**N-my whole myc gene product expression in neuroblastoma**

H HASHIMOTO,* Y DAIMARU,* M ENJOJI,* A NAKAGAWARA

*From the* *Second Department of Pathology and the Department of Paediatric Surgery, Faculty of Medicine, Kyushu University, Fukuoka, Japan*

**SUMMARY** The presence and distribution of *N-my whole myc* gene product were studied in 13 neuroblastomas and five ganglioneuroblastomas, using immunohistochemical techniques. Nine tumours (eight neuroblastomas and one ganglioneuroblastoma of composite type) contained neuroblastoma cells with positive nuclei for *N-my whole myc* protein. Microscopic examination showed that most of the positive neuroblastoma cells seemed to be immature, with no apparent neuronal differentiation. Nine of 11 tumours with amplified *N-my whole myc* gene copies exhibited tumour cells with positive immunostaining for the *N-my whole myc* gene product, while none of the seven non-*N-my whole myc* amplified cases contained immunoreactive tumour cells. The survival of the patients positive for *N-my whole myc* protein was significantly low compared with that of the negative ones. It is concluded that immunohistochemical staining for the *N-my whole myc* gene product will facilitate prediction of the prognosis of patients with neuroblastoma.

The *N-my whole myc* gene was originally identified in human neuroblastoma cell lines and was grouped as a proto-oncogene because of its partial homology to the proto-oncogene c-myc. Close correlation between genomic amplification of the *N-my whole myc* gene in neuroblastomas and the stage and aggressiveness of the tumours was first reported by Brodeur *et al.* and Seeger *et al.* The key role of *N-my whole myc* gene amplification in determining the biological behaviour of neuroblastomas has also been given further support. The enhanced expression of *N-my whole myc* has also been detected in relation to malignant progression, using Northern blot analysis or in situ hybridisation techniques. Recently, antibodies against genetically engineered *N-my whole myc* gene products were prepared and these tools paved the way for retrospective immunohistochemical studies on the *N-my whole myc* gene product. We used immunohistochemical techniques to evaluate distribution of the *N-my whole myc* gene product on paraffin wax sections of neuroblastomas and compared our findings with the number of *N-my whole myc* gene copies and how this affected prediction of the prognosis of the patients.

**Material and methods**

Between 1979 and 1986 13 cases of neuroblastoma and five of ganglioneuroblastoma (two composite, two well differentiated, one poorly differentiated) were selected for study. In all tumours the number of copies of *N-my whole myc* gene had been determined by Southern blot analysis, using radiolabelled *N-my whole myc* (NB-19-21), the details of which have been reported elsewhere.

Five micron sections of 10% formalin fixed, paraffin wax embedded material were mounted on slides coated with poly(L-lysine) for immunohistochemical study. After the tissue sections had been deparaffinised and the endogenous peroxidase activity blocked by using a short 10 minute oxidation procedure in 0.5% periodic acid, followed by treatment with a freshly prepared 0.02% sodium borohydrate for 30 minutes, the avidin-biotin peroxidase complex (ABC) method of Hsu *et al.* was used to determine localisation of the *N-my whole myc* gene product. The slides were immersed in 10% normal goat serum for 30 minutes to eliminate the non-specific binding of immunoglobulins and then incubated overnight at 4°C with the affinity purified anti-human *N-my whole myc* gene product rabbit IgG (MBL, Nagoya, Japan) diluted 1/10. The antibody was produced against the genetically engineered human *N-my whole myc* gene product. After a buffer wash 50 μg/ml of the biotinylated anti-rabbit goat immunoglobulin was applied for 30 minutes. The avidin-biotin peroxidase complex (Vector, Burlingame, California, USA) was applied for an additional 60 minutes at room temperature. The site of localisation of the peroxidase label was then identified by formation of a brown reaction product, using hydrogen peroxidase and diaminobenzidine. These sections were lightly counterstained with methyl green.

