Fiber Composition of the Grasscutter (Thryonomys swinderianus, Temminck 1827) Thigh Muscle: An Enzyme-histochemical Study

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Abstract

The aim of this study was to describe de fiber composition in the thigh muscles of grass cutter (Thryonomys swinderianus, Temminck 1827). Ten 4 to 6-month-old (3 to 4 kg) male grasscutter were used in this study. Eleven skeletal muscles of the thigh [M. biceps femoris (BF), M. rectus femoris (RF), M. vastus lateralis (VL), M. vastus medialis (VM), M. tensor fasciae latae (TFL), M. semitendinosus (ST), M. semimembranosus (SM), M. semimembranosus accessorius (SMA), M. Sartorius (SRT), M. pectineus (PCT), M. adductor magnus (AM)] were collected after animals euthanasia and examined by light microscopy. Three muscle fiber types (I, IIB and IIA) were found in these muscles using enzyme histochemical techniques [myosine adenosine triphosphatase (ATPase) and nicotinamide adenine dinucleotide tetrazolium reductase (NADH-TR)]. Ten of these eleven muscles are composed by 89% to 100% of fast contracting fibers (types IIA and IIB), while the SMA was almost exclusively formed by slow contracting fibers.

Keywords

Skeletal muscle; Muscle fiber types; Histochemistry; Grasscutter

Introduction

The grasscutter (Thryonomys swinderianus, Temminck 1827) is a hystricomorph rodent whose breeding in close captivity grows increasingly in West and Central Africa for both

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food speculation and wildlife resource management [1-5]. In fact, grasscutter meat is highly
valued by African game consumers, whether urban or rural for its organoleptic qualities.

The meat corresponds to striated skeletal muscles of meat animals, delicatessen and game
meat. Because of their contractile properties, they can engage anatomical structures to which
they are attached to, and are as such active motion organs.

Structurally, vertebrate striated skeletal muscle is mainly composed of fiber muscles
arranged in clusters through the connective tissue. In the conjunctiva frame we can find
neurovascular structures and clusters of adipocytes that form intramuscular adipose tissue
[6].

Within a given muscle, the fibers differ in their morphological, physiological and
biochemical characteristics. This heterogeneity of muscle tissue reflects its high degree of
specialization and is the basis of its diversity [7].

In rabbits, a phylogenetically close species to the grass cutter, there are in fact two broad
categories of muscle fibers in terms of the speed of contraction: some slow (type I fibers)
and others fast (fiber Type IIA, IIB and IIX) [8]. The energy required for the contraction of
the muscle fibers comes from the hydrolysis of adenosine di - phosphate (ADP) catalysed by
the myofibrillar ATPase. Contractile properties of muscle fibers depend on the ATPase type
carried by myosin heads [9], the main protein constituent of myofibrils. The myosin
molecule consists of two polymorphs of heavy chains (200 kDa), the actual carriers of the
ATPase activity, to which four light chains (16-30 kDa) are associated. The speed of
contraction of muscle fibers thus depends directly on their specific isoform composition of
slow myosin heavy chain (type I) and fast (type IIa, IIX and IIB) [10].

To ensure their effective functioning, the muscle fibers have permanent enzymatic
equipment for regenerating the hydrolyzed ATP during muscle contraction. ATP synthesis is
provided by the catabolism of energy substrates such as glucose and its storage form,
glycogen and fat. The stock of ATP can be reconstituted anaerobically (glycolytic) and/or
aerobic (oxidative). The measurement of the activity of certain enzymes [Succinate
Dehydrogenase (SDH) and Cytochrome oxidase nicotinamide adenine dinucleotide-
tetrazolium reductase (NADH-TR)] involved in either of these two channels allows the
assessment of their respective importance and thus to distinguish the glycolytic metabolism
fibers from the oxydo-glycolytic or oxidative ones.

The purpose of this study is to present the composition type of muscle fibers of grasscutter
thigh (*Thryonomys swinderianus*, Temminck, 1827) using histochemical techniques.

