Expression of NM23 in human melanoma progression and metastasis

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Summary NM23 is a putative metastasis-suppressor gene for some human cancers. Here we have studied NM23 expression during melanoma progression using Northern blotting and immunocytochemistry. There was no significant difference in the average amounts of NM23 mRNA between cell lines derived from metastatic and primary melanomas. The level of NM23 mRNA was also determined for three pairs of poorly metastatic parental (P) and their highly metastatic variant (M) cell lines; the ratios for M/P were 1.2, 0.98 and 0.80. Next we used immunocytochemistry to study NM23 protein in normal skin, benign naevi and primary and metastatic melanomas. Melanocytes in all normal skin and benign samples were positive for NM23; however most primary melanomas (7/11) were not stained by the antibody. All metastatic melanoma samples (5/5) were positively stained. Findings were similar with an antiseraum reactive with both forms of NM23 (H1 and H2), and with an antibody specific for NM23-H1. No relationship was apparent between NM23 immunoreactivity in primary tumours and their aggressiveness or prognosis. Hence, in contrast to the situation described for murine melanoma, the amount of NM23 mRNA or protein in human melanoma did not correlate inversely with metastasis.

Keywords: human melanoma; metastasis; NM23

Successful metastasis is a complex series of biological events (MacDonald and Steeg, 1993; Dorudi and Hart, 1993). Acquisition of metastatic capacity by a cell appears to involve both positive and negative changes in gene expression (Liotta et al., 1991). Hence subtractive and differential cDNA hybridisation methods provide approaches for the isolation of relevant genes (Hart and Easty, 1991), starting from closely related tumour cell lines selected for differing metastatic behaviour in experimental assays (Fidler and Radinsky, 1990). Thus a panel of subclones from mouse melanoma K1735 was used to isolate the metastasis-suppressor gene nm23 on the basis of a lower level of expression in highly metastatic compared with poorly metastatic clones (Steeg et al., 1988). The human homologues of the gene, NM23-H1 and NM23-H2, both mapping to chromosome 17q21 (Chang et al., 1994). They encode the monomers of nucleoside diphosphate kinases (NDPK)-A and -B respectively. Both are homohexamers, except that hybrid forms containing both NDPK-A and -B subunits have been found in erythrocytes (Gilles et al., 1991). It is not known whether these mixed forms occur elsewhere. These proteins may also have other functions, notably a DNA-binding activity (Postel et al., 1993) or a protein kinase activity (MacDonald et al., 1993; Bertheau et al., 1994). Suppressor activity was further indicated when transfection of murine nm23 into the highly metastatic melanoma subline K1735 TK resulted in significantly reduced metastasis (Leone et al., 1991).

More recent studies however have not shown a universal pattern of deficient expression of this gene in metastatic tumours. In other mouse melanoma sublines (Parker and Sherbet, 1992), further clones derived from line K1735, and human tumour cell clones (Radinsky et al., 1992), no correlation was detected between nm23 transcription levels and metastatic potential. In studies of various human cancers, an inverse correlation of NM23 mRNA or protein expression with metastatic potential has been described in some cases (Bevilacqua et al., 1989; Hennessy et al., 1991; Hirayama et al., 1991; Flørenes et al., 1992), but not others (Sastre-Garau et al., 1992; Radinsky et al., 1992; Higashiyama et al., 1992; Sawan et al., 1994). Moreover, groups who also examined related normal or benign tissues generally reported that expression of NM23 was lower than typically seen in malignant lesions (e.g. Lacombe et al., 1991; Hirayama et al., 1991; Hallat et al., 1991; Sawan et al., 1994). The amount of NM23-H1 mRNA in human metastatic melanoma has been studied by two groups (Flørenes et al., 1992; Xerri et al., 1994). Both groups reported generally decreased amounts in more aggressive tumours. Flørenes and colleagues found that metastases appearing soon after diagnosis of the primary tumour had significantly lower levels of NM23 mRNA than metastases that developed after a longer time. Similarly Xerri et al. (1994) when studying patients with melanoma metastasis confined to regional lymph nodes, found significantly longer survival associated with higher levels of NM23 mRNA in the resected nodes. Conversely Flørenes et al. (1992), also found lower levels of NM23 mRNA in benign naevi than in metastases, and speculated that NM23-H1 might act as a suppressor of differentiation in melanocytes, as previously suggested for NM23-H2 in lymphocytes (Okabe-Kado et al., 1992).

