Abl expression, tumour grade, and apoptosis in chondrosarcoma

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Abstract
Aims—To determine whether Abl immunoreactivity correlates with grade and cell kinetics (apoptosis and mitosis) in chondrosarcoma.

Methods—Sections from 16 chondrosarcomas were stained immunohistochemically using a polyclonal antibody to the c-Abl/Bcr–Abl oncoprotein. Apoptotic indices and mitotic indices were assessed in all tumours. Sections from 24 paraffin wax blocks of human fetal rib (gestational ages, 15–42 weeks) were also stained to determine whether the Abl protein is synthesised consistently throughout endochondral ossification.

Results—Abl staining in immature fetal rib chondrocytes at all stages of development was predominantly nuclear, and 70% of cells showed moderate to strong staining. Abl immunoreactivity was minimal or absent in hypertrophic chondrocytes about to undergo apoptosis at the growth plate. There was strong Abl staining in grade 1 and grade 2 chondrosarcomas but staining was greatly reduced or absent in grade 3 chondrosarcomas. There was a very significant linear correlation between apoptotic index (mean, 0.68%; range, 0–3.2%) and mitotic index (mean, 0.23%; range, 0–0.9%), and both indices were significantly lower in grade 1 than in grade 2 and grade 3 chondrosarcomas.

Conclusions—These data suggest that abl gene expression is associated with differentiation and apoptosis inhibition in fetal and neoplastic chondrocytes. However, these putative effects cannot be ascribed solely to the Abl protein, because several additional factors contribute to the regulation of both differentiation and apoptosis.


Keywords: abl gene expression; chondrosarcoma; apoptosis

The abl oncogenes are cellular homologues of the v-abl oncogene of the Abelson murine leukaemia retrovirus. They encode nuclear and cytoplasmic protein tyrosine kinases that function in signal transduction, cell cycle dependent and DNA damage induced gene expression, and apoptosis inhibition. The biological effects of c-Abl vary in different cell types and may depend on an equilibrium between nuclear and cytoplasmic c-Abl. The bcr–abl fusion gene occurs in 90% of cases of chronic myeloid leukaemia, as a result of reciprocal translocation of c-abl (chromosome 9) to the breakpoint cluster region, bcr on chromosome 22. bcr–abl encodes the chimaeric protein Bcr–Abl, which has enhanced protein tyrosine kinase activity and suppresses apoptosis in the indolent phase of chronic myeloid leukaemia.

Recently, it has been reported that c-Abl proteins, encoded by the ubiquitous c-abl proto-oncogene, also inhibit apoptosis. We have previously reported variable Abl immunoreactivity in a wide range of normal human fetal and adult tissues and tumours. Intense Abl immunoreactivity was seen in fetal and adult chondrocytes and in low grade chondrosarcomas.

The purpose of our study was to determine whether Abl is: (1) synthesised consistently in sequential stages of fetal chondrocyte maturation, and (2) related to grade and tumour cell kinetics (apoptosis and mitosis) in chondrosarcoma.

Materials and methods
TISSUE SAMPLES
Twenty four formalin fixed paraffin wax embedded blocks of normal fetal rib taken at postmortem were obtained from the files of the department of histopathology, Rotunda Hospital, Dublin. Specimens were selected to encompass gestational ages from 15 to 42 weeks. The material was well fixed, showing minimal autolysis.

Chondrosarcomas were retrieved from the files of the departments of histopathology, St James’s Hospital and Mater Misericordiae Hospital, Dublin. Of 20 cases diagnosed from 1989 to 1997, 16 (femur (four), rib (three), ilium (two), humerus (two), scapula (two), fibula (one), sites not recorded (two)) had sufficient well preserved formalin fixed tissue for our study. Tumours were graded using standard criteria recently summarised by Bjornsson and colleagues as follows: six grade 1, five grade 2, and five grade 3 (including one dedifferentiated chondrosarcoma). From each case, one paraffin wax block was selected for Abl immunostaining and assessment of apoptosis and mitosis.

IMMUNOHISTOCHEMISTRY
Paraffin wax sections (4 µm thick) were cut and mounted on 3-aminopropyltriethoxysilane coated slides, oven dried, dewaxed in xylene, and rehydrated in alcohol. The immunostaining protocol has been described previously. Briefly, sections were incubated with a polyclonal sheep anti-c-Ab/Bcr–Abl oncprotein antibody (Serotec, Oxford, UK; 1/200 dilution), for one hour at room temperature, then incubated with biotinylated rabbit anti-sheep antibody (Serotec; 1/400), followed by peroxidase conjugated
streptavidin (1/400; Dako, Glostrup, Denmark). The reaction product was visualised with 3,3'-diaminobenzidine and a haematoxylin counterstain. Appropriate positive and negative controls were used, as described previously. Abl immunoreactivity was assessed using a three tiered scale: + (focal or diffuse weak staining), ++ (moderately intense staining), and +++ (intense staining).

