Introduction

Overexpression of the Ha-ras\textsuperscript{val12} oncogene has been observed in primary and metastatic human carcinomas. Alterations in Ha-ras per se or its expression have also been reported in metastatic tumors (1). Gallick et al. examined the expression of the c-ras gene in fresh biopsies from primary and secondary colon carcinomas and found that overexpression of p21\textsuperscript{ras} was common in the primary tumors of Dukes’ B and C stages when compared to the surrounding normal tissue (2). In contrast, Sukumar et al. demonstrated that about one-third of N-methyl-N-nitrosourea-induced rat carcinomas expressed the activated c-Ha-ras gene (3). The tumor metastatic process is complex, requiring cells to move, invade tissues and basement membranes, avoid immunological responses, and colonize distant sites within the organism (4, 5). However, the mechanism of tumor cell-induced metastasis is poorly understood and the identification of the genes involved is still ongoing. The discovery of such genes may lead to new strategies for diagnostic and therapeutic approaches that help mitigate metastatic events. High nm23 expression indicated a relatively low metastatic potential in many cell types (1). Although eight human nm23 homologs have been identified, only H1 and H2 have been extensively studied for their metastasis-related properties. It was shown that aurora-A associates with NM23-H1 and that NM23-H1 further phosphorylates the kinase repressor of Ras, a negative modulator of Ras downstream of the MEK/MAPK pathway (6, 7). The above data suggest that NM23 may be involved in Ha-ras-related metastasis process. In addition, overexpression of the NM23-H1 protein may indicate poor survival for metastatic tumors.

Materials and Methods

The stable NIH/3T3 cell line harboring an inducible Ha-ras\textsuperscript{val12} oncogene (designated as 7-4) and small interfering RNAs (siRNAs) were used to clarify the inverse correlation between nm23 and Ha-ras expression both in vitro and in vivo. A derivative 7-4/Z-3 cell line harboring a β-galactosidase reporter gene was used to trace cell metastasis in a murine tumor model. The data presented here reveal that Ha-ras\textsuperscript{val12} is able to cause cell morphological changes, induce tumor formation, and promote metastasis of tumor cells to the lungs. In mice with metastases, the immune surveillance against tumor formation was suppressed by Ha-ras\textsuperscript{val12} overexpression through an increase in T-reg cells and a decrease of cytotoxic T lymphocytes and natural killer cell populations. Our results suggest that the Ha-ras oncogene regulates morphogenesis, tumorigenesis, and metastasis through suppressing nm23 expression and modulation of immune cell function.

Results

Effect of Ha-ras\textsuperscript{val12} on nm23 Expression, Tumor Formation and Metastasis of the Transformants, and Immunomodulation in Tumor-bearing Mice

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Abstract. Overexpression of the Ha-ras\textsuperscript{val12} oncogene has frequently been detected in primary and metastatic carcinomas. NM23 is a metastasis inhibition factor and plays a suppressive role in metastasis in many types of cancer. In this study, a stable NIH/3T3 cell line harboring an inducible Ha-ras\textsuperscript{val12} oncogene (designated as 7-4) and small interfering RNAs (siRNAs) were used to clarify the inverse correlation between nm23 and Ha-ras expression both in vitro and in vivo. A derivative 7-4/Z-3 cell line harboring a β-galactosidase reporter gene was used to trace cell metastasis in a murine tumor model. The data presented here reveal that Ha-ras\textsuperscript{val12} is able to cause cell morphological changes, induce tumor formation, and promote metastasis of tumor cells to the lungs. In mice with metastases, the immune surveillance against tumor formation was suppressed by Ha-ras\textsuperscript{val12} overexpression through an increase in T-reg cells and a decrease of cytotoxic T lymphocytes and natural killer cell populations. Our results suggest that the Ha-ras oncogene regulates morphogenesis, tumorigenesis, and metastasis through suppressing nm23 expression and modulation of immune cell function.

Keywords: Ha-ras, nm23, metastasis, immunomodulation.
cervical cancer patients (8). However, the relationships of these proteins and their role in metastasis have not yet been determined.