Accepted for publication 21 July 1988
N-myc gene product expression in neuroblastoma

Table  Clinical profiles and immunohistochemistry for N-myc gene product

<table>
<thead>
<tr>
<th>Case No</th>
<th>Age</th>
<th>Sex</th>
<th>Primary site</th>
<th>Stage</th>
<th>Histology</th>
<th>No of N-myc copies</th>
<th>N-myc gene product</th>
<th>Follow up</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>11 y</td>
<td>M</td>
<td>Adrenal gland</td>
<td>III</td>
<td>Neuroblastoma</td>
<td>130</td>
<td>++ +</td>
<td>Died 7 m</td>
</tr>
<tr>
<td>2</td>
<td>2y 9m</td>
<td>F</td>
<td>Adrenal gland</td>
<td>III</td>
<td>Neuroblastoma</td>
<td>113</td>
<td>++</td>
<td>Died 6 m</td>
</tr>
<tr>
<td>3</td>
<td>4y</td>
<td>M</td>
<td>Adrenal gland</td>
<td>IVA</td>
<td>Neuroblastoma</td>
<td>112</td>
<td>++ + +</td>
<td>Died 4 m</td>
</tr>
<tr>
<td>4</td>
<td>1y 7m</td>
<td>F</td>
<td>Adrenal gland</td>
<td>IVA</td>
<td>Neuroblastoma</td>
<td>92</td>
<td>+</td>
<td>Died 9 m</td>
</tr>
<tr>
<td>5</td>
<td>1y 9m</td>
<td>M</td>
<td>Adrenal gland</td>
<td>II</td>
<td>Neuroblastoma</td>
<td>71</td>
<td>+</td>
<td>Died 2 5 y</td>
</tr>
<tr>
<td>6</td>
<td>2y 6m</td>
<td>F</td>
<td>Adrenal gland</td>
<td>IVA</td>
<td>Neuroblastoma</td>
<td>37</td>
<td></td>
<td>Died 5 m</td>
</tr>
<tr>
<td>7</td>
<td>1y 1m</td>
<td>M</td>
<td>Adrenal gland</td>
<td>III</td>
<td>Neuroblastoma</td>
<td>37</td>
<td>+</td>
<td>Died 4 m</td>
</tr>
<tr>
<td>8</td>
<td>1y 3m</td>
<td>F</td>
<td>Adrenal gland</td>
<td>IVA</td>
<td>Neuroblastoma</td>
<td>36</td>
<td>+</td>
<td>Died 8 m</td>
</tr>
<tr>
<td>9</td>
<td>1y 9m</td>
<td>F</td>
<td>Adrenal gland</td>
<td>III</td>
<td>Ganglioneuroblastoma, composite</td>
<td>32</td>
<td>+</td>
<td>Died 1 7 y</td>
</tr>
<tr>
<td>10</td>
<td>1y</td>
<td>F</td>
<td>Adrenal gland</td>
<td>IVA</td>
<td>Neuroblastoma</td>
<td>8</td>
<td>+</td>
<td>Alive 3 7 y</td>
</tr>
<tr>
<td>11</td>
<td>5y 8m</td>
<td>F</td>
<td>Adrenal gland</td>
<td>IVA</td>
<td>Ganglioneuroblastoma, composite</td>
<td>6</td>
<td></td>
<td>Alive 4 5 y</td>
</tr>
<tr>
<td>12</td>
<td>7m</td>
<td>F</td>
<td>Retroperitoneum</td>
<td>IVA</td>
<td>Ganglioneuroblastoma, poorly differentiated</td>
<td>1</td>
<td>-</td>
<td>Alive 3 6 y</td>
</tr>
<tr>
<td>13</td>
<td>3y</td>
<td>F</td>
<td>Mediastinum</td>
<td>II</td>
<td>Ganglioneuroblastoma, well differentiated</td>
<td>1</td>
<td>-</td>
<td>Alive 3 5 y</td>
</tr>
<tr>
<td>14</td>
<td>9y</td>
<td>F</td>
<td>Pelvic cavity</td>
<td>III</td>
<td>Ganglioneuroblastoma, well differentiated</td>
<td>1</td>
<td>-</td>
<td>Alive 3 1 y</td>
</tr>
<tr>
<td>15</td>
<td>7m</td>
<td>F</td>
<td>Mediastinum</td>
<td>II</td>
<td>Neuroblastoma</td>
<td>1</td>
<td>-</td>
<td>Alive 2 2 y</td>
</tr>
<tr>
<td>16</td>
<td>4m</td>
<td>M</td>
<td>Retroperitoneum</td>
<td>III</td>
<td>Neuroblastoma</td>
<td>1</td>
<td>-</td>
<td>Alive 2 2 y</td>
</tr>
<tr>
<td>17</td>
<td>6y</td>
<td>M</td>
<td>Adrenal gland</td>
<td>IVA</td>
<td>Neuroblastoma</td>
<td>1</td>
<td>-</td>
<td>Alive 1 8 y</td>
</tr>
<tr>
<td>18</td>
<td>6m</td>
<td>M</td>
<td>Adrenal gland</td>
<td>I</td>
<td>Neuroblastoma</td>
<td>1</td>
<td>-</td>
<td>Alive 1 6 y</td>
</tr>
</tbody>
</table>

