Rat diaphragm contractility and histopathology are affected differently by low dose treatment with methylprednisolone and deflazacort

P.N.R. Dekhuijzen, G. Gayan-Ramirez, A. Bisschop, V. de Bock, R. Dom, R. Bouillon, M. Decramer

Recent animal studies have demonstrated that administration of corticosteroids affects diaphragm structure and function [1-6]. Fluorinated steroids (e.g. triamcinolone 0.5-3 mg·kg⁻¹ i.m. and dexamethasone 1-4 mg·kg⁻¹ i.m. administered during 1-7 weeks) caused severe loss of body weight and muscle mass up to 40%. This was accompanied by type IIb fibre atrophy in the diaphragm as well as in peripheral skeletal muscles [1-5]. These morphological alterations were reflected in changes in the physiological properties of isolated diaphragm bundles, which showed a shift towards the contractile profile of slow muscle. In contrast, the nonfluorinated steroid prednisolone, administered in a dose of 5 mg·kg⁻¹ i.m. daily during 4 weeks, did not cause loss of body mass [4]. Histological examination revealed myogenic changes in the diaphragm, including greater than normal variation of the diameter of all fibre types, scattered necrotic fibres, excess nuclei, and an increased amount of connective tissue, but without fibre atrophy. In line with these alterations, an increased tendency towards fatigue was observed when examining the contractile characteristics of the diaphragm bundles.

In the above mentioned studies, moderate to high doses of corticosteroids were administered. It remains unclear whether lower doses may cause alterations in animal diaphragm. In patients with chronic obstructive pulmonary disease (COPD) there is evidence to suggest that treatment with low doses of methylprednisolone (MP) affects respiratory and peripheral muscle force [7].

The aim of the present study, therefore, was to investigate whether in rats contractile properties of the diaphragm (DIA) and morphological characteristics of DIA and gastrocnemius muscle (GA) were affected by treatment with a low dose (0.4 mg·kg⁻¹ i.m. daily) of MP, theoretically comparable to doses commonly used for therapeutic purposes in humans. In addition, we studied...
the effects of an equipotent dose of deflazacort (DF), 0.5 mg·kg⁻¹ i.m. daily, an oxazoline derivative of prednisolone with similar anti-inflammatory and immunosuppressive activity [8], but less side-effects on bone structure and carbohydrate metabolism compared to prednisolone [9, 10].

**Methods**

**Study design, animals and treatment**

Twenty six adult male Wistar rats, aged 18 weeks, weighing 350–400 g, were randomized into one of three treatment groups: 1) control (C), (n=10), saline 0.05 ml daily i.m.; 2) methylprednisolone (MP), (n=10), 0.4 mg·kg⁻¹ daily i.m.; and 3) deflazacort (DF), (n=6), 0.5 mg·kg⁻¹ daily i.m.

The dosages of MP and DF were chosen in order to administer an equivalent anti-inflammatory dose to the animals. During 6 weeks, the animals were injected daily in the left hind limb. They were fed ad libitum. All animals were weighed thrice weekly. After the treatment period, contractile properties of DIA and histo­

**Contractile properties**

Twenty four hours after the last injection, rats were anaesthetized with sodium pentobarbital (Nembutal), 60 mg·kg⁻¹ i.p. Animals were tracheotomized and a tra­

The diaphragm was quickly removed through a lap­

**Histological and histochemical procedures**

**MUSCLE STRIPS**

Muscle strips obtained from the costal region of the diaphragm and from the mid-belly of the GA of the right hind limb were prepared for histopathological examination. Muscle samples were put into "tissue glue" (TissueTek, Elkhard, IN, USA) on a cork holder, with the muscle fibres orientated perpendicularly to the surface of the cork. Proper orientation of the bundles was controlled by using magnifying glasses. Subsequently, these specimens were quickly frozen in isopentane cooled with liquid N₂. Serial cross-sections were cut at 10 μm thickness with a cryostat kept at -20°C, parallel to the cork. Sections of each DIA and GA were taken for routine haematoxylin and eosin staining.