**Materials and Methods**

**Ethics consideration**

The study was approved by the Ethics Committee of Ecole Inter-Etats des Sciences et
Médecine Vétérinaires (EISMV) of Dakar (Senegal).
**Animal source**

Ten 4 to 6-month-old (3 to 4 kg) male grasscutters were used in this study. The animals were purchased from different grass cutter farms in the peri-urban area of Abidjan (Côte d’Ivoire). They were transferred into standard grasscutter breeding cages at the Department of Natural Sciences, Nangui Abrogua University, Abidjan, Côte d’Ivoire and fed with grass and commercial feed for a while before sacrifice. Water was given *ad libitum* during the period.

**Experimental design**

Sample collection: Each animal was weighed alive and sacrificed after anaesthesia with a mixture of equal volumes of xylazin hydrochlorate (Rompun ND) and ketamine hydrochlorate (Imalgen 1000 ND), at 0.1 ml/kg of body weight.

Eleven muscles [M. biceps femoris (BF), M. rectus femoris (RF), M. vastus lateralis (VL), M. vastus medialis (VM), M. tensor fasciae latae (TFL), M. semitendinosus (ST), M. semimembranosus (SM), M. semimembranosus accessorius (SMA), M. sartorius (SRT), M. pectineus (PCT), M. adductor magnus (AM)] of the thigh were removed 15 minutes after the sacrifice (Table 1).

**Histochemical processing**

A transverse section (about 0.5 cm) was obtained from the muscle midbelly and frozen for 30 s at −80°C in isopentane previously cooled with liquid nitrogen. Sections at 12 μm were cut using a cryostat and then treated with histochemical techniques. Nicotinamide adenine tetrazolium-reductase (NADH-TR) reaction according to Dubowitz [11] to characterise fiber metabolism and myofibrillar, adenosine triphosphatase (ATPase) reaction after acid pre-incubation (pH 4.35 and 4.6) and alkaline pre-incubation (pH 10.4) according to Guth and Samaha [9] to detect ATPase activity within fibers.

**Morphometric analysis**

The different types of fibers were determined according to the classification of Brooke and Kaiser [12].

Morphometric parameter was assessed using a semi-automatic image analysis system (NIS – Elements AR 3,1 image analyser, Nikon Laboratory Imaging, Prague, Czech Republic). Thus, the relative proportion of the different fibers types (I, IIA and IIB) in thigh muscles was determined from an average count of 300 fibers in 6 random fields. Determination of the proportion of different muscle fiber types were performed on sections of frozen muscles treated with acid ATPase (pH 4.35) [13-15].

**Statistical analysis**

Conventional statistical procedures have been used to calculate mean and standard deviation (S.D.). The coefficient of variation for observational bias (repeatability) was about 3%.

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Results

Morphology

Most of the muscles studied were characterized by a mosaic pattern of polygonally shaped fibers with the exception of M. semimembranosus accessorius, which exhibited an exclusively slow-twitch fibers pattern. We found no evidence of fiber type grouping or marked regionalisation of distinct fiber types in all the other muscles studied (Figure 1).

Fiber type

Data on the composition of the fiber type in the muscles of the grasscutter thigh are shown in Table 1.

The reaction to the myofibrillar ATPase acid (pH 4.35) revealed three types of fibers (I, IIA and IIB) in most of the muscles of the grasscutter thigh (Figure 1). Thus, in all muscles examined, M. semimembranosus accessorius is the only homogeneous fiber containing only type I fibers. Mm. rectus femoris and sartorius are almost exclusively composed of type IIB fibers. The other muscles of the grasscutter thigh (M. adductor magnus, M. biceps femoris, M. pectineus, M. semimembranosus, M. semitendinosus, M. tensor fascia latae, M. vastus lateralis and M. vastus medialis) are heterogeneous composed of the three fiber types I, IIA and IIB.

In heterogeneous muscles type IIB fibers are the most numerous (76.48% to 99%). Type IIA fibers come in second place and represent 18.88%, 8.58% and 8.04% in Mm adductor magnus, biceps femoris and vastus lateralis respectively. Finally, type I fibers are fewer in all heterogeneous muscles of the grasscutter thigh except for M. pectineus where they account for 11.78% of total fibers.

