Effects of Melanocyte-stimulating Hormone on Plasma Levels of Testosterone and Estradiol Hormones in Alloxan-Induced Diabetic Rats

1Mahmoud Abu-Samak, 2Fahmi Mahmoud, 1Moayad Khataibeh,
3Suhail Hamdan and 4Aurelia Crevoi
1Department of Medical Technology, Faculty of Allied Medical Sciences,
Applied Science University, Amman, Jordan
2Faculty of Pharmacy, Al-Zaytoonah University of Jordan, Amman Jordan
3Medical School, Al-Quds University, Al-Quds, Palestine
4Department of Human and Animal Physiology, Moldova State University, Moldova

Abstract: This study was designed to investigate the effects of Melanocyte stimulating hormone MSH on serum testosterone T and estradiol E2 hormones concentrations in alloxan-induced diabetic rats. Eighty male and female Sprague Dawley rats, weighing 180-200 g, were divided into four groups of normal rats and four groups of alloxan-induced diabetic rats were given intraperitoneally (i.p.) a daily injection of 20 mg alloxan solution/100 g of body weight for 10 days. Two groups, male and female from the normal and 2 diabetic groups served as controls and did not inject with MSH 2 groups, male and female from the normal rats and 2 groups from the diabetic rats injected (i.p) daily with MSH at a dose of 2-microg/100 g of body weight, for 10 days. The control group was only injected with the same volume of normal saline. Serum glucose concentrations were higher and serum insulin, testosterone and estradiol concentrations were lower in diabetic rats than those in the control groups. MSH administration decreased the elevated blood glucose concentrations of the diabetic rats to the normal levels and decreased estradiol concentration in female normal rats while increased the testosterone concentration in male normal rats. Present findings indicate that MSH plays adaptive role during early stages of alloxan induced-diabetes mellitus. Further studies are needed to identify the mechanism.

Key words: MSH, testosterone, estradiol, sex hormone, alloxan induced-diabetic rats, Type 2 diabetes mellitus

INTRODUCTION

Melanocyte Stimulating Hormone (MSH) has modulatory role in food intake, body fat and glucose metabolism (Biebermann et al., 2006; Fehm et al., 2001; Costa et al., 2006), but little is known about the effects of MSH on gonadal function during diabetes mellitus and practically there are no literature relating to the MSH influence on sex hormones levels during diabetes mellitus. A number of investigators reported on the relationship between gonadal dysfunction and Diabetes Mellitus (DM) in humans and animals (Matsushita et al., 2005, Komaki et al., 2005). Male and female reproductive alterations have been widely reported in individuals with diabetes including decreases in testicular testosterone production, reduced serum LH and male and female estradiol (Ballester et al., 2004; Steger and Rabb, 1997; Sanguinetti et al., 2004). Although several investigators have reported serum E2 free Testosterone and LH not affected in diabetic rats (Komaki et al., 2005) and diabetes-related effects on testicular function have been attributed to the lack of insulin. The regulatory action of this hormone is known and observations of a direct effect on both Leydig cells (Khan et al., 1992; Hurtado de Catalfo et al., 1998) and Sertoli cells (Berland et al., 1984; Mita et al., 1985) have been reported. Nonetheless, the data are confusing and the exact role that insulin plays in the regulation of the male reproductive function is still unclear, even though the melanocortins are known to mediate some of the effects of both leptin and insulin (Benoit et al., 2004; O'Shaughnessy et al., 2003; Costa et al., 2006).

Therefore the aim of the present research is to investigate whether MSH can stimulate the gonadal axis by altering glucose homeostasis or sex hormones dynamic in diabetic rats.

Corresponding Author: Dr. Mahmoud Abu-Samak, Department of Medical Technology, Faculty of Allied Medical Sciences, Applied Science University, Amman 11931, Jordan Fax: 8232899
1350
MATERIALS AND METHODS

Animals: This research was carried out at the laboratories of Department of Medical Technology, Applied Science University from September 2005 to January 2006. Eighty male and female Sprague Dawley rats, were housed in our laboratory for at least one week before the experiments. The rats were maintained in a temperature-controlled room (22-25°C). Food and water were available for the animals all the time and without any restrictions. The weights of rats were taken on the day of the experiment and only those weighing 180-250 g were used in this study.

The animals were divided into two sections according to the sex of the animal and the animals of each section were divided into four treatment groups: control, MSH injected rats (MSH), Alloxan-induced diabetic rats (Allox) and Alloxan-diabetic rats injected with MSH (Allox+MSH). All the animals were given daily injections of following preparations, for 10 days: Control rats injected with 1 mL of normal saline, (MSH) and (Allox+MSH) Rats injected (i.p. daily with alpha melanocyte stimulating hormone (Sigma, Firm, USA) at a dose of 2 microg (dissolved in 1 mL of normal saline)/100 g of body weight. (Allox) and (Allox+MSH) rats were given intraperitoneally (i.p) a daily injection of 20 mg alloxan (dissolved in 1 mL of normal saline) solution/100 g of body weight (Sigma, Firm, USA).

At the end of the experiment, all rats were fasted for 12 h before they were sacrificed and blood collected.

Serum analysis: The serum was isolated by centrifugation and analyzed by radioimmunoassay for serum insulin, estradiol and testosterone using a rat insulin, estradiol and testosterone radioimmunoassay kit (Ceni-Jeford Firm, France). Serum glucose concentrations were measured by the glucose oxidase method using a spectrophotometer (Cecil ce 1010 England).

Statistical analysis: Data were expressed as means±SE and were analyzed with a two-way ANOVA followed by LSD multiple comparison test, using Statistica Software (OK, USA). Differences were considered significant at p<0.05.

RESULTS AND DISCUSSION

Daily MSH administration for 10 days did not alter insulin levels but normalized blood glucose in the Allox rats, compared with control group (Table 1). Diabetic male rats showed a low weight gain and decreased levels of plasma insulin and testosterone when compared with healthy controls (Table 1). Diabetes also induced hyperglycemia (Table 1 and 2).

Allox-induced diabetes increased estradiol concentration, while decreased testosterone concentration compared with control groups (Fig. 1 and 2). MSH administration decreased estradiol concentration and increased testosterone concentration in diabetic groups (Fig.1 and 2).

Present results indicate that MSH plays modulatory role on sex hormones levels in diabetic rats. MSH may exert this role through glucose homeostasis peripheral pathway because there was no change in insulin concentration while it normalized blood sugar in both male and female diabetic rats. Costa et al. (2006) reported that MSH has reciprocal effects in which MSH appears to increase sensitivity to insulin when present in the CNS, while MSH in the periphery is necessary for insulin resistance. Their results are in agreement with our results in that MSH plays modulatory role in diabetes mellitus type 2. Hochgeschwender et al. (2003) stated that the regulation of glucose homeostasis requires the integration of both central and peripheral melanocortin signalling systems. Furthermore, Abu-Samak and Savontaus et al. (2004) reported that MSH could serve as a potential strategy for anti-obesity and/or antidiabetic therapy.

Also the clear differences in sex hormones levels observed between male and female rats suggest that MSH has contrasting effects on male and females rats. This is supported by the reports of Kim et al. (1999), Robertson et al. (2003). There are researches indicating that estrogen normally acts in the brain to reduce food intake and body weight in female individuals (Clegg et al., 2003) and other reports show that MSH mimics these effects (Kim et al., 1999; Robertson et al., 2003; Fan et al., 1997). On the other hand, numerous experiments have reported that MSH has been shown to stimulate gonadotropin secretion in humans (Celis, 1985), in which an elevation of plasma testosterone hormone levels in male group injected with MSH have regulatory effects on the physiological function of rats. This is supported by previous lines of evidence that indicate that elevation of testosterone support the concept of a functional continuity between the central and peripheral actions of melanocortins in regulating certain physiological functions like sexual behavior (Van der Kraan et al., 1998). Further studies are needed to identify the mechanism underlying this enhanced sensitivity to peripherally administered MSH. There are
Table 1: Effects of melanocyte stimulating hormone and diabetes on physical and seric parameters

<table>
<thead>
<tr>
<th></th>
<th>Control</th>
<th>MSH</th>
<th>Allox</th>
<th>MSH+ Allox</th>
</tr>
</thead>
<tbody>
<tr>
<td>Initial body wt. (g)</td>
<td>202.5±8.24</td>
<td>236.5±11.2</td>
<td>226.4±13.3</td>
<td>192.6±7.66</td>
</tr>
<tr>
<td>Final body wt. (g)</td>
<td>216.5±9.88</td>
<td>228.5±0.8</td>
<td>208.9±13.1</td>
<td>177.6±6.28</td>
</tr>
<tr>
<td>Serum glucose, (mmol L⁻¹)</td>
<td>5.3±0.25</td>
<td>5.4±0.35</td>
<td>8.42±1.3</td>
<td>5.5±0.65</td>
</tr>
<tr>
<td>Serum insulina, (µIU L⁻¹)</td>
<td>1.7±0.22</td>
<td>2.7±0.30</td>
<td>0.5±0.08</td>
<td>0.5±0.10</td>
</tr>
</tbody>
</table>

Table 2: Significant differences (p-values) between female estradiol (*) and male testosterone (**) concentrations

<table>
<thead>
<tr>
<th></th>
<th>Control</th>
<th>MSH</th>
<th>Allox</th>
<th>MSH+ Allox</th>
</tr>
</thead>
<tbody>
<tr>
<td>Female Rats</td>
<td>0.910159</td>
<td>0.001987</td>
<td>0.2084</td>
<td></td>
</tr>
<tr>
<td>MSH</td>
<td>0.001452</td>
<td>0.17165</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Allox</td>
<td>0.0473</td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th></th>
<th>Control</th>
<th>MSH</th>
<th>Allox</th>
<th>MSH+ Allox</th>
</tr>
</thead>
<tbody>
<tr>
<td>Male Rats</td>
<td>0.445</td>
<td>0.000</td>
<td>0.000432</td>
<td></td>
</tr>
<tr>
<td>MSH</td>
<td>0.000</td>
<td>0.000044</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Allox</td>
<td>0.0142</td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

Fig. 1: Effects of peripheral injection of MSH for 10 days on estradiol serum concentration. Values are means±SE

Fig. 2: Effects of peripheral injection of α-MSH (2 microg/100 g bw) for 10 days on Testosterone serum concentration in the male rats. Values means±SE

therapeutics is specifically active in the CNS or peripheral circulation for the treatment of type 2 gonadal dysfunction.

REFERENCES


