	.370438	.503078	.1033	.7795
ExtConPkTqNorm	Normal	17	1.281353	.5213672	.1264501	1.013291	1.549415	.4480	2.1245
	Valgus	14	1.244378	.4238226	.1132714	.999670	1.489086	.6017	1.8479
	Total	31	1.264655	.4723985	.0848453	1.091378	1.437932	.4480	2.1245
ExtEccPkTqNorm	Normal	17	.806239	.4145150	.1005347	.593115	1.019363	.2827	1.8846
	Valgus	14	.730298	.3345796	.0894202	.537118	.923479	.1661	1.3295
	Total	31	.771943	.3763289	.0675907	.633905	.909982	.1661	1.8846
AbConPkTqNorm	Normal	17	.438308	.1749017	.0424199	.348381	.528234	.0543	.6667
	Valgus	14	.326804	.1420702	.0379699	.244775	.408833	.1269	.5404
	Total	31	.387951	.1680570	.0301839	.326308	.449595	.0543	.6667
AbEccPkTqNorm	Normal	17	.508174	.2769238	.0671639	.365793	.650555	.0497	1.0031
	Valgus	14	.501200	.2165695	.0578806	.376157	.626244	.1142	.8333
	Total	31	.505025	.2474600	.0444451	.414256	.595794	.0497	1.0031

ANOVA

		Sum of Squares	df	Mean Square	F	Sig.
ERConPkTqNorm	Between Groups	.015	1	.015	1.309	.262
	Within Groups	.323	29	.011		
	Total	.337	30			
EREccPkTqNorm	Between Groups	.018	1	.018	.514	.479
	Within Groups	1.011	29	.035		
	Total	1.029	30			
IRConPkTqNorm	Between Groups	.000	1	.000	.000	.985
	Within Groups	1.895	29	.065		
	Total	1.895	30			
IREccPkTqNorm	Between Groups	.002	1	.002	.060	.808
	Within Groups	.979	29	.034		
	Total	.981	30			
ExtConPkTqNorm	Between Groups	.010	1	.010	.046	.833
	Within Groups	6.684	29	.230		
	Total	6.695	30			
ExtEccPkTqNorm	Between Groups	.044	1	.044	.305	.585
	Within Groups	4.204	29	.145		
	Total	4.249	30			
AbConPkTqNorm	Between Groups	.095	1	.095	3.682	.065
	Within Groups	.752	29	.026		
	Total	.847	30			
AbEccPkTqNorm	Between Groups	.000	1	.000	.006	.939
	Within Groups	1.837	29	.063		
	Total	1.837	30			

Oneway

Descriptives

	N	Mean	Std. Deviation	Std. Error	95% Confidence Interval for Mean		Minimum	Maximum
					Lower Bound	Upper Bound		
ERConTmPkT Normal	17	3.352941	1.6349808	.3965411	2.512312	4.193571	1.3333	6.3333
Valgus	14	2.976200	1.7805440	.4758704	1.948144	4.004256	1.0000	8.0000
Total	31	3.182800	1.6839899	.3024535	2.565107	3.800493	1.0000	8.0000
EREccTmPkT Normal	16	3.229169	1.2395829	.3098957	2.568642	3.889696	1.0000	6.0000
Valgus	14	3.119050	1.8332602	.4899594	2.060557	4.177543	1.3333	6.6667
Total	30	3.177780	1.5180521	.2771571	2.610930	3.744630	1.0000	6.6667
IRConTmPkT Normal	17	2.294112	1.1110650	.2694729	1.722855	2.865369	1.0000	4.3333
Valgus	14	3.214279	1.1810949	.3156609	2.532335	3.896222	1.0000	5.3333
Total	31	2.709671	1.2163714	.2184668	2.263502	3.155840	1.0000	5.3333
IREccTmPkT Normal	15	2.755553	1.3479478	.3480386	2.009085	3.502022	1.0000	4.6667
Valgus	14	2.857143	2.1630790	.5781072	1.608218	4.106068	1.0000	8.6667
Total	29	2.804597	1.7559904	.3260792	2.136654	3.472540	1.0000	8.6667
ExtConTmPkT Normal	17	4.490194	1.7642959	.4279046	3.583077	5.397311	2.3333	9.3333
Valgus	14	4.904764	1.5821169	.4228385	3.991277	5.818251	2.0000	6.6667
Total	31	4.677419	1.6699658	.2999347	4.064871	5.289968	2.0000	9.3333
ExtEccTmPkT Normal	17	2.509806	1.9759540	.4792392	1.493864	3.525748	1.0000	7.0000
Valgus	13	3.307685	1.9122394	.5303598	2.152130	4.463239	1.0000	8.3333
Total	30	2.855553	1.9567736	.3572563	2.124882	3.586225	1.0000	8.3333
AbConTmPkT Normal	17	1.176471	.5787576	.1403693	.878901	1.474040	1.0000	3.3333
Valgus	14	1.619050	.7937605	.2121414	1.160746	2.077354	1.0000	3.0000
Total	31	1.376345	.7083765	.1272282	1.116511	1.636180	1.0000	3.3333
AbEccTmPkT Normal	17	2.725488	1.4153696	.3432775	1.997772	3.453204	1.0000	6.0000
Valgus	14	3.357157	1.2227414	.3267914	2.651167	4.063147	1.0000	5.0000
Total	31	3.010758	1.3484797	.2421941	2.516132	3.505384	1.0000	6.0000

