Overexpression of Numb suppresses tumor cell growth and enhances sensitivity to cisplatin in epithelioid malignant pleural mesothelioma

YANMENG KANG¹, MINGJIE DING², GUANGYAN TIAN³, HAISHENG GUO⁴, YUNYAN WAN¹, ZHOUHONG YAO¹, BIN LI¹ and DIANJIE LIN¹

¹Department of Respiratory Medicine, Shandong Provincial Hospital, Shandong University, Jinan, Shandong 250021; Departments of ²Respiratory Medicine, and ³Neurology, Jinan Children’s Hospital, Jinan, Shandong 250022; ⁴Department of Oncology, Dongying People’s Hospital, Dongying 257091, P.R. China

Received March 2, 2013; Accepted April 11, 2013

DOI: 10.3892/or.2013.2429

Abstract. Malignant pleural mesothelioma (MPM) is a highly aggressive and conventional treatment-resistant tumor with a dismal prognosis. Among the three histological subtypes of MPM, the epithelioid is the most common type. Numb is considered as a tumor suppressor playing a critical role in controlling asymmetric cell division, maintenance of stem cell compartments, ubiquitination of specific substrates and regulating Notch-, Hedgehog- and TP53-activated pathways. The present study was designed to analyze the role of Numb in epithelioid MPM. We investigated the expression of Numb in 39 epithelioid MPM and 22 normal pleural tissues by immunohistochemistry. Furthermore, we overexpressed Numb in NCI-H2452, an epithelioid human MPM cell line, and investigated the effect of Numb overexpression on the proliferation, apoptosis and sensitivity to cisplatin in cells. The expression of Numb was significantly lower in MPM compared to the control group and Numb had an inverse correlation with the Ki-67 labeling index. Loss of Numb expression was associated with poor prognosis in epithelioid MPM. Overexpression of Numb in NCI-H2452 cells significantly inhibited proliferation, promoted apoptosis and enhanced sensitivity to cisplatin. Moreover, Numb overexpression activated caspase-9 and caspase-3 through release of cytochrome c as well as downregulation of XIAP and survivin. We speculate that cytochrome c/caspase signaling is a possible mechanism through which Numb enhances the apoptosis of NCI-H2452 cells. These results suggest that Numb may be involved in epithelioid MPM development, and its upregulation may confer sensitivity to cisplatin, suggesting potential therapeutic options for MPM.

Introduction

Malignant pleural mesothelioma (MPM) is a highly aggressive and conventional treatment-resistant tumor with a dismal prognosis. Its incidence has been increasing over the past two decades in industrialized countries and is expected to peak during the period 2010-2020 (1,2). There are three distinct histologic subtypes of MPM and the epithelioid type is the most common one which comprises 50-70% of all cases. Asbestos exposure, non-asbestos mineral fibers, organic chemicals, radiation and simian virus 40 (SV40) have been suggested as risk factors for mesothelioma (3). These etiologic agents can cause chronic tissue injury and tissue repair in the long latency period. The misappropriating homeostatic mechanisms that govern tissue repair and stem cell self-renewal may lead to carcinogenesis (4). Targeting the genes which play a part in the determination and maintenance of the stem cell compartment may contribute to identifying more accurate prognostic factors and developing effective therapies.

Numb is an evolutionarily conserved protein that plays a critical role in cell fate determination. Numb was originally identified in Drosophila (5). It has been reported to be involved in the control of asymmetric cell division (6), maintenance of stem cell compartments (7), regulation of cell polarity and adhesion (8), migration (9) and ubiquitination of specific substrates (10-12). Subversion of Numb has been linked to many pathological mechanisms, including cancer. Numb plays an important role in the transformation target and the plasticity of the stem cell compartment (13), which are closely related to tumor formation. In particular, the plasticity of the stem cell compartment is related to epithelial-mesenchymal transition (EMT), which contributes to the acquirement of invasive properties as well as resistance to cell apoptosis and chemotherapy (14,15). Subversion of Numb is also predicted to have a major impact on the homeostasis of endocytosis (16,17) and the alteration of many polarity functions through connecting with the PAR complex (18,19), which both facilitate transformation events of tumors. In the absence of Numb, both Notch (10,20,21) and Hedgehog (Hh) signaling (11) are augmented, with pro-proliferative...
and anti-differentiation effects, while the suppressor TP53 signaling is attenuated (12).

