Cytidine deaminase activity as a measure of acute inflammation in rheumatoid arthritis

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SUMMARY Cytidine deaminase (CD), a cytoplasmic enzyme, is thought to leak out of damaged cells and can be measured in fluids by a simple biochemical assay. This study has shown that serum CD activity is raised in rheumatoid arthritis (RA) compared with osteoarthritis (OA). Synovial fluid (SF) CD activity was always less than the corresponding serum activity (mean SF/serum ratio=0.6) in OA but up to 22 times greater than the corresponding serum activity in RA (mean SF/serum ratio=13.1), suggesting CD production in inflammatory joints. Evidence to support the SF neutrophil as a cell of CD origin is provided by the CD gradient running from cells to SF to synovium. The close correlation between SF CD activity and neutrophil count (r=0.93) indicates that SF CD activity is an accurate measure of acute synovial inflammation. Weak correlation of serum CD activity with erythrocyte sedimentation rate (ESR) (r=0.44) and C-reactive protein (CRP) (r=0.49) implies that CD estimations supply different though related information about rheumatoid disease activity. We suggest that CD released from damaged neutrophils diffuses from all inflamed joints into the blood, so that serum CD activity may provide an integrated measure of joint inflammation more specific than traditional measures such as the ESR.

Key words: cytidine aminohydrolase, nucleoside deaminases, nucleoside aminohydrolases, joint inflammation.

Cytidine deaminase (CD) is a cytoplasmic enzyme that catalyses the hydrolytic deamination of cytidine, deoxycytidine, and their analogues.

\[
\text{cytidine} + \text{H}_2\text{O} \xrightarrow{\text{CD}} \text{uridine} + \text{NH}_3
\]

Although its exact physiological role is uncertain, it may provide a salvage pathway for pyrimidine nucleosides produced during the breakdown of nucleic acids.1

Clinical interest in the enzyme developed when some patients with acute myeloblastic leukaemia proved resistant to treatment with cytosine arabinoside, an antileukaemic drug known to be degraded by CD.2 The finding that their marrow cells had raised levels of CD led to the suggestion that CD inhibition might be used to augment treatment.3

A simple assay has been developed1,4 and the distribution of CD in human tissues determined (Fig. 1).1

In a similar manner to the release of creatine phosphokinase from cardiac muscle during myocardial infarction CD is thought to leak out of damaged cells and can be measured quantitatively in serum. Serum CD levels are raised in a variety of conditions, including rheumatoid arthritis (RA) (Table 1).1

The relationship between serum CD levels and the development of pre-eclamptic toxaemia is now well established, though the organ of origin of CD is uncertain. Estimation of maternal serum CD activity has been recommended as a screening test and is in routine use at several centres.4

This paper reports the results of a study designed to investigate the site of origin and significance of CD activity in RA patients.
Materials and methods

LONDON

Samples of blood, synovial fluid (SF), and synovium were collected from patients with osteoarthritis (OA) and definite or classical RA during routine venepuncture, joint aspiration, or joint replacement at The London Hospital.

Blood was taken into ethylenediaminetetra-acetic acid (EDTA) or plain glass tubes. SF samples collected without anticoagulant were centrifuged at 1000 g for 10 minutes and supernatant separated. In some cases a recorded volume of SF was centrifuged and the pellet which adhered to the bottom of the tubes separated from the supernatant.

Loss of serum enzyme activity has been shown to be minimal if serum is frozen within four hours, but in order to test if the same were true for SF a number of samples were stored at 22°C and aliquots taken at intervals, centrifuged, and the supernatant collected.

All samples were stored at −20°C within four hours of collection. Samples were coded, packed in solid carbon dioxide, and sent from London to Aberystwyth once per fortnight. In all cases samples arrived still frozen within 24 hours of departure.

Synovial fluid leucocyte counts were performed with a haemocytometer (Hawksley BS 748) on samples collected into EDTA, and differential cell counts performed on stained smears of EDTA SF by

### Table 1 Serum CD activity in disease

<table>
<thead>
<tr>
<th>Raised CD activity</th>
<th>Normal CD activity</th>
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<tr>
<td>Septicaemia</td>
<td>Influenza</td>
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<td>Pyelonephritis</td>
<td>Cystitis</td>
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<td>Cellulitis</td>
<td>Polymyositis</td>
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<td>Pneumonia</td>
<td>Obstructive jaundice</td>
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<td>Ulcerative colitis</td>
<td>Pancreatitis</td>
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<td>Rheumatoid arthritis</td>
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<td>Pre-eclampsia</td>
<td>Essential hypertension of pregnancy</td>
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<tr>
<td>Intrauterine death</td>
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<td>Viral hepatitis</td>
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<td>Liver metastases</td>
<td></td>
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<td>Osteomyelitis</td>
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a single observer without knowledge of the fluid CD activity.

Estimations of the erythrocyte sedimentation rate (ESR) were made by the modified Westergren method.

Aberystwyth

Samples were thawed and processed immediately without knowledge of the patients’ diagnosis or condition.

Synovium and cellular pellets were homogenised with distilled water in a tissue grinder. The extract was ultrasonicated for 10 minutes at 8 kHz and then centrifuged to remove debris.1

Estimations of CD activity were carried out with cytidine as a substrate, and the ammonia liberated was measured by a modified Berthelot reaction without protein precipitation. Details of this technique have been published and show high inter- and intrabatch reproducibility (coefficient of variation 1.6% and 2.8% respectively).1

Enzyme activity is expressed as the amount of ammonia produced per unit time. One unit being equivalent to the release of 10⁻⁴ μmol ammonia per minute. For fluids the activity is expressed as units/ml, for tissues as units/g wet weight, and for cells as units/10⁶ cells.

