Effect of long term Treatment of Chlorpromazine and Diazepam on the activity of acetyl cholinesterase in the chicks’ blood

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Received 5/2/2006 Accepted 4/9/2006

Abstract
Administration of chlorpromazine at (50 mg / kg / day ) orally for a period of 21 days resulted in a significant increase in chick blood plasma and RBC acetyl cholinesterase activity by 48 % and 62% respectively, where as only a slight and non significant increase by 9% and 13% respectively was observed when diazepam was given at (30 mg / kg / day ) orally in comparison with control group .The result suggest that chronic treatment with chlorpromazine may stimulate blood acetyl cholinesterase activity in chicks.

Introduction
Acetyl cholinesterase ( Ache ) is a cell microsomal membrane bound enzyme found in different tissues and play a responsible role for regulation of most physiological event, involving the turnover of acetylcholine( 1,2,3 ). A number of psychotropic drugs that change blood and/or 'true' Ache activity is known to affect various neurophysiologic and metabolic functions at cellular and molecular levels (1, 4).
Administration of chlorpromazine to chicks and/or rabbits increases the concentration of Ache in the brain and cerebrospinal fluid (5, 6). And the increase in levels of enzyme activity by chlorpromazine in certain brain regions depends on the duration of drug administration (6, 7). Long – term treatment with chlorpromazine resulted in significant increase in chick and rat 'true' Ache activity (3, 6). Diazepam in rats and /or chicks change (non significantly) the level of Ache in The brain (3, 6, 17). The present study was undertaken in chicks to examine Ache activity following chronic administration of chlorpromazine and diazepam.

Materials and Methods
Twenty- one chicks 2 – 2.5 months old and weighing 300 – 350 g were used. They were housed under standard condition of temperature (27°C) and humidity, and had free access to food (Commercial chick ration, Nebrese Co., Mosul, Iraq.) and water. The birds were divided into three groups of 7 chicks each. Control chicks were treated with physiological saline solution at 1ml/kg/day. The remaining two groups were treated with chlorpromazine (May and Baker Co., England) at 50 mg/kg/day orally and diazepam (Arab Pharmaceutical Manufacturing Co., Jordan) at 30 mg/kg/day orally. The volume of administration of both drugs was 1 ml / kg body weight. Treatment was continued daily for 21 day. At the end of experimental period, chicks were sacrificed after ether anesthesia, and blood was rapidly collected. Blood samples were deranged by using treated heparinized syringe (Braun, Germany) and separated plasma from RBC by centrifugation in (3000 rpm) for 10 min. Samples of blood stored in -20°C until the time of estimation of the enzyme activity (7 – 10 days). 0.2 ml from plasma and RBC samples was used for Ache activity measurement using acetylthiocholine iodide (7.5 %) (Fluka, Switzerland) (8). All assays were done in duplicate. The statistical significance of the differences between mean values were analyzed by student’s paired ‘t’ test (9). The level of significance was at p < 0.01.

Results and Discussion
Chronic chlorpromazine administration to chicks at 50 mg/kg/day significantly increases plasma and RBC Ache activity by 48% and 62% respectively in comparison with the control group as shown in table. While chronic diazepam treatment at 30 mg/kg/day caused only a slight but non significant increase in both plasma and RBC enzyme activity which was about 9% and 13% respectively of the control group. Estimation of acetyl cholinesterase activity is of importance in studies concerning enzyme synthesis mechanisms (10-12). In the present study, we found that long-term administration of chlorpromazine increased blood Ache activity. This effect seemed to be due to accelerated turnover of Ach (3, 12, 13). Possibly through change in cell membrane associated
properties (3,5,14,15), any drug which alter the composition of cell microsomal membrane would also affect the enzyme–bound activity (1,3,5,14,15). On the other hand, increased Ach turnover might cause adaptive change in the enzyme activity for Ach synthesis and degradation (3,14). Similarly, 21 day of haloperidol treatment is associated with cell microsomal membrane enzyme-bound activity (3,7,14). Stimulation of the enzymatic activity observed in the present study may be partly due to cell membrane effect of this drug (2-4, 16). The non significant change of enzyme activity by diazepam is in accordance with other reports (3, 4, 17).

References

Table: blood acetyl cholinesterase activity in chicks treated orally with chlorpromazine and diazepam at 50 and 30 mg /kg / day, respectively for 21 days.

<table>
<thead>
<tr>
<th>Treatment</th>
<th>Δ pH/ 30 min</th>
<th>Ache activity</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Plasma</td>
<td>% Increase</td>
</tr>
<tr>
<td>Saline (control)</td>
<td>0.33 ± 0.01</td>
<td></td>
</tr>
<tr>
<td>Chlorpromazine</td>
<td>0.49 ± 0.02</td>
<td>48% *</td>
</tr>
<tr>
<td>Diazepam</td>
<td>0.36 ± 0.01</td>
<td>9%</td>
</tr>
</tbody>
</table>

* P < 0.01 Versus Saline control group. The values are mean ± SE of 7 chicks / group.