ANOVA

		Sum of Squares	df	Mean Square	F	Sig.
ERConTmPkTq	Between Groups	1.090	1	1.090	.376	.544
	Within Groups	83.985	29	2.896		
	Total	85.075	30			
EREccTmPkTq	Between Groups	.091	1	.091	.038	.847
	Within Groups	66.739	28	2.384		
	Total	66.830	29			
IRConTmPkTq	Between Groups	6.501	1	6.501	4.976	.034
	Within Groups	37.886	29	1.306		
	Total	44.387	30			
IREccTmPkTq	Between Groups	.075	1	.075	.023	.880
	Within Groups	86.263	27	3.195		
	Total	86.338	28			
ExtConTmPkTq	Between Groups	1.320	1	1.320	.465	.501
	Within Groups	82.344	29	2.839		
	Total	83.664	30			
ExtEccTmPkTq	Between Groups	4.690	1	4.690	1.235	.276
	Within Groups	106.350	28	3.798		
	Total	111.040	29			
AbConTmPkTq	Between Groups	1.504	1	1.504	3.218	.083
	Within Groups	13.550	29	.467		
	Total	15.054	30			
AbEccTmPkTq	Between Groups	3.063	1	3.063	1.725	.199
	Within Groups	51.489	29	1.775		
	Total	54.552	30			

General Linear Model

Within-Subjects Factors

Measure: MEASURE_1

phase	Dependent Variable
1	GMaxDes AmpNorm
2	GMaxAsc AmpNorm

Between-Subjects Factors

	Value Label	N
Group 1.0000	Normal	16
2.0000	Valgus	13

Descriptive Statistics

	Group	Mean	Std. Deviation	N
GMaxDesAmpNorm	Normal	10.984009	12.5250030	16
	Valgus	11.370089	5.6519150	13
	Total	11.157080	9.8878250	29
GMaxAscAmpNorm	Normal	11.521487	8.0315153	16
	Valgus	11.395781	5.7234116	13
	Total	11.465136	6.9713218	29

Multivariate Tests

Effect		Value	F	Hypothesis df	Error df	Sig.	Partial Eta Squared	Noncent. Parameter	Observed Power ^a
phase	Pillai's Trace	.001	.024 ^b	1.000	27.000	.877	.001	.024	.053
	Wilks' Lambda	.999	.024 ^b	1.000	27.000	.877	.001	.024	.053
	Hotelling's Trace	.001	.024 ^b	1.000	27.000	.877	.001	.024	.053
	Roy's Largest Root	.001	.024 ^b	1.000	27.000	.877	.001	.024	.053
phase * Group	Pillai's Trace	.001	.020 ^b	1.000	27.000	.888	.001	.020	.052
	Wilks' Lambda	.999	.020 ^b	1.000	27.000	.888	.001	.020	.052
	Hotelling's Trace	.001	.020 ^b	1.000	27.000	.888	.001	.020	.052
	Roy's Largest Root	.001	.020 ^b	1.000	27.000	.888	.001	.020	.052

a. Computed using alpha = .05

b. Exact statistic

c.

Design: Intercept+Group

Within Subjects Design: phase

Mauchly's Test of Sphericity

Measure: MEASURE_1

		Approx. Chi-Square	df	Sig.	Epsilon ^a		
					Greenhouse-Geisser	Huynh-Feldt	Lower-bound
Within Subjects Effects	Mauchly's W						
phase	1.000	.000	0	.	1.000	1.000	1.000

Tests the null hypothesis that the error covariance matrix of the orthonormalized transformed dependent variables is proportional to an identity matrix.

a. May be used to adjust the degrees of freedom for the averaged tests of significance. Corrected tests are displayed in the Tests of Within-Subjects Effects table.

b.

Design: Intercept+Group

Within Subjects Design: phase

Tests of Within-Subjects Effects

Measure: MEASURE_1

Source		Type III Sum of Squares	df	Mean Square	F	Sig.	Partial Eta Squared	Noncent. Parameter	Observed Power ^a
phase	Sphericity Assumption	1.137	1	1.137	.024	.877	.001	.024	.053
	Greenhouse-Geisser	1.137	1.000	1.137	.024	.877	.001	.024	.053
	Huynh-Feldt	1.137	1.000	1.137	.024	.877	.001	.024	.053
	Lower-bound	1.137	1.000	1.137	.024	.877	.001	.024	.053
phase * Group	Sphericity Assumption	.939	1	.939	.020	.888	.001	.020	.052
	Greenhouse-Geisser	.939	1.000	.939	.020	.888	.001	.020	.052
	Huynh-Feldt	.939	1.000	.939	.020	.888	.001	.020	.052
	Lower-bound	.939	1.000	.939	.020	.888	.001	.020	.052
Error(phase)	Sphericity Assumption	1257.722	27	46.582					
	Greenhouse-Geisser	1257.722	27.000	46.582					
	Huynh-Feldt	1257.722	27.000	46.582					
	Lower-bound	1257.722	27.000	46.582					

a. Computed using alpha = .05

Tests of Within-Subjects Contrasts

Measure: MEASURE_1

Source	phase	Type III Sum of Squares	df	Mean Square	F	Sig.	Partial Eta Squared	Noncent. Parameter	Observed Power ^a
phase	Linear	1.137	1	1.137	.024	.877	.001	.024	.053
phase * Group	Linear	.939	1	.939	.020	.888	.001	.020	.052
Error(phase)	Linear	1257.722	27	46.582					

a. Computed using alpha = .05

Tests of Between-Subjects Effects

Measure: MEASURE_1

Transformed Variable: Average

Source	Type III Sum of Squares	df	Mean Square	F	Sig.	Partial Eta Squared	Noncent. Parameter	Observed Power ^a
Intercept	7349.919	1	7349.919	69.890	.000	.721	69.890	1.000
Group	.243	1	.243	.002	.962	.000	.002	.050
Error	2839.411	27	105.163					

a. Computed using alpha = .05

Estimated Marginal Means

1. Grand Mean

Measure: MEASURE_1

Mean	Std. Error	95% Confidence Interval	
		Lower Bound	Upper Bound
11.318	1.354	8.540	14.096