APOPTOSIS AND MITOSIS ASSESSMENT
The extent of apoptosis was assessed in all 16 tumours by counting tumour cells in 10 high power fields using the ×40 objective with a 100 square graticule. We decided to use haematoxylin and eosin stained sections, and not in situ end labelling of fragmented apoptotic cell DNA, because apoptosis and mitosis could be assessed in identical microscopic fields. The apoptotic index assigned to each sample represents the total number of apoptotic tumour cells and apoptotic bodies expressed as a percentage of the total number of tumour cells in 10 fields. The mitotic index was determined using the same procedure.

From raw data, the relation between the apoptotic index and the mitotic index was analysed using parametric (Pearson) linear correlations (Instat, Graphpad Software, San Diego, California, USA). From contingency tables, Fisher’s exact test was used to compare grade 1 chondrosarcomas and grades 2/3 chondrosarcomas (rows) with apoptotic index = 0 and apoptotic index > 0 (columns), and also with mitotic index = 0 (rows) and mitotic index > 0 (columns).

Results
FETAL CARTILAGE
Abl staining was predominantly nuclear, although cytoplasmic or cell membrane staining was found occasionally. Most epiphyseal reserve chondrocytes and proliferating immature chondrocytes in fetuses of all ages showed moderate to strong Abl immunoreactivity (fig 1A). In contrast, hypertrophic chondrocytes about to undergo apoptosis during enchondral ossification showed minimal or no Abl staining (fig 1B). There was intense Abl staining of adjacent proliferating osteoblasts and invading metaphyseal blood vessels, as seen previously.

ABL EXPRESSION, GRADE, APOPTOSIS, AND MITOSIS IN CHONDROSARCOMA
In all six grade 1 chondrosarcomas, more than 70% of tumour cells showed intense (3+) nuclear Abl staining (table 1). Grade 2 chondrosarcomas showed variably intense nuclear Abl staining: 3+ in one tumour and 2+ in the other four samples (fig 1C). In grade 3 chondrosarcomas, Abl staining was negative (fig 1D) or, in one case, very weak (1+). The dedifferentiated chondrosarcoma showed intense (3+) Abl immunoreactivity in the low grade component, but no staining in the high grade component (fig 1E).

Mean apoptotic indices and mitotic indices (and ranges) for the 16 chondrosarcomas (table 1) were 0.68% (0–3.2%) and 0.23% (0–0.9%), respectively. In all tumours except one the number of mitoses was less than or equivalent to the number of apoptotic tumour cells. There was a very significant linear correlation between the apoptotic and mitotic indices (p < 0.0090; r = 0.6294).

Figure 1 (A) Immature chondrocytes in the developing human fetal rib show nuclear, cell membrane, or cytoplasmic immunoreactivity for Abl. (B) Hypertrophic chondrocytes show minimal or no Abl staining in sites of enchondral ossification, whereas osteoblasts and proliferating blood vessels (below) are stained intensely. (C) Variably intense Abl immunoreactivity in grade 2 chondrosarcoma. (D) Abl immunoreactivity is absent in grade 3 chondrosarcoma. (E) Numerous apoptotic tumour cells in the high grade component of dedifferentiated chondrosarcoma.
The extent of both apoptosis and mitotic activity correlated with chondrosarcoma grade (fig 1E; table 1). There was no apoptosis or mitotic activity in grade 1 chondrosarcomas. Grade 2 chondrosarcomas (apoptotic index, 0.5%; range, 0–1.9%; mitotic index, 0.16%; range, 0–0.6%) and grade 3 chondrosarcomas (apoptotic index, 1.7%; range, 0.3–3.2%; mitotic index, 0.6%; range: 0.3–0.9%) combined had significantly higher apoptotic indices (p < 0.0114) and mitotic indices (p < 0.0114) than grade 1 tumours.

<table>
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<th>Case no.</th>
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In summary, our results suggest that Abl immunoreactivity correlates with differentiation in both fetal chondrocytes and chondrosarcomas. Absent Abl staining correlates with terminal differentiation and apoptosis in developing cartilage and with high chondrosarcoma grade. However, the anti-apoptotic effects of Abl cannot be ascribed solely to Abl, because several additional factors contribute to the regulation of both differentiation and apoptosis.
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