Designing A Model of Induced-Benzhexol Addiction In Male Rats

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Abstract
Background: Benzhexol hydrochloride (Trihexyphenidyl) is a potent anticholinergic agent. Abuse of benzhexol has been noted with increasing frequency in recent years. By reviewing the relevant literature, no model for benzhexol addiction in rats was found.
Aim of the study: to designing a model of induced benzhexol addiction in male rats.
Materials and methods: Thirty two male, adult, Albino rats were enrolled in this experiments. The animals were randomly divided into 4 groups, eight rats for each group. Each rat of group 1 received normal saline (N.S.) in equal volume to the benzhexol dose, twice daily (b.d.), orally (p.o.) by gastric tube for 10 days. Each rat of group 2, 3 and 4 received benzhexol 0.1mg, 0.2mg and 0.4mg respectively, b.d., p.o. by gastric tube for 10 days. On the 11th day, each rat was placed in the open-field chamber for 15 minutes and all behaviors (withdrawal symptoms) of the animals were recorded by video camera.
Results and conclusion: Withdrawal symptoms such as rooming, wet dog shakes, head shakes, abdominal constriction, facial fasciculation, ptosis and piloerection significantly increased (P<0.05) in group 3 and 4 as compared with control group and model of benzhexol addiction in male rats can be induced by administration of 0.2 or 0.4 mg/kg of benzhexol, p.o, for 10 days.
Keywords: Benzhexol, withdrawal symptoms, grooming, wet dog shakes, head shakes, abdominal constriction, facial fasciculation, ptosis, piloerection and irritability.

تصميم موديل لادمان البنزهكسول المستحدث في ذكور الجرذان

الخلاصة

يعتبر البنزهكسول هايدروكلاوريد (الأزيين) دواءً فعالًا مضادًا للكلورونيك. وان اسعا استعمال هذا الدواء زاد بشكل مطرد في السنوات الأخيرة. حيث تشير البيانات المتوفرة إلى عدم وجود موديل للادمان على هذا الدواء في الجرذان. هدفت هذه الدراسة إلى تصميم موديل لادمان البنزهكسول المستحدث في ذكور الجرذان. تم استعمال اثنان وثلاثون جرذًا ذكرًا وسموا عشوائيًا إلى أربعة مجاميع. ثمانية حيوانات في كل مجموعة. أظهرت المجموعة الأولى نموذج سلبيًا. أما المجموعات الثانية والثالثة والرابعة فقد أعطيت عقار البنزهكسول جرعة 0.1 و 0.2 و 0.4 ملغ/كلغ حسب التوزيع مرتين يوميا لمدة 10 أيام عن طريق الفم. ان نتائج هذه الدراسة تشير إلى حصول زيادة بالغة الأهمية بالاعراض الانسحابية كالفصل، انقطاعية الكتل، اهتزاز الرأس، تقلصات عضلات
Introduction

Benzhexol hydrochloride (Trihexyphenidyl) is a potent anticholinergic agent [1]. It causes reversible blockade of cholinomimetic action at muscarinic receptors [2]. It characterized by a higher affinity for the neuronal (M1) muscarinic receptor than other subtypes of (M2 and M3) muscarinic receptors [3]. Benzhexol can be used for Parkinson's disease [4,5], extrapyramidal side effects, such as tardive dyskinesia, occurring during antipsychotic treatment [6,7,8], dystonia [9,10], tremor [11,12], excessive salivation [13,14], spasmodic torticollis [15,16].

Abuse of benzhexol has been noted with increasing frequency in recent years [17,18,19,20]. By reviewing the relevant literature, no model for benzhexol addiction in rats was found.

Two drugs, opioids and alcohol, provide classic examples of rodent somatic signs of withdrawal and served as models for measures of withdrawal per se. The somatic signs of withdrawal are an index of dependence and provide a quantifiable measure by which to assess the level of dependence [21].

Opioid withdrawal signs in rodents include weight loss, grooming, wet dog shakes, abdominal constrictions, facial fasciculation/teeth chattering, ptosis, abnormal posture, diarrhea and irritability [22].

For alcohol, withdrawal signs characterized by hyperactivity, tail tremors, tail stiffness, head tremors, general tremors, wet shakes, teeth chattering, akinesia, spastic rigidity, and convulsions [23].

Aim of the study: to design a model of induced benzhexol addiction in male rats.

Materials and methods

1- Animals:
Thirty two male, adult, Albino rats were enrolled in this experiments. Their weights were 150-200 g. The rats were housed in the Animal House of the College of Medicine/Babylon University, and kept on 25°C and 12 hours light–dark cycle with water and food ad libitum. After two weeks of adaptation, the animals were randomly divided into 4 g.oups, eight rats for each group.

2- Drug
Benzhexol HCL (Parkizol tablet, 5mg, PHARMALINE, Lebanon) was dissolved in 50 ml of normal saline (N.S.), so the final product contained 0.1 mg of benzhexol in each ml.

