# Introduction

Glucocorticoids mediate a number of metabolic processes regarding muscle whether produced endogenously or taken exogenously in the form of prescription medications. The estimated prevalence of oral glucocorticoids usage in the United States is 1.2% for a variety of health concerns including asthma, chronic obstructive pulmonary disease, COPD, and a range of autoimmune disorders (2). Elevated levels of glucocorticoids within the human body have shown to cause skeletal muscle atrophy. This muscle atrophy stems from an upregulation of atrogenes, or genes that cause muscle atrophy.

Skeletal muscle is vital for most everyday basic functions and maintenance of health. However, many factors including poor nutrition, lack of exercise, and a myriad of diseases can lead to loss of skeletal muscle. This loss can be measured, in most cases, by evaluating muscle size via mass or cross-sectional area and muscle function. Muscle mass can be quantified as the volume of muscle within the body while strength refers to contractile ability of a given muscle. In both humans and animals, muscle mass can be measured quite accurately using Magnetic Resonance Imaging while strength is assessed dependent on the specific muscle group. Grip strength in human and mice is simple and easy way to measure limb strength. Advantageously using animal models, we are able to evaluate muscle strength using direct muscle and nerve stimulation (1). In order to properly study skeletal muscle atrophy, we must quantify size and strength as well as the factors that induce such atrophy.

Glucocorticoids are steroid hormones that function through a Glucocorticoid Receptor (GR) or NR3C1, which can act directly on gene transcription . (Patel et al. 2014) Exogenous glucocorticoid consumption has been shown to induce muscle atrophy through increased muscle proteolysis and inhibition of protein synthesis in lean mice (3). Muscle proteolysis may be caused by glucocorticoid induction of atrogenes, including MuRF1 and Atrogin-1 through the FOXO pathway. (Kang et al., 2017) The inhibition of protein synthesis is believed to be directed by inhibition of the mTOR pathway associated with muscle growth by glucocorticoids (3).

Preliminary research conducted at the Bridges lab shows the both lean and obese mice develop significant reductions in lean mass, muscle mass, and grip strength when treated with dexamethasone. Treated-mice had temporary induction of muscle degradation transcripts including Fbxo32 and Trim63, (Atrogin-1 and Murf1 respectively) and their upstream regulator Foxo3.

# Methods

## Animals

C57BL/6J adult male mice were purchased from The Jackson Laboratory at 9 weeks of age and randomly caged. All animals were on a light/dark cycle of 12:12 hours and housed at 22°C. At 10 weeks of age, mice were placed on a high-fat diet (HFD; 45% fat from lard, 35% carbohydrate mix of starch, maltodextrin, and sucrose, and 20% protein from casein) or a normal chow diet (NCD; 13% fat, 57% carbohydrate, and 30% protein) for 12 weeks. At 22 weeks, mice were either treated with with vehicle (water) or 1 mg/kg/d of dexamethasone dissolved in their drinking water. All mice were provided with access to food and their respective waters *ad libitum* throughout the study. Food and both waters were measured weekly to determine the concentration of dexamethasone consumed per cage and volumes were averaged per mouse per cage.

## Grip Strength

C57BL/6J mice were tested during treatment with vehicle (water) or dexamethasone using a grip strength meter with a Chatillon digital force gauge (AMETEK). These mice were treated for six weeks with their respective waters. A grip strength baseline was established per mouse and all measurements were reported in torque(N). Mice were placed on a grid attached to the meter and once all four paws had contact with the grid, the mice were slowly pulled backwards by the tail until they left the grid. Each mouse was tested five times and given approximately 10 seconds rest in between each test. Final measurements for grip strength were assessed by taking the average of the five trials and reported as average peak torque (N).

## Contractile Measurements

Mice were anesthetized using XXX in a XXX . All contractile properties were measured for gastrocnemius muscles *in situ*. After the mouse was anesthetized, the right gastroc muscle was carefully isolated and a 4–0 silk suture was tied around the distal tendon. After the tendon was secured, the tendon was cut so the hindlimb could be secured at the knee to a fixed post. Animals were placed on a temperature-controlled platform with continual drip of saline over the gastroc at approximately 37°C to keep with muscle warm and moist. The distal tendon of the gastroc muscle was tied to the lever arm of a servomotor (model 305B, Aurora Scientific).