Experimental evidence indicates that Numb has the potential to function as a tumor suppressor. Studies of mutant Numb in Drosophila have shown neuroblast overproliferation and tumor formation, which strongly supports the hypothesis that impairment of asymmetric cell division in stem cells results in an imbalance in self-renewal and differentiation which leads to cancer (22). In addition, in vivo RNA interference of Numb in a model of mouse lymphomagenesis was found to accelerate the onset of lymphomas (23). Loss of Numb expression has also been reported in some types of human cancers, such as breast cancers, non-small cell lung carcinomas, salivary gland carcinomas and medulloblastomas (20,21,24,25). In human astrocytomas, there is a trend of higher Numb expression in more malignant tumors (26). Whether Numb functions as a tumor suppressor in MPM or not has not been clarified.

In the present study, we analyzed the expression of Numb in epithelioid MPM and investigated the potential role of Numb in MPM cells. The aim of our study was to clarify the role of Numb in epithelioid MPM and to explore novel strategies for drug-resistant MPM therapy.

Materials and methods

Patients and specimens. Tissue samples of 39 MPM patients who underwent medical thoracoscopy procedures (Department of Respiratory Medicine, Provincial Hospital, Shandong University, China), from 2007 to 2010, were analyzed and 22 normal pleural samples as a control group were recruited. The MPM diagnosis was based on WHO criteria (27) and confirmed in all instances by clinical, morphologic and immunohistochemical data. Specimens were reviewed by two pathologists, and the histological type was confirmed as epithelioid cell type according to the 2004 WHO classification of pleural tumors (27). All procedures were approved by the Ethics Committee of Shandong University.

Immunohistochemistry. Formalin-fixed and paraffin-embedded tissue sections (4 µm) were processed for immunohistochemical investigation. Tissue sections were first deparaffinized, followed by rehydration with serially decreased concentrations of ethanol. The following antigen retrieval was carried out in 0.01 M citrate buffer (pH 6.0) at 96˚C for 15 min in a thermostat controlled water bath. The tissue sections were then immersed in 3% H2O2 for 30 min and were blocked with normal serum at 37˚C for 30 min. The primary antibodies anti-Numb antibody (1:200, rabbit polyclonal; Abcam, Cambridge, UK) and anti-Ki-67 antibody (1:150, rabbit polyclonal; Santa Cruz Biotechnology, Santa Cruz, CA, USA) were incubated at 4˚C overnight. The following biotinylated secondary antibodies (Beijing Zhongshan Golden Bridge Biotechnology Co., Ltd., Beijing, China) were added at 37˚C for 30 min. For detection of the binding sites, a streptavidin-biotin system (Beijing Zhongshan Golden Bridge Biotechnology) was used. Diaminobenzidine (DAB) was used as the enzyme substrate to observe the specific antibody localization, and nuclei were counterstained using hematoxylin. Negative controls were performed in all cases by omitting the primary antibodies. Observation was carried out under a light microscope (Leica DM4000, Wetzlar, Germany) and the images were processed with image processing software (Leica IM45).

Histological evaluation. Two independent observers blinded to the clinical data were assigned to evaluate all samples. Quantitation of cells staining positivity for Numb in the cytoplasm or membrane was recorded. Sections with staining in ≥10% of cells were considered as positive. Immunohistochemical reactivity was graded on a scale of 0-3 according to the percentage of immunopositive cells as follows: grade 0, no staining or <10%; grade 1, 10-30%; grade 2, 30-50%; grade 3, >50% positive cells. For Ki-67, the percentage of positively stained nuclei out of the total cancer cells was calculated, and we used 25% as the cut-off point for categorizing.

