Expression of Müllerian inhibiting substance type II receptor and antiproliferative effects of MIS on human cervical cancer

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Abstract. This study aimed to analyze expression of Müllerian inhibiting substance type II receptor (MISRII) protein and mRNA in cervical neoplasia, to demonstrate the growth inhibition of cervical cancer cells by administration of highly purified recombinant human Müllerian inhibiting substance (MIS) and, furthermore, to evaluate the clinical significance of MIS as a biological modifier for MIS receptor expressing tumors. Reverse transcriptase polymerase chain reaction (RT-PCR) was used for MISRII mRNA expression, and \textit{in situ} hybridization and immunohistochemistry were used to observe expression, location of MISRII mRNA and protein, respectively. To demonstrate the effect of MIS on the viability of cervical cancer cells, methyl thiazole tetrazolium (MTT) assay was performed. Flow cytometry was used to evaluate the cell cycle distribution after exposure to MIS in cervical cancer cells, and the annexin-V-FITC staining method was performed to demonstrate apoptosis by MIS in cervical cancer cells. Expression of MISRII protein and mRNA were observed in all normal cervical and cervical carcinoma tissues. There was no significant difference in expression of MISRII protein and MISRII mRNA between normal cervical and cervical carcinoma tissues. MTT assay showed negative correlation between MIS exposure time and the viability of cervical cells (P=0.008). The changes in cell cycle distribution after MIS exposure suggest that MIS plays an important role in inducing cellular apoptosis by causing arrest at the G\textsubscript{1} phase and increasing cells at sub-G\textsubscript{0}/G\textsubscript{1} phase. Annexin-V-FITC staining methods showed that cellular apoptosis was, respectively, 10.44 and 12.89% after 24 and 48 h of MIS exposure in cervical carcinoma cells. There was a negative correlation between cellular survival and MIS exposure time. This study demonstrates that MISRII is present on normal cervical and cervical carcinoma tissues, and MIS shows receptor-mediated antiproliferative effect on cervical cells \textit{in vitro}. These data suggest that MIS may be used as a biological modifier or therapeutic modulator on MISRII-expressing tumors in the future.

Introduction

In 1946, Alfred Jost observed that the regression of the Müllerian duct in the rabbit embryo was not due to testosterone after removing undifferentiated gonad and inserting testosterone crystals in its place. He named the critical agent produced by testis, Müllerian inhibiting substance (MIS) (1), also known as anti-Müllerian hormone (AMH). Following the discovery, an \textit{in vitro} MIS bioassay was developed (2), bovine MIS was purified (3) and an immunoassay was created employing monoclonal antibody (4). The human gene was cloned and recombinant human MIS (rhMIS) was purified (5) soon thereafter a human specific ELISA came to use (6–8). MIS is a 140-kilodalton (kDa) glycoprotein homodimeric glycoprotein that is a member of TGF-\(\beta\) multigene family (9). The human MIS gene is on short arm of chromosome 19 the encoded protein produces a 25-kDa carboxy-terminus by proteolytic cleavage. This domain is sufficient to induce Müllerian duct regression \textit{in vitro}, whereas 110 kDa amino-terminal extends the half-life of MIS (10) and may facilitate MIS activity. The MIS receptor is expressed in all tissues of Müllerian duct origin. It is a heteromeric complex consisting of single membrane spanning type I and II serine threonine kinases. The MIS carboxy-terminus binds to a unique MISRII which in turn phosphorylates one of several MIS type I receptors and produces signal transduction via a cascade of the intracellular Smad pathway originally found to be used in BMP signaling (11,12). Other studies suggest that MIS inhibits the cell growth by different cellular signaling pathways involving cyclin-dependent kinase inhibitor (CDKI) (19,22,23) or nuclear factor-\(\kappa\)B (NF\(\kappa\)B) (13). In addition to being an inhibiting factor associated with sex differentiation and reproductive physiology, the MIS receptor was identified in tissues other than the reproductive system, therefore, MIS could be a multifunctional cellular modulator (14).

