INCREASED plasma histamine levels were associated with significantly lowered diamine and type B monoamine oxidase activities in platelet-rich plasma of atopic eczema (AE) patients. The diamine oxidase has almost normal cofactor levels (pyridoxal phosphate and Cu^{2+}) but the cofactor levels for type B monoamine oxidase (flavin adenine dinucleotide and Fe^{2+}) are lowered. The biogenic amines putrescine, cadaverine, spermidine, spermine, tyramine and serotonin in the sera, as well as dopamine and epinephrine in EDTA-plasma were found to be normal. It is unlikely, therefore, that these amines are responsible for the decreased activities of monoamine and diamine oxidase in these patients. The most likely causative factors for the inhibition of the diamine oxidase are nicotine, alcohol, food additives and other environmental chemicals, or perhaps a genetic defect of the diamine oxidase.

Key words: Atopic eczema, Biogenic amines, Diamine oxidase, Histamine, Monoamine oxidase

Introduction

Many attempts have been made to use the activity of the mitochondrial B-type monoamine oxidase of platelets for the diagnosis of mental disease. However, the results remain contradictory. All attempts to use diamine oxidase for the diagnosis of certain diseases have been unsuccessful until now, owing to an uncertain discrimination between the normal and pathological range. The preferred substrates of type B monoamine oxidase are benzylamine, 2-phenylethylamine, dopamine, tyramine and, with lower activity, tryptamine. N-methylhistamine, histamine methylated by N-methyl-transferase, is also oxidized by type B monoamine oxidase. This enzyme is therefore important in catabolizing histamine. Diamine oxidase is active on short-chain aliphatic diamines, such as putrescine and cadaverine, and is also the first and major histamine catabolizing enzyme. The purpose of this study was to evaluate the activity and the cofactor levels of the diamine oxidase in platelets of atopic dermatitis patients and healthy controls.

Materials and Methods

Twenty-one patients with clinically proved atopic eczema (age 15-44 years) and 13 healthy volunteers with no allergic history (age 16-39 years) gave their consent to participate. All patients avoided any steroid or antihistaminic treatment for at least 3 months before admission.

Platelet-rich plasma (PRP) was obtained by centrifuging the stabilized (EDTA) blood at 53 × g for 10 min at 20°C. The oxidases were measured by the method of Köchli and Wartburg with minor modifications. An aliquot (0.6 ml) of peroxidase buffer (8.3 mg peroxidase in 100 ml 0.1 M sodium phosphate, pH 7.15), 0.2 ml PRP and 10 µl 10% Triton X-100 were mixed. After 5 min 0.2 ml of 0.25 mM 2',7'-dichloro-fluorescein diacetate dissolved in 0.01 N NaOH and 10 µl 50 mM benzylamine (for monoamine oxidase) or 10 µl 50 mM putrescine (for diamine oxidase) were added and mixed. The absorbance at 502 nm was recorded at 15 to 25 min, at 20°C, on a Shimadzu UV-160 spectrophotometer.

Histamine was measured in EDTA plasma by the method of Shore using a Perkin-Elmer LS-2 filter fluorimeter. Serum copper, serum pyridoxal phosphate, serum iron, EDTA-blood haemoglobin and flavin adenine dinucleotide (FAD) in EDTA-blood were measured by routine procedures as outlined.

After deproteinisation of the sera with sulfosalicylic acid, putrescine and cadaverine were isolated from the supernatant by solid phase extraction. Dansyl derivatives formed after dansyl chloride treatment were extracted with ethyl acetate in the eluted liquid. The organic phase was dried and the residue solved in acetonitrile. Putrescine and cadaverine concentrations were measured fluorometrically after their separation using a HPLC system with gradient elution. The system contained an internal standard. Spermidine and spermine concentrations were measured in a similar way, but without the solid phase extraction at the beginning of the whole procedure. Tyramine was isolated from the sera by solid phase extraction on Bond...
Elut RP-C18 columns. It was measured after separation using an HPLC system with electrochemical detection.20,21 This system also contained an internal standard. Serum serotonin was separated from accompanying materials by HPLC and measured fluorometrically.22 Dopamine, epinephrine and norepinephrine concentrations in EDTA-plasma were determined by reverse phase HPLC with electrochemical detection.23

EDTA-blood samples were frozen twice and thawed for the induction of haemolysis. The haemolysate was diluted 1:3 with bi-distilled water. The analysis of the biogenic amines was then performed as described for the sera.18,19

The blood samples for atopic eczema patients and control persons were drawn at the same time of the day, although a diurnal variation in the DAO/MAO-B levels was not seen.

