Apoptosis of Skeletal Muscle on Steroid-Induced Myopathy in Rats

Recently apoptotic cell death has been reported in differentiated skeletal muscle, where apoptosis was generally assumed not to occur. To investigate whether apoptosis may contribute to the steroid-induced myopathy, rats treated with triamcinolone acetonide (TA) for 9 days were sacrificed for detecting apoptosis by in situ end labeling (ISEL) and electron microscopy in the soleus muscles. Immunohistochemical stainings of Fas antigen and p53 protein were performed to examine whether apoptosis-related proteins were present in the myopathy. Muscle fiber necrosis and apoptotic myonuclei appeared in the soleus muscles following administration of TA, while control muscles showed no evidences for apoptosis. Fas antigen was not detected in control muscles, but expressed in the soleus muscles of steroid-induced myopathy. Some of the Fas antigen-expressing muscle fibers were positive for ISEL. p53 protein was not detected in any muscle fibers. These findings indicate that TA can induce apoptosis in differentiated skeletal muscles, and Fas antigen might be partly related to apoptotic muscle death in steroid-induced myopathy.

Key Words: Myopathy, Steroid-Induced; Apoptosis; Fas Antigens, CD95; Protein p53; Muscular Diseases

INTRODUCTION

Long-term administration of glucocorticoids in human and animals has been reported to induce steroid myopathy (1-3). Although any of the commonly available glucocorticoid preparations can cause myopathy, the fluorinated steroids, e.g., triamcinolone, betamethasone, and dexamethasone, seem more likely to produce muscle weakness. The typical findings of steroid myopathy are selective atrophy of type II muscle fibers and necrotic changes (4, 5). It has been suggested that glucocorticoid-induced mitochondrial damage can lead to muscle fiber necrosis (4, 6-8).

It is widely accepted that there are two major distinct modes of death in eukaryotic cells, i.e., necrosis and apoptosis (9-11). Apoptosis in response to a variety of injurious stimuli has been demonstrated in several organs (12-16). In the lymphoid tissue and brain, glucocorticoids are known to easily induce apoptosis (17-19). Although differentiated skeletal muscles usually undergo necrotic death in response to injury, apoptotic cellular death occurs in dystrophin-deficient muscle (20, 21) and differentiated skeletal muscle after treatment with anabolic steroids (22).

Apoptosis is stimulated or inhibited by several mediator proteins. Fas antigen, which mediates apoptosis of lymphoid cells (23, 24), has been identified in diseased muscles (25-27). p53 protein was proven to stimulate apoptosis (28, 29).

This study investigated whether apoptosis might contribute to the death of differentiated skeletal muscle after administration of glucocorticoid. Two assays for the detection of apoptosis were employed, i.e., in situ end labeling (ISEL) and electron microscopic examinations (30, 31). Immunohistochemical stainings of Fas antigen and p53 protein were performed to examine the roles of these proteins in apoptosis of skeletal muscle in steroid-induced myopathy.

MATERIALS AND METHODS

Development of steroid-induced myopathy and tissue sampling

Twenty female Sprague-Dawley rats weighing between 180-210 g used in this study were maintained under standard condition. Rats were divided into two groups. Each group was given daily intraperitoneal injection of either physiologic saline (n=5) or triamcinolone acetonide (TA) at a dose of 5 mg/kg body weight for 9 days (n=15). Ten days after the completion of the injections, the soleus muscles from both legs were taken under ether anesthesia, and mounted on cork in a manner such that the muscle fibers were perpendicular to the cork surface. The sections were quick-
frozen in isopentane cooled by liquid nitrogen. The tissues were cut in 10 μm thick serial sections in a cryostat maintained at -20°C and mounted on grooved microscopic slides (FisherBiotech, Pittsburgh, PA, U.S.A.). The frozen sections were used for histological studies, ISEL, and immunohistochemical studies. Remnant of the soleus muscles were used for electron microscopic examination.

In situ end labeling (ISEL)

Apoptotic cells were visualized using the ApopTag® Plus In Situ Apoptosis Detection Kit (Oncor, Gaithersburg, MD, U.S.A.). Briefly, frozen sections were fixed in 10% neutral buffered formalin for 10 min and postfixed in ethanol: acetic acid (2:1) for 5 min at -20°C. After washing in phosphate-buffered saline (PBS; 50 mM sodium phosphate, pH 7.4, 200 mM NaCl), sections were digested with proteinase K (20 μg/ml in PBS; Sigma, St. Louis, MO, U.S.A.) for 15 min at room temperature and washed with distilled water. Slides were then put into 3% H2O2 for 5 min to quench the endogenous peroxidase activity and washed with PBS. After adding the equilibration buffer for 1 min, terminal deoxynucleotidyl transferase (TdT) enzyme was pipetted onto the sections, which were then incubated in a humidified chamber at 37°C for 1 hr. The reaction was stopped by putting sections in a stop/wash buffer. After washing, anti-digoxigenin-peroxidase was added to the slides. Slides were washed, stained with diaminobenzidine substrate, and counterstained with hematoxylin. A specimen known to be positive for apoptotic cells was used as a positive control for staining. Substitution of TdT with distilled water was used as a negative control. Positive muscle fibers were counted in all sections.

