Perineal rhabdomyosarcoma in a newborn child: pathological and biochemical studies with emphasis on contractile proteins

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SUMMARY Histological and ultrastructural studies have been undertaken on a perineal rhabdomyosarcoma from a newborn child. The spontaneous tumour has the typical feature of mesenchymoma. The recurrent tumour, however, displays some rhabdopoietic characteristics. The myosin of the recurrent tumour has been extracted and compared with human fetal myosin. These two myosins are identical in their synthetic filaments and their light-chain composition. Nevertheless, whereas the ATPase activity of fetal myosin can be stimulated considerably by increasing the Ca²⁺ concentration, that of tumoral myosin remains very low. These results show that there are isoenzymes of myosins and there must be differences in the myosin heavy chains, particularly in the active sites. These findings are identical with those seen in experimental rhabdomyosarcoma.

Human rhabdomyosarcoma is considered to be a relatively rare tumour. Investigation of this tumour during the last 15 to 20 years has shown, however, that embryonic and alveolar rhabdomyosarcomas are the two soft tissue tumours seen most frequently in children. In childhood, rhabdomyosarcoma accounts for 12 to 56% of all soft tissue tumours. The tumours may appear at any age in childhood, but the majority are seen in infants from 3 to 4 years of age. In children aged 10 to 12 years, rhabdomyosarcomas are less common, but they occur more frequently in adolescents 15 to 18 years old. Rhabdomyosarcomas are exceptional in the newborn. The high incidence of this tumour in these reported series reflects the greater confidence with which the diagnosis can now be made. This is to a great extent the result of electron microscopy. Normal myoblasts grown in vitro display mobility and contract. This is not so with rhabdomyoblasts grown in culture; although one of the most typical characteristics of the tumour is lack of contractility, little attention has been paid to their contractile proteins. In previous investigations, tumoral myosins isolated from NlaS₂-induced rhabdomyosarcomas in rats and rabbits were analysed and compared with normal and fetal skeletal myosins. Tumoral myosin was found to be a new isoenzyme with oncofetal character. The present report deals with the study of myosin from a spontaneous human tumour and compares it to myosin from experimentally induced tumours.

Case report

The tumour was seen in a full-term newborn girl weighing 3 kg at birth. She was the first child, and the parents had no genetic abnormalities. A vulvo-perineal tumour, 10 cm in diameter, was found at birth. The tumour pushed back the labia majora. In addition, the left tibia showed a focus of osteolysis. The tumour was excised on the fifth day of life. It was hard and vascularised and had a lobulated structure. During surgery the bladder, vagina, and rectum were completely freed from the tumour which weighed 180 g. The infant died aged 3 months with a local recurrence of the same size as the primary tumour, which was excised immediately after death.

Material and methods

LIGHT MICROSCOPY

Portions of tissue were taken and fixed in Dubosc-
Brasil's fluid and in formaldehyde. The fixed material was embedded in paraffin, and sections 5-8 μm thick were stained with haematoxylin and eosin, Masson's trichrome, Mallory's phosphotungstic acid haematoxylin, Wilder's reticulin, and Heidenhain's azan.

**Electron Microscopy**

The primary tumour was cut into serial sections which were fixed in formalin for 72 hours. Small pieces were then minced into 1 mm cubes and placed in 10% Millonig's buffered formalin. Small tissue portions (about 1 mm³) of the recurrent tumour were fixed for 1 hour in 5% glutaraldehyde in Millonig's buffer. The samples of both tumours were then postfixed for 1 hour in 2% Millonig's buffered OsO₄, dehydrated in a graded series of ethanols, and embedded in styrene-methacrylate as described by Stockem and Komnick. Thin sections were cut on a LKB ultratome III and placed on Pioloform F coated copper grids. After staining with uranyl acetate and lead citrate, sections were examined with a Philips 300 electron microscope at an accelerating voltage of 80 kV.

**Myosin Preparations**

Normal adult rabbit muscles as controls were taken from the thigh of healthy animals. For comparative purposes, fetal muscles were taken from 3-month-old human fetuses. Normal, fetal, and tumoral muscles were fragmented in a Waring-Blender homogeniser for 2 minutes at high speed. The myosins were extracted with a modified Hasselbach-Schneider solution at high ionic strength and in the presence of 40 mM K₃P₂O₇. They were then purified by suspension-reprecipitation cycles and by ammonium sulphate fractionation using the 40 to 50% saturated ammonium sulphate fraction. All details were described in a previous investigation.

Protein concentrations were determined by the biuret method as well as with the molar extinction coefficient E₁% = 5-60. The latter values were compared with those obtained by the biuret method and found to be accurate when the A₂₈₀nm/A₂₆₀nm ratio was 1-40 or greater.

Reprecipitated or synthetic filaments were obtained by rapid dilution from 500 mM KCl, 10 mM MgCl₂, pH 6-8, to 100 mM KCl, 10 mM MgCl₂, pH 6-8, and examined in a Philips 300 electron microscope after negative staining with uranyl acetate.