MIS is produced by the embryonic testes and responsible for the regression of the Müllerian duct by binding the receptor...
in mesenchyme surrounding the Müllerian duct. Thus blocking its differentiating into the oviduct, uterus, upper vagina and the outer lining of the ovary. MIS is also associated with fetal lung maturation (15), and differentiation of the ovary and testis (16). The serum concentration of MIS is 10-70 ng/ml in neonatal boys, it slightly elevates before puberty then declines after puberty, and maintains a level of 2-5 ng/ml (6,17). In females, MIS is not produced by the ovarian granulosa cells to measurable levels in serum or follicular fluids until after puberty. The concentration of MIS in females is similar to male (2-5 ng/ml). In postmenopausal woman, MIS is not produced. This allows it to be used as a parameter of an ovarian function (6,9,17). In addition, MIS regulates recruitment and maturation of follicle, folliculogenesis and acts as an inhibiting factor of steroid hormone production, in autocrine, paracrine and endocrine manner (9,17). MIS not only inhibits differentiating into the oviduct, uterus and upper vagina in male, but also inhibits meiosis of oocyte in late prophase from 3 days after birth to puberty before ovulation in female. In male, MIS inhibits meiosis of germ cell in prophase, although varying in degree according to the timing. MIS acts as a regulatory factor of cell division both in male and female (16,18). MIS-deficient male mice had Leydig cell hyperplasia and hyperandrogenism while MIS-deficient female mice showed accelerated oocyte degeneration and increased follicular atresia (16,18-21).

Not only is MIS an inhibiting factor of a sexual development and reproductive physiology, it is established as a regulator of cell growth. After MIS was proposed as possible tumor growth suppressor, it has been investigated intensively, since the early 1980's. The ovarian surface epithelium morphologically simulates the lining cells of the fallopian tube, which are derivatives of the coelomic epithelium of the urogenital ridge, which invaginates to form a Müllerian duct early in embryonic life (11). Several studies provide physiologic confirmation of the Müllerian origin of serous ovarian tumors and suggest that MIS may ultimately prove to be effective in its therapy. The results of the studies presented show that the growth of ovarian tumor cell lines and ovarian tumors were suppressed by MIS concentration, especially effective in highly purified rhMIS (22-25).

It has also been demonstrated in previous studies that proliferation of cervical, endometrial and breast cancer cells are suppressed by MIS treatment (13,26,27). Many of these studies mentioned the potential of MIS as a treatment of MIS-Receptor expressing tumors, however, in-depth discussion or investigation has yet to come (14). MIS inhibits proliferation of cervical cancer cell lines (CaSki, SiHa, C33A) (26). Previously, we demonstrated the expression of MIS type II receptor (MISRII) in the human papilloma virus (HPV)-16 related cervical cancer cell lines (CaSki, SiHa) and a non-HPV-related cervical cancer cell line (C33A) (28). We also showed that MIS inhibited growth of cervical cancer cells, and induced cellular apoptosis of C33A. In addition, we identified characteristic molecular signature of MIS in CaSki cells by using whole genome expression analysis (28). However, evidenced based data on the potential benefit of MIS against human cervical cancers requires further study.

Therefore, this study is aimed to analyze expression of MISRII protein and mRNA in cervical neoplasia, to demonstrate the growth inhibition of cervical cancer cells by administration of highly purified MIS, and, furthermore, to evaluate the clinical significance of MIS as a biological modifier for MIS receptor expressed tumors.

Materials and methods

Clinical specimens. Paraffin-fixed 12 normal cervical tissues and 30 cervical cancer tissues (6 squamous cell carcinoma in situ, 2 microinvasive squamous carcinoma, 11 squamous cell carcinoma, 2 adenocarcinoma in situ, 2 microinvasive adenocarcinoma, 7 adenocarcinoma) were obtained through St Mary's Hospital Tissue Banks in the Clinical Pathology Department from September 2006 to May 2010. Fifteen fresh human tissues (4 normal cervical tissue, 9 squamous carcinoma, 2 adenocarcinoma) were obtained from discarded tissues of patients undergoing surgery at the hospital. Among four fresh tissues of normal cervix, there was one case infected by HPV 51. On the other hand, there were 12 cases infected by HPV 16 (7 cases), HPV 18 (2 cases), HPV 31 (1 case), HPV 33 (1 case), HPV 52 (1 case) among 11 fresh tissues of cervical cancer. One fresh tissue of squamous cancer was co-infected by HPV 16 and HPV 18. The patients ranged in age from 34 to 68 years (mean 45.4 years). This study was approved by the Institutional Review Board-Human Research Committee at the Hospital (no. SCMC07BR22). Informed consent was obtained from each patient.