The evasion of immunological surveillance is one of the primary strategies tumor cells utilize in becoming metastatic. Immunological modulation of T lymphocytes by their surface markers may be responsible for the regulation of tumor metastasis (9, 10). The population of regulatory CD4+ T-cells expressing the interleukin-2 (IL-2) receptor α-chain (CD25) plays a key role in controlling immunosuppressive capacity both in vitro and in vivo (11, 12). In this study, we clarified the relationship between Ras and nm23 and the role of nm23 in Ras-related tumor formation and metastasis. The effects of the Ha-ras oncogene on modulation of immune cells and the ability of Ha-ras-transformed cells to suppress immune surveillance and induce metastasis were also investigated.

Materials and Methods

Cell culture. NIH/3T3 and its derivatives, 7-4 and 7-4/Z-3, were maintained in minimum essential medium (MEM) containing 10% calf serum. The 7-4 cells contained the lacI and Ha-ras transgenes and the expression of Ha-rasval12 was regulated by the administration of isopropyl β-D-1-thiogalactopyranoside (IPTG) (13). After IPTG induction, the cells were observed at different intervals and the relative expression levels of the Ha-ras and nm23 genes in NIH/3T3 and 7-4 cells were measured by Northern and Western blotting. The 7-4/Z-3 cells harboring the Ha-ras oncogene and β-galactosidase reporter gene were detected by histochemical staining using X-gal (14, 15).

Northern blot analysis. Total RNA (20 μg) prepared from the cells was denatured using glyoxal (1.0 M). The denatured RNA was separated in 1% agarose at 80 V for 4 h in a 10 mM circulated sodium phosphate (NaPi) buffer system and then transferred to a Hybond-N membrane (Amersham, USA) using 25 mM NaPi buffer. A 4.7 kb Ha-ras probe, digested from the pSVlacOraS plasmid, was used to detect the expression of the transcreated Ha-ras construct (13). The nm23 primers (nm23-1a, 5'-CAGTCGCAGCCGGC GTFAAAG-3' and nm23-1b, 5'-GCAGCTAACACTGCACA-3') were used to amplify a 636 bp fragment by PCR, which was used as a probe to detect the endogenous nm23 RNA expression. The radioactive probes were hybridized with the RNA-bound membrane and the signals were detected following exposure to Kodak X-OMAT AR film. The quantification of the autoradiograph was done using scanning densitometry (420oe: PDI Inc, USA).

Western blot analysis. Protein expression of Ras in 7-4 cells was induced by different concentrations of IPTG. The cell lysates (50 μg) were separated in a 12% polyacrylamide gel and subsequently transferred to a PVDF membrane (Millipore, USA). After the membranes were blocked with 5% non-fat milk, Ras, NM23, and β-actin proteins were detected using anti-Ras (Oncogene, USA), anti-NM23-H1/H2 (Cell Signaling, UK), and anti-β-actin (Sigma, USA) antibodies, respectively. The reaction was followed by probing with peroxidase-coupled secondary antibodies and then detection using enhanced chemiluminescence (Amersham Pharmacia, USA).

Small interfering RNA (siRNA) transfection. The Ha-rasval12-specific siRNA was designed and synthesized by Qiagen, Inc (www.qiagen.com). The sense sequence was as follows: 5'-CCGGCAAGAGTGCGCTGACCACCTCAGTGATGGGTC AGCGCAC TCTGGTTTTTG-3'. The cells (5×10^5) were seeded in a 6-cm^2 cell culture dish and incubated in Dulbecco’s modified Eagle’s medium (DMEM) without antibiotics overnight in a 5% CO2 incubator. siRNA was transiently transfected into cells using Lipofectamine™ 2000 reagent (Invitrogen Inc, Carlsbad, CA, USA). Briefly, siRNA was diluted in 500 μl of serum-free DMEM to the assigned concentrations (0, 2, 4, 6, and 8 μM). The Lipofectamine-siRNA solution was mixed gently and added to the cells. The cells were incubated at 37°C in a 5% CO2 incubator for 4 h. The media were then replaced with normal media and the cells were incubated for an additional 48 h before protein extraction.