The CD activity in the cellular component of a unit volume of SF was calculated by the following equation:

\[
\text{CD activity in the cells from } 1 \text{ ml SF} = \frac{\text{total CD of SF cell sample}}{\text{volume of SF in ml}}
\]

In an attempt to exclude enzyme release from liver or kidney estimations of γ-glutamyltransferase (γ-GT) and creatinine were made on the serum samples.

C-reactive protein (CRP) was measured by the turbidimetric method with a Hyland laser nephelometer, creatinine by the Technicon AA11 continuous flow system using alkaline picrate, and γ-GT by the calorimetric method as supplied by Boehringer.

Analysis of data employed the techniques of linear correlation and Student’s t test.

Results

Serum

There was a large variation in serum CD activity among the 88 RA patients tested (Fig. 2). The mean CD activity (5.5 units/ml) was significantly higher than in both OA patients (3.0 units/ml) and a reference population of Welsh students1 (2.4 units/ml) (p<0.001).

Fig. 3 CD activity during short term storage of synovial fluid at 22°C.

Fig. 4 Synovial fluid CD activity in OA and RA patients. OA – Mean (SD)=1.8 (0.8) units/ml. RA – Mean (SD)=52.3 (40.0) units/ml. p<0.001.
SYNOVIAL FLUID

From Fig. 3 it can be seen that short term storage had minimal effect on SF CD activity.

SF CD activity was markedly raised in RA patients compared with OA patients (Fig. 4) and correlated well with SF neutrophil count (r=0.93) (Fig. 5).

The CD activity of the cellular component of a unit volume of RA SF (mean=89 units/ml) was greater than the corresponding supernatant CD activity (mean=53 units/ml) (p<0.05), and the cellular and supernatant CD levels correlated well (r=0.86) (Fig. 6).

SYNOVIAL FLUID/SERUM RATIO

SF CD activity was lower than the corresponding serum activity for the OA patients (mean SF/serum ratio=0.6, n=11) but up to 22 times higher for the RA patients (mean SF/serum ratio=13.1, n=16). Fig. 7 shows weak positive correlation between serum and SF CD levels in RA patients (r=0.45).

SYNOVIIUM

For the nine samples of rheumatoid synovium that were processed the mean (SD) CD activity was 53 (48) units/g and the mean (SD) CD activity of the corresponding SF was 25 (34) units/ml.

COMPARISON WITH ESR AND CRP

Comparison of serum CD activity with ESR and CRP was made in 49 cases of RA. In one case the serum γ-GT was raised at 86 IU/l but in all other cases γ-GT and creatinine measurements fell within.

Fig. 6 Unit volume of RA synovial fluid – scattergram. CD cells versus CD supernatant. r=0.86, n=23.
CD Supernatant – Mean (SD)=53 (42) units/ml, n=23.
CD Cells – Mean (SD)=89 (107) units/ml, n=23.

SYNOVIAL FLUID – CD*

Fig. 7 RA patients – scattergram. Serum CD versus synovial fluid CD. r=0.45, n=16.
Cytidine deaminase activity in rheumatoid arthritis

The cellular components of RA SF contained higher CD activity than the corresponding supernatant, but RA synovial CD activity was no greater than that of 15 normal postmortem specimens previously studied (mean (SD)=45 (24) units/g). This suggests a CD concentration gradient running from cells to fluid to synovium. Neutrophils are the predominant cell type in inflammatory SF. It has been shown that their half life is about four hours and estimated that in an inflamed rheumatoid knee the daily neutrophil breakdown in the synovial cavity might well exceed one billion cells. Neutrophils contain 20 times more CD than lymphocytes (Fig. 1), so it seems likely that CD originates from SF neutrophils.

As the SF neutrophil count has long been considered the marker of acute inflammation, the close correlation between it and CD levels (r=0.93) implies that SF CD activity is an accurate measure of the acute inflammatory activity of a joint.

Synovial permeability can be assessed by the ratio of SF/serum concentrations of molecules that are not produced or destroyed in joints. Inflammation increases synovial permeability, so that the ratio of SF/serum for transferrin, a protein manufactured solely by the liver and of similar molecular weight to CD, varied from 0.56 in OA patients to 0.84 in RA patients in one study. Our value of 0.6 for the CD SF/serum ratio of OA patients agrees well with this, suggesting diffusion of CD from blood to SF. In contrast, the SF/serum CD ratios for RA patients (mean=13.1) reflect the high levels of CD activity in SF compared with serum and suggest that CD originates from inflamed joints and diffuses into the blood.

Such diffusion is reflected by the correlation found between CD activity in serum and SF (Fig. 7). Close correlation would not be expected because of the varying contributions from other joints in different patients.

If it is assumed that all synovial joints react to inflammation in a similar manner and that background CD production, metabolism, and excretion occur at constant rates, then serum CD activity will be directly proportional to total neutrophil turnover.

Serum CD levels correlated weakly but significantly with ESR (r=0.45) and rather better with CRP (r=0.49). The relationship is, however, not so strong as to suggest that CD estimation is merely a further reflection of the acute phase response but close enough to support the hypothesis that serum CD activity is a specific measurement of inflammation.

Thus serum CD estimation may offer an integrated measure of acute inflammation in rheumatoid arthritis. Such a laboratory measure would reflect an entirely different dimension of rheumatoid disease activity compared with indexes such as the ESR.

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