2. Group

Measure: MEASURE_1

Group	Mean	Std. Error	95% Confidence Interval	
			Lower Bound	Upper Bound
Normal	11.253	1.813	7.533	14.972
Valgus	11.383	2.011	7.256	15.509

3. phase

Measure: MEASURE_1

phase	Mean	Std. Error	95% Confidence Interval	
			Lower Bound	Upper Bound
1	11.177	1.880	7.321	15.034
2	11.459	1.325	8.739	14.178

4. Group * phase

Measure: MEASURE_1

Group	phase	Mean	Std. Error	95% Confidence Interval	
				Lower Bound	Upper Bound
Normal	1	10.984	2.517	5.820	16.148
	2	11.521	1.775	7.880	15.163
Valgus	1	11.370	2.792	5.641	17.099
	2	11.396	1.969	7.356	15.436

General Linear Model

Within-Subjects Factors

Measure: MEASURE_1

phase	Dependent Variable
1	GMedDes AmpNorm
2	GMedAsc AmpNorm

Between-Subjects Factors

	Value Label	N
Group 1.0000	Normal	16
2.0000	Valgus	13

Descriptive Statistics

	Group	Mean	Std. Deviation	N
GMedDesAmpNorm	Normal	14.985443	10.0408630	16
	Valgus	19.024160	9.7204132	13
	Total	16.795903	9.9339103	29
GMedAscAmpNorm	Normal	15.499042	11.0197818	16
	Valgus	19.028615	9.7469128	13
	Total	17.081264	10.4384500	29

Multivariate Tests

Effect		Value	F	Hypothesis df	Error df	Sig.	Partial Eta Squared	Noncent. Parameter	Observed Power ^a
phase	Pillai's Trace	.004	.101 ^b	1.000	27.000	.753	.004	.101	.061
	Wilks' Lambda	.996	.101 ^b	1.000	27.000	.753	.004	.101	.061
	Hotelling's Trace	.004	.101 ^b	1.000	27.000	.753	.004	.101	.061
	Roy's Largest Root	.004	.101 ^b	1.000	27.000	.753	.004	.101	.061
phase * Group	Pillai's Trace	.004	.098 ^b	1.000	27.000	.757	.004	.098	.060
	Wilks' Lambda	.996	.098 ^b	1.000	27.000	.757	.004	.098	.060
	Hotelling's Trace	.004	.098 ^b	1.000	27.000	.757	.004	.098	.060
	Roy's Largest Root	.004	.098 ^b	1.000	27.000	.757	.004	.098	.060

a. Computed using alpha = .05

b. Exact statistic

c.

Design: Intercept+Group

Within Subjects Design: phase

Mauchly's Test of Sphericity

Measure: MEASURE_1

Within Subjects Effect	Mauchly's W	Approx. Chi-Square	df	Sig.	Epsilon ^a		
					Greenhouse-Geisser	Huynh-Feldt	Lower-bound
phase	1.000	.000	0	.	1.000	1.000	1.000

Tests the null hypothesis that the error covariance matrix of the orthonormalized transformed dependent variables is proportional to an identity matrix.

a. May be used to adjust the degrees of freedom for the averaged tests of significance. Corrected tests are displayed in the Tests of Within-Subjects Effects table.

b.

Design: Intercept+Group

Within Subjects Design: phase

Tests of Within-Subjects Effects

Measure: MEASURE_1

Source		Type III Sum of Squares	df	Mean Square	F	Sig.	Partial Eta Squared	Noncent. Parameter	Observed Power ^a
phase	Sphericity Assumed	.962	1	.962	.101	.753	.004	.101	.061
	Greenhouse-Geisser	.962	1.000	.962	.101	.753	.004	.101	.061
	Huynh-Feldt	.962	1.000	.962	.101	.753	.004	.101	.061
	Lower-bound	.962	1.000	.962	.101	.753	.004	.101	.061
phase * Group	Sphericity Assumed	.930	1	.930	.098	.757	.004	.098	.060
	Greenhouse-Geisser	.930	1.000	.930	.098	.757	.004	.098	.060
	Huynh-Feldt	.930	1.000	.930	.098	.757	.004	.098	.060
	Lower-bound	.930	1.000	.930	.098	.757	.004	.098	.060
Error(phase)	Sphericity Assumed	257.040	27	9.520					
	Greenhouse-Geisser	257.040	27.000	9.520					
	Huynh-Feldt	257.040	27.000	9.520					
	Lower-bound	257.040	27.000	9.520					

a. Computed using alpha = .05

Tests of Within-Subjects Contrasts

Measure: MEASURE_1

Source	phase	Type III Sum of Squares	df	Mean Square	F	Sig.	Partial Eta Squared	Noncent. Parameter	Observed Power ^a
phase	Linear	.962	1	.962	.101	.753	.004	.101	.061
phase * Group	Linear	.930	1	.930	.098	.757	.004	.098	.060
Error(phase)	Linear	257.040	27	9.520					

a. Computed using alpha = .05

Tests of Between-Subjects Effects

Measure: MEASURE_1

Transformed Variable: Average

Source	Type III Sum of Squares	df	Mean Square	F	Sig.	Partial Eta Squared	Noncent. Parameter	Observed Power ^a
Intercept	16845.691	1	16845.691	85.005	.000	.759	85.005	1.000
Group	205.414	1	205.414	1.037	.318	.037	1.037	.166
Error	5350.643	27	198.172					

a. Computed using alpha = .05

Estimated Marginal Means

1. Grand Mean

Measure: MEASURE_1

Mean	Std. Error	95% Confidence Interval	
		Lower Bound	Upper Bound
17.134	1.858	13.321	20.947