Discussion

In general, the muscles of grasscutter thigh, has a fiber composition similar to that of the pelvic limb of other rodents in particular the guinea pig (Cavia porcellus) and rat (Rattus rattus) and the rabbit (Oryctolagus cuniculus) a lagomorph which is a phylogenetically close species [10,15-21]. Indeed, the reaction to the myofibrillar ATPase revealed the existence of three types of fibers which according to the classification of Brooke and Kaiser [12] correspond to fiber types I, IIA and IIB. However, the distribution of these three types of muscle fibers varies from one to another and enables to distinguish three main types of muscles in the grasscutter thigh: (i) a homogeneous muscle exclusively composed of type I fibers (SMA), (ii) almost exclusively composed of type IIB fibers (SM and RF) muscles, (iii) and heterogeneous muscles composed of fibers type I, IIA and IIB (AM, BF, PCT, SM, ST, TFL, VL and VM).

Except for M. adductor magnus, the results of this study clearly show that the muscles of grasscutter thigh are mostly composed of rapid and glycolytic fibers (type IIB). For example, M. rectus femoris, M. Sartorius, M. vastus medialis, M. semimembranosus and M. tensor fascia latae are composed of fast glycolytic fibers with an aspect ratio of 99 ± 1%, 98.1 ± 1.43%, 97.5 ± 1.97%, 96.64 ± 2.17% and 94.02 ± 2.19% respectively. This high
proportion of fast glycolytic fibers in the thigh muscles of the grasscutter is consistent with the observations reported by some authors in the rabbit [8,15]. Furthermore, it is well established that the presence of this high proportion of type IIB fibers correlates with the means of locomotion of the rabbit, a species that moves using successive leaps because of the greater length of its hind limbs compared to its fore limbs. The grasscutter is a digitigrade rodent that is nonetheless capable of making spectacular leaps [22]. Also, type IIB fibers that make up his thigh muscles could explain this ability of this species. Among the thigh muscles studied, M. semimembranosus accessorius is entirely composed of oxidative slow-twitch fibers (type I). The M. semimembranosus accessorius in the grasscutter is a fusiform muscle housed within the M. semimembranosus with which it shares the same origin (ischiatric tuberosity) but ends on the medial side of the distal end of the femur [23]. M. semimembranosus unlike M. semimembranosus accessorius contains very little type I fibers (1.22 ± 1.14%). Our results are opposed to those reported by Bacou et al. [17] in rabbits. Indeed, these authors reported that M. semimembranosus proprius contains 100% of oxidative slow-twitch fibers (type I), while it’s dependent, the M. semimembranosus accessorius is composed of fast glycolytic fibers (type IIB) and oxidative - glycolytic (type IIA). Although the reaction to the myofibrillar ATPase is widely used for typing of muscle fibers, this method does not always manage to highlight all type II fibers [24]. This is why the use of immunohistochemistry, which the benefit lies in the fact that the isoforms may be detected by appropriate antibody, regardless of their phenotypic characteristics must complete this last method for a better typing of muscle fiber [25].

Different types observed in the thigh muscles of the grasscutter are widely represented both in mammals and in birds [13,25]. They reflect the phenomenon of “fibers regionalization” described for the first time by Gordon and Phillips in 1955 [26]. According to these authors, the distribution of fibers in skeletal muscles of mammals is “regionalized” with slow fibers located in deep region near the bone and fast fibers at the surface area (subcutaneous). This assertion is consistent with our results. Indeed, the deep muscles of the thigh as Mm. semimembranosus accessorius, pectineus, and adductor magnus are those that contain the most slow-twitch oxidative ie respectively, 100%, 11.78 ± 3.27% and 4.64 ± 2.17%. Mechanisms and functional implications of the phenomenon of regionalization remain to this day not very well known. These mechanisms seem to take place during the embryonic development of muscle fibers [27-29]. Thus, in the muscles of the pelvic limb in rats [28], cattle [27] pig and guinea pigs [29], the first generation of myotubes give rise to slow type primary fibers. They provide a subsequent frame for the formation of a second generation of myotubes, a larger number that will give fast type secondary fibers. This arrangement in concentric bundles slow and fast fibers persists during the postnatal growth of the pig [29]. However, during fetal development and during part of the postnatal growth, some secondary fibers (fast) change type and become slow and other new bundles of slow fibers appear. In addition, Condon et al. [28], explains the phenomenon of regionalization by the presence of “fast or slow gradients morphogenetic” that result in the transition from slow to fast isoform of the myosin and vice versa.