Data from Northern blotting using fresh tumour samples will be influenced by any non-neoplastic cells present (including stromal cells and any inflammatory infiltrate), and also will not provide information regarding tumour heterogeneity. To circumvent these problems we have used: (1) Northern blotting to study 30 cell lines derived from lesions at various stages of melanoma progression; and (2) immunocytochemistry to study human biopsy material. To ensure that data from cell lines were relevant to metastasis we included three pairs of poorly metastatic parental lines and their highly metastatic variant lines. These human cell lines were previously selected and assayed in nude (thymus-deficient) mice (Ormerod et al., 1986, Herlyn et al., 1990), and so resemble the murine melanoma cell lines used initially to isolate nm23 (Steeg et al., 1988).

The amount of NM23 mRNA (by Northern blotting) has been shown to correlate well with protein expression assessed with an anti-NM23 polyclonal antiseraum (Sawan et al., 1994). We have used the same antiseraum in an immunohistochemical study of the expression of NM23 in pigmented lesions at various stages of tumour progression.

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Materials and methods

Patients and Tumours

Five biopsies of normal skin and biopsies of five benign compound naevi, five dysplastic melanocytic lesions, five melanomas in situ, five radial growth phase (RGP) melanomas, six vertical growth phase (VGP) melanomas (Clark et al., 1989) and five metastatic melanomas were obtained from the archives of the Department of Histopathology, St. George’s Hospital. Clinical follow-up was available in every case. Tissue had been fixed in 10% neutral buffered formalin, routinely processed and embedded in paraffin wax. Sections (5 μm) were cut and floated onto to poly-L-lysine coated glass slides for immunohistochemistry. Before inclusion in the study, haematoxylin and eosin stained sections were reviewed by the dermatopathologist (MEF) to verify the pathological diagnosis.

Immunohistochemistry

An alkaline phosphatase-conjugated second antibody was used for immunostaining (Warburton et al., 1982). This method produces a red product that is easily distinguished from melanin pigment. Mounted sections were dewaxed, taken to water and preincubated in 5% goat serum in PBSA (Dulbecco’s phosphate-buffered saline lacking calcium and magnesium chlorides), to block non-specific binding. They were next incubated with anti-NM23 antiserum (Sastre-Garau et al., 1992) that had been affinity purified using recombinant NDPK-A (Sawan et al., 1994). This antiserum has a preference for NDPK-A but also cross-reacts with NDPK-B (Sawan et al., 1994). The serum was diluted 1:500 in PBSA with 5% normal human serum. Incubation was overnight at 4°C. Sections were washed in PBSA and incubated with alkaline phosphatase-conjugated goat anti-mouse antisera (Sigma) diluted 1:60, for 1 h at room temperature. They were then washed in PBSA and the antibody visualised by developing in substrate buffer: naphthol AS-BI phosphate (sodium salt) and fast red TR salt in veronal acetate buffer, pH 9.2, containing levamisole to inhibit endogenous alkaline phosphatase activity as previously described (Warburton et al., 1982). Sections were counterstained with haematoxylin. For negative controls the primary antibody was replaced by normal rabbit serum.

Representative sections (more than half of the total stained previously with the rabbit antibody and additional new blocks) were also stained with a monoclonal antibody to NM23-H1 that does not cross-react with NM23-H2 (nm23-H1/NM301, Santa Cruz Biotechnology). This was diluted 1:50 and detection was with an alkaline phosphatase-conjugated rabbit anti-mouse antisera (Sigma). Antibody incubation and substrate development were as described above.