In order to measure force generated at the nerve, a bipolar platinum wire electrode was used to stimulate the muscle at the tibial nerve. The voltage of the electrode pulses was incrementally adjusted to find maximum isometric twitch and the muscle length was altered to find the optimal length (Lo). Optimal length is the length of the muscle in which the maximal twitch force was obtained.

Once Lo was found, gastrocs were kept at that length (Lo) and the frequency of pulses was increased in increments of 300-ms to obtain maximum isometric tetanic force (Po).

In order to measure force generated at the muscle, an electrode cuff was placed around the mid-belly of gastroc for muscle stimulation. The process was then repeated as done for the nerve.  
After all force measurements, both gastrocnemius and quadricep muscles were dissected, weighed, and snap frozen in liquid nitrogen. Mice were sacrificed under anesthesia and muscles were stored at -80℃.

## Histology

Quadriceps were collected and snap frozen in 2methyl-butane. Quadricep samples were sectioned at -20 degree celsius with a thickness of 10um through the mid-belly and mounted on SuperFrost glass slides. For analysis of fiber cross-sectional area (CSA), fibers were identified by hematoxylin and eosin (H&E staining) and the area of 200 individual fibers were averaged per mouse quadricep. For analysis of fiber-type composition, fibers were stained using β-Nicotinamide adenine dinucleotide, reduced disodium salt hydrate or NADH-NBT staining. Sections were imaged using 20x objective of an EVOS XL digital inverted microscope and measured or quantified using ImageJ.

## Cell Culture

C2C12 cells were cultured in 10% Fetal Bovine Serum (FBS), Dulbecco's Modification of Eagle's Medium (DMEM; 4.5 g/L D- glucose; Fisher Scientific; catalog #11965118) with penicillin, streptomycin and glutamine (PSG). Cells were split at approximately 75% confluency and differentiated using DMEM, 1x PSG with 2% Horse serum until myotubes were obtained. Media was replenished as needed until myotube differentiation was complete around one week. Myotubes were treated with 250nm dexamethasone for either 2, 4, 8, 12, or 24 hours or left untreated. All cells will kept in a 5% CO2 regulated incubator at 37 °C. After treatment, cells were homogenized in TRIZol using a TissueLyser II (Qiagen) and prepared for RNA extraction using a PureLink RNA kit (catalog no. 12183025; Life Technologies).

## RNA

Cells and tissues were lysed in TRIzol using a TissueLyser II (Qiagen) and RNA was extracted using a PureLink RNA kit (catalog no. 12183025; Life Technologies). Complementary DNA (cDNA) was synthesized using the High Capacity cDNA Reverse Transcription Kit without RNAse inhibitor (catalog no. 4368813; Life Technologies). Quantitative Real-Time Polymerase Chain reaction (qPCR) was performed using a QuantStudio 5 (Thermo Fisher Scientific) with primers, complementary DNA, and Power SYBR Green PCR Master Mix (catalog no. 4368708; Life Technologies) per recommended guidelines. Messenger RNA (mRNA) expression levels were normalized to *XXX* and analyzed.

LIST OF PRIMERS

|  |  |  |
| --- | --- | --- |
| **Gene** | **Forward 5’-3’ Sequence** | **Reverse 5’-3’ Sequence** |
| *Fbxo32* | CTTCTCGACTGCCATCCTGG | GTTCTTTTGGGCGATGCCAC |
| *Trim63* | GAGGGCCATTGACTTTGGGA | TTTACCCTCTGTGGTCACGC |
| *Foxo1* | AGTGGATGGTGAAGAGCGTG | GAAGGGACAGATTGTGGCGA |
| *Foxo3* | AAACGGCTCACTTTGTCCCA | ATTCTGAACGCGCATGAAGC |

## Data Analysis and Statistics

All results are represented as mean ± SEM. Two-Way ANOVA analyses were performed to test for significance and determine interactions between diet and dexamethasone treatment, if applicable. Pairwise testing was performed to check for normality and equal of variances. If Shapiro-Wilks passed, a Levene’s tests was performed and followed by either a Welch’s or Student’s *t*-test. Any P-value under p=0.05 was considered significant. All statistical tests were conducted using R software.