O-phthalaldehyde, histamine, benzylamine, putrescine, Triton X-100 were obtained from Sigma, München, Germany; 2',7'-dichlorofluorescein diacetate from Serva Heidelberg, Germany; and horse-radish peroxidase from Boehringer-Mannheim, Germany. Other chemicals were reagent grade quality.

**Results and Discussion**

The investigations showed reduced type B monoamine (MAO) and diamine oxidase (DAO) activities in platelet-rich plasma of atopic eczema patients when compared with control subjects (Tables 1 and 2). The difference was highly significant for the first step histamine catabolizing enzyme, diamine oxidase, (p < 0.001) and significant for the methyl histamine catabolizing enzyme monoamine oxidase B, (p < 0.05). Concomitantly with reduction of monoamine and diamine oxidase activities plasma histamine levels were increased (Table 3). High plasma histamine levels in atopic eczema have been reported in fasting subjects as well as after food ingestion.24,25 Low monoamine and diamine oxidase activities may account for increased histamine levels of endogenous or exogenous origin in atopic eczema patients.

The cofactor levels for monoamine oxidase, flavin adenine dinucleotide (FAD) and iron,1 were lowered in AE patients when compared with control subjects (Table 1). Iron values were at the bottom of the normal range. Despite the lowered iron concentrations haemoglobin values were almost normal (Table 1), suggesting that AE patients may have a small iron storage deficiency. FAD in these patients was significantly lowered (p < 0.005) when compared with the control group (Table 1). Monoamine oxidase activity can be restored by iron and vitamin B$\text{$_{2}$}$ supplementation.

The cofactor levels for diamine oxidase, pyridoxal phosphate and copper,26 were almost normal in the atopic group (Table 2). It was therefore concluded that, contrary to monoamine oxidase activity, a reason other than the cofactor deficiency must be responsible for the lowered activity of diamine oxidase. Biogenic amines, food additives, or drugs27,28 are the most probable candidates for inhibition of the activity. Drugs have

### Table 1. Monoamine oxidase activity, cofactor level and haemoglobin concentration in atopic eczema

<table>
<thead>
<tr>
<th></th>
<th>Type B monoamine oxidase (mmol/min/l)</th>
<th>FAD (µg/l)</th>
<th>Iron (µg/dl)</th>
<th>Haemoglobin (g/dl)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Atopic eczema</td>
<td>0.223 ± 0.110 (19)</td>
<td>79.6 ± 31.1 (16)</td>
<td>men 94.6 ± 25.8 (7)</td>
<td>men 13.6 ± 1.5 (7)</td>
</tr>
<tr>
<td>Patients (n)</td>
<td></td>
<td></td>
<td>women 73.6 ± 27.0 (13)</td>
<td>women 13.2 ± 0.9 (13)</td>
</tr>
<tr>
<td>Controls (n)</td>
<td>0.371 ± 0.086 (11)</td>
<td>132 ± 42 (11)</td>
<td>men 130 ± 50*</td>
<td>men 15.5 ± 2.5*</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>women 110 ± 50</td>
<td>women 14 ± 2</td>
</tr>
<tr>
<td>Significance, Student's t-test</td>
<td>p &lt; 0.05</td>
<td>p &lt; 0.005</td>
<td>NS</td>
<td>NS</td>
</tr>
</tbody>
</table>

* Generally accepted normal values.

### Table 2. Diamine oxidase activity and cofactor level in atopic eczema patients and healthy controls

<table>
<thead>
<tr>
<th></th>
<th>Diamine oxidase (mmol/min/l)</th>
<th>Copper (µg/dl)</th>
<th>Pyridoxal phosphate (µg/l)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Atopic eczema</td>
<td>0.270 ± 0.089 (18)</td>
<td>125.3 ± 28.9 (18)</td>
<td>12.1 ± 9.4 (17)</td>
</tr>
<tr>
<td>Patients (n)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Controls (n)</td>
<td>0.511 ± 0.125</td>
<td>115 ± 50*</td>
<td>10.8 ± 7.2 (11)</td>
</tr>
<tr>
<td>Significance, Student's t-test</td>
<td>p &lt; 0.001</td>
<td>NS</td>
<td>NS</td>
</tr>
</tbody>
</table>