2. Group

Measure: MEASURE_1

Group	Mean	Std. Error	95% Confidence Interval	
			Lower Bound	Upper Bound
Normal	15.242	2.489	10.136	20.348
Valgus	19.026	2.761	13.362	24.691

3. phase

Measure: MEASURE_1

phase	Mean	Std. Error	95% Confidence Interval	
			Lower Bound	Upper Bound
1	17.005	1.848	13.213	20.797
2	17.264	1.955	13.252	21.276

4. Group * phase

Measure: MEASURE_1

Group	phase	Mean	Std. Error	95% Confidence Interval	
				Lower Bound	Upper Bound
Normal	1	14.985	2.475	9.907	20.064
	2	15.499	2.618	10.127	20.871
Valgus	1	19.024	2.746	13.390	24.658
	2	19.029	2.905	13.069	24.989

General Linear Model

Within-Subjects Factors

Measure: MEASURE_1

phase	Dependent Variable
1	AddDes AmpNorm
2	AddAsc AmpNorm

Between-Subjects Factors

	Value Label	N
Group 1.0000	Normal	16
2.0000	Valgus	13

Descriptive Statistics

	Group	Mean	Std. Deviation	N
AddDesAmpNorm	Normal	29.680102	39.0152292	16
	Valgus	18.848112	14.2616026	13
	Total	24.824383	30.5398423	29
AddAscAmpNorm	Normal	18.024013	7.6244966	16
	Valgus	35.598841	33.7813601	13
	Total	25.902384	24.4814316	29

Multivariate Tests

Effect		Value	F	Hypothesis df	Error df	Sig.	Partial Eta Squared	Noncent. Parameter	Observed Power ^a
phase	Pillai's Trace	.007	.196 ^b	1.000	27.000	.661	.007	.196	.071
	Wilks' Lambda	.993	.196 ^b	1.000	27.000	.661	.007	.196	.071
	Hotelling's Trace	.007	.196 ^b	1.000	27.000	.661	.007	.196	.071
	Roy's Largest Root	.007	.196 ^b	1.000	27.000	.661	.007	.196	.071
phase * Group	Pillai's Trace	.184	6.095 ^b	1.000	27.000	.020	.184	6.095	.663
	Wilks' Lambda	.816	6.095 ^b	1.000	27.000	.020	.184	6.095	.663
	Hotelling's Trace	.226	6.095 ^b	1.000	27.000	.020	.184	6.095	.663
	Roy's Largest Root	.226	6.095 ^b	1.000	27.000	.020	.184	6.095	.663

a. Computed using alpha = .05

b. Exact statistic

c.

Design: Intercept+Group

Within Subjects Design: phase

Mauchly's Test of Sphericity

Measure: MEASURE_1

		Approx. Chi-Square	df	Sig.	Epsilon ^a		
					Greenhouse-Geisser	Huynh-Feldt	Lower-bound
Within Subjects Effects	Mauchly's W						
phase	1.000	.000	0	.	1.000	1.000	1.000

Tests the null hypothesis that the error covariance matrix of the orthonormalized transformed dependent variables is proportional to an identity matrix.

a. May be used to adjust the degrees of freedom for the averaged tests of significance. Corrected tests are displayed in the Tests of Within-Subjects Effects table.

b.

Design: Intercept+Group

Within Subjects Design: phase

Tests of Within-Subjects Effects

Measure: MEASURE_1

Source		Type III Sum of Squares	df	Mean Square	F	Sig.	Partial Eta Squared	Noncent. Parameter	Observed Power ^a
phase	Sphericity Assumed	93.081	1	93.081	.196	.661	.007	.196	.071
	Greenhouse-Geisser	93.081	1.000	93.081	.196	.661	.007	.196	.071
	Huynh-Feldt	93.081	1.000	93.081	.196	.661	.007	.196	.071
	Lower-bound	93.081	1.000	93.081	.196	.661	.007	.196	.071
phase * Group	Sphericity Assumed	2893.880	1	2893.880	6.095	.020	.184	6.095	.663
	Greenhouse-Geisser	2893.880	1.000	2893.880	6.095	.020	.184	6.095	.663
	Huynh-Feldt	2893.880	1.000	2893.880	6.095	.020	.184	6.095	.663
	Lower-bound	2893.880	1.000	2893.880	6.095	.020	.184	6.095	.663
Error(phase)	Sphericity Assumed	12819.595	27	474.800					
	Greenhouse-Geisser	12819.595	27.000	474.800					
	Huynh-Feldt	12819.595	27.000	474.800					
	Lower-bound	12819.595	27.000	474.800					

a. Computed using alpha = .05

Tests of Within-Subjects Contrasts

Measure: MEASURE_1

Source	phase	Type III Sum of Squares	df	Mean Square	F	Sig.	Partial Eta Squared	Noncent. Parameter	Observed Power ^a
phase	Linear	93.081	1	93.081	.196	.661	.007	.196	.071
phase * Group	Linear	2893.880	1	2893.880	6.095	.020	.184	6.095	.663
Error(phase)	Linear	12819.595	27	474.800					

a. Computed using alpha = .05

Tests of Between-Subjects Effects

Measure: MEASURE_1

Transformed Variable: Average

Source	Type III Sum of Squares	df	Mean Square	F	Sig.	Partial Eta Squared	Noncent. Parameter	Observed Power ^a
Intercept	37421.498	1	37421.498	37.394	.000	.581	37.394	1.000
Group	163.050	1	163.050	.163	.690	.006	.163	.068
Error	27020.104	27	1000.745					

a. Computed using alpha = .05

Estimated Marginal Means

1. Grand Mean

Measure: MEASURE_1

Mean	Std. Error	95% Confidence Interval	
		Lower Bound	Upper Bound
25.538	4.176	16.969	34.107