Construction of tissue microarray block. For the construction of the tissue microarray block (TMB), H&E-stained sections from all specimens were carefully reviewed. Tissue cylinders 3 mm in diameter were punched from carefully selected histologically representative regions of each paraffin-embedded donor tissue block and brought into a recipient paraffin block using a tissue punch instrument. In constructed TMB, clinical samples were arrayed in 5 rows x 6 columns position, and control samples were positioned in the 7th column.

Tissue preparation. In order to extract the RNA for reverse transcriptase polymerase chain reaction (RT-PCR), 4 normal cervical tissues and 11 cases of cervical cancer were frozen and stored in liquid nitrogen. For in situ hybridization and immuno-histochemistry, 5 μm-thick sections from TMB and some paraffin-embedded donor blocks were prepared on the slides (Probe on slide, Fisher Scientific Co., Pittsburgh, PA, USA).

The expression of MISR II mRNA by RT-PCR. Total RNA was isolated from 4 normal cervical tissues and 11 cases of cervical cancer using RNA tissue kit (Boehringer-Mannheim GmbH, Mannheim, Germany). First-strand cDNA was reverse-transcribed from the total RNA using RT kit (Boehringer-Mannheim) according to the manufacturer's instruction. The each reaction mixture was as followed; 10X reverse transcription buffer 2 μl, MMLV reverse transcriptase 0.8 μl, dNTP mix 2 μl, 25 mM MgCl 4 μl, random hexamer 2 μl, RNase inhibitor 1 μl, total RNA 2 μg, and total volume was adjusted to 20 μl with Dnase water. The reaction mixtures were incubated for 10 min at 25°C, and then for 60 min at 42°C followed by RT enzyme inactivation step for 10 min at 70°C. The reaction products were then stored at -20°C before the next procedure.

RT-PCR. cDNA products were amplified using by Takara PCR amplification kit (Takara Shuzo Corp., Shiga, Japan). PCR mixture consisted of 10X PCR buffer 3 μl, dNTP 2.4 μl,
downstream primer (10 pmol) 1 µl, upstream primer (10 pmol) 1 µl, Taq polymerase 0.4 µl, cDNA 1 µl, and DEPC water 21.2 µl. Amplification started with denaturation at 94°C for 4 min followed by 30 cycles of 94°C for 1 min, 58°C for 1 min and 72°C for 1 min. The final extension was made at 72°C for 10 min. The sequences of PCR primers were as follows; upstream primer 5'-ccctgcagctagaaac-3' (MISR II cDNA; Gene Bank, accession no. AF172932; sequence 581-600), and downstream primer 5'-tgggcatagttggcagc-3' (sequence 921-941). PCR products (4 µl) were separated by electrophoresis on a 1% agarose gel, and the band size (361 bp) was examine compared with DNA ladder.

**Sequence protocol.** Sequencing reactions were performed in an MJ Research PTC-225 Peltier Thermal Cycler using an ABI PRISM Big Dye Terminator Cycle Sequencing Kits with AmpliTaq DNA polymerase (FS enzyme) (Applied Biosystems, Foster City, CA, USA), following the protocols supplied by the manufacturer. Single-pass sequencing was performed on each template using primer. The fluorescent-labeled fragments were purified from the unincorporated terminators with an ethanol precipitation protocol. The samples were resuspended in distilled water and subjected to electrophoresis in an ABI 3730xl sequencer (Applied Biosystems).

**Immunohistochemistry.** Tissues for immunochemical detection of the MIS type II receptor were processed using a fast temperature-controlled machine, microprobe immunostaining station (Biomedica Co., Foster City, CA, USA). The immunostaining procedure was as follows: TMB slides were preheated at 85°C for 30 min, dewaxed with xylene (2 min, 4 times) and re-hydrated with water. In order to retrieve antigenic sites, the slides were autoclaved at 121°C for 10 min in citrate buffer (Zymed Lab. Inc., San Francisco, CA, USA), and cooled for 20 min at room temperature. The slides were then treated with 3% H₂O₂ at 45°C for 5 min to eliminate endogenous peroxidase activity followed by T-TBS wash 4 times. After treatment with normal rabbit serum (Zymed Lab. Inc.) at 45°C for 5 min to block the non-specific protein binding, the slides were incubated with rabbit polyclonal anti-human MIS type II receptor antiserum (provided by Dr David T. MacLaughlin, Massachusetts General Hospital, Boston, MA, USA) as primary antiserum at 4°C overnight. The slides were rinsed in T-TBS 4 times, and incubated with biotinylated anti-rabbit IgG (Zymed Lab. Inc.) as the second antibody at 45°C for 7 min. After T-TBS rinse 4 times, Streptavidin HRP detection system (Zymed Lab. Inc.) was applied to the slides at 45°C for 5 min to induce the biotin-avidin binding reaction. The slides were treated with 3-amino-9-ethylcarbazole (AEC) for 10 min at room temperature, counterstained with hematoxylin, and then mounted with glycerol gel.