The other serial sections were stained for myofibrill­

**Study design, animals and treatment**

Twenty six adult male Wistar rats, aged 18 weeks, weighing 350–400 g, were randomized into one of three treatment groups: 1) control (C), (n=10), saline 0.05 ml daily i.m.; 2) methylprednisolone (MP), (n=10), 0.4 mg·kg⁻¹ daily i.m.; and 3) deflazacort (DF), (n=6), 0.5 mg·kg⁻¹ daily i.m.

The dosages of MP and DF were chosen in order to administer an equivalent anti-inflammatory dose to the animals. During 6 weeks, the animals were injected daily in the left hind limb. They were fed ad libitum. All animals were weighed thrice weekly. After the treatment period, contractile properties of DIA and histo­

**Contractile properties**

Twenty four hours after the last injection, rats were anaesthetized with sodium pentobarbital (Nembutal), 60 mg·kg⁻¹ i.p. Animals were tracheotomized and a tra­

The diaphragm was quickly removed through a lap­

**Histological and histochemical procedures**

**MUSCLE STRIPS**

Muscle strips obtained from the costal region of the diaphragm and from the mid-belly of the GA of the right hind limb were prepared for histopathological examination. Muscle samples were put into "tissue glue" (TissueTek, Elkhard, IN, USA) on a cork holder, with the muscle fibres orientated perpendicularly to the surface of the cork. Proper orientation of the bundles was controlled by using magnifying glasses. Subsequently, these specimens were quickly frozen in isopentane cooled with liquid N₂. Serial cross-sections were cut at 10 μm thickness with a cryostat kept at -20°C, parallel to the cork. Sections of each DIA and GA were taken for routine haematoxylin and eosin staining.

The other serial sections were stained for myofibrill­

**Study design, animals and treatment**

Twenty six adult male Wistar rats, aged 18 weeks, weighing 350–400 g, were randomized into one of three treatment groups: 1) control (C), (n=10), saline 0.05 ml daily i.m.; 2) methylprednisolone (MP), (n=10), 0.4 mg·kg⁻¹ daily i.m.; and 3) deflazacort (DF), (n=6), 0.5 mg·kg⁻¹ daily i.m.

The dosages of MP and DF were chosen in order to administer an equivalent anti-inflammatory dose to the animals. During 6 weeks, the animals were injected daily in the left hind limb. They were fed ad libitum. All animals were weighed thrice weekly. After the treatment period, contractile properties of DIA and histo­

**Contractile properties**

Twenty four hours after the last injection, rats were anaesthetized with sodium pentobarbital (Nembutal), 60 mg·kg⁻¹ i.p. Animals were tracheotomized and a tra­

The diaphragm was quickly removed through a lap­

**Histological and histochemical procedures**

**MUSCLE STRIPS**

Muscle strips obtained from the costal region of the diaphragm and from the mid-belly of the GA of the right hind limb were prepared for histopathological examination. Muscle samples were put into "tissue glue" (TissueTek, Elkhard, IN, USA) on a cork holder, with the muscle fibres orientated perpendicularly to the surface of the cork. Proper orientation of the bundles was controlled by using magnifying glasses. Subsequently, these specimens were quickly frozen in isopentane cooled with liquid N₂. Serial cross-sections were cut at 10 μm thickness with a cryostat kept at -20°C, parallel to the cork. Sections of each DIA and GA were taken for routine haematoxylin and eosin staining.

The other serial sections were stained for myofibrill­

**Study design, animals and treatment**

Twenty six adult male Wistar rats, aged 18 weeks, weighing 350–400 g, were randomized into one of three treatment groups: 1) control (C), (n=10), saline 0.05 ml daily i.m.; 2) methylprednisolone (MP), (n=10), 0.4 mg·kg⁻¹ daily i.m.; and 3) deflazacort (DF), (n=6), 0.5 mg·kg⁻¹ daily i.m.