2. Group

Measure: MEASURE_1

Group	Mean	Std. Error	95% Confidence Interval	
			Lower Bound	Upper Bound
Normal	23.852	5.592	12.378	35.326
Valgus	27.223	6.204	14.494	39.953

3. phase

Measure: MEASURE_1

phase	Mean	Std. Error	95% Confidence Interval	
			Lower Bound	Upper Bound
1	24.264	5.712	12.544	35.984
2	26.811	4.336	17.914	35.709

4. Group * phase

Measure: MEASURE_1

Group	phase	Mean	Std. Error	95% Confidence Interval	
				Lower Bound	Upper Bound
Normal	1	29.680	7.649	13.986	45.374
	2	18.024	5.807	6.110	29.938
Valgus	1	18.848	8.486	1.437	36.259
	2	35.599	6.442	22.381	48.817

General Linear Model

Within-Subjects Factors

Measure: MEASURE_1

board	phase	Dependent Variable
1	1	GMaxDes AmpNorm
	2	GMaxAsc AmpNorm
2	1	GMaxDes Ampboard Norm
	2	GMaxAsc Ampboard Norm

Group = Valgus

Multivariate Tests^{b,c}

Effect		Value	F	Hypothesis df	Error df	Sig.
board	Pillai's Trace	.011	.134 ^a	1.000	12.000	.721
	Wilks' Lambda	.989	.134 ^a	1.000	12.000	.721
	Hotelling's Trace	.011	.134 ^a	1.000	12.000	.721
	Roy's Largest Root	.011	.134 ^a	1.000	12.000	.721
phase	Pillai's Trace	.113	1.525 ^a	1.000	12.000	.241
	Wilks' Lambda	.887	1.525 ^a	1.000	12.000	.241
	Hotelling's Trace	.127	1.525 ^a	1.000	12.000	.241
	Roy's Largest Root	.127	1.525 ^a	1.000	12.000	.241
board * phase	Pillai's Trace	.106	1.430 ^a	1.000	12.000	.255
	Wilks' Lambda	.894	1.430 ^a	1.000	12.000	.255
	Hotelling's Trace	.119	1.430 ^a	1.000	12.000	.255
	Roy's Largest Root	.119	1.430 ^a	1.000	12.000	.255

a. Exact statistic

b.

Design: Intercept

Within Subjects Design: board+phase+board*phase

c. Group = Valgus

Mauchly's Test of Sphericity^c

Measure: MEASURE_1

Within Subjects Effect	Mauchly's W	Approx. Chi-Square	df	Sig.	Epsilon ^a		
					Greenhouse-Geisser	Huynh-Feldt	Lower-bound
board	1.000	.000	0	.	1.000	1.000	1.000
phase	1.000	.000	0	.	1.000	1.000	1.000
board * phase	1.000	.000	0	.	1.000	1.000	1.000

Tests the null hypothesis that the error covariance matrix of the orthonormalized transformed dependent variables is proportional to an identity matrix.

a. May be used to adjust the degrees of freedom for the averaged tests of significance. Corrected tests are displayed in the Tests of Within-Subjects Effects table.

b.

Design: Intercept

Within Subjects Design: board+phase+board*phase

c. Group = Valgus

Tests of Within-Subjects Effects

Measure: MEASURE_1

Source		Type III Sum of Squares	df	Mean Square	F	Sig.
board	Sphericity Assumed	2.550	1	2.550	.134	.721
	Greenhouse-Geisser	2.550	1.000	2.550	.134	.721
	Huynh-Feldt	2.550	1.000	2.550	.134	.721
	Lower-bound	2.550	1.000	2.550	.134	.721
Error(board)	Sphericity Assumed	228.463	12	19.039		
	Greenhouse-Geisser	228.463	12.000	19.039		
	Huynh-Feldt	228.463	12.000	19.039		
	Lower-bound	228.463	12.000	19.039		
phase	Sphericity Assumed	20.663	1	20.663	1.525	.241
	Greenhouse-Geisser	20.663	1.000	20.663	1.525	.241
	Huynh-Feldt	20.663	1.000	20.663	1.525	.241
	Lower-bound	20.663	1.000	20.663	1.525	.241
Error(phase)	Sphericity Assumed	162.606	12	13.550		
	Greenhouse-Geisser	162.606	12.000	13.550		
	Huynh-Feldt	162.606	12.000	13.550		
	Lower-bound	162.606	12.000	13.550		
board * phase	Sphericity Assumed	19.829	1	19.829	1.430	.255
	Greenhouse-Geisser	19.829	1.000	19.829	1.430	.255
	Huynh-Feldt	19.829	1.000	19.829	1.430	.255
	Lower-bound	19.829	1.000	19.829	1.430	.255
Error(board*phase)	Sphericity Assumed	166.412	12	13.868		
	Greenhouse-Geisser	166.412	12.000	13.868		
	Huynh-Feldt	166.412	12.000	13.868		
	Lower-bound	166.412	12.000	13.868		