Cell culture and plasmid transfection. Human MPM cell line NCI-H2452 was stemmed from the American Type Culture Collection (ATCC, Manassas, VA, USA), cultured in RPMI-1640 medium supplemented with 10% fetal calf serum, 1.5 g/l NaHCO3, 2.5g/l glucose, 0.11 g/l sodium pyruvate and maintained at 37˚C in a humidified atmosphere with 5% CO2. We selected the HEK-293T cell line which has been proven to express Numb as a control (11). HEK-293T cells were cultured in Dulbecco’s modified Eagle’s medium (DMEM) containing 10% fetal bovine serum (FBS). Plasmid pcDNA3.1-Numb was synthesized by Shanghai GenePharma Co., Ltd. (Shanghai, China). First, 1x106 cells in a 60-mm dish were transfected with either the empty vector (mock) or pcDNA3.1-Numb (Numb) using Lipofectamine™ 2000 (Invitrogen) according to the manufacturer’s guidelines. After 24 h, the cells were split into three sets: one set was used for cell viability assay; one was used to analyze protein expression by western blot analysis; and one was used for cell apoptosis analysis.

Real-time quantitative reverse transcription-polymerase chain reaction. Primer sequences used for Numb were: 5’-CAATCTTCTACCTTCCAAGGG-3’ and 5’-CGGACGCC TCTTAGACACCTC-3’. For GAPDH, 5’-TGTTGACACCA GGCTCCTGCT-3’ and 5’-AGCTTCCGGTCTACAGCCTT-3’. All primers were synthesized by Takara Co., Ltd. (Dalian, China).

Total RNA from cells was isolated using TRIzol (Invitrogen) and reverse-transcribed by the ReverTra Ace qPCR RT kit (Toyobo, Osaka, Japan). qRT-PCR was performed using SYBR Green Real-Time PCR Master Mix (Toyobo) following the manufacturer’s instructions and on the LightCycler 480 System (Roche, Germany). Each reaction was performed in triplicate and analysis was performed by the 2-ΔΔct method.

Western blot analysis. At 48-72 h after transfection, cells were harvested for western blot analysis. Total protein was extracted by RIPA lysis buffer. To analyze cytochrome c expression, cytosolic and mitochondrial fractions were prepared using the Mitochondria Extraction kit (Nanjing KeyGen Biotech., Co., Ltd., Nanjing, China). Proteins were transferred to a PVDF membrane and subjected to the antibodies including Numb (1:1,000; Abcam), caspase-3, caspase-9, cytochrome c, XIAP and survivin (1:500; Santa Cruz Biotechnology). Peroxidase-labeled anti-mouse or anti-rabbit antibodies were
used as secondary antibodies (Zhongshan Goldenbridge Biotechnology). Blots were developed using enhanced chemiluminescence detection reagent (Applygen Technologies Inc., Beijing, China).

Cell viability assay. We used an MTT (3-[4,5-dimethylthiazol-2-yl]-2,5-diphenyltetrazolium bromide) assay to evaluate cell viability. Cells (5x10^3/well) were grown in a 96-well plate and were tested at 24, 48 and 72 h after transfection. The cells (1x10^3/well) were grown in a 96-well plate and incubated for another 24 h, and these cells were then treated with different concentrations of cisplatin ranging from 1 to 10 µmol/l for 24 h in triplicate. Twenty microliters of 5 mg/ml MTT was added into each well and incubated for 4 h at 37°C in a culture hood. Media were carefully removed, and 150 µl DMSO was added. The absorbance was measured at a wavelength of 490 nm from which the background was subtracted. The cell survival value index was calculated by the formula: \[\frac{A_{490 (+\text{cisplatin})}}{A_{490 (-\text{cisplatin})}} \times 100\%\].

Cell apoptosis analysis. The morphological changes in apoptotic cells were investigated using Hoechst 33258 staining (Solarbio, Shanghai, China) and fluorescence microscopy. The apoptosis rate was examined by two-color analysis of Annexin V/FITC binding and propidium iodide (PI) uptake using flow cytometry. Cells (1x10^5) were seeded into 6-well plates and were tested at 72 h after transfection. Cells (1x10^5) were then seeded into 6-well plates and incubated for another 24 h, and these cells were then treated with 5 µmol/l cisplatin for 24 h. FITC-conjugated Annexin V was added at a concentration of 0.5 µg/ml. After incubation for 20 min at room temperature in the dark, PI was added at 1 µg/ml, and the samples were immediately analyzed by fluorescence-activated cell sorting (FACS).