Immunohistochemistry for Fas antigen and p53 protein

Frozen sections were fixed with the ethanol/acetic acid fixative solution for 10 min. The slides were immunostained with the avidin-biotin-peroxidase complex method (32). The endogenous peroxidase activity was blocked by incubating 1.5% H2O2 in PBS. The slides were treated in the 10% normal in PBS for 30 min in order to block charged sites on tissue surfaces and then incubated with anti-Fas antigen (titer 1:50; done U-2; MBL, Watertown, MA, U.S.A.) or anti-p53 protein (titer 1:500; done Ab-7; Oncogene, Cambridge, MA, U.S.A.) overnight at 4°C. The streptavidin-horseradish peroxidase (Research Genetics, Huntsville, Alabama, U.S.A.) detection system was then applied. After treatment with 1% avidin-biotinylated horseradish peroxidase for 1 hr at room temperature, the tissue sections were prepared for chromogen reaction with 3-aminon-9-ethyl carbazole (Biomeda, Foster, CA, U.S.A.). The sections were counterstained with hematoxylin and mounted on Crys-tal/Mount (Biomeda). Positive muscle fibers were counted in all sections.

Electron microscopy

The soleus muscles used for electron microscopy were initially kept in a slightly stretched state, via pins passed through their tendons into dental wax, while they were fixed for 1 hr in 2.5% glutaraldehyde in 0.2 M sodium cacodylate buffer, pH 7.2. Three pieces of muscle, measuring 2 × 2 × 3 mm, were then excised and left for several hours in fresh fixative. They were then postfixed in 1% osmium tetroxide for 1 hr, dehydrated in graded alcohols, cleared in propylene oxide, and embedded in epoxy resin. Sections were cut 80 nm in thickness by an ultramicrotome, collected on uncoated grids, stained with uranyl acetate and lead citrate, and examined with a JEM 1200 EX II (JEOL).

RESULTS

Gross and histologic examinations of myopathy

Control rats gained weight during the experimental periods. In contrast, rats treated with TA showed loss of body weight (average weight loss, 26%). The excised soleus muscle in TA-treated rats weighed 11% less than in control rats.

In light microscopic examinations, the muscle fibers of control rats were arranged in bundles or fascicles, surrounded by perimysial connective tissue. There was little variation in muscle fiber size or shape. Cytoplasmic staining was uniform, and small peripherally located nuclei were abundant (Fig. 1A). Variation in fiber size and shape was seen in the TA-treated soleus muscles. Atrophic fibers were noted occasionally (Fig. 1B). Some fibers had pale-staining cytoplasm. Necrotic fibers invaded by numerous macrophages were seen (Fig. 1C). A few small eosinophilic fibers with increased nuclei, indicative of atrophic change, were observed (Fig. 1D).

In situ end labeling (ISEL) and electron microscopy for apoptosis

No ISEL-positive signal was noted in the soleus muscles of control group (Fig. 2A). The TA-treated soleus muscles had 1 to 4 muscle fibers having apoptotic myonuclear (Fig. 2B). Myonuclei labeled with ISEL were found in muscle fibers, which had pale or intact staining cytoplasm (Fig. 2C and D). Necrotic fibers were negative for ISEL (Fig. 2E and F).

In electron microscopic examinations, the nuclei of control soleus muscles appeared just beneath the plasma mem-
brane. They were elongated and ovoid in shape, with the chromatin distributed towards the periphery (Fig. 3A).

In the TA-treated soleus muscles, a few apoptotic myonuclei characterized by irregularly condensed chromatin near the myonuclear membrane with intact sarcoplasm were noted (Fig. 3B). There were no discernible signs of degeneration or necrosis in fibers that contained apoptotic nuclei. Some muscle fibers contained swollen sarcoplasmic reticulum and mitochondria. Necrotic muscle fibers, with focal loss of myofilaments and swollen nuclei containing dispersed chromatin, were also observed.

Immunohistochemical staining of Fas antigen and p53 protein

Immunohistochemical staining of Fas antigen was not detected in the normal soleus muscles (Fig. 4A). Fas antigen was detected in the TA-treated soleus muscles and was mainly localized in the cytoplasm with granular pattern (Fig. 4B). Number of Fas antigen-positive muscle fibers ranged from 1 to 8 per each muscle bundle. Some of the Fas antigen-expressing muscle fibers were positive for ISEL (Fig. 4C and D). Necrotic muscle fibers invaded by...
DISCUSSION

Several of synthetic glucocorticoids have been developed and used in a wide range of medical and surgical fields. Despite their therapeutic benefits, the long-term administration of glucocorticoids in humans has often resulted in progressive muscle weakness and steroid myopathy. Characteristic biopsy findings from patients with steroid myopathy are selective atrophy of type II muscle fibers and occasional random fiber necrosis (1-3). As in human disease, steroid myopathy has been induced in a variety of experimental animals using a number of different steroids. The fluorinated steroids, e.g., triamcinolone, betamethasone, and dexamethasone, appear more likely to produce steroid myopathy (4, 5). Necrotic lesions are easily induced in the rat soleus muscle by steroid administration (4). In this experiment, steroid-induced myopathies in rats were devel-

macrophages demonstrated no expression of Fas antigen. p53 protein was not detected in any muscle fibers of the normal or TA-treated soleus muscles.