The dosages of MP and DF were chosen in order to administer an equivalent anti-inflammatory dose to the animals. During 6 weeks, the animals were injected daily in the left hind limb. They were fed ad libitum. All animals were weighed thrice weekly. After the treatment period, contractile properties of DIA and histo­

**Contractile properties**

Twenty four hours after the last injection, rats were anaesthetized with sodium pentobarbital (Nembutal), 60 mg·kg⁻¹ i.p. Animals were tracheotomized and a tra­

The diaphragm was quickly removed through a lap­

**Histological and histochemical procedures**

**MUSCLE STRIPS**

Muscle strips obtained from the costal region of the diaphragm and from the mid-belly of the GA of the right hind limb were prepared for histopathological examination. Muscle samples were put into "tissue glue" (TissueTek, Elkhard, IN, USA) on a cork holder, with the muscle fibres orientated perpendicularly to the surface of the cork. Proper orientation of the bundles was controlled by using magnifying glasses. Subsequently, these specimens were quickly frozen in isopentane cooled with liquid N₂. Serial cross-sections were cut at 10 μm thickness with a cryostat kept at -20°C, parallel to the cork. Sections of each DIA and GA were taken for routine haematoxylin and eosin staining.

The other serial sections were stained for myofibrill­
oxidative), or type IIb (fast-twitch glycolytic) fibres [15]. Slides preincubated at pH 4.5 offered the best separation of different fibre types, and were subsequently used for further analysis.

Morphometric examination was carried out with a Leitz microscope (Wetzlar, Germany) at ×25 magnification, connected to a digitizing board (Numonics 2207; Montgomeryville, PA, USA). Areas in which fibre orientation was not transverse to the long axis of the fibre were not analysed. Boundaries of individual muscle fibres were delineated, and fibre CSA was determined from the number of pixels within the outlined fibre. At least 150 fibres of each DIA and deep (red) part of the GA were used to calculate mean CSA of all fibre types.

**Data analysis**

Two diaphragm bundles were obtained from each animal. Since variation between animals was as large as within animals, all bundles were used for statistical analysis as independent cases. Data from the different treatment groups were compared, using two-way analysis of variance. Differences between means were assessed using Duncan’s multiple range test. A chi-squared likelihood ratio test was used to detect differences in the distribution of maximal tetanic tensions between the three treatment groups. Statistical significance was set at a value of p less than 0.05. All analyses were performed using the statistical Package for the Social Sciences (SPSS)/PC+ package [16]. Means±SD are represented in text, tables and figures, unless otherwise specified.

**Results**

**Body, muscle and adrenal weight**

Body weight at the start of the study was not different between the three groups (fig. 1). During the treatment period, weight increased by 10.9±1.7% in C, and by 6.6±1.4% in MP-treated animals, whereas it decreased by 4.2±1.1% in the DF group (p<0.05 compared to C).

Muscle masses in the DF group were significantly reduced compared to C (DIA, scalenus and GA) and MP (scalenus and GA) (table 1). No changes occurred in the parasternal muscles, probably because the major part of the weight consisted of the ribs. In addition, the mass of the adrenal glands was decreased compared to C and MP. The ratio of DIA mass to total body weight was similar among the three groups (C 0.16±0.03%, MP: 0.15±0.02%, DF: 0.17±0.02%).

**Diaphragmatic contractile properties**

**Bundle dimensions.** Bundle dimensions are shown in table 2. The DIA bundles in the DF group tended to be smaller, but no significant differences were found compared to the other groups.