Standardisation of immunostaining was made possible by inclusion of normal skin in each experiment. Normal skin consistently contained intensely stained sweat and sebaceous glands and weakly stained or unstained epidermis. Normal skin present at the edges of some test sections gave a further control. Sections were graded using a five point scale as previously described (Sawan et al., 1994). Absence of staining (0) and weak, equivocal staining (+/−) were classed as negative, while unequivocal weak (+), moderate (+++) and intense (+++) staining were classed as positive.

Culture of melanocytes and melanomas

Normal melanocytes and melanomas were cultured as previously described (Easty et al., 1993). Biological details of most of the lines used here are tabulated in Easty et al., (1995a,b). Melanoma cell line RPMI-7932 and all COLO lines were gifts from Professor G. Moore (Department of Health, Denver, CO, USA) (Morse and Moore, 1993). Lines ME1402 and ME1402 were gifts from Dr P Parsons (Queensland Institute of Medical Research, Brisbane, Australia). 451LU and all WM lines were gifts from Professor M Herlyn. Cell line 451LU is a highly metastatic variant cell line derived from the parental cell line WM164 which has only a very low rate of spontaneous metastasis (Herlyn et al., 1990). DX3 and SKMEL23 cells were donated by Dr T Albino (Memorial Sloan-Kettering Cancer Center, New York, USA). DX3LT5.1 is a highly metastatic variant cell line that was derived from the poorly metastatic parental line DX3 following brief exposure to 5-azacytidine and repeated passage through the lungs of nude mice by the intravenous route (Ormerod et al., 1986). Similarly A375M is a highly metastatic variant line that was derived from the poorly metastatic parental line A375P in the same way. Thus, three pairs of poorly metastatic parental lines and their highly metastatic variants were examined for NM23 expression. The metastatic behaviour of these variants is stable (Ormerod et al., 1986; Herlyn et al., 1990); nevertheless the earliest available passage was used for Northern blotting to minimise clonal drift. All melanoma lines were cultured in Dulbecco’s modification of Eagle’s medium with 5% fetal calf serum, except for WM1650, a line derived from the radial growth phase of a primary melanoma, which was grown in melanocyte medium (Easty et al., 1993).

Northern blotting analysis

Total RNA was isolated and poly (A) + RNA selected by oligo(dT)-cellulose chromatography as described previously (Easty et al., 1993). Aliquots (10 μg) of poly(A)-enriched RNA were separated by electrophoresis in agarose gels containing formaldehyde and transferred to nitrocellulose membranes. Prehybridisation (4 h) and hybridisation (18 h) were at 42°C in 5 × SSPE, 5 × Denhardt’s solution, 50% w/v formamide, 500 μg ml−1 salmon sperm DNA and 10% w/v sodium dodecyl sulphate (SDS). The final wash was in 0.5 × SSPE and 0.1% SDS at 65°C. After washing, filters were autoradiographed for up to 1 h. The probe used was an NM23-H1 sequence, a gift from Dr P Steeg (Steeg et al., 1988). This would be expected to detect NM23-H2 as well, because of the high homology of the two coding sequences. Probes were labelled by random hexamer priming (Feinberg and Vogelstein, 1984) to a specific activity of about 2 × 107 d.p.m. μg−1 DNA. For normalisation of RNA loading, blots were rehybridised with a probe for housekeeping gene glyceraldehyde phosphate dehydrogenase (GAPDH), a gift from Professor M Clemens, St. George’s Hospital Medical School. Quantitation of autoradiographs was by laser densitometry, using Image Quant software (Molecular Dynamics, Sunnyvale, CA, USA). The relative level of expression was calculated as the ratio of signal in the area of the NM23 band to that for GAPDH mRNA in the same cell line. Figures were then normalised to the level in one cell line (DX3LT5.1), defined as 100 as described (Easty et al., 1993).