Sodium dodecyl sulphate (SDS)-polyacrylamide gel electrophoresis using Laemmli's buffer system and Kerckhaert's apparatus was performed on 10 to 25% acrylamide gradient slab gel (15 x 10 x 0.1 cm).

The conditions for Ca²⁺ATPase activity measurements were those described previously.

**Results**

**Morphology of Primary Tumour**

Histologically, the neoplasm displayed mesenchymatous features. Most of the tumour was composed of undifferentiated spindle-celled elements (Fig. 1), the character of which could not be determined: possibly fibroblastic cells, tumour cells of smooth muscle origin, transformed Schwann cells, or elongated rhabdomyoblasts. After exhaustive examination of histological sections, different regions were found which indicated a rhabdomyosarcomatous origin. There were areas with an appearance of undifferentiated embryonic rhabdomyosarcoma and others of the botryoid type (Fig. 2). The low frequency of rhabdomyoblasts was confirmed by electron microscopy, where only few cells with myofibrillar differentiation were found (Fig. 3).

![Fig. 1 General histological appearance of the primary tumour. 90% of the examined surface displays undifferentiated spindle-like cells. Haematoxylin and eosin × 200.](http://jcp.bmj.com/).
MORPHOLOGY OF RECURRENT TUMOUR

The same histological features occurred in the recurrent tumour but in different proportions. Botryoid areas of embryonic rhabdomyosarcoma were particularly developed at the periphery of the tumour; in the centre, the undifferentiated type, characterized by the presence of long fusiform cells (Masson's myocytes) and large polymorphic cells (Masson's myoblasts) (Fig. 4), was prominent. The latter cells displayed a fibrillar network in their cytoplasm.

Electron microscopy revealed the existence of myofibrillar structures in practically all the cells. The most differentiated stage that could be observed showed only myofilament alignment in the form of myofibrils without precise orientation. Z-lines were very hypertrophied. Sarcomeres were very irregular and generally found to be in a supercontracted state. M-lines were not observed. Large quantities of glycogen were present, especially in the vicinity of Z-lines. The nuclei often contained nuclear bodies of fibrillar and/or granular structure. An electron dense, amorphous material was frequently observed in the sarcoplasmic reticulum.

Fig. 2 Botryoid type region of embryonic rhabdomyosarcoma in the primary tumour. H and E × 100.

Fig. 3 Rare rhabdomyoblast of the primary tumour with myofibrillar differentiation. Myofilaments have no precise orientation. × 13700.
TUMORAL AND FETAL MYOSINS

Tumoral myosin was extracted from 80 g of the recurrent tumour. Fetal myosin was isolated from 60 g fresh human fetal muscle.

The synthetic filaments

In classical terminology, reprecipitated myosin filaments are called synthetic filaments. The myosins were precipitated by rapid dilution in the absence and in the presence of 10 mM MgCl₂. The two myosins formed in the absence of MgCl₂ were seen to be exclusively short bipolar filaments (Fig. 5a, b) such as myosin of normal skeletal muscle of rat and rabbit precipitated under the same conditions. The filaments varied in length from 0.3 to 0.6 μm.

The width was about 15 nm at the bare zone and approximately 50 nm at the extremities. In the presence of MgCl₂, however, the tumoral and fetal myosins precipitated essentially as long fusiform filaments with an average width of 25 nm (Fig. 5c, d). The length of the filaments was quite variable for both myosins. The smallest filaments measured 0.6 μm, the longest 10 μm or greater. The majority of the filaments did not exceed 2 μm. There was no possibility of isolating myosin from normal human adult skeletal muscle.

Analytical gel electrophoresis

The tumoral and fetal human myosins were analysed by SDS-gel electrophoresis and compared with rabbit white skeletal muscle myosin.

In the gel of rabbit myosin (Fig. 6a), myosin heavy chains, M-proteins, C-protein, actin, the three light chains of 25 000, 18 000, and 16 000 daltons, and troponin-T, tropomyosin, and troponin-I as contaminants could be recognised. The human tumoral myosin (Fig. 6b) as well as the human fetal myosin (Fig. 6c) contained only two light chains. Other proteins, such as β-actin and troponin C, could be observed in these gels. The 27 000 daltons protein of human tumoral myosin and the 17 000 daltons protein of human fetal myosin could not be identified.

Furthermore, the gel of tumoral human myosin (Fig. 6b) revealed various proteins, the molecular size of which was found to be between 140 000 and 45 000 daltons and between 34 000 and 25 000 daltons. These proteins were probably polypeptid fragments of the heavy chains. Indeed, many proteolytic enzymes are known to exist in tumours.