a. Group = Valgus

Tests of Within-Subjects Contrasts

Measure: MEASURE_1

Source	board	phase	Type III Sum of Squares	df	Mean Square	F	Sig.
board	Linear		2.550	1	2.550	.134	.721
Error(board)	Linear		228.463	12	19.039		
phase		Linear	20.663	1	20.663	1.525	.241
Error(phase)		Linear	162.606	12	13.550		
board * phase	Linear	Linear	19.829	1	19.829	1.430	.255
Error(board*phase)	Linear	Linear	166.412	12	13.868		

a. Group = Valgus

Tests of Between-Subjects Effects^a

Measure: MEASURE_1

Transformed Variable: Average

Source	Type III Sum of Squares	df	Mean Square	F	Sig.
Intercept	7002.385	1	7002.385	59.407	.000
Error	1414.464	12	117.872		

a. Group = Valgus

Estimated Marginal Means

1. Grand Mean^a

Measure: MEASURE_1

Mean	Std. Error	95% Confidence Interval	
		Lower Bound	Upper Bound
11.604	1.506	8.324	14.885

a. Group = Valgus

2. board^a

Measure: MEASURE_1

board	Mean	Std. Error	95% Confidence Interval	
			Lower Bound	Upper Bound
1	11.383	1.577	7.946	14.820
2	11.826	1.667	8.194	15.457

a. Group = Valgus

3. phase^a

Measure: MEASURE_1

phase	Mean	Std. Error	95% Confidence Interval	
			Lower Bound	Upper Bound
1	10.974	1.450	7.815	14.133
2	12.235	1.718	8.491	15.978

a. Group = Valgus

4. board * phase^a

Measure: MEASURE_1

board	phase	Mean	Std. Error	95% Confidence Interval	
				Lower Bound	Upper Bound
1	1	11.370	1.568	7.955	14.786
	2	11.396	1.587	7.937	14.854
2	1	10.578	1.403	7.520	13.636
	2	13.074	2.386	7.875	18.273

a. Group = Valgus

General Linear Model

Within-Subjects Factors

Measure: MEASURE_1

board	phase	Dependent Variable
1	1	GMedDes AmpNorm
	2	GMedAsc AmpNorm
2	1	GMedDes Ampboard Norm
	2	GMedAsc Ampboard Norm

Group = Valgus

Descriptive Statistics^a

	Mean	Std. Deviation	N
GMedDesAmpNorm	19.024160	9.7204132	13
GMedAscAmpNorm	19.028615	9.7469128	13
GMedDesAmpboard Norm	18.142014	9.4341316	13
GMedAscAmpboardNorm	20.258858	11.8490756	13

a. Group = Valgus

Multivariate Tests^a

Effect		Value	F	Hypothesis df	Error df	Sig.	Partial Eta Squared	Noncent. Parameter	Observed Power ^a
board	Pillai's Trace	.002	.022 ^b	1.000	12.000	.885	.002	.022	.052
	Wilks' Lambda	.998	.022 ^b	1.000	12.000	.885	.002	.022	.052
	Hotelling's Trace	.002	.022 ^b	1.000	12.000	.885	.002	.022	.052
	Roy's Largest Root	.002	.022 ^b	1.000	12.000	.885	.002	.022	.052
phase	Pillai's Trace	.145	2.034 ^b	1.000	12.000	.179	.145	2.034	.259
	Wilks' Lambda	.855	2.034 ^b	1.000	12.000	.179	.145	2.034	.259
	Hotelling's Trace	.169	2.034 ^b	1.000	12.000	.179	.145	2.034	.259
	Roy's Largest Root	.169	2.034 ^b	1.000	12.000	.179	.145	2.034	.259
board * phase	Pillai's Trace	.140	1.955 ^b	1.000	12.000	.187	.140	1.955	.251
	Wilks' Lambda	.860	1.955 ^b	1.000	12.000	.187	.140	1.955	.251
	Hotelling's Trace	.163	1.955 ^b	1.000	12.000	.187	.140	1.955	.251
	Roy's Largest Root	.163	1.955 ^b	1.000	12.000	.187	.140	1.955	.251

a. Computed using alpha = .05

b. Exact statistic

c.

Design: Intercept

Within Subjects Design: board+phase+board*phase

d. Group = Valgus

Mauchly's Test of Sphericity^b

Measure: MEASURE_1

Within Subjects Effects	Mauchly's W	Approx. Chi-Square	df	Sig.	Epsilon ^a		
					Greenhouse-Geisser	Huynh-Feldt	Lower-bound
board	1.000	.000	0	.	1.000	1.000	1.000
phase	1.000	.000	0	.	1.000	1.000	1.000
board * phase	1.000	.000	0	.	1.000	1.000	1.000

Tests the null hypothesis that the error covariance matrix of the orthonormalized transformed dependent variables is proportional to an identity matrix.

a. May be used to adjust the degrees of freedom for the averaged tests of significance. Corrected tests are in the Tests of Within-Subjects Effects table.

b.