Statistical analysis. All analyses were performed using SPSS 17.0 software package (SPSS Inc., Chicago, IL, USA). Chi-square test was used to compare the expression of Numb in the MPM and control groups. Correlations of protein expression (grade scores) with clinical and pathological parameters of the tumors were evaluated with the Mann-Whitney test for ordinal variables. Overall survival (OS) was defined as the time of diagnosis to the time of death from cancer-related cause or to the date the patient was last known to be alive. The univariate analysis of OS was carried out by the Kaplan-Meier method using the log-rank test. All cell culture experiments were conducted at least three times. Data are shown as means ± SD and were analyzed using the Student’s t-test for two samples. All statistical tests were two-sided, and probability values <0.05 and <0.01 were defined as being statistically significant.

Results

Expression of Numb in MPM tissues. The characteristics of the 39 epithelioid MPM patients enrolled in the present study were as follows. The median age for the patients was 55 years (range, 32-77). Nineteen patients were males and 20 were females, with a male to female ratio of 1:1.05. Patients were divided into two groups according to the Eastern Cooperative Oncology Group (ECOG) performance status. Twenty-nine patients were in the grade <2 group and 10 patients were in the grade ≥2 group (Table I).

![Figure 1. Immunostaining of Numb and Ki-67.](image)

Figure 1. Immunostaining of Numb and Ki-67. (A) Strong positive staining in the control group compared to (B) negative staining of Numb in epithelioid MPM. (C) Representative weak staining of Numb in MPM and the corresponding high Ki-67 labeling index (>25%); (D), magnification, x200.

Table I. Expression of Numb in epithelioid MPM and its correlation with clinical features.

<table>
<thead>
<tr>
<th>Variables</th>
<th>Numb expression</th>
<th>N</th>
<th>Grade 0/1/2/3</th>
<th>P-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>MPM (epithelioid) total</td>
<td>39</td>
<td>20/10/6/3</td>
<td>0.783</td>
<td></td>
</tr>
<tr>
<td>Age (years)</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>≤55</td>
<td>20</td>
<td>10/5/3/2</td>
<td>0.025</td>
<td></td>
</tr>
<tr>
<td>&gt;55</td>
<td>19</td>
<td>10/5/3/1</td>
<td>0.258</td>
<td></td>
</tr>
<tr>
<td>Gender</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Male</td>
<td>19</td>
<td>8/5/5/1</td>
<td>0.159</td>
<td></td>
</tr>
<tr>
<td>Female</td>
<td>20</td>
<td>12/5/1/2</td>
<td>0.011</td>
<td></td>
</tr>
<tr>
<td>ECOG performance status</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>≤2</td>
<td>11</td>
<td>4/3/2/2</td>
<td>0.011</td>
<td></td>
</tr>
<tr>
<td>&gt;2</td>
<td>28</td>
<td>16/7/4/1</td>
<td>0.001</td>
<td></td>
</tr>
<tr>
<td>Ki-67 index</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>≤25</td>
<td>26</td>
<td>10/7/6/3</td>
<td>0.258</td>
<td></td>
</tr>
<tr>
<td>&gt;25</td>
<td>13</td>
<td>10/3/0/0</td>
<td>0.258</td>
<td></td>
</tr>
</tbody>
</table>

Oncology (ECOG) performance status. Twenty-nine patients were in the grade <2 group and 10 patients were in the grade ≥2 group (Table I).

Numb localization on the membrane or in the cytoplasm was demonstrated in tumor samples (51.3% (20/39) positive) with a slighter intensity compared with [86.4% (19/22)] the normal pleural specimens (P<0.05; Fig. 1A and B). A marked inverse correlation was found between Numb expression levels and Ki-67 labeling index (P<0.05; Fig. 1C and D, Table I). There was no correlation between loss of Numb
expression and gender, age and ECOG performance status (P>0.05).