Abbreviations: Prevalence of positive tumour cells: ++ + +, almost all; ++ , a moderate number; +, a few; -, none.

We examined the kidneys, adrenal glands, and sympathetic ganglia from three patients with neuroblastoma as controls. The N-myc gene product did not stain any of the following: nephrons; parenchymal cells of the adrenal cortex; ganglion cells and stromal elements such as vascular endothelial cells; smooth muscle cells; fibroblasts; and Schwann cells.

Results

The staining results for N-myc gene product, together with the clinical data and the number of N-myc gene copies are given in the table. In nine of the 18 tumour cells were positive for the N-myc gene product. Of these nine, two were positive in almost all of the tumour cells (fig 1) and four contained a moderate number of positive cells (fig 2). In the remaining three positive cells were rare. Brown benzidine products in the immunoperoxidase reaction for N-myc gene product were observed only in the nuclei of the neuroblastoma cells. Almost all of the positive neuroblastoma cells were immature neuroblasts, and no ganglion cells and few differentiated neuroblasts cells with nuclear enlargement and cytoplasmic eosinophilia and enlargement with distinct cell border13 were immunoreactive to the N-myc gene product. Except for one ganglioneuroblastoma of a composite type with a few positive immature

![Fig 1](http://jcp.bmj.com/)

Almost all neuroblastoma cells are positive for N-myc protein. Immunoreaction product is confined to the nuclei of those tumour cells. (ABC technique.)

![Fig 2](http://jcp.bmj.com/)

Scattered neuroblastoma cells strongly positive for N-myc protein. (ABC technique.)
neuroblasts, tumours in the positive cases were classic neuroblastomas.

Among 11 with amplified N-myc gene copies, nine had tumour cells positive for the N-myc protein. The remaining two were stage IVA neuroblastoma and stage IVA ganglioneuroblastoma of the composite type, respectively, both with no positive cells. These two tumours were composed of more differentiated neuroblastic cells than the other amplified ones. In contrast, there were no immunoreactive tumour cells in the seven non-amplified cases.

Tumour cells positive for N-myc protein were not observed in any of the four patients under 1 year of age, but such cells were present in nine (64%) of 14 patients over 1 year of age. Of nine with N-myc positive tumour cells in the tumour, all but one died within one year after surgery. The one exception was a patient with a stage IVS tumour; the child was still alive 44 months after surgery. Of the nine with an N-myc protein negative tumour, one died and tumour recurred in the other. The remaining seven were alive at the time of writing with no evidence of disease, with a follow up ranging from 19 to 54 months. There was a significant difference in the survival curves (generalised Wilcoxon test) between patients with an N-myc protein positive neuroblastoma and those with an N-myc protein negative tumour (p < 0.01).