Design: Intercept

Within Subjects Design: board+phase+board*phase

c. Group = Valgus

Tests of Within-Subjects Effects

Measure: MEASURE_1

Source	Type III Sum of Squares	df	Mean Square	F	Sig.	Partial Eta Squared	Noncent. Parameter	Observed Power ^a
board	Sphericity Assumed	.394	1	.394	.022	.885	.002	.052
	Greenhouse-Geisser	.394	1.000	.394	.022	.885	.002	.052
	Huynh-Feldt	.394	1.000	.394	.022	.885	.002	.052
	Lower-bound	.394	1.000	.394	.022	.885	.002	.052
Error(board)	Sphericity Assumed	215.282	12	17.940				
	Greenhouse-Geisser	215.282	12.000	17.940				
	Huynh-Feldt	215.282	12.000	17.940				
	Lower-bound	215.282	12.000	17.940				
phase	Sphericity Assumed	14.625	1	14.625	2.034	.179	.145	.259
	Greenhouse-Geisser	14.625	1.000	14.625	2.034	.179	.145	.259
	Huynh-Feldt	14.625	1.000	14.625	2.034	.179	.145	.259
	Lower-bound	14.625	1.000	14.625	2.034	.179	.145	.259
Error(phase)	Sphericity Assumed	86.289	12	7.191				
	Greenhouse-Geisser	86.289	12.000	7.191				
	Huynh-Feldt	86.289	12.000	7.191				
	Lower-bound	86.289	12.000	7.191				
board * phase	Sphericity Assumed	14.502	1	14.502	1.955	.187	.140	.251
	Greenhouse-Geisser	14.502	1.000	14.502	1.955	.187	.140	.251
	Huynh-Feldt	14.502	1.000	14.502	1.955	.187	.140	.251
	Lower-bound	14.502	1.000	14.502	1.955	.187	.140	.251
Error(board*phase)	Sphericity Assumed	89.006	12	7.417				
	Greenhouse-Geisser	89.006	12.000	7.417				
	Huynh-Feldt	89.006	12.000	7.417				
	Lower-bound	89.006	12.000	7.417				

a. Computed using alpha = .05

b. Group = Valgus

Tests of Within-Subjects Contrasts

Measure: MEASURE_1

Source	board phase	Type III Sum of Squares	df	Mean Square	F	Sig.	Partial Eta Squared	Noncent. Parameter	Observed Power ^a
board	Linear	.394	1	.394	.022	.885	.002	.022	.052
Error(board)	Linear	215.282	12	17.940					
phase	Linear	14.625	1	14.625	2.034	.179	.145	2.034	.259
Error(phase)	Linear	86.289	12	7.191					
board * phase	Linear Linear	14.502	1	14.502	1.955	.187	.140	1.955	.251
Error(board*ph	Linear Linear	89.006	12	7.417					

a. Computed using alpha = .05

b. Group = Valgus

Tests of Between-Subjects Effects

Measure: MEASURE_1

Transformed Variable: Average

Source	Type III Sum of Squares	df	Mean Square	F	Sig.	Partial Eta Squared	Noncent. Parameter	Observed Power ^a
Intercept	18996.771	1	18996.771	49.171	.000	.804	49.171	1.000
Error	4636.130	12	386.344					

a. Computed using alpha = .05

b. Group = Valgus

Estimated Marginal Means

1. Grand Mean^a

Measure: MEASURE_1

Mean	Std. Error	95% Confidence Interval	
		Lower Bound	Upper Bound
19.113	2.726	13.175	25.052

a. Group = Valgus

2. board^a

Measure: MEASURE_1

board	Mean	Std. Error	95% Confidence Interval	
			Lower Bound	Upper Bound
1	19.026	2.699	13.145	24.908
2	19.200	2.874	12.938	25.463

a. Group = Valgus

3. phase^a

Measure: MEASURE_1

phase	Mean	Std. Error	95% Confidence Interval	
			Lower Bound	Upper Bound
1	18.583	2.642	12.826	24.340
2	19.644	2.855	13.422	25.865

a. Group = Valgus

4. board * phase^a

Measure: MEASURE_1

board	phase	Mean	Std. Error	95% Confidence Interval	
				Lower Bound	Upper Bound
1	1	19.024	2.696	13.150	24.898
	2	19.029	2.703	13.139	24.919
2	1	18.142	2.617	12.441	23.843
	2	20.259	3.286	13.099	27.419

a. Group = Valgus

General Linear Model

Within-Subjects Factors

Measure: MEASURE_1

board	phase	Dependent Variable
1	1	AddDes AmpNorm
	2	AddAsc AmpNorm
2	1	AddDes Ampboard Norm
	2	AddAsc Ampboard Norm

Group = Valgus

Descriptive Statistics^a

	Mean	Std. Deviation	N
AddDesAmpNorm	18.848112	14.2616026	13
AddAscAmpNorm	35.598841	33.7813601	13
AddDesAmpboardNorm	18.090004	14.0836079	13
AddAscAmpboardNorm	28.494032	20.9464810	13

a. Group = Valgus

Multivariate Tests^a

Effect		Value	F	Hypothesis df	Error df	Sig.	Partial Eta Squared	Noncent. Parameter	Observed Power ^a
board	Pillai's Trace	.166	2.390 ^b	1.000	12.000	.148	.166	2.390	.296
	Wilks' Lambda	.834	2.390 ^b	1.000	12.000	.148	.166	2.390	.296
	Hotelling's Trace	.199	2.390 ^b	1.000	12.000	.148	.166	2.390	.296
	Roy's Largest Root	.199	2.390 ^b	1.000	12.000	.148	.166	2.390	.296
phase	Pillai's Trace	.477	10.951 ^b	1.000	12.000	.006	.477	10.951	.859
	Wilks' Lambda	.523	10.951 ^b	1.000	12.000	.006	.477	10.951	.859
	Hotelling's Trace	.913	10.951 ^b	1.000	12.000	.006	.477	10.951	.859
	Roy's Largest Root	.913	10.951 ^b	1.000	12.000	.006	.477	10.951	.859
board * phase	Pillai's Trace	.124	1.700 ^b	1.000	12.000	.217	.124	1.700	.225
	Wilks' Lambda	.876	1.700 ^b	1.000	12.000	.217	.124	1.700	.225
	Hotelling's Trace	.142	1.700 ^b	1.000	12.000	.217	.124	1.700	.225
	Roy's Largest Root	.142	1.700 ^b	1.000	12.000	.217	.124	1.700	.225

a. Computed using alpha = .05

b. Exact statistic

c.