Inoculation of tumor cells into mice. Male, 6-week-old BALB/c and SCID mice were obtained from the National Laboratory Animal Breeding and Research Center, Taipei, Taiwan. The experimental protocol adhered to the rules of the Animal Protection Act of Taiwan and was approved by the Laboratory Animal Care and Use Committee of the National Cheng Kung University. Mice were randomly divided into the three groups. Using a 28-gauge needle, the mice were injected subcutaneously (s.c.) in the back with 0.2 ml of NIH3T3, 7-4 or 7-4/Z-3 cells (2.0×10^6 cells/each side). Briefly, group A was inoculated with PBS alone, group B was inoculated with NIH3T3 cells and group C was inoculated with 7-4/Z-3 cells. Each group comprised eight mice and four mice were housed per cage and provided with sterile food and water ad libitum. Tumor sizes were measured using Vernier calipers to calculate the mean diameter at the second and third weeks.

Tumor embedding, sectioning, and X-gal staining. Two weeks post-inoculation with PBS, NIH3T3 or 7-4/Z-3 cells, the lungs and tumors were removed from the injected mice immediately after sacrifice and embedded in cryoress. The frozen embedded tumors were processed by cryosection using a microtome (Shandon, Pittsburgh, USA). The sections were rinsed with Dulbecco’s modified phosphate-buffered saline (DPBS) and fixed for 1 h at 4°C with 2% (v/v) formaldehyde and 0.2% (v/v) glutaraldehyde in DPBS. The fixed sections were rinsed with DPBS three times and then stained with staining solution (20 mM potassium ferricyanide, 2 mM MgCl2, 0.02% [v/v] Nonidet-P40, and 0.01% sodium deoxycholate [w/v] in HEPES buffered DPBS) containing X-gal (1 mg/ml) at 37°C overnight (16).

Lymphocyte preparation from tumor-draining lymph node (TDLN) and splenocytes. Two weeks post-inoculation with PBS, NIH3T3 or 7-4/Z-3 cells, mice were sacrificed by injecting an overdose of sodium pentobarbital. The spleen and superficial inguinal TDLNs of each mouse were collected, weighed, minced finely, and passed through a stainless mesh in RPMI-1640 with 10% fetal bovine serum (FBS). Any residual tissue was further disintegrated by passage through a 21-gauge needle. Erythrocytes in the spleen were lysed for 5 min with 2 ml of lysis solution (0.15 mM NH4Cl, 10 mM KHCO3, and 0.1 mM EDTA, pH 7.2). The cells were washed in Hank’s balanced salt solution (HBSS), and the number of viable cells was measured using the trypan blue dye exclusion method.
**Flow cytometric analysis of splenocytes.** The phenotypes of the different lymphocyte populations were determined by the detection of surface markers with specific monoclonal antibodies (mAbs). Briefly, cell suspensions (5.0×10^5) were incubated on ice for 30 min in the dark in the presence of fluorescein, phycoerythrin, or PerCP-conjugated mAb (PharMingen, San Diego, USA). Cytofluorometric analysis was performed by laser excitation at 488 nm on the lymphocyte fractions. A computer system (CellQuest; BD Biosciences, Mountain View, CA, USA) was used for data acquisition and analysis. Percentages were calculated on the basis of the number of lymphocytes found in each quadrant. T-Cells were identified by expression of the surface marker CD3ε (clone 145-2C11), T-helper cells by CD3ε/CD4 (clone RM4-5), T cytotoxic/suppressor cells by CD3ε/CD8 (clone 53-6.7), B-cells by CD19 (clone 1D3), and natural killer (NK) cells by PanNK (clone DX5). Identification of natural T-reg cells by CD25 (clone PC61) and CD4 markers indicates the cell surface with the interleukin-2 receptor α-chain in T-helper cells (17).

**Statistical analysis.** The weights of tumor and spleen, and subpopulations of splenocytes are presented as the mean±SD. One-way analysis of variance (ANOVA) was used for statistical analysis. The mean values of two groups were considered significantly different when \( P < 0.05 \).
Figure 1. Time course of mRNA and protein expression of Ha-ras\textsuperscript{val12} and nm23 genes in 7-4 cells. A: The 7-4 cells were induced with IPTG (20 mM) continuously for 6 days. B: The 7-4 cells were induced with IPTG for one day and then grown under normal conditions without IPTG for the remaining days. The total RNA was extracted from treated cells at each time point followed by Northern blotting using the γP32-labeled probes of Ha-ras\textsuperscript{val12} and nm23; 28S and 18S were used as the loading control. C: Total protein was extracted from 7-4 cells in the presence of various dosages of IPTG for 4 days. D: Total protein was extracted from 7-4 cells after treatment with Ras si-RNA at various dosages for 48 h. Antibodies for Ha-ras, NM23 and β-Actin were used to detect each individual gene. E: The 7-4 cells with or without IPTG (5 mM) treatment were injected subcutaneously into SCID mice. Protein was extracted from the tumors to detect Ha-ras and NM23 by Western blotting. β-Actin was used as the internal control. F: Quantification of Ha-ras and NM23 expression levels in Figure 1E as well as the tumor weight in mice injected with 7-4 cells treated with or without IPTG.
Results