Results

Northern blotting

By Northern analysis a 0.8 kb NM23 transcript was detected in all cell lines studied. This mRNA was highly expressed (easily detected by autoradiography in 30 min), although with an 11-fold range of level (Figure 1). The levels of NM23 mRNA in cell lines derived from primary and metastatic melanomas were compared (Figures 1 and Table 1). Median mRNA indices were 73.5, 55 and 85 in normal melanocytes, primary and metastatic melanomas respectively. Levels ranged widely in the tumour cells and there was an apparent trend towards higher mRNA levels in metastatic than in primary melanoma lines (Table I), although this was not statistically significant. Cell lines WM239A and WM115 were from the same patient and were both a metastasis and a primary melanoma respectively; WM239A had a 5-fold higher level of NM23 mRNA than WM115.
Three pairs of poorly metastatic cell lines (P) and highly metastatic variants (M) were examined by Northern blotting (Table I). The ratios M/P for NM23 mRNA were: 1.20 (DX3LT5.1/DX3), 0.98 (A375M/A375P), and 0.80 (451LU/ WM164). Thus there was no obvious relationship between the level of NM23 transcription in these cell lines and their metastatic behaviour in nude mice. Nor could we detect any strong relationship with cell differentiation. The normal melanocytes and two of the melanoma lines used, WM1650 and SKMEL23, were morphologically differentiated, pigmented and non-tumorigenic in nude mice; however there was an 8-fold difference between the NM23 mRNA indices for these two melanoma lines and the values for melanocytes were intermediate (Table I).

**Immunohistochemistry**

Initially, a rabbit anti-NM23 polyclonal antibody was used for immunohistochemistry. In normal skin this antiserum bound strongly (+ + +) to some epidermal adnexal components such as sweat gland coils and sebaceous glands, whereas the epidermis was unstained or only weakly

![Figure 1](https://example.com/figure1.png)

**Figure 1** Northern blot analysis of poly(A)-enriched RNA from cultured melanocytes and cell lines from primary melanomas (WM115, MM485) and melanoma metastases (all other lines): 10 μg was loaded per lane. Filters were hybridised to probes for NM23-H1 and GAPDH and autoradiographic signals were quantitated by laser densitometry. To control for loading the ratio of signal for NM23 to that for GAPDH was calculated for each cell line and then normalised to the ratio for one cell line (LT5.1). Both primary and metastatic lines show a range of levels of NM23 mRNA; see Table I for details.

![Figure 2](https://example.com/figure2.png)

**Figure 2** Northern blot analysis of poly(A)-enriched RNA from poorly metastatic parental melanoma cell lines (WM164, DX3, A375P), and their closely related highly metastatic variant sublines (451LU, LT5.1, A375M): 10 μg was loaded per lane. Probes were for NM23-H1 and GAPDH. Quantitation of autoradiographs was by laser densitometry; see Table I for details.

| Normal melanocytes mRNA index | Primary melanoma lines mRNA index | Metastatic melanoma lines mRNA index |
|-------------------------------|----------------------------------|------------------------------------|
| M/P                           | M/P                              | M/P                                |
| WM115                         | 65                               | 23                                 |
| ME10538                       | 451LU                            | 22                                 |
| WM98-1                        | 50                               | WM852                              |
| MM485                         | 60                               | WLM164                             |
| ME1402                        | 67                               | C32TG                              |
| WM1158                        |                                  | WM1158                             |
| COLO 679                      | 66                               |                                   |
| COLO 792                      | 68                               |                                   |

|mRNA levels were quantitated by Northern blotting and laser densitometry of autoradiographs (Materials and methods). Levels are relative to GAPDH mRNA levels in each line and the index is obtained by dividing the value for one line (DX3LT5.1) and multiplying by 100 (Easy et al., 1993). *These lines were obtained from different metastatic sites in the same patient. Thus they were not independent and were represented by a single median value (121) for statistical purposes. The dashed line represents the median value for normal melanocytes. The difference between primary and metastatic lines was not significant by the χ² test on the numbers of values falling above and below this line (P<0.1), not by Mann–Whitney rank test.
stained (+) (Figure 3). This provided an internal control for NM23 reaction in pathological material that contained adjacent normal skin. Inflammatory and stromal cells in lesions (including lymphocytes, macrophages and fibroblasts) were generally unstained.