Moreover, according to Brandstetter et al. [27], innervation and muscular work influence the regionalization of fiber types acquired during embryonic life in adult muscles. Many works carried out on the neuromuscular system, studied the interactions between muscle and motor...
nerve and tried to explain the specificity of the association between neurons and fibers of the same type [30]. In this work, two different but non-exclusive hypotheses have been advanced. The first is that the characteristics of the muscle fibers are genetically different and neurons recognize each type of fiber, the second is that the fibers are genetically equipotential and their specificity is determined by the nature of the neuron that innervates them [31].

Adult skeletal striated muscles have no autonomous activity and they cannot be dissociated from motor nerves that control their activity. They form with the motoneuron, which innervates them the drive unit representing the functional entity of the muscular contraction.

According to the theory of Henneman [32], motor units formed by oxidative slow fibers are recruited for movements that require a slow speed and low contraction force (standing, walking). Motor units formed by the fast oxidative-glycolytic fibers are recruited for movements that require high speed and high force contraction (race). Finally, the motor units formed by fast glycolytic fibers are recruited for fast and powerful movements (jumping). Thus, the balance of glycolytic fibers in Mm. rectus femoris, sartorius, vastus medialis and semimembranosus observed in our study, have a functional explanation. Furthermore, in mammalian quadrupeds, chest members which are closer to the centre of gravity of the animal are mostly sustainers, while the pelvic limbs are mostly thrusters. The muscles of the thoracic limb will therefore be involved in the coordination of body balance during the execution of a movement. For this reason, they are composed by motor units formed mainly by oxidative fibers. However, the muscles of the pelvic limb involved in all movements requiring great strength (jumping, rearing, startup, race) consist of drive units formed mostly by glycolytic fibers.

In conclusion, the type of fiber composition of the grasscutter thigh muscles is directly related to the functions of the pelvic limb region muscles.

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Figure 1.
Light photomicrographs demonstrating the features of different muscles use in our study. Cross-sections from rectus femoris (A), Sartorius (B), biceps femoris (C), vastus medialis (D), adductor magnus (E), semimembranosus accessorius (F), pectineus (G) muscles stained by myofibrillar ATPase after acid preincubation (pH 4.2). These muscles are mixed and composed of type I (dark), type IIB (light) and type IIA (intermediate) fibres. Cross section...
from vastus lateralis muscle stained by NADH-Tetrazolium reductase (SO: Slow oxidative fibres; FOG: Fast oxydo-glycolytic fibres; FG: fast glycolytic fibres).
Table 1

Proportion (means ± S.D.) of type I, type IIA and type IIB, expressed as a percentage of 300 fibres counted in thigh muscle of grass cutter (*Thryonomys swinderianus*, Temminck 1827).

| Muscle | n | Type I     | Type IIA    | Type IIB    |
|--------|---|------------|-------------|-------------|
| AM     | 10| 4.64 ± 0.99| 18.88 ± 6.20| 76.48 ± 5.37|
| BF     | 10| 2.90 ± 1.35| 8.58 ± 1.18 | 88.52 ± 0.52|
| PCT    | 10| 11.78 ± 3.27| 3.72 ± 0.46 | 84.50 ± 3.57|
| RF     | 10| 1.00 ± 1.00 | 0.00 ± 0.00 | 99.00 ± 1.00|
| SRT    | 10| 1.90 ± 1.43 | 0.00 ± 0.00 | 98.10 ± 1.43|
| SM     | 10| 1.22 ± 1.14 | 2.14 ± 1.62 | 96.64 ± 2.17|
| SMA    | 10| 100 ± 0.00  | 0.00 ± 0.00 | 0.00 ± 0.00 |
| ST     | 10| 1.62 ± 0.61 | 4.68 ± 0.63 | 93.70 ± 0.70|
| TFL    | 10| 1.82 ± 1.31 | 4.16 ± 1.41 | 94.02 ± 2.19|
| VL     | 10| 5.24 ± 1.78 | 8.04 ± 3.26 | 86.72 ± 3.33|
| VM     | 10| 1.42 ± 1.41 | 1.08 ± 0.72 | 97.50 ± 1.97|