nm23 gene expression inversely correlates with Ha-ras oncogene expression in Ha-ras transformants. At the transcriptional level, high expression of nm23 was observed on the first day post-IPTG induction in 7-4 cells. The expression levels of nm23 in the cells that continuously overexpressed Ha-ras<sup>Val12</sup> gradually declined, falling to half of the amount at day six as compared to the level on the first day post-IPTG induction (Figure 1A). In contrast, the expression levels of nm23 gradually increased from day one to day six if IPTG was removed one day after induction (Figure 1B). Consistently, IPTG induced Ha-ras overexpression at the translational level, in a dose-dependent manner, and NM23 protein expression was inversely suppressed (Figure 1C). Ha-ras siRNA suppressed Ras expression in a dose-dependent manner, and the expression of NM23 protein increased conversely (Figure 1D).

The 7-4 cells with or without IPTG (5 mM) treatment were subsequently injected s.c. into SCID mice. Tumor formation was investigated over the subsequent 17 days, and the tumor proteins were then extracted for evaluation of specific protein expression. Our data showed that Ha-ras expression was significantly higher in the tumors from the 7-4 cells with IPTG treatment relative to the tumors derived from 7-4 cells without IPTG treatment (Figure 1E and 1F). Consistently, the 7-4 cells with IPTG treatment also induced significantly larger tumor formation compared to 7-4 cells without IPTG treatment (Figure 1F). Conversely, the expression of NM23 was lower in tumor from 7-4 cells with IPTG-induction when compared to tumors derived from 7-4 cells without IPTG treatment (Figure 1E and 1F). In summary, our data demonstrate a positive correlation between Ha-ras expression and tumor formation and an inverse correlation between Ha-ras and NM23 both in vitro and in vivo.

Ha-ras overexpression causes morphological changes and cell aggregation. The 7-4 cells became round and formed aggregates in the presence of IPTG for 3 days (Figure 2C). In contrast, these same cells without IPTG induction, or incubated in the presence of IPTG for only one day and in the absence of IPTG for the following 7 days, did not
become round or form aggregates (Figure 2A and 2B). Our data indicate that Ha-ras overexpression changes the cell morphology and causes substantial cell aggregation.

The 7-4 and 7-4/Z-3 cells cause tumor growth and lung metastasis in BALB/c Mice. To determine the potential of the Ha-ras transformants for tumor growth and metastasis, 7-4, 7-4/Z-3, control NIH/3T3 cells, or PBS only were injected s.c. into BALB/c mice. In the mice injected with 7-4/Z-3 cells, tumors were detected and the size was measured (see Table I). Metastasis of the ras transformants was determined by the detection of cells positively stained for X-gal in lung cryosections as well as nodule formation in the lung (Figure 3B and 3D). Our results revealed that 7-4 and 7-4/Z-3 cells harboring Ha-rasval12 or Ha-rasval12 and β-galactosidase genes form tumors and also readily migrated from the initial tumor site to the lungs of the inoculated mice.

Ha-ras overexpression affects a subpopulation of the immune cells in the spleen and TDLN of the tumor-bearing BALB/c mice. The weight, number of cells, and the lymphocyte subpopulations in the spleen and TDLN of the different groups of mice are shown in Table II. The 7-4/Z-3 tumor-bearing mice (Group C) showed significant increases in spleen weight and the number of splenocytes. The percentage of activated T-reg cells (CD4+/CD25+) of the splenocytes and TDLNs was significantly higher in the 7-4/Z-3 tumor-bearing mice when compared to the control group (Group A). In contrast, the percentage of cytotoxic T-cells (CD3ε+/CD8+) and natural killer cells (PanNK+) was significantly lower in these mice. Together, these data indicate that the Ha-ras transformed cell line (7-4/Z-3) modulated the immunity against Ha-ras transformants in tumor-bearing mice.