## Protein- ARE WE INCLUDING IDK

Tissues were lysed in RIPA buffer (50 mM Tris, pH 7.4, 0.25% sodium deoxycholate, 1% NP40, 150 mM sodium chloride, 1 mM EDTA, 100 uM sodium orthovanadate, 5 mM sodium fluoride, 10 mM sodium pyrophosphate and 1x protease inhibitor) and 1x protease inhibitor using a TissueLyser II (Qiagen) and centrifuged at 14,000 rpm for 10 minutes at 4°C. Loading buffer was added to the lysates and heated at approximately 85°C. Proteins were separated by SDS-PAGE gel electrophoresis (Life Technologies) and gels were transferred onto nitrocellulose overnight. Nitrocellulose blots were stained with Revert Total Protein Stain (catalog no. 926-11011; Li-Cor Biosciences) for total protein. Anti-mouse and anti-rabbit fluorescent-conjugated antibodies (Invitrogen) were used to detect secondary antibodies.

Antibody list...

A Odyssey CLx image scanner (Li-Cor Biosciences) was used to visualize the images. All calculation used to quantify protein concentrations were performed using version 5.2 Image Studio software (Li-Cor Biosciences) and first normalized to Revert Total Protein stained blots.

# Results

## Greater Losses in Grip Strength in Obese-Dexamethasone Mice

As a test to assess the effect of glucocorticoids on muscle strength, we treated lean and obese male mice with dexamethasone for five weeks and measured four-paw grip strength. Dexamethasone treatment resulted in synergistic reductions in grip strength in both lean and obese mice when compared to their counterparts (Figure 1a-b). This result is consistent with studies in adult humans in which elevated salivary cortisol had a significantly higher risk of loss of grip strength than their peers(Peeters et al., 2008). In addition, obese dexamethasone-treated mice had greater overall losses in grip strength when compared to the lean animals. For mean grip strength, we saw a -0.4% reduction in lean animals (P=0.007) but a -0.9% reduction in grip strength for obese animals (P=3.62e-05).

## Reductions in Strength are Related to Smaller CSA

In order to expand upon these results, we measured the force generated by stimulation nerve and gastrocnemius muscle. Though we are specifically interested in muscle strength, stimulating the XX nerve and assessing the resulting muscle force allows us to evaluate the potential effect of neuropathy. In NCD animals, force generated by nerve stimulation was reduced X% when treated with dexamethasone. However in HFD animals, force generated by nerve stimulation was reduced X% when treated with dexamethasone. There was a significant interaction between diet and treatment, P=.009 (Figure 1c). These results are consistent with direct muscle stimulation where a electrode cuff is placed around the mid-belly of the gastrocnemius muscle in order to stimulate the whole muscle. In NCD animals, force generated by direct muscle stimulation was reduced X% when treated with dexamethasone. While in HFD animals, the force generated by direct muscle stimulation was reduced X% when treated with dexamethasone. There was a significant interaction between diet and treatment P=.024(Figure 1d).

In order to examine whether changes in muscle strength were correlated to changes in or declined in proportion to muscle size, we plotted a regression of force (mN) versus whole-muscle cross-sectional area (CSA). Mice appeared to follow a linear pattern in which animals with a larger CSA generated proportionally larger force for both nerve and direct muscle stimulation (Figure 1e-f). Equation of the line idk idk idk

Reduction is strength are significantly correlated to smaller whole-muscle CSA, P=.003. Report the adjusted R^2 for the model

Can explain percent of variance- of strength by knowing CSA

## Enhanced Muscle Loss in Obese Mice

To determine whether obesity interacts with long-term glucocorticoid treatment to induce losses in muscle mass, we treated lean and obese male mice with dexamethasone for five weeks. Dexamethasone caused a reduction in total body lean mass in both lean and obese mice. Consistent with losses in strength, obese-dexamethasone treated mice had greater losses in lean mass as assessed by echoMRI(Figure 2a). This loss in lean mass is consistent with previously reported effects of glucocorticoids on muscle atrophy (Pleasure, Walsh, and Engel 1970). This loss was also observed in specific hindlimb muscles. At sacrifice, the NCD animals quadricep and tricep surae weights were smaller by X percent in the dexamethasone treated. While in HFD animals, quadricep and tricep surae weights were smaller by X percent in the dexamethasone treated (Figure 2b).