Ca²⁺ATPase activity

Measurements of the specific Ca²⁺ATPase activity of the two different human myosins were performed and the results compared with the specific Ca²⁺ATPase activity of rabbit white muscle myosin and of rat red muscle myosin. The comparison with rat red muscle myosin seemed to be necessary as human muscle is also a red muscle. Human fetal myosin has a specific activity of Ca²⁺ATPase somewhat lower than that of rat myosin (Table). The specific Ca²⁺ATPase activity of human tumoral myosin was very much decreased. These results are in agreement with the values found for Ca²⁺ATPase activity of rabbit and rat tumoral myosins. 14, 15

Discussion

In the rhabdomyosarcoma examined in this investigation, the essential morphological characteristics that have already been described thus far for chemically induced muscle tumours in rats 26–28 and rabbits 29 or for spontaneous human rhabdomyosarcomas 30 have been observed.

In contrast to the recurrent tumour, myofibrils were extremely rare in the primary tumour, and only a careful examination of many histological sections established its rhabdomyosarcomatous nature. Light microscopy and the ultrastructure of the recurrent tumour confirmed its nature. Nevertheless it is difficult to understand why in the recurrent tumour only one predominant cell type, muscle cells, can be observed.

As described previously in chemically induced rhabdomyosarcomas, irregularly arranged myofibrils with supercontracted sarcomeres were obser-
Fig. 5 Synthetic filaments of human fetal and tumoral myosins (negative staining). In the absence of MgCl₂, fetal (a) and tumoral (b) myosins form bipolar filaments. When MgCl₂ is added, fetal (c) as well as tumoral (d) myosins precipitate as long fusiform filaments. All micrographs: original magnification × 120 000.
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Maximal values of Ca\(^{2+}\) ATPase activity of different normal, fetal, and tumoral myosins

<table>
<thead>
<tr>
<th>Material</th>
<th>Myosins</th>
<th>Maximal values (nmol Pi/mg/min)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Human</td>
<td>Fetal</td>
<td>414 ± 10</td>
</tr>
<tr>
<td></td>
<td>Tumoral</td>
<td>90 ± 8</td>
</tr>
<tr>
<td>Rat*</td>
<td>Normal</td>
<td>450 ± 60</td>
</tr>
<tr>
<td></td>
<td>Fetal</td>
<td>470 ± 21</td>
</tr>
<tr>
<td>Rabbit*</td>
<td>Normal</td>
<td>546 ± 48</td>
</tr>
<tr>
<td></td>
<td>Fetal</td>
<td>510 ± 35</td>
</tr>
<tr>
<td></td>
<td>Tumoral</td>
<td>30 ± 10</td>
</tr>
</tbody>
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*From Hildebrand et al.*

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The absence of the M-line, necessary for the maintenance of the sarcomere, revealed a deficiency of M-protein synthesis. The function of the granular precipitate in the sarcoplasmic reticulum was not clear. The precipitate was believed to be a calcium carbonate or phosphate salt and has been observed in other rhabdomyosarcomas as well as in Ni\(_{3}\)S\(_{2}\)-induced leiomyosarcomas. The presence of this substance in the cisternae may be related to a defect in calcium transport which prevents normal muscle contractile function. It is noteworthy that Gilman has demonstrated that myoblasts grown in vitro from nickel-induced rhabdomyosarcoma fail to achieve contractility. The role of calcium in triggering or potentiating normal and neoplastic cell division in relation to an alternative role in neuromyal transmission and contraction was shown previously by Balk.

Human fetal and tumoral myosins can be compared with those of fetuses and Ni\(_{3}\)S\(_{2}\)-induced rhabdomyosarcomas of rat and rabbit. The same synthetic filaments are observed in both myosins precipitated as short bipolar filaments without MgCl\(_{2}\) in the precipitating buffer but as long fusiform filaments when MgCl\(_{2}\) is added. Moreover, analogies are found in the light-chain composition: the two myosins possess two light chains with a molecular size of 25,000 and 18,000 daltons, whereas normal muscle myosin has three light chains. A clear difference is finally observed between the specific activities of Ca\(^{2+}\) ATPase of human fetal and tumoral myosins. Thus myosin of human spontaneous rhabdomyosarcomas can be considered as an isoenzyme comparable to the myosins of experimentally induced rhabdomyosarcomas of rat and rabbit.

The experimental model seems to be convenient for further studies in molecular biology on the control of synthesis of tumoral myofibrillar proteins. On a more general level, one can believe that the molecular mechanisms occurring in spontaneous or chemically induced tumoral transformation are comparable.

It is noteworthy that these biochemical results can be obtained in less than three days, and they may be useful for diagnostic purposes, especially in embryonic rhabdomyosarcomas, the diagnosis of which is often rather difficult. In addition, the method may be used to distinguish undifferentiated rhabdomyosarcomas from fibrosarcomas because of the different light chain composition of the myosins of these tumours: rhabdomyosarcoma myosin possesses two light chains of skeletal myosin (25,000 and 18,000 daltons); fibrosarcoma myosin, however, has two light chains completely different (20,000 and 15,000 daltons) similar to fibroblast myosin.

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Contractile proteins of human rhabdomyosarcoma

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