Collagenase digestion of bone marrow trephine biopsy specimens: an important adjunct to haematological diagnosis when marrow aspiration fails

Z T Maung, N P Bown, P J Hamilton

Abstract

Failure to obtain sufficient material from marrow aspiration (dry tap) posed a diagnostic problem in two patients with pancytopenia. By using collagenase digestion of the trephine biopsy specimen, a precise diagnosis was reached. This technique is very useful because it permits flow cytometric and immunocytochemical analyses of cell suspensions obtained after collagenase digestion of the trephine biopsy specimen core. Acute leukaemia presenting with a dry tap can therefore be accurately immunophenotyped. The technique is easy to perform and merits wider use.

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Conventional cytomorphological analysis of Romanowsky stained bone marrow smears is frequently complemented by cytochemical, immunophenotypic, and cytogenetic studies for complete diagnostic, therapeutic, and prognostic information. But sometimes bone marrow aspiration fails and a dry tap results. Collagenase digestion of a trephine biopsy specimen is easily performed in such situations and results in the garnering of enough cells in a suspension to permit meaningful cytomorphology and immunophenotyping and rapid arrival at a precise diagnosis.1

Methods

A modification of the original method of Mononen and Jansson was used.1 A 2 mm long piece of bone marrow trephine obtained using a Jamshidi needle was cut off the parent biopsy specimen core and incubated for 1 hour at 37°C with 2 units of collagenase (Sigma C2799) in 400 μl of RPMI 1640 medium (Northumbria Biologicals) containing L-glutamine 2 mM, fetal calf serum 10% (v/v), adjusted to pH 7.4-7.6 by 1M HCl. The liberated cells were resuspended in phosphate buffered saline. The cell suspension is suitable for flow cytometry and, after cytocentrifugation, for a range of cytochemical investigations.

Case reports

Case 1
A 51 year old man presented with epistaxis and widespread purpura. A blood count showed that the haemoglobin was 59 g/l, mean cell volume 101 fl, white cell count 2.9 × 10⁹/l, and platelets 8 × 10⁹/l. There were no blasts in the peripheral blood smear. Plasma B₁₂, folate, and red cell folate concentrations were high normal. Dry taps were obtained from both iliac crests. A drop of marrow was expelled from the needle with the stylet and used for cytogenetic analysis. A trephine biopsy specimen was obtained and imprints of the core showed only bare nuclei with no morphological detail. The core was bisected, one part sent for histological examination and the other for collagenase digestion. The digest was cytocentrifuged and stained with May-Grünwald-Giemsa (MGG). This showed multinuclearity and delayed nuclear condensation of erythroid precursors. Myelopoesis was morphologically unremarkable with no excess of blasts. Histological examination of the trephine biopsy specimen showed hypercellular marrow with expansion of red and white cell precursors; no megakaryocytes were identified. Reticulin fibres were greatly increased. In the context of severe peripheral pancytopenia and a hypercellular marrow with no evidence of infiltration, the dysplastic, albeit non-specific, appearance of the erythroid cells suggested a primary myelodysplastic syndrome (MDS) associated with increased marrow fibrosis.2 The diagnosis of MDS was supported by the cytogenetic findings which included trisomy 6, 5q– and 7p– and 21p+, well established markers of MDS and myeloid leukaemia. Over the next four months the refractory anaemia evolved with up to 10% myeloblasts appearing in the blood, the spleen enlarged, and he died from complications of bone marrow failure.

Case 2
A 28 year old man presented with purpura. A full blood count showed that the haemoglobin has 45 g/l, white cell count 6.4 × 10⁹/l with blasts 0.1-1 × 10⁹/l, and platelets 10 × 10⁹/l. Bone marrow aspirate was difficult, yielding a minute quantity of marrow barely sufficient for one smear for MGG staining and a few drops for cytogenetic studies. The aspirate smear showed a heavy monotonous infiltration of undifferentiated blasts. Histological examination of the trephine biopsy specimen confirmed acute leukaemia but could not distinguish lymphoblastic from myeloid form. An aliquot of cells liberated by collagenase digestion of the core was analysed by flow cytometry (FACScan, Becton-Dickinson) using a panel of monoclonal antibodies. The
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Collagenase digestion of bone marrow trephine biopsy specimens has been of value in diagnosis of idiopathic myelofibrosis, myelodysplastic syndromes, myeloproliferative disorders, and chronic myelogenous leukaemia. The digestion allows re-examination of the biopsy specimen, and increased yields of information are obtained. It is important to re-emphasise the value of this technique. It is always difficult to distinguish fibrotic MDS from idiopathic myelofibrosis by inspection of decalcified, paraffin wax embedded biopsy specimens. Characteristic features of MDS, such as dysmegalakoytopoiesis and abnormal localisation of immature precursors (ALIP), have been described, but they are neither restricted to various FAB subtypes of MDS nor easily detected other than in plastic embedded biopsy specimens. The demonstration of dysplastic erythropoiesis by cytology allowed a diagnosis of MDS to be made in case 1. The clonal nature of the disorder was confirmed by the cytogenetic findings and the haematological progression indicated the original diagnosis. It has been reported that cells treated with collagenase retain their cytochemical reactivity, allowing a rapid definitive diagnosis of leukaemia. In case 2 successful use of flow cytometry and immunocytochemistry on collagenase-treated cells made it possible to make a firm diagnosis of pre-B ALL.

Cytogenetic analysis is important for diagnosis, classification, and prognosis of most clonal haematological disorders. Successful cytogenetic analysis of cells from bone marrow biopsy cores has been reported, but the yield of dividing cells suitable for chromosomal analysis has not been high in our hands. In both cases reported here we obtained cytogenetic information from culturing of what little marrow was present in our aspirate needle, relying on obtaining sufficient cells from the collagenase digest for morphological and immunphenotypic studies. The knowledge that useful morphological and immunphenotypic information can be obtained from the digested cells provides an opportunity for more flexible use of the small amount of marrow sometimes obtained in difficult aspirates. There is now little justification for accepting failure to establish a firm diagnosis of leukaemia on the grounds that a difficult aspirate yielded insufficient cells.

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