* Generally accepted normal values.
Platelet histamine oxidases

<table>
<thead>
<tr>
<th>Table 3. Biogenic amine levels in sera and blood of atopic eczema patients and healthy controls</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Putrescine</strong> (μg/l)</td>
</tr>
<tr>
<td>------------------------</td>
</tr>
<tr>
<td><strong>Sera</strong></td>
</tr>
<tr>
<td>Atopic eczema patients (n = 21)</td>
</tr>
<tr>
<td>Controls (n = 13)</td>
</tr>
<tr>
<td>Significance, Student’s t-test</td>
</tr>
<tr>
<td><strong>Blood</strong></td>
</tr>
<tr>
<td>Atopic eczema patients (n = 12)</td>
</tr>
<tr>
<td>Controls (n = 13)</td>
</tr>
<tr>
<td>Significance, Student’s t-test</td>
</tr>
</tbody>
</table>

to be excluded as the reason for lowered DAO activities, as the patients in our study were free of any drug. However, in spite of normal pyridoxal phosphate levels we found that decreased diamine oxidase activity may be reversed by vitamin B6 supplementation.11

The studies show that the sera of atopic eczema patients have normal biogenic amine (putrescine, cadaverine, spermidine, spermine and tyramine) concentrations, when compared with control individuals (Table 3). In atopic eczema the dopamine (21.2 ± 18.4 vs. control value 16.6 ± 13.7, NS),23 epinephrine (36.4 ± 17.5 vs. control value 43.1 ± 22.2, NS)23 and serotonin (148 ± 81 vs. control value 125 ± 75, NS) (unpublished) concentrations are also normal. The only measured biogenic amine with elevated concentration was norepinephrine (401.3 ± 164.5 vs. control value 174.3 ± 55.8, p < 0.005). The possible reasons for the elevated norepinephrine values have been discussed.23,29 In the same patients reduced MAO-B and DAO activities in platelet-rich plasma (Tables 1 and 2) are paralleled by concomitantly increased histamine concentrations (Table 3) (see also References 32 and 33). The type A and B monoamine oxidases have only small differences in their substrate specificity. Some of the preferred substrates of type A monoamine oxidase are dopamine, norepinephrine, epinephrine, tyramine and serotonin. Type A monoamine oxidase activity might also be lowered. Because of limitations in the sensitivity of the method used for the measurements it was not possible for this to be evaluated directly.

However, the biogenic amines described above are not responsible for the low oxidase activities in these patients, since their concentrations in the sera were normal. Furthermore we found almost normal cofactor levels for DAO but lowered cofactor concentrations for MAO-B (see also Reference 33). This means that the lowered MAO-B activities may be explained by the lowered cofactor concentrations, but this explanation does not hold for the measured DAO values. The remaining candidates for the inhibition of DAO are nicotine, alcohol, food additives27 or other environmental factors. These powerful oxidase inhibitors can also induce a release of histamine (pseudoallergic reaction). The result is a further increase of histamine concentration and an oxidase overload.

Although nicotine or alcohol are able to inhibit DAO activity, the general low activities of patients’ DAO in our study cannot be caused by these drugs, because patients did not smoke or drink any more alcohol during treatment than the control patients.

A short time overload of the oxidases with concomitant competitive inhibition may also be caused by food containing high amounts of biogenic amines. Nevertheless allergic reactions are also responsible for a rise of the histamine levels.

Although mercurials may induce changes in the oxidase activities,34 the blood mercurial concentrations of our patients are identical to values measured in the control persons (0.8 ± 0.5 vs. control value 1.2 ± 0.9 μg/l, NS) and therefore not responsible for the oxidase changes measured in these patients.

Circulating immune complexes and IgE in the patients’ blood activates the coagulation system with elevation of platelet aggregation35 and histamine release with further enhancement of the aggregation—a process probably related to the DAO activities of platelets. A puzzling question may then also be answered: why is the histamine level in the patients’ sera elevated but not the level of the other normally present biogenic amines? The other biogenic amines are presumably catabolized by the amine oxidases outside the platelets. Platelet amine oxidases, essentially DAO, may then serve especially to detoxify high histamine concentrations. Human blood, with 140 to 440 platelets/ml, is the major ‘organ’ for metabolizing circulating histamine. Most other biogenic amines are
catabolized by MAO-A or MAO-B—oxidases that are present in sufficient amounts in, for instance, nerve endings near their target cells. It cannot be ruled out that a genetic defect (including enzyme repression) may be responsible for the present results. Further investigations concerning platelet DAO inhibitory factors are in progress in the authors’ laboratory.

References


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