Discussion

In 1986 Ikegaki et al identified and characterised the N-myc gene product in human neuroblastoma cells, using monoclonal antibodies with defined specificities. These were prepared against a bacterially expressed fusion protein containing a portion of the N-myc sequence.\(^\text{10}\) The relative level of the polypeptides was roughly proportional to the level of N-myc transcripts present in a panel of neuroblastoma cell lines. We applied an affinity purified rabbit antibody against the genetically engineered human N-myc oncogene product prepared by Shimatake et al\(^\text{11}\) to paraffin wax sections. Those authors reported that; (i) Western blot analysis showed the presence of the N-myc protein (63 kd) in the nuclear fraction of the neuroblastoma cells examined; (ii) immunohistochemically, the N-myc protein was distributed in the nuclei of neuroblastoma cells obtained from various culture cell lines, a xenograft, and a surgical specimen; (iii) the anti-N-myc antibody did not react with cells producing c-myc in the enzyme linked immunosorbent assay (ELISA), or in immunohistochemical tests. In nine of our 18 tumours, there were tumour cells positive for the N-myc gene product. The immunoreaction products were recognised only in the nuclei of neuroblastoma cells, with virtually no staining in the cytoplasm of these cells or normal ganglion cells, nephrons, and vascular or other stromal elements. Almost all the positive cells seemed to be primitive neuroblasts and were not observed among neuroblastic cells showing differentiation, even in the same tumour. A close relation of N-myc oncogene positivity to undifferentiated phenotypes in neuroblastoma cells is compatible with the finding that in situ hybridisation to sections of neuroblastomas showed a high expression of N-myc, predominately in the undifferentiated neuroblasts.\(^\text{17}\) The expression of N-myc was found to be greatly decreased before morphological evidence of differentiation when primitive human neuroblastoma cell lines were differentiated in vitro by the addition of retinoic acid.\(^\text{14,15}\) The N-myc oncogene may therefore have an important role in the oncogenicity, proliferating ability, and regulation of cell differentiation of neuroblastomas.

Increased production of N-myc protein seems to correlate closely with N-myc gene amplification and a poor prognosis. Nine of the 11 neuroblastic tumours with the amplified N-myc gene copies contained N-myc protein positive tumour cells, while none of the seven non-amplified cases showed immunostaining for N-myc. Because the N-myc gene product has been reported to have a short half life (30–40 minutes),\(^\text{10}\) the negativity for the N-myc protein in the two cases with amplified N-myc gene copies may have been the result of a loss of intranuclear N-myc protein by the prolonged lapse of time until fixation or delayed permeation of fixatives. Even though there may be some loss of the antigen, patients with an N-myc immunoreactive neuroblastoma had a significantly poorer prognosis than did those with a neuroblastoma negative for N-myc protein. These results suggest that amplification and consequent increased transcripts of the N-myc oncogene may be related to the aggressiveness of neuroblastomas. Immunohistochemical examination for the N-myc gene product in neuroblastomas is thus useful for predicting the prognosis of these patients.

We thank Mariko Ohara for pertinent advice.

References

5 Nakagawara A, Ikeda K, Higashi K. N-myc oncogene amplification.


N-myc gene product expression in neuroblastoma


Requests for reprints to: Dr Munetomo Enjoji, Second Department of Pathology, Faculty of Medicine, Kyushu University 60, 3-1-1 Maidashi, Higashi-ku, Fukuoka 812, Japan.
N-myc gene product expression in neuroblastoma.

H Hashimoto, Y Daimaru, M Enjoji and A Nakagawara

doi: 10.1136/jcp.42.1.52

Updated information and services can be found at:
[http://jcp.bmj.com/content/42/1/52](http://jcp.bmj.com/content/42/1/52)

**Email alerting service**

*These include:*

Receive free email alerts when new articles cite this article. Sign up in the box at the top right corner of the online article.

**Notes**

To request permissions go to:
[http://group.bmj.com/group/rights-licensing/permissions](http://group.bmj.com/group/rights-licensing/permissions)

To order reprints go to:
[http://journals.bmj.com/cgi/reprintform](http://journals.bmj.com/cgi/reprintform)

To subscribe to BMJ go to:
[http://group.bmj.com/subscribe/](http://group.bmj.com/subscribe/)