Design: Intercept

Within Subjects Design: board+phase+board*phase

d. Group = Valgus

Mauchly's Test of Sphericity^a

Measure: MEASURE_1

		Approx. Chi-Square	df	Sig.	Epsilon ^a		
					Greenhouse-Geisser	Huynh-Feldt	Lower-bound
board	Mauchly's W	.000	0	.	1.000	1.000	1.000
phase		.000	0	.	1.000	1.000	1.000
board * phase		.000	0	.	1.000	1.000	1.000

Tests the null hypothesis that the error covariance matrix of the orthonormalized transformed dependent variables is proportional to an identity matrix.

a. May be used to adjust the degrees of freedom for the averaged tests of significance. Corrected tests are in the Tests of Within-Subjects Effects table.

b.

Design: Intercept

Within Subjects Design: board+phase+board*phase

c. Group = Valgus

Tests of Within-Subjects Effects^a

Measure: MEASURE_1

Source		Type III Sum of Squares	df	Mean Square	F	Sig.	Partial Eta Squared	Noncent. Parameter	Observed Power ^b
board	Sphericity Assumed	200.933	1	200.933	2.390	.148	.166	2.390	.296
	Greenhouse-Geisser	200.933	1.000	200.933	2.390	.148	.166	2.390	.296
	Huynh-Feldt	200.933	1.000	200.933	2.390	.148	.166	2.390	.296
	Lower-bound	200.933	1.000	200.933	2.390	.148	.166	2.390	.296
Error(board)	Sphericity Assumed	1008.777	12	84.065					
	Greenhouse-Geisser	1008.777	12.000	84.065					
	Huynh-Feldt	1008.777	12.000	84.065					
	Lower-bound	1008.777	12.000	84.065					
phase	Sphericity Assumed	2396.488	1	2396.488	10.951	.006	.477	10.951	.859
	Greenhouse-Geisser	2396.488	1.000	2396.488	10.951	.006	.477	10.951	.859
	Huynh-Feldt	2396.488	1.000	2396.488	10.951	.006	.477	10.951	.859
	Lower-bound	2396.488	1.000	2396.488	10.951	.006	.477	10.951	.859
Error(phase)	Sphericity Assumed	2626.032	12	218.836					
	Greenhouse-Geisser	2626.032	12.000	218.836					
	Huynh-Feldt	2626.032	12.000	218.836					
	Lower-bound	2626.032	12.000	218.836					
board * phase	Sphericity Assumed	130.912	1	130.912	1.700	.217	.124	1.700	.225
	Greenhouse-Geisser	130.912	1.000	130.912	1.700	.217	.124	1.700	.225
	Huynh-Feldt	130.912	1.000	130.912	1.700	.217	.124	1.700	.225
	Lower-bound	130.912	1.000	130.912	1.700	.217	.124	1.700	.225
Error(board*phase)	Sphericity Assumed	924.341	12	77.028					
	Greenhouse-Geisser	924.341	12.000	77.028					
	Huynh-Feldt	924.341	12.000	77.028					
	Lower-bound	924.341	12.000	77.028					

a. Computed using alpha = .05

b. Group = Valgus

Tests of Within-Subjects Contrasts^a

Measure: MEASURE_1

Source		Type III Sum of Squares	df	Mean Square	F	Sig.	Partial Eta Squared	Noncent. Parameter	Observed Power ^b
board	Linear	200.933	1	200.933	2.390	.148	.166	2.390	.296
Error(board)	Linear	1008.777	12	84.065					
phase	Linear	2396.488	1	2396.488	10.951	.006	.477	10.951	.859
Error(phase)	Linear	2626.032	12	218.836					
board * phase	Linear Linear	130.912	1	130.912	1.700	.217	.124	1.700	.225
Error(board*phase)	Linear Linear	924.341	12	77.028					

a. Computed using alpha = .05

b. Group = Valgus

Tests of Between-Subjects Effects^b

Measure: MEASURE_1

Transformed Variable: Average

Source	Type III Sum of Squares	df	Mean Square	F	Sig.	Partial Eta Squared	Noncent. Parameter	Observed Power ^a
Intercept	33173.597	1	33173.597	20.711	.001	.633	20.711	.986
Error	19220.971	12	1601.748					

a. Computed using alpha = .05

b. Group = Valgus

Estimated Marginal Means

1. Grand Mean^a

Measure: MEASURE_1

Mean	Std. Error	95% Confidence Interval	
		Lower Bound	Upper Bound
25.258	5.550	13.165	37.350

a. Group = Valgus

2. board^a

Measure: MEASURE_1

board	Mean	Std. Error	95% Confidence Interval	
			Lower Bound	Upper Bound
1	27.223	6.541	12.972	41.475
2	23.292	4.696	13.060	33.524

a. Group = Valgus

3. phase^a

Measure: MEASURE_1

phase	Mean	Std. Error	95% Confidence Interval	
			Lower Bound	Upper Bound
1	18.469	3.921	9.926	27.012
2	32.046	7.393	15.939	48.153

a. Group = Valgus

4. board * phase^a

Measure: MEASURE_1

board	phase	Mean	Std. Error	95% Confidence Interval	
				Lower Bound	Upper Bound
1	1	18.848	3.955	10.230	27.466
	2	35.599	9.369	15.185	56.013
2	1	18.090	3.906	9.579	26.601
	2	28.494	5.810	15.836	41.152

a. Group = Valgus

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