Discussion

Using rat embryo fibroblast (REF) cells, Steeg demonstrated that adenovirus early region IA (EIA) suppressed the metastatic potential of REFs overexpressing the Ha-ras oncogene and the expression of NM23 was also increased (16). This indicates that adenovirus EIA can induce NM23 expression which is involved in the suppression of metastasis. The present study demonstrates that increased tumor formation and lung metastasis correlate with lower nm23 gene expression. The expression of nm23 at the transcriptional and translational levels was decreased while Ha-ras was continuously overexpressed for 6 days (Figure 1). In this study, the anti-NM23 antibody detects both NM23-H1 and NM23-H2. At least three mechanisms are thought to contribute to the metastasis-suppressive effect of NM23-H1: Firstly, NM23-H1 demonstrates histidine kinase activity toward ATP-citrate lyase, aldolase C, and the kinase suppressor of ras (KSR), with the latter inactivating mitogen-activated protein kinase signaling. Secondly, binding proteins that reduce the availability of free NM23-H1 can inhibit its ability to suppress metastasis. Thirdly, NM23-H1 can alter downstream gene expression and demonstrates an inverse association with the lysophosphatidic acid receptor endothelial differentiation gene-28 (EDG2) (18). NM23-H2 is a negative regulator of epidermal growth factor (EGF)- and Ras-mediated cellular proliferation via its effect on the
extracellular signal-regulated protein kinase (ERK) pathway (19). The underlying mechanism and signaling pathway(s) that Ha-ras influences to suppress NM23 remain to be determined.

The morphology of the transformed cells reverted to the non-transformed form after withdrawal of IPTG (Figure 2, Group B). The nm23 levels decreased while Ha-ras was overexpressed and the cells became round and formed aggregates. These aberrant cell morphological changes were detected under continuous IPTG induction. We have demonstrated that Ha-rasVal12-induced morphological change is via microtubule disruption and via the activation of MEK/ERK signaling pathway that post-translationally down-regulates Stat3 expression (20). In our murine metastasis model, initially, we demonstrated that all the injected mice formed tumors and in the Ha-ras overexpressed tumor, the NM23 level was reduced (Figure 1E and 1F). Furthermore, we revealed that 7-4 and 7-4/Z-3 (harboring a β-galactosidase reporter gene) cells migrated to the lung and formed tumors as demonstrated by tumor nodules and blue cells labeled by X-gal staining (Figure 3B and 3D). Collectively, we reveal a relationship between Ha-rasVal12 overexpression and increased tumor formation, decreased nm23 expression, cell morphological changes, as well as the metastasis of the transformants both in vitro and in vivo.

Immunomodulation of T lymphocytes plays an important role in immune surveillance against tumor cell invasion and metastasis. In this study, we monitored a population of immune cell subtypes in the spleen and TDLNs using immunofluorescence to assess lymphocyte surface antigens. In 7-4/Z-3 tumor-bearing mice, cytotoxic T-cell (CD3e+/CD8+) and NK (PanNK+/CD45+) populations in the spleen and TDLNs were significantly increased at 7-4/Z-3 bearing mice. In contrast, CD25+/CD4+ T-reg cells in the spleen and TDLNs were significantly decreased. This suggests that the increase in T-reg cell numbers might be due to down-regulation of tumor-specific CD8+ cytotoxic T lymphocytes (CTLs) and general cytotoxic NK cells, as previously reported. Suppression of CD8+ CTLs and NK cell populations reduces their cytolytic activity and may enhance the initial activation of tumor cells and help sustain tumor growth (21, 22). Clark et al. demonstrated the presence of a reduced population of CD8+ CTLs in highly invasive pancreatic cancer (23). Such a microenvironment contributes to the great reduction of tumor-associated myeloid-derived suppressor cells and creates the opportunity for tumor cells to escape from immunosurveillance (24).

In conclusion, nm23 appears to be responsible for the suppression and regression of tumors, and continuous Ha-ras overexpression and tumor formation correlates with down-regulation of nm23 expression. Our findings shed the light on the regulation between nm23 and oncogenic Ha-ras-associated tumorigenesis, as well as metastasis. Ha-ras may also increase the metastatic potential of tumor cells through regulation of the activation of T-reg cells and suppressing the immune surveillance by CTLs and NK cells.

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