Most pathological specimens showed heterogeneous staining of cells forming the lesion. Staining was generally cytoplasmic. There was intense staining of melanocytes in junctional nests of compound naevi (+ + +) with some more weakly stained cells (+ to + +) in the dermis (Figure 3); there was some indication of reduced staining with increasing depth in the dermis (from + + + to + +). Similar patterns were observed in compound naevi with melanocytic dysplasia (not shown). In contrast, primary melanomas (both RGP and VGP) were more weakly stained, areas within a lesion varying from 0 to + or occasionally + +. Lesions were scored as positive if they contained any clearly stained malignant cells. Some primary melanomas (7/11) were unstained (0) or gave at best an equivocal reaction (+/-), and so were classed as negative (Figure 4). Melanoma metastases also tended to react heterogeneously for NM23. However, in all samples from metastases the majority of melanoma cells were moderately or highly stained and lesions were thus scored as + + or + + + (Figure 5). Nuclear staining was observed in some cases.

Similar patterns of staining were obtained when the anti-NM23-H1 monoclonal antibody was used, except that in some compound naevi immunostaining was only moderate (+ to + +), and occasionally restricted to one part of the naevus in contrast with the situation with the rabbit anti-NM23 antibody which bound to all parts. The reaction in melanoma metastases was intense (+ + +) and the pattern of staining closely resembled that found previously with the rabbit antiserum.

There was no apparent relationship between NM23 immunoreactivity and prognosis. Positive NM23 staining was seen in 3/6 primary melanomas from which metastasis was subsequently detected (median time from resection of the primary tumour, 7 months, range 1-24 months), and in 1/5

![Figure 3](image1.png) **Figure 3** Immunohistochemical staining for NM23 in compound naevi using (a) a rabbit polyclonal antibody, and (b) a mouse monoclonal antibody. There is an intense reaction in naevus cells (solid arrow), whereas staining in the epidermis (open arrow) is weak to negative. Individual positive cells in the basal epidermis near the naevus are likely to be naevus cells.

![Figure 4](image2.png) **Figure 4** Lack of immunohistochemical staining for NM23 in a VGP primary melanoma using a rabbit polyclonal antibody. There was no clear staining in tumour cells. Brown melanin can be seen in the cytoplasm of some melanoma cells.

![Figure 5](image3.png) **Figure 5** Immunohistochemical staining for NM23 in a melanoma metastasis to a lymph node. (a) Rabbit polyclonal antibody and haematoxylin counterstain. (b) Mouse monoclonal antibody. There is intense staining of trabeculae of tumour cells (solid arrow). Stromal cells and lymphocytes are negative (open arrow).
Table II Immunohistochemical staining for NM23 in melanocytic lesions

| Lesion              | Percentage of lesions expressing NM23 | Comments                      |
|---------------------|---------------------------------------|-------------------------------|
| Compound naevi      | 100 (5/5)                             | Generally heterogeneous       |
|                     |                                       | labelling of cells:          |
|                     |                                       | junctional nests, + + +;     |
|                     |                                       | cells in dermis, + to + +    |
| Atypical naevi      | 100 (5/5)                             | As for compound naevi        |
| Primary melanomas   | 36 (4/11)                             | Heterogeneous labelling       |
|                     |                                       | (4/4): to - or to + +        |
| Metastatic melanomas| 100 (5/5)                             | Heterogeneous labelling       |
|                     |                                       | in all lesions, - to + + or  |
|                     |                                       | to + + +                     |

*Immunohistochemical staining with a rabbit polyclonal anti-NM23 antibody. 2Primary melanomas positive for NM23 were 3 VGP and 1 RGP. After resection and follow-up, one of these patients had no evidence of secondary disease and three patients developed metastases; times from excision of primary to first metastasis were 6, 8 and 14 months. Primary melanomas negative for NM23 were 3 VGP and 4 RGP; three such patients had no evidence of secondary disease whereas four patients developed metastases after 1, 2, 18 and 24 months.