We then evaluated short-term dexamethasone treated animals by placing male mice on vehicle or dexamethasone for two weeks. These are the same animals and muscles used for all isometric force testing. The obese-dexamethasone treated animals had the smallest gastrocnemius weights and whole-muscle CSA (Figure 2c-d). At sacrifice, the NCD animals gastrocnemius weights were smaller by X percent in the dexamethasone treated. While in HFD animals, gastrocnemius weights were smaller by X percent in the dexamethasone treated. There was a significant interaction between diet and treatment, P=.021.

## Obesity and Dexamethasone Treatment Resulted in Muscle Fiber Degradation

In order to assess changes in muscle size at the fiber-level, we sectioned the 5-week dexamethasone-treated mice quadriceps at the mid-belly and H&E stained the sections(Figure 2e). The NCD animal’s muscle fibers were smaller by X percent in the dexamethasone treated and in HFD animals muscle fibers were smaller by X percent in the dexamethasone treated. There was a significant interaction between diet and treatment, P=.001 (Figure 2f).

## Dexamethasone did not Induce Changes Fiber-Type Composition

In order to assess any changes in the ratio of oxidative versus non-oxidative fiber-types, we stained muscle sections and quantified the muscle fibers based upon their oxidative capacity. These are the same section from the quadriceps used ….. Mice skeletal muscle is made up Type I, Type IIa, Type IIb, and Type IIx fibers. Oxidative fibers or Type I fibers stained the darkest (Figure 2g). We found no significant change in the ratio of oxidative to total fibers in the mice quadriceps in lean or obese. (Figure 2h) This result may be different in other types of muscles within the hindlimb because quadriceps have fairly high oxidative capacity due their mitochondrial content (Jacobs et al., 2012).

## Dexamethasone Reduced Type II Fiber CSA

Though we did not see changes in composition of fiber types, we wanted to investigate changes in fiber-type size. In order to evaluate fiber-type specific atrophy, we labeled fiber types based on their stained color and measured their CSA. Type I fibers stained darkest, Type IIb stained intermediate, and Type IIa stained the lightest (Figure 2g). Dexamethasone-treatment reduced Type IIa fibers CSA in lean and obese mice by X% and X% respectively. Dexamethasone-treatment also reduced Type IIb fibers CSA in lean and obese by X% and X% respectively. As for Type I Fibers, dexamethasone treatments only reduced fiber CSA in NCD animals. There was no significant effect of dexamethasone on Type I fibers in obese mice (Figure 2i). This outcome is consistent with previous data shown in which plantares muscles from mice treated with dexamethasone for 13 days showed significant atrophy in Type IIb and Type IIa and not in Type I fibers (Rouleau et al., 1987).

## Short-term Dexamethasone-Treatment Induced Muscle Degradation Transcripts as seen *in vitro*

It is well established that dexamethasone treatment induces expression of muscle atrophy-related genes (Sandri et al., 2004; Waddell et al., 2008; Kang et al., 2017). We chose to treat C2C12 myotubes with dexamethasone overtime in order to assess the expression of Foxo1, Foxo3, and well-established atrogenes, Murf1 and Atrogin-1 (encoded by *Trim63* and *Fbxo32* respectively) *in vitro.* Relative expression of all genes were elevated after 2 hours of treatment with dexamethasone (Figure 3a). Only Murf1 and Atrogin-1 reached peak expression at the end of the time-course at 24 hours of dexamethasone.

To evaluate the molecular effects of dexamethasone *in vivo*, we treated lean and obese mice with dexamethasone and evaluated atrogene expression. After one week of dexamethasone treatment, we observed a greater induction of both *Foxo3* and the atrogenes, *Trim63* and *Fbxo32*, in obese mice as compared to their lean counterparts (Figure 3b). The expression of *Trim63, Fbxo32,* and *Foxo3* was elevated by approximately X%, X% and X% respectively in obese mice than their lean counterparts*.* However we need not see an increase in Foxo1 or Ncr31, glucocorticoid receptor.