**In situ hybridization.** Five µm-thick sections from TMB were preheated at 85°C in oven for 30 min, dewaxed with xylene (2 min, 4 times). After dewaxing, slides were rinsed with 100% alcohol, and then dried at 40°C in oven for 1 h. The dried sections were treated in 0.2 N HCl for 20 min and incubated in 20 µg/ml pepsin (0.1 N HCl) for 20 min at room temperature, which was followed by three washes with diethyl pyrocarbonate-treated PBS. The sections were dehydrated with ethanol and dried. Prehybridization and hybridization steps were carried out at 53°C for 2 and 15 h, respectively. The prehybridization buffer was composed of 50% formamide, 4X SSC, 10% dextran sulfate, 1X Denhardt's solution, and 1 mg/ml salmon sperm DNA. The hybridization buffer was identical to the prehybridization buffer except that salmon sperm DNA was substituted with 200 ng/ml MISR II riboprobe. After post-hybridization washing, sections were incubated with anti-digoxigenin antiserum conjugated with alkaline phosphatase (Boehringer-Mannheim), and histochemical detection was then performed using 4-nitroblue tetrazolium and 5-bromo-4-chloro-3-indoly-phosphate (Boehringer-Mannheim).

**Cervical cancer cell culture and addition of MIS.** Fresh human uterine cervix cancer specimens were processed as quickly as possible after collection. The tissues were rinsed twice with phosphate-buffered saline (PBS) and were cut into ~1-3 mm³ sized pieces. The portions of minced tumor were then placed into 50 ml conical tube containing 20 ml of enzyme solution (0.14% collagenase type I (Sigma, St. Louis, MO, USA) and 0.01% DNaseI (Sigma, 20,000 U/ml in Dulbecco's modified Eagle's medium (Gibco, Grand Island, NY, USA)), and incubated on a magnetic stirring apparatus for 2 h at 37°C. Enzymatically dissociated tumor was then filtered through 100 mm nylon mesh to generate a single cell suspension. The resultant cell suspension was then washed twice in Dulbecco's modified Eagle's medium (Gibco) containing 10% fetal bovine serum (FBS), 100 U/ml penicillin and 100 U/ml streptomycin. The pieces were subsequently incubated at 37°C in humid air with 5% CO₂, rhMIS (71 nM) was added on the cervical cancer cell cultures.

**The measurement of cell viability by MTT assay.** Three thousand cells/well were seeded in 96-well plates (Costar, Cambridge, MA, USA) in DMEM with 10% FBS. After 24 h the cells were exposed to 71 nM of MIS for 24 and 48 h. The treated cells were washed with PBS and 100 µl of MTT solution (5 mg/ml MTT stock in PBS diluted to 1 mg/ml with 10% DMEM) was added to each well. Cells were incubated for 4 h at 37°C at the end of which time 200 µl DMSO (Sigma) was added and incubated further for 30 min at room temperature in the dark. Optical densities at 550 nm were measured using an ELISA plate reader (Bio-Tek Instruments, Winooski, VT, USA).

**Cell cycle analysis.** Cells were exposed to MIS for 24 and 48 h, collected by trypsinization, fixed with methanol and stored for 30 min at 20°C with the equivalent volume of PBS (pH 7.4) vehicle control. The cells were washed with PBS and resuspended in 1 ml DNA staining solution (20 µg/ml propidium
iodide, 200 µg/ml DNase free RNase) and incubated in the dark at 37˚C for 30 min. The cells were analyzed on a FACSVatage SE flow cytometer (Becton-Dickison, San Jose, CA, USA). The results were analyzed using Cell Quest™ software and Modfit LT 3.0 program (Verity Software House, Topsham, ME, USA).