**Twitch characteristics and maximal tetanic tension.** The data of the contractile properties of the DIA are summarized in table 3. Pt, TPT, 1/2RT and Po/Po were similar in the three groups. Po (expressed both as absolute values (kg) and corrected per CSA (kg-cm²)) was decreased in the DF group (p<0.05). When the values were divided in two classes (Po <2.0 and >2.0 kg-cm²), more bundles in the DF-treated animals generated tetanic tensions below 2.0 kg-cm² (37%, compared to 10% in the other two groups; χ²=8.712, p<0.05).

**Force-frequency curve.** The response of diaphragm strips to increasing stimulus frequencies is shown in figure 2. When expressed in absolute values (kg-cm²), a significant

<table>
<thead>
<tr>
<th>Treatment</th>
<th>Diaphragm (DIA) mg</th>
<th>Parasternals mg</th>
<th>Scalenus mg</th>
<th>Gastrocnemius (GA) mg</th>
<th>Soleus mg</th>
<th>Adrenal glands mg</th>
</tr>
</thead>
<tbody>
<tr>
<td>Control (C)</td>
<td>663 (67)</td>
<td>3233 (311)</td>
<td>578 (56)</td>
<td>2238 (179)</td>
<td>163 (17)</td>
<td>45 (3)</td>
</tr>
<tr>
<td>Methylprednisolone (MP)</td>
<td>626 (40)</td>
<td>3171 (167)</td>
<td>580 (53)</td>
<td>2053 (137)*</td>
<td>165 (12)</td>
<td>42 (6)*</td>
</tr>
<tr>
<td>Deflazacort (DF)</td>
<td>597 (39)*</td>
<td>3176 (299)</td>
<td>504 (46)*#</td>
<td>1872 (132)*#</td>
<td>155 (15)</td>
<td>29 (7)*#</td>
</tr>
</tbody>
</table>

Data are presented as mean, and sd in parenthesis. *: p<0.05 compared to control; #: p<0.05 compared to methylprednisolone.
Table 2. – Diaphragm bundle dimensions

<table>
<thead>
<tr>
<th>Treatment</th>
<th>Length (mm)</th>
<th>Thickness (mm)</th>
<th>Width (mm)</th>
<th>Weight (mg)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Control</td>
<td>21.2 (1.9)</td>
<td>0.51 (0.13)</td>
<td>1.62 (0.20)</td>
<td>28.5 (6.2)</td>
</tr>
<tr>
<td>Methylprednisolone</td>
<td>20.3 (2.2)</td>
<td>0.53 (0.12)</td>
<td>1.62 (0.24)</td>
<td>27.1 (5.3)</td>
</tr>
<tr>
<td>Deflazacort</td>
<td>19.8 (2.5)</td>
<td>0.46 (0.11)</td>
<td>1.60 (0.28)</td>
<td>25.2 (7.1)</td>
</tr>
</tbody>
</table>

Data are presented as mean.

downward shift of the DF bundles at 50, 80 and 120 Hz was observed (p<0.05) (fig. 2a). A nonsignificant decrease in force generation was observed in the MP bundles. When expressed as a percentage of the 160 Hz stimulations before and after each stimulus frequency, no differences were observed among the three groups (fig. 2b).

Fatigue properties. In all groups, force generated at 160 Hz decreased during the force-frequency stimulation procedure. In the C, MP and DF groups, the values for percentage decrease relative to initial \( P_0 \) were 8.0±1.9, 15.4 ±2.6, and 19.6±5.8 (mean±SD), respectively. This decrease in generated force in the MP and DF groups was significantly higher than in C (p<0.05).

During the low-frequency fatigue run, stress generation in absolute values (kg-cm\(^2\)) was similar in all groups (fig. 3a). When expressed as a percentage of initial stresses, this ratio (fatigue/fresh, a dimensionless index) was higher in the DF bundles (fig. 3b), (p<0.05).

Histology and histochemistry

Histological examination of haematoxylin and eosin stained slides showed a normal muscular pattern of DIA and GA in the three groups.