### Obese Dexamethasone-Treated Mice are Profoundly Insulin Resistant

Since we have highlighted that obesity can enhance skeletal muscle atrophy generated by glucocorticoids, we wanted to take a look at the role of insulin and insulin sensitivity in this muscle atrophy model. Insulin is central to glucose-uptake in skeletal muscle, where the majority of all glucose uptake occurs within the body. Both dexamethasone and obesity are known to cause insulin resistance (references) In order to measure insulin sensitivity during dexamethasone-treatment, we treated lean and obese mice with dexamethasone and measured their fasting blood glucose(FBG). In lean animals, there was no significant change in FBG between treatment groups however there was a X% increase in obese animals given dexamethasone (Figure 4a). There was a synergistic effect of obesity and dexamethasone on fasted blood glucose, P=.033.

In order to evaluate whether the dexamethasone-treated animals were insulin resistant, we treated the same lean and obese mice with insulin and monitored with blood glucose overtime. Insulin doses were given dependent on their lean mass composition to account for their difference in size between chow-fed and high-fat diet fed mice. In both NCD and HFD animals, dexamethasone induced insulin resistance (Figure 4b). This is consistent with prior research in which insulin-stimulated glucose-uptake is significantly reduced in lean and obese dexamethasone-treated muscles when compared to controls.

# Discussion

Dexamethasone-treatment reduced muscle strength and size in diet-induced obese mice. This is supported by that fact that muscle weakness is a common side effect of exogenous glucocorticoid consumption as well as continually elevated levels of endogenous hormones By evaluating the effects of glucocorticoids on muscles within obese animals, this may give us insights into reviewing muscle atrophy in humans afflicted with obesity. This research could be particularly important because those suffering with obesity are shown to have reduced muscle function (Maffiuletti et al. 2007; Blimkie, Sale, and Bar-Or 1990; Hulens et al. 2001; Zoico et al. 2004). Therefore we may to able to more effectively care for individuals on glucocorticoids suffering from obesity by better understanding this animal model.

Not only are glucocorticoids shown to induce changes in type II muscle fibers, there are a variety of conditions and lifestyle factors such a bed-rest that also lead to other significant myofiber changes. For instance, disuse atrophy as a result of denervation or immobilization of a limb, is shown to reduce type I fiber size and muscle mass (Nicks et. al., 1989, Rouleau et al., 1987, Trappe et al., 2004). These physiological changes are consistent with our results in the way that we saw reductions in whole muscle mass, CSA and muscle fiber CSA, but no decrease in total number of fibers. Disuse atrophy has even been shown to generate fiber plasticity where type I fibers or oxidative fibers take on characteristics of fast-twitch, non-oxidative fibers or change completely to type II fibers. (Debré et al., 2016, Stevenson et al., 2003) Even though we saw no change in fiber composition when treating animals with dexamethasone, other studies have shown that dexamethasone reduces both that quantity and size of Type 2 fibers in muscles in mice and rats (Ma et al., 2003). This difference could be due to quantity of dexamethasone provided to the animals; our dose which is consistent with human under constant stress or taking a high prescription dose.

It is also important to note that glucocorticoids induce muscle atrophy in a muscle specific manner. Researchers often test on mouse hindlimb muscle because they are fairly large and accessible load bearing muscles. Specifically, type II fibers are more prone to the effect of glucocorticoids (Falduto et al., 1990, Kelly and Goldspink, 1982, Livingstone et al., 1981), so it is possible that muscles with higher concentrations of type II fibers may be more vulnerable to atrophy. For example, rats treated with dexamethasone for two weeks had no significant reduction in mean fiber CSA in their solei but had significant reduction in their plantares muscles, which have higher type II fiber composition (Rouleau et al., 1987). We evaluated fibers within the mouse quadriceps. It also possible that we would see more dramatic changes in the forces generated by other muscles, than the gastrocnemius we tested, that have higher in type II fiber content. Studies have shown that increased losses in contraction force of muscle wither higher percentages of type IIa fibers (Mänttäri et al., 2005)

Though it is well known that glucocorticoids induce muscle atrophy, we do not understand how obesity interacts with glucocorticoids to exaggerate muscle loss. One possible theory is that glucocorticoids increase sensitivity to pro-inflammatory cytokines and separately glucocorticoids and excess adiposity exacerbate levels of these cytokines circulating in the body and act upon skeletal muscle. For example leptin-receptor deficient obese mouse mice were found to have a lowered threshold for release of pro-inflammatory cytokines in the brain but an increase in pro-inflammatory cytokines such as interleukin 1beta and tumor necrosis factor alpha. (Dey at al. 2014) It is possible that glucocorticoids and proinflammatory factors could coregulate genes associated with atrophy that are unknown.