**Annexin-V analysis.** MIS treated cells were stained for annexin-V and propidium iodide (PI) using the Annexin-V-FITC Apoptosis Detection Kit I (BD Biosciences, San Diego, CA, USA) according to the manufacturer’s protocol. Briefly, following drug treatment, 1x10^5 cells were pelleted and washed once with PBS and resuspended in 100 ml of binding buffer [10 mM HEPES (pH 7.4), 150 mM NaCl, 5 mM potassium chloride, 1 mM MgCl₂, and 2 mM calcium chloride]. Subsequently, 5 µl of annexin V-FITC and PI was added to the cells that were then incubated for 15 min at room temperature in the dark. After incubation, 400 µl of binding buffer was added to the stained cells, and cells were analyzed using a FACSVatage SE Flow Cytometer (Becton-Dickison). Data analysis was conducted using Cell Quest™ software.

**Expression scoring system and statistical analysis.** The intensity of immunohistochemistry and in situ hybridization staining were assessed independently by 2 pathologists on a scale of increasing intensity, 0 (no staining), 1 (weak), 2 (moderate), and 3 (strong staining). Data were analyzed by using Wilcoxon rank sum test and Wilcoxon signed rank test. A P<0.05 was considered statistically significant.

**Figure 1.** RT-PCR of human MISRII mRNA from human cervix and cervical cancers. Bands of 361 bp are detected in normal cervix (A and B), squamous cell carcinoma of cervix (C), and adenocarcinoma of cervix (D). Lane E is the DNA ladder.

**Figure 2.** Light micrography of human uterine cervix. (A) The squamous epithelium moderately express MISRII in the cell membrane. In this and all subsequent panels, the right lower boxed area is of higher magnification (x400). Chromogen, AEC. Magnification x200. (B) In carcinoma in situ (CIS) of the cervix cells moderately express MISRII. (C) Microinvasive squamous cell carcinoma of the cervix moderately express MISRII in the cell membrane. (D) Squamous cell carcinoma, well differentiated of uterine cervix, moderately express MISRII in the cell membrane. (E) Moderately differentiated carcinoma of uterine cervix moderately expresses MISRII. (F) Poorly differentiated cells moderately express MISRII. (G) Human adenocarcinoma in situ (AIS) strongly express MISRII. (H) Microinvasive adenocarcinoma diffusely and strongly express MISRII. (I) Adenocarcinoma, well differentiated diffusely and strongly express MISRII. (J) Adenocarcinoma, moderately differentiated diffusely and strongly express MISRII. (K) Adenocarcinoma, poorly differentiated diffusely and strongly express MISRII. (L) Human liver specimen used as negative control. No immunoreactivity for MISRII is detected.
Results

Expression of MISRII mRNA by RT-PCR. We analyzed MISRII mRNA expression in 2 normal cervical tissues, a squamous cell carcinoma of cervix, and an adenocarcinoma of cervix by RT-PCR and all tissues showed 361 bp band which was confirmed to be identical to a segment of human MISRII cDNA sequence (581-941) (Gene Bank, accession no. AF172932) (Fig. 1).

Expression of MISRII protein by immunohistochemistry. In all twelve normal human uterine cervical samples, the squamous epithelium moderately express MISRII protein in the cell membrane (2.33±0.19) (Fig. 2A). The cancer cells in cervical carcinoma in situ (6 cases) moderately express MISRII protein (2.17±0.31) (Fig. 2B). The cancer cells in microinvasive squamous carcinoma (2 cases) moderately express MISRII protein (2.50±0.50) (Fig. 2C). In 11 cases of invasive squamous carcinoma, the cancer cells diffusely and moderately express MISRII protein in the cell membrane with similar intensity according to cancer differentiation (2.27±0.19) (Fig. 2D-F).

In adenocarcinoma in situ (2 cases) and microinvasive adenocarcinoma (2 cases), the cancer cells strongly express MISRII protein (2.50±0.5 and 2.50±0.5, respectively) (Fig. 2G and H). In adenocarcinoma (7 cases), the cancer cells strongly express MISRII protein in the cell membrane with similar intensity according to cancer differentiations (2.43±0.20) (Fig. 2I-K). In human liver specimen, negative control for MISRII immunostain, no immunoreactivity is detected (Fig. 2L).

Expression of MISRII mRNA by in situ hybridization. In all twelve normal human uterine cervical samples, the squamous epithelium moderately express MISRII mRNA in the cell membrane (2.25±0.18) (Fig. 3A). The cancer cells in cervical carcinoma in situ (6 cases) moderately express MISRII mRNA (2.50±0.22) (Fig. 3B). The cancer cells in microinvasive squamous carcinoma (2 cases) moderately express MISRII mRNA (2.00±0.00) (Fig. 3C). In 11 cases of invasive squamous carcinoma, the cancer cells diffusely and moderately express MISRII mRNA in the cell membrane with similar intensity according to cancer differentiation (2.27±0.14) (Fig. 3D-F).
carcinoma (2 cases), the cancer cells strongly express MISR II mRNA (2.50±0.5 and 2.50±0.5, respectively) (Fig. 3G and H). In adenocarcinoma (7 cases), the cancer cells strongly express MISR II mRNA in the cell membrane with similar intensity according to cancer differentiation (2.28±0.18) (Fig. 3I-K). In human liver specimen, used as negative control, no in situ signal is shown (Fig. 3L).