Discussion

Some previous studies have concluded that the expression of NM23 in certain human cancers correlates inversely with the incidence of metastases (Hennessy et al., 1989; Hirayama et al., 1991; Florenes et al., 1992; MacDonald and Steig, 1993). The approach has been often to determine the level of NM23 expression in primary tumours in comparison with the incidence of metastasis (or prognosis) in the patients. Alternatively, Florenes et al. (1992) and Xerri et al. (1994) determined the level of NM23 mRNA in melanoma metastases and compared this with the aggressiveness of the primary tumour. Here we examined primary melanomas and related the expression of NM23 to the same measure of tumour aggressiveness (the interval between diagnosis of primary and secondary tumours) as used in the earlier study (Florenes et al., 1992).

The NM23-H1 and -H2 transcripts are 88% homologous, and Northern analyses that use full-length NM23 probes will detect mRNA for both genes (Leone et al., 1993). Such studies include the present work and two previous reports on NM23 in human melanoma (Florenes et al., 1992; Xerri et al., 1994). These genes are not always similarly regulated and the H1 gene product may be a better marker for metastasis in at least some tumours (Tokunaga et al., 1993). Nevertheless, both previous studies of human melanoma found reduced amounts of total NM23 mRNA in more aggressive tumours. For the present study we used a similar probe, and a rabbit polyclonal antibody that cross-reacts with both gene products. For comparison we also included an antibody specific for NM23-H1.

We did observe generally decreased NM23 protein levels in primary melanomas as compared with melanocytes in naevi. More than half the primary melanomas were unstained (Table II). However, no relationship was apparent between the amount of NM23 in the primary melanomas and the subsequent incidence of metastasis for this small group of patients. Immunostaining for NM23 in these lesions was generally heterogeneous. Tumour heterogeneity for metastatic propensity has been well documented (e.g. Fidler and Radinsky, 1990), and it is possible that metastases arose selectively from cells lacking NM23 within these lesions even when the lesion overall was classified as positive. However, if so, one might expect that melanoma metastases would have at least as low a level of NM23 expression as primary tumours, and that primary tumours expressing no NM23 would have the poorest prognosis. Neither was the case: we found high levels of NM23 protein in all metastatic melanomas studied.

With the polyclonal antibody there was intense immunohistochemical staining for NM23 in benign naevi. This apparently contrasts with the finding of low NM23-H1 mRNA levels in naevi by Florenes et al. (1992). Naevi are generally small, thin, discontinuous lesions and biopsies are likely to contain epidermis and dermis, which express little NM23 and might dilute NM23 mRNA from naevus cells. Another possible source of discrepancy might be high expression of NM23-H2/NDPK-B in naevi. This might be detected to a lesser extent by the NM23-H1 gene probe than by the rabbit antibody. Consistent with this, the NM23-H1-specific antibody reacted only weakly with some naevi. However, other naevi were strongly stained, indicating high expression of NM23-H1 protein in at least some naevi. Moreover, from results with this antibody, NM23-H1 was highly expressed in all melanoma metastases tested.

NM23 mRNA levels in cultured melanoma cells ranged from lower than those in cultured normal melanocytes to several fold higher; there was no significant difference between the distributions of amounts in metastatic and primary melanoma cell lines. Using immunohistochemistry we found more NM23 protein in metastases than in primary melanomas. This tendency seems surprising given that it was in melanoma that a metastasis-suppressor activity of nm23 was first demonstrated. Recalling the often high expression of NM23-H1 mRNA in metastatic cell lines, it is possible that an abnormal form of NM23-H1 is being overexpressed. It was recently reported that point mutations of NM23-H1 can be coupled with amplification of the gene in neuroblastoma (Chang et al., 1994). This is reminiscent of findings with the p53 tumour-suppressor gene: either (1) inactivation and silencing of the normal p53 gene; or (2) overexpression of a mutated form of the gene can be oncogenic (Levine, 1993). We are currently examining this further by sequencing NM23-H1 genes from metastatic melanomas. Our results raise the possibility that, while in primary human melanoma expression of the NM23-H1 gene tends to be repressed, overexpression of an altered gene becomes a more common mechanism during progression and metastasis.

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