Fig. 2. In situ end labeling (ISEL) for apoptotic cells. None of myonuclei in control muscles show positive reaction (A, ×200). Positive staining in myonuclei of TA-treated soleus muscle is seen (B, ×200). Muscle fiber having pale cytoplasm (C, ×400) is positive by ISEL (D). (Fig. 2 continued next)
oped by daily intraperitoneal injections of TA for 9 days. In the steroid-induced myopathy, necrotic changes were found in frozen sections prepared for histological study and in electron microscopical examination. We thought that our experimental protocol (i.e. species, drug, duration) might be suitable for the study of this specific muscle lesion. Glucocorticoid-induced mitochondrial damage leading to muscle fiber necrosis has been suggested in both ultrastructural and biochemical studies (4, 6-8). The soleus muscle, which showed necrotic change in this experiment, consists mainly of type I fibers, being rich in mitochondrial oxidative enzymes (33).

Apoptosis is a type of cell death that has been shown to differ from necrosis by defined ultrastructural and biochem-

Fig. 2. (Continued from the previous page) In situ end labeling (ISEL) for apoptotic cells. Necrotic muscle fiber invaded by macrophages (E, H&E, ×400) is negative for ISEL (F).

Fig. 3. Electron micrograph of a nucleus of the normal soleus muscle displays condensed chromatin along the nuclear membrane. Other organelles in the sarcoplasm appear normal (A, ×4,000). Apoptotic change in the TA-treated soleus muscle shows irregularly clumped chromatin of the nucleus without sarcoplasmic degeneration (B, ×10,000).
ical features (9-11). Through the implementation of ISEL and electron microscopy, we examined the possibility that apoptosis might contribute to the death of differentiated skeletal muscles of steroid myopathy. The technique of ISEL introduced by Gavrieli et al. (30) detects fragmented DNA via TdT reactions. This technique not only labels apoptotic cells already in progress but also potentially detects any cells where DNA strand breaks occurred without morphological changes. In addition, this technique can be performed on histologic sections so that both the quantification of apoptosis and the location of apoptotic nuclei can be determined. Morphological changes in cell structure representing apoptotic cell death can be identified by electron microscopy (31).

Glucocorticoids are known to easily induce apoptosis in some tissue. Apoptosis of thymocytes (17) and the intestinal intraepithelial lymphocytes (18) has been experimentally induced by treatment with glucocorticoids. Hassan et al. have also demonstrated the exacerbation of apoptosis in the dentate gyrus of aged rats induced by dexamethasone (19).

Differentiated skeletal muscle cells of rats underwent apoptosis in response to glucocorticoid injury. Some myonu-

Fig. 4. Immunohistochemical staining of Fas antigen and ISEL. Fas antigen is not detected in control muscle (A, ×200), and is positive in the cytoplasm of the TA-treated soleus muscle (B, ×400). Fas antigen-positive muscle fiber (C, ×400) observed in the TA-treated soleus muscle is also positive by ISEL (D, ×400).
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were stained positively by ISEL. Peripheral condensation of nuclear chromatin, which appears at early stages of apoptosis and is distinct from necrotic cells, was observed by electron microscopy. Contrary to a widespread belief that differentiated skeletal muscle undergoes only necrotic death, apoptotic cellular death has been demonstrated in dystrophin-deficient muscle (20, 21) and differentiated skeletal muscle after treatment with anabolic steroids (22).

Apoptosis in any particular cell lineage is mediated by exogenous influences, such as survival factors and the genes of the susceptible cell. One signal which leads to cell death by apoptosis is the Fas gene product (23, 24). The Fas molecule, synonymously referred to as APO-1, contains a cytoplasmic “death domain” shared with the type I tumor necrosis factor receptor (34). To date, apoptosis induced by Fas antigen has been extensively investigated in the lymphoid systems (23, 24). Immunohistochemical studies demonstrated that Fas antigen was expressed on muscle fibers from patients with various muscle wasting diseases, but not in normal muscle cells (26, 27). However, whether the muscle fiber that expressed the Fas antigen is involved in apoptotic processes has not been fully investigated. In this study, Fas antigen was expressed only in the TA-treated soleus muscles. Fas antigen was mainly detected in the cytoplasm of muscle fibers. Some Fas antigen-positive muscle fibers showed positive reaction for ISEL, indicating nuclear chromatin, which appears at early stages of apoptosis (30), the small number of apoptotic nuclei in this study is not an indicator of insignificance of apoptosis in steroid myopathy. The apoptotic cell death in steroid myopathy is mediated by Fas antigen expression in this study.

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