Effect of MIS on the viability of cervical cancer cells. Cells were treated with 71 nM MIS at 24 h after plating. At 24 and 48 h after treatment, cells were incubated with MTT, and the absorbance was read at 550 nm. Results are presented as percentage of control which was calculated using the equation: (mean absorbance of treated cells/mean absorbance of control cells) x 100. Data are expressed as mean ± standard deviation (SD) from 5 independent experiments. *P<0.01 as compared to corresponding control cells.

In order to find out the changes in cell cycle distribution after MIS exposure, cells were treated with 71 nM MIS for 24 and 48 h and histograms of cellular DNA content were obtained by flow cytometry (Fig. 5). In control incubations the proportion of cellular DNA of G0/G1, S, G2/M and sub-G0/G1 phase was 46.43, 23.83, 26.49 and 3.25%, respectively. At 24 h after treatment with MIS, the proportions were 48.43, 19.09, 22.14 and 10.34%, respectively.

The proportion of cellular DNA of G0/G1, S, G2/M and sub-G0/G1 phase in controls for the 48 h MIS exposure experiment were 47.08, 23.20, 25.21 and 4.51%, respectively. At 48 h after treatment with MIS, the proportions were 52.43, 14.95, 19.55 and 13.07%. In short, the changes in cell cycle distribution after MIS exposure at 24 and 48 h demonstrated that S and G2/M phases were shortened, G0/G1 and sub-G0/G1 phases were lengthened. Therefore, MIS plays an important role in inducing cellular apoptosis by causing arrest at G1 phase and increasing cells at sub-G0/G1 phase.

Induction of apoptosis by MIS in cervical cancer cells. Cells were treated with 71 nM MIS for 24 or 48 h. For apoptosis, the externalization of phosphatidylserine was assessed by measuring annexin-V-FITC binding using propidium iodide as a counterstain (Fig. 6). The proportion of surviving cells/ early apoptotic cells/late apoptotic cells and necrotic cells on annexin-V-FITC staining were 92.13/1.78/1.04% in controls and changed to 78.85/10.44/6.32% after 24 h with treatment of MIS.
The proportions were 90.11/2.47/3.53% in control and changed to 73.84/12.89/9.92% after 48 h, respectively. Annexin-V-FITC staining method showed that cellular apoptosis was 10.44 and 12.89% after 24 and 48 h of MIS exposure in cervical carcinoma cells. There was a negative correlation between cellular survival and MIS exposure time.

Discussion

A considerable body of *in vitro* and *in vivo* animal evidence supports the conclusion that recombinant human MIS may affect cancer cell growth, particularly those originating from Müllerian tissue including endometrial, cervical, fallopian tubal, and ovarian, perhaps even cancer stem cells (13,14,22-37). Data presented in the current study confirm and extend the fact that human cervical cancers are also potential MIS targets. The usual treatment of malignant female reproductive tract tumors of surgery followed by chemotherapy or radiation therapy has resulted in serious systemic toxicities including induced infertility. Additionally, relapse is common due to drug resistance. Because MIS mechanism is different from the traditional chemotherapy, MIS treatment combined with chemotherapy holds promise in controlling the tumor size more effectively, with lower dose of chemotherapy agent, leading to less systemic toxicity, decreasing the prevalence of the disease, and widening the optimal therapeutic dose range of the chemoagent (26). Moreover, as seen in granulosa cell tumor patients, high MIS concentration up to 1000-fold of normal adult female and over 70-fold concentration in normal male infant are noted without adverse effect. Therefore, unlike chemotherapy or radiation therapy, high concentrations of MIS are expected to be non-toxic, when MIS only or MIS combined with chemotherapy is administered (14,25).

It is noteworthy that our study shows, for the first time, the expression pattern of MISRII protein and mRNA in various cervical carcinomas. MISRII protein and mRNA expression and intensity were noted by immunohistochemistry and *in situ* hybridization for each