Eosinophilic leukaemia in association with a double Philadelphia chromosome

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Summary
A case of eosinophilic leukaemia in association with chromosomal abnormalities including a double Philadelphia chromosome is reported. Comment is also made on the cardiological problems which arise in this condition.

Introduction
Eosinophilic leukaemia (EL) is a rare disorder. It is associated with a marked peripheral blood and marrow eosinophilia and almost invariably with myocardial involvement, frequently leading to cardiac failure. A close affinity with chronic granulocytic leukaemia (CGL) has been suggested and although cytogenetic abnormalities have been recorded in only a limited number of patients, 7 have had a Philadelphia chromosome (Ph' positive). A case of EL with a double Ph' chromosome and an abnormal chromosome 12 is now reported.

Case report
A 59-year-old male presented with a 3-month history of excessive tiredness and weight loss of 14 kg over one year. Physical examination revealed marked hepato-splenomegaly. Investigations: Hb 10.0 g/dl; WCC 27.2 x 10^9/l (neutrophils 6%,

Fig. 1. Karyotype of a bone marrow cell following Trypsin-Giemsa banding: 48,XY, t(9;22) (q34;q11), r(12) (p12q22), +del(22) (q11), +22.
lymphocytes 7%, monocytes 6%, eosinophils 78%, basophils 1%, myelocytes 1%); platelets 30 × 10^9/l. Blood film revealed many large eosinophils with large sparsely scattered granules, hypersegmented nuclei and cytoplasmic vacuolation. Marrow aspirate was hypercellular with 80% cells being eosinophilic in origin but no increase in blast cells, the predominant cell being the eosinophilic myelocyte. Antinuclear factor and rheumatoid factor were both negative, as were chest X-ray and ECG. The echocardiograph, however, showed slight thickening of the left ventricular endocardium. Immunoglobulins including IgE were normal. Neutrophil alkaline phosphatase (NAP) score was normal. Eosinophil function studies showed normal Fc and C3 receptors, which responded normally to enhancement by chemotactic factors.

Chromosome analysis on a marrow aspirate showed 2 Ph' chromosomes, one resulting from a classical 9;22 translocation and one derived from the 22q- translocation chromosome. In addition, one of the no. 12 chromosomes was in a ring form and an extra chromosome 22 was present, giving a karyotype of 48,XY,t (9;22) (q34;q11), r(12) (p12 q22), +del(22) (q11), +22 (Fig. 1).

In view of the possibility of this ‘mature cell’ EL being a variant of CGL, treatment was started with a combination of 6-mercaptopurine 50 mg daily, busulphan 2 mg daily and allopurinol 300 mg daily. This resulted in a rapid reduction in the eosinophil count, eventually down to normal levels, the absolute neutrophil count being relatively unaffected (Fig. 2).

Hepatosplenomegaly regressed but unfortunately, as the eosinophil count was falling, the patient developed left ventricular failure (A on Fig. 2) which proved extremely slow to respond to therapy. Chemotherapy was discontinued at this point but had to be re-instituted some time later after development of a haemorrhagic pleural effusion and rising eosinophil count (B on Fig. 2). Repeat chromosome analysis on bone marrow after 3 months’ therapy showed reversion to a completely normal male karyotype 46,XY. Chemotherapy in a reduced dosage maintained a normal eosinophil count for 12 weeks, but the patient then entered an accelerated phase with a rising eosinophil count and increasing splenomegaly which proved refractory to further treatment, and he required considerable support in terms of blood and platelet transfusions. He developed lobar pneumonia and cardiac failure preterminally, and died 11 months after presentation.

Post-mortem examination confirmed the presence of widespread leukaemic deposits in bone, lymph nodes, spleen and muscle. The heart showed left ventricular dilatation with extensive endomyocardial fibrosis (Fig. 3) and thrombotic vegetations overlying this fibrosis and on the mitral valve. Leukaemic deposits were also noted in the myocardium of the left ventricle.

**Discussion**

It is now accepted that EL exists as a distinct entity and represents the more aggressive end of the spectrum of disease covered by the term ‘hypereosinophilic syndrome’ (Hardy and Anderson, 1968). It has been classified into 3 morphological categories.
—acute blastic, immature, and mature (Benvenisti and Ultman, 1969). The present authors consider that this case falls into the 'mature' category. Although cytogenetic studies have been performed on only a limited number of patients with EL, abnormalities have been frequently uncovered (Weinfeld, Westin and Swolin, 1977). Seven cases have been found with Ph' chromosome and this has led to the suggestion that EL is a variant of CGL and that EL could be similarly classified into PH' negative and Ph' positive groups. This patient had a double Ph' chromosome—a feature frequently found in transforming CGL, and occasionally at the time of diagnosis (International Workshop on Chromosomes in Leukaemia, 1978). In addition, he had an abnormal C group chromosome—12. C group chromosome abnormalities are a frequent finding in addition to the Ph' in classical CGL, and imply a poorer prognosis (Sakurai, Hayata and Sandberg, 1976). This may also be the case in EL. However, the normal NAP score, normal neutrophil count and severe cardio-respiratory problems encountered in this patient are features not usually associated with CGL. Similarly, the striking effect of therapy on the eosinophils with sparing of neutrophils supports the claim of existence of separate precursor cells for both eosinophils and neutrophils. The response to this combination therapy recently shown to be effective in CGL (Allan, Duvall and Stockdill, 1978) was initially good but the normal karyotype encountered during the 'remission' phase with loss of Ph' positivity was unexpected as it is an extremely uncommon occurrence in classical CGL. This suggests that in this case the chromosome abnormalities were confined to the eosinophil precursors.

Endomyocardial fibrosis, a feature associated with hypereosinophilia in other diseases (Chusid et al., 1975) and problems of cardiac failure are known to occur in eosinophilic leukaemia. The rapid lysis of eosinophils achieved by treatment in this case may have precipitated the severe cardiac failure encountered, and emphasizes the necessity for careful reduction of the hypereosinophilia in such cases.

References
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*Postgrad Med J* 1980 56: 268-270
doi: 10.1136/pgmj.56.654.268

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