Complete clinical follow-up data were obtained from 32 patients allowing retrospective survival analysis. At the completion of the study in July 2011, 30 patients had died, with a median follow-up of 11.3 months. Univariate analysis indicated that overall survival was influenced by Numb expression (log-rank test P<0.05; Fig. 2), comparing negativity (grade 0; median survival, 10.3 months; 95% CI, 9.1-11.5) vs. weak-moderate-strong expression (grade 1-3; median survival, 11.7 months; 95% CI, 11.1-12.3).

Numb protein expression is increased in NCI-H2452 cells by transfection. Before transfection, NCI-H2452 cells displayed levels of Numb mRNA expression comparable to that detected in HEK-293T cells, but low protein expression was noted (Fig. 3A and B). The NCI-H2452 cells transfected with pcDNA3.1-Numb showed higher Numb protein expression compared with the mock-transfected control (Fig. 3C).

Numb inhibits proliferation and promotes apoptosis in the NCI-H2452 cells. The absorbance of Numb-transfected cells, as indicated by MTT, was markedly lower at 48 and 72 h compared to that of the mock-transfected cells (P<0.05, P<0.01; Fig. 4A), which indicates that Numb overexpression inhibited the growth of NCI-H2452 cells. There were no significant differences between the two groups at 24 h (P>0.05). Apoptosis was investigated at 72 h after transfection. Hoechst staining showed that Numb overexpression increased the number of apoptotic cells, which were characterized by several unique morphological nuclear changes, such as chromatin condensation and nuclear fragmentation (Fig. 5). The apoptosis rate was assayed using flow cytometry. The percentage of early and late apoptotic cells was notably higher in the Numb-transfected (19.88±3.31%) when compared to the percentage in the mock-transfected cells (9.41±2.34%, P<0.05; Fig. 4B and C).

Overexpression of Numb induces activation of caspase-3 and caspase-9, as well as cytochrome c release and XIAP and survivin degradation. At 48-72 h after transfection, we analyzed various proteins related to apoptosis in the NCI-H2452 cells by western blot analysis. Numb-transfected cells showed increased activated caspase-3 (17 kDa) and caspase-9 (35 kDa) bands compared with the mock-transfected cells. In addition, we found release of cytochrome c into the cytoplasm as well as downregulated levels of XIAP and survivin in the Numb-transfected cells (Fig. 6).
Numb sensitizes NCI-H2452 cells to cisplatin. MTT assay was used to determine the cell growth and proliferation of the NCI-H2452 cells after treatment with cisplatin for 24 h. Significant decreases in cell viability were observed in Numb-transfected cells treated with cisplatin at concentrations of 1-10 µmol/l for 24 h compared with the mock-transfected cells (P<0.05, **P<0.01). There were no significant differences between the two groups at 24 h. (B and C) Numb induced apoptosis at 72 h after transfection in NCI-H2452 cells as detected by Annexin V-PI analysis. The apoptosis rate (the percentage of Annexin V/PI double-positive cells) was notably higher in Numb-transfected (19.88±3.31%) than that in the mock-transfected cells (9.41±2.34%; P<0.05).

Discussion

In the present study, we reported the role of Numb as a tumor suppressor in epithelioid MPM, and revealed that it enhances the sensitivity of the human MPM cell line NCI-H2452 to cisplatin.

Our data demonstrated that Numb staining in normal pleural specimens was intense, but in tumors it was weak and in many cases completely absent. In accordance with the present study, the expression of Numb was found to be completely lost in breast cancers and non-small cell lung carcinomas (NSCLCs) (20,21). In addition, Numb expression was inversely correlated with the Ki-67 labeling index, an indicator of tumor aggressiveness. This finding is consistent with a study concerning breast cancer (20). High Ki-67 index is the marker of rapid cell proliferation which is related to high malignant potential, high recurrence rate and poor survival (28). This suggests that loss of Numb expression is involved in the formation and development of epithelioid MPM. Negative Numb immunohistochemistry with poor prognosis was previously reported in human salivary gland carcinomas and breast tumors (12,24). Our data also showed that the reduction in Numb expression was associated with poor long term outcome in epithelioid MPM patients. Numb plays an important role in tumorigenesis...
due to its crucial feature of asymmetric cellular distribution. Moreover, due to its ability to function as an adaptor for E3-enzymes, Numb is involved in a complex network of ubiquitin-regulated pathways which are related to the development of cancer. Numb does not live up to its name as ‘fate determinant’.