# References

1. **Larkin LM**, **Davis CS**, **Sims-Robinson C**, **Kostrominova TY**, **Remmen H V.**, **Richardson A**, **Feldman EL**, **Brooks S V.** Skeletal muscle weakness due to deficiency of CuZn-superoxide dismutase is associated with loss of functional innervation. *AJP Regul Integr Comp Physiol* 301: R1400–R1407, 2011.

2. **Overman RA**, **Yeh JY**, **Deal CL**. Prevalence of oral glucocorticoid usage in the United States: A general population perspective. *Arthritis Care Res* 65: 294–298, 2013.

3. **Schakman O**, **Kalista S**, **Barbé C**, **Loumaye a**, **Thissen JPP**. Glucocorticoid-induced skeletal muscle atrophy. *Int J Biochem Cell Biol* 45: 2163–2172, 2013.

# Figure Legends

**Figure 1. Obese-Dexamethasone Treated Mice Lost Significant Muscle Strength**

Grip strength (N) in lean (a) and obese (b) male mice over the course of six weeks of dexamethasone treatment. N=4-8 per group. Data collected by Innocence Harvey. \*=Significance identified by Student’s T-Test. Force (mN) generated by nerve stimulation (c) and by direct muscle gastrocnemius stimulation (d) in lean and obese mice treated with dexamethasone for 15-21 days. Force plotted by whole gastrocnemius CSA (e-f). N=5-8 per group. \*=Diet-Treatment interaction identified by Two-Way ANOVA.

**Figure 2. Obese-Dexamethasone Treated Mice had Reduced Lean Mass, Muscle Weights, and Type II Fiber CSA.**

Lean mass via EchoMRI (a) and muscle weights (b) in lean and obese mice following 6 weeks of dexamethasone treatment. N=8-22 per group. Data collected by Innocence Harvey. Gastrocnemius weIghts (c) and CSA (d) in lean and obese mice treated with dexamethasone for 15-21 days. N=5-8 per group. H&E stained section of quadriceps (e) from mice treated with vehicle (water) or dexamethasone for six weeks. N=4 quadricep sections per group. Average fiber CSA (f) from 200 fibers per quadricep section. NADH-NBT stained section of quadriceps (g) from mice treated with vehicle (water) or dexamethasone for six weeks. Percent of oxidative or type I fibers to total mouse fibers (h). N=4 quadricep sections per group. Average fiber CSA by NADH-NBT staining density (i). \*=Diet-Treatment interaction identified by Two-Way ANOVA.

**Figure 3. Short-term Dexamethasone Treatment Induced Muscle Degradation Transcripts unlike Chronic or Long-Term Treatment**

Relative atrogene (*Fbxo32, Trim63, Foxo1* and *Foxo3*) expression in C2C12 myotubes treated with 250 nm dexamethasone for 2, 4, 8,12, or 24 hours or left untreated(a). After treatment, cells were homogenized and prepared for RNA extraction.

Atrogene expression in mice treated for either 72 hours, one week, or two weeks with vehicle(water) or 1mg/kg/d dexamethasone (b). RNA was extracted from the quadriceps. \*=Significance identified by Student’s T-Test and in mice treated for six weeks with vehicle (water) or dexamethasone. \*=Diet-Treatment interaction identified by Two-Way ANOVA. N=8 per group.

**Figure 4. Dexamethasone Treatment Induced Insulin Resistance** Blood glucose values taken from the tail vein in lean and obese male mice after a 6-hour fast and two weeks of dexamethasone or vehicle (water) treatment (a). \*=Diet-Treatment interaction identified by Two-Way ANOVA. N=4 mice per group. Glucose values after insulin administration at time 0, following a 6-hour fast (b). Insulin was given via intraperitoneal injection at .75g/kg lean mass for lean mice and 1.5g/kg for obese mice.N=4 mice per group.