3- Experimental protocol
a- Each rat of group 1 received N.S. in equal volume to the benzhexol dose, twice daily (b.d.), orally (p.o.) by gastric tube for 10 days
b- Each rat of group 2,3 and 4 received benzhexol 0.1mg, 0.2mg and 0.4mg respectively, b.d., p.o. by gastric tube for 10 days.
c- On the 11th day, each rat was placed in the open-field chamber for 15 minutes and all behaviors (withdrawal symptoms) of the animals were recorded by video camera.

4- Open-field box
A glass chamber (40 cm x 30 cm x 30 cm) was made by the researcher (Picture 1). The rats were placed in this chamber and their activities were recorded by video camera.
5- **Video camera**
Digital Video Camera, (SONY, Japan) was used in this experiment to record the behaviors of animals.

6- **Statistical data**
The results were expressed as mean ± standard error of the mean (SEM). Statistical analysis was carried out by using one way ANOVA. Differences were considered statistically significant if the p value is lower than 0.05. The data of the research were carried out by using 17th edition of IBM® SPSS® statistics for Windows® 7.

**Results**

1- **Number of grooming**
There is no significant difference (P>0.05) in the number of grooming (Pictures 2 and 3) between control group and group 2, while number of grooming significantly increased (P<0.05) in group 3 and 4 as compared with control group. Number of grooming in group 3 and 4, also, significantly increased (P<0.05) as compared with group 2, while there is no significant increase between groups 3 and 4 (P>0.05) (Figure 1).
Figure 1: Mean of the grooming, wet dog shakes, and head shakes of all groups for 10 days following benzhexol withdrawn; group 1 (control group) and group 2 (benzhexol 0.1 mg/kg), group 3 (benzhexol 0.2 mg/kg), and group 4 (benzhexol 0.4 mg/kg).

Picture 2: Forelimbs grooming

Picture 3: Penile grooming
mg/kg).

2- **Number of wet dog shakes**

There is no significant difference (P>0.05) in the number of wet dog shakes between control group and group 2, while number of wet dog shakes significantly increased (P<0.05) in group 3 and 4 as compared with control group. Number of wet dog shakes in group 3 and 4, also, significantly increased (P<0.05) as compared with group 2, while there is no significant increase between groups 3 and 4 (P>0.05) (Figure 1).

3- **Number of head shakes**

There is no significant difference (P>0.05) in the number of head shakes between control group and group 2, while number of head shakes significantly increased (P<0.05) in group 3 and 4 as compared with control group. Number of head shakes in group 3 and 4, also, significantly increased (P<0.05) as compared with group 2, while there is no significant increase between groups 3 and 4 (P>0.05) (Figure 1).

4- **Other withdrawal symptoms**

There is no significant difference (P> 0.05) in the percentage of occurrence of abdominal constriction, facial fasciculation, Piloerection (Picture 4) and ptosis (Picture 5) between group 2 and control group, while they significantly increased in groups 3 and 4 as compared with control group.

Also there is no significant difference (P> 0.05) in the percentage of occurrence of abdominal constriction, facial fasciculation, ptosis, piloerection, and irritability between groups 3 and 4, and control group (Table 1).
Table 1: Percentage of occurrence of abdominal constriction, facial fasciculation, ptosis, and piloerection in different groups (group 1: control group; group 2: benzhexol 0.1 mg/kg; group 3: benzhexol 0.2 mg/kg; and group 4: benzhexol 0.4 mg/kg).

Picture 4: Piloerection

Picture 5: Ptosis
### Table

<table>
<thead>
<tr>
<th>Group</th>
<th>Abdominal constriction</th>
<th>Facial Fasciculation</th>
<th>Ptosis</th>
<th>Piloerection</th>
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<td>0%</td>
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<td>0%</td>
</tr>
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<td>100% *</td>
<td>87.5% *</td>
<td>75%</td>
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<tr>
<td>Group 4</td>
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<td>100% *</td>
<td>100% *</td>
<td>75%</td>
</tr>
</tbody>
</table>

*P<0.05

### Discussion

With this study, statistical analysis revealed that withdrawal symptoms such as grooming, wet dog shakes, head shakes, abdominal constriction, facial fasciculation, ptosis, piloerection, and irritability did not occurred after termination of 0.1mg/kg of benzhexol, pd, for 10 days, but occurred after termination of 0.2 mg/kg and 0.4 mg/kg of benzhexol, pd, for 10 days. These withdrawal symptoms are the classic withdrawal symptoms of other drugs in rodents [21] (Koob, 2011).

By reviewing the relevant literature, no specific animal model for benzhexol addiction was found. Benzhexol withdrawal symptoms which occur in this study agree with those reported in opioids and alcohol withdrawal symptoms [21,22] (Koob, 2011; Gellert and Holtzman, 1978).

### Conclusion

Model of benzhexol addiction in male rats can be induced by administration of 0.2 or 0.4 mg/kg of benzhexol, pd, po, for 10 days.

### References


6- Bartzokis G., Cummings J.L., Freidengerg D.L. and Wirshing W.C.