We selected NCI-H2452, an epithelioid human MPM cell line to investigate the expression and function of Numb in vitro. We detected comparable levels of Numb mRNA in NCI-H2452 cells vs. HEK-293T cells, and this contrasted with the markedly different levels of Numb protein in the same cultures. Thus, we speculated that loss of Numb expression in NCI-H2452 cells is determined at the posttranslational level, through enhanced protein degradation, similarly to what was shown in breast cancers (20) and NSCLCs (21), but the exact mechanism needs to be further explored.

In Numb-negative breast tumor cells, re-expression of Numb was found to selectively suppress growth (20). Our results also showed that overexpression of Numb inhibits proliferation and promotes apoptosis in NCI-H2452 cells. Malignant cells are frequently resistant to apoptotic stimuli, and the inhibition of apoptosis may enhance tumor development. Caspase-3 activity is often used as a definite marker for apoptosis, and cytochrome c/caspase-9 activation is an initial event in the intrinsic pathway for apoptosis. In the present study, activated caspase-3 and caspase-9 bands as well as release of cytochrome c to the cytoplasm were observed in Numb-transfected cells. This suggests that overexpression of Numb induced apoptosis perhaps through cytochrome c/caspase-9/caspase-3 signaling. Inhibitors of apoptosis proteins, IAPs, are a family of proteins that regulate the cytochrome c/caspase activating pathway (29). XIAP is involved in the inactivation of caspase-3, -7 and -9 and has been found to be elevated in mesothelioma (30,31). Several studies have now indicated that higher levels of XIAP may contribute to chemotherapy resistance in mesothelioma cells (32,33). Survivin was also found actively expressed in MPM (32,34) and it has an important role in maintaining apoptosis resistance. It can partially block activation of caspases due to its ability to bind to activated caspases (35,36). Survivin inhibition increases the rates of both spontaneous and radiation-induced apoptosis (37) as well as decreased survival of MPM cells (38). In our study, Numb overexpression decreased the expression of XIAP and survivin, yet the mechanism in unknown. It has been shown that IAPs contain a RING finger domain which confers ubiquitin protease ligase (E3) activity, and these IAPs having E3 activity are able to catalyze their own ubiquitination and degradation. As we know, Numb is at the intersection of a complex network of E3-ligases for it contains a functional phosphotyrosine-binding (PTB) domain. We hypothesized that Numb downregulates IAPs directly through promoting ubiquitination and protease degradation, but further study is needed to explore the exact mechanism. However, we speculate that IAPs play important roles in the apoptosis promoted by Numb. Overexpression of Numb to downregulate IAPs may be an important strategy for overcoming chemotherapy and apoptosis resistance in MPM.

The standard first-line treatment used for MPM is cisplatin-based chemotherapy, either as a single agent or in combination with pemetrexed. Its cytotoxicity is mediated through platinum-DNA adducts and subsequent induction of apoptosis (39). However, in MPM the chemotherapy resistance leads to low efficacy and high doses of cisplatin cause prohibitive toxicity such as nephrotoxicity (40). In the present study, we showed that overexpression of Numb significantly enhanced sensitivity to cisplatin-induced apoptosis and suppressed growth in NCI-H2452 cells. These data indicate that Numb may be involved in conferring cisplatin sensitivity to MPM cells. Decreased caspase-3 and caspase-9 activation have been described as an additional mechanism of cisplatin resistance in various types of cancers (41-44). To confirm whether or not Numb sensitizes MPM cells to cisplatin through the caspase-dependent apoptotic pathway, further investigation is needed.

In conclusion, our results revealed that Numb was frequently downregulated in epithelioid MPM cases, which was also associated with poor prognosis. Overexpression of Numb in NCI-H2452 cells inhibited proliferation and promoted apoptosis, and we speculated that cytochrome c/caspase signaling was a possible mechanism through which Numb enhanced apoptosis. Moreover, Numb sensitized NCI-H2452 cells to
Acknowledgements

We thank the doctors from the Department of Respiratory Medicine and the Department of Pathology of Shandong Provincial Hospital for their technical support.

References