Analysis of ATPase stainings showed that fibre type distribution was not changed by the different treatments. For groups taken together, the DIA consisted of 40.4±5.3% type I fibres, 29.7±6.1% type IIa fibres, and 29.9±4.2% type IIb fibres. The red (deeper) part of the GA consisted of 30.1±4.5% type I fibres, 25.3±5.1% type IIa fibres, and 44.6±6.5% type IIb fibres. Dimensions of the three fibre types in both DIA and GA were similar in C and MP groups (fig. 4). Treatment with DF, however, was accompanied by a decrease of diaphragmatic type I, IIa, and IIb fibre cross sectional area (p<0.05 compared to the other groups) (fig. 4a). Similar changes were observed in the GA (fig. 4b).

Table 3. – Diaphragmatic contractile properties

<table>
<thead>
<tr>
<th>Treatment</th>
<th>( P_t ) (kg-cm(^2))</th>
<th>TPT (ms)</th>
<th>1/2RT (ms)</th>
<th>( P_0 ) (kg)</th>
<th>( P_0 ) (kg-cm(^2))</th>
<th>( P_t/P_0 )</th>
</tr>
</thead>
<tbody>
<tr>
<td>Control</td>
<td>0.72 (0.19)</td>
<td>19 (2)</td>
<td>24 (2)</td>
<td>0.033 (0.009)</td>
<td>2.57 (0.45)</td>
<td>0.28 (0.04)</td>
</tr>
<tr>
<td>Methylprednisolone</td>
<td>0.70 (0.15)</td>
<td>20 (2)</td>
<td>24 (4)</td>
<td>0.031 (0.009)</td>
<td>2.46 (0.33)</td>
<td>0.28 (0.05)</td>
</tr>
<tr>
<td>Deflazacort</td>
<td>0.62 (0.16)</td>
<td>19 (2)</td>
<td>25 (2)</td>
<td>0.028 (0.008)*</td>
<td>2.32 (0.45)*</td>
<td>0.27 (0.06)</td>
</tr>
</tbody>
</table>

Data are presented as mean (SD). \( P_t \): twitch tension; TPT: time to peak tension; 1/2RT: half-relaxation time; \( P_0 \): tetanic tension; \( P_t/P_0 \): twitch/tetanic ratio. *: p<0.05 compared to control.
Fig. 3. - a) Fatigue curve of diaphragm bundles. b) Fatigue curve of diaphragm bundles, expressed as a percentage of initial value. --- : control; —O— : methylprednisolone; ■— : deflazacort; —— : pooled sd. *: deflazacort compared to control; p<0.05; #: p<0.05 deflazacort compared to control and methylprednisolone.

Fig. 4. - a) Fibre cross-sectional area in diaphragm; and b) gastrocnemius. □ : control; ■ : methylprednisolone; ■■ : deflazacort. *: p<0.05 deflazacort compared to control and methylprednisolone.

Discussion

The present study shows that low dose MP administered during 6 weeks hardly affects the contractile properties of the rat diaphragm, if at all. Only subtle changes were found, such as slightly increased fatigability during high-frequency stimulation and a nonsignificant downward shift of the force-frequency curve. MP did not alter diaphragm or gastrocnemius morphology. Treatment with DF, however, was associated with more pronounced changes in the contractile characteristics of the diaphragm. These alterations consisted of a decrease in maximal tetanic tension, a downward shift of the force-frequency curve and decreased maximal tension generation during this protocol. In contrast to MP, DF caused mild atrophy of all fibre types, both in the diaphragm and in the gastrocnemius. In line with these findings, treatment with DF was accompanied by slight loss of body and muscle mass.

Interpretation of the results

Side-effects, such as changes in bone and carbohydrate metabolism, associated with the use of DF appear to be less than with prednisolone [17–20]. Comparisons between DF and MP have not been made to our knowledge, nor have animal studies been published regarding the effects of DF on respiratory and peripheral skeletal muscles.

One early study has been published regarding the effects of MP on skeletal muscles in dogs [21]. In this study, the effects of (among other steroids) prednisolone 25 mg i.m. and MP 20 mg i.m. daily during 5 weeks were compared. Body weight decreased by ±10% in both groups. Prednisolone caused mild atrophy (not specified for different fibre types) in the quadriceps muscle, whereas MP did not change fibre size. This is in line with the morphological findings in the present study.

The pattern of fibre atrophy associated with DF in the
Mechanisms of steroid-induced changes in striated muscle

The mechanism of steroid-induced changes in striated muscle is complex and in part unknown. Protein wasting is due to a reduction in protein synthesis and to an increase in intracellular proteolysis. Inhibition of protein synthesis occurs primarily in type II muscle fibres, mainly by affecting the activity of factors involved in peptide initiation [25]. Increased muscle cytoplasmatic protease activity, associated with steroid treatment, results in myofibrillar destruction [26]. Carbohydrate metabolism within muscle fibres is also affected by corticosteroids. Muscle glycogen content is increased by increased muscle glycogen synthetase activity and decreased glycogen utilization [27]. Mitochondrial alterations in muscles affected by steroids have suggested that impaired oxidative respiration may also be an important factor in pathogenesis [28].

There is no clear explanation for the difference in response of striated muscle to different subgroups of glucocorticoids, such as fluorinated and nonfluorinated steroids. A difference in binding of triamcinolone, dexamethasone, and cortisol to cytoplasmatic proteins may play a role [26]. In the case of DF, another mechanism has been proposed, associated with its chemical structure. DF is characterized by an oxazoline group at C-17, which is believed to sterically hinder reactivity of the side chain [29]. This results in a reduced lipid solubility compared to prednisolone. Therefore, it has been postulated that DF is less extractable from serum by cells with a limited degree of exposure to the circulation, such as osteoblasts, accounting for its favourable effects on bone metabolism [8]. Such a mechanism, however, may be expected to protect organs with an abundant blood supply, such as the dia-phragm, less, as suggested by the findings in the present study. Assandri et al. [30] showed that the concentration of deflazacort in striated muscle in rats after oral administration of 5 mg·kg\(^{-1}\) deflazacort was low. Data on the diaphragm were not presented.

Potential clinical significance of steroid-induced alterations

In previous studies, the doses of steroids administered were high compared to current use in clinical practice. In the present study, MP and DF were administered in low doses, comparable with the dose sometimes administered to patients with asthma or COPD. In these patients, diaphragm function is compromised by malnutrition [31, 32], hyperinflation [33], disturbances in blood gases [34], and cardiac failure [35, 36]. It may be speculated that the effects of steroids on already impaired diaphragmatic function, as in patients with COPD, might be more pronounced.

Zanotti et al. [37] showed that respiratory muscle strength was similar in patients with COPD treated with DF, 15 mg daily for at least four consecutive months, compared to COPD patients matched for age and severity of airflow obstruction. Firm conclusions from this study, however, are precluded since it was performed in a retrospective and cross-sectional design.

In a recent study, eight out of 21 consecutive patients who were admitted to hospital with acute exacerbations of COPD or asthma, suffered from generalized muscle weakness [7]. In seven of these eight patients the average daily dose of methylprednisolone during the last 6 months exceeded 4 mg, whilst this was the case in only three of the 13 patients with normal muscle strength. Although the therapeutic efficacy of corticosteroids in COPD is at least controversial [38], this observation indicates the potential effect of low-dose treatment with methylprednisolone on muscle strength.

In summary, this study shows that low dose of MP cause only subtle changes in contractile properties of rat diaphragm, without alterations in morphology. In contrast, more pronounced abnormalities in contractile
properties are observed in the DF treated bundles, which are accompanied by generalized muscle fibre atrophy in both diaphragm and gastrocnemius.

Acknowledgement: The authors thank N. Plaisance-Buts for expert technical assistance.

References


