**IS FAST FOOD SPEEDING UP THE AGING PROCESS? COMPARING THE SKELETAL MUSCLE INFLAMMATORY ENVIRONMENT IN DIET INDUCED OBESITY AND OLD RATS**

Charmi Dholakia, Christine Waters-Banker, Geoffrey A. Power, Walter Herzog

Faculty of Kinesiology, Human Performance Lab, University of Calgary

Charmi Figure for JURAcharmidholakia4@gmail.com

**INTRODUCTION**

Obesity, linked to various deleterious health problems such as heart disease and Type II diabetes, is becoming a public health concern and economic burden worldwide [1]. Loss of skeletal muscle content, or atrophy, and fat infiltration are often reported in obese individuals [2,3]. In a separate population, elderly adults also exhibit a loss of muscle mass commonly referred to as sarcopenia [4]. Similarly, both of these populations are characterized by systemic inflammation [5,6]. However, it is not yet known whether skeletal muscle inflammation is similar in obese and aging individuals, or if obesity may be an accelerated model of aging. Therefore, the purpose of this study was to compare the general cellular infiltration associated with inflammation in the tibialis anterior (TA) muscle of a diet induced obesity model, and an established aging model, against a young healthy model, in rats.

**METHODS**

Six male Sprague Dawley rats provided with a high fat/high sucrose diet for 6 months were analyzed for this study. Additionally, The TA muscles from the non-perturbed contralateral limb of 12 Fisher 344 x Brown Norway rats (old (n=6) and young (n=6)), were collected following mechanical testing. All TA muscles were harvested and flash frozen in liquid nitrogen. Four 8μm cryosections were stained with Hematoxylin and Eosin (H&E), and imaged using light microscope. Because skeletal muscle consists of resident ‘anti-inflammatory’ macrophages, stereological point counting techniques were used to count clusters of immune cells likely associated with a ‘pro-inflammatory’ response. These clusters were normalized per 100 muscle fibres. All procedures were performed according to the guidelines of the Canadian Council on Animal Care and were approved by the Animal Care Committee of the University of Calgary. A Kruskal-Wallis One-Way ANOVA was used to determine differences between groups.

**RESULTS**

Obese rats had a significantly greater number of immune cell clusters per 100 muscle fibres compared to Young rats

(p=0.027)(**Figure 1)**. No statistical differences were observed

**Figure 1.** Average immune cell clusters normalized to skeletal muscle fibre number with representative H&E histology. Black boxes indicate immune cell clusters.

between Obese and Old rats.

**DISCUSSION AND CONCLUSIONS**

The inflammatory environment within the skeletal muscle of Obese rats more closely resembled Old than the Young rats. This was surprising considering the average age difference between the Obese and Young rats at time of sacrifice was only one month (9 and 8 months, respectively), compared to Old rats at 36 months. Additionally, similar gross morphological differences were observed in the Old and Obese rats, such as inflammatory cell mediated fibre breakdown and fat infiltration, not observed in the Young tissue (**Figure 1**). The inflammatory process in skeletal muscle is crucial to the maintenance, repair, and regeneration of skeletal muscle. Dysregulation of this response may have severe consequences on skeletal muscle health, especially as young obese individuals progress into old age.

**REFERENCES**

1. Popkin & Slining. *Obes Rev.* **2**:11-20,2013.
2. Hilton TN, et al. *Phys Ther.* **88**:1336-1344, 2008.
3. Sishi B, et al. *Exp Physiol.* **96**:179-193, 2011.
4. Power GA, et al. *Age.* **36**:1377-1388, 2014.
5. Roubenoff R. *Curr Opin Clin Nutr Metab Care.* **6**:295-299, 2003.
6. Fontana L, Eagon JC, Trujillo ME, Scherer PE, Klein S. *Diabetes.* **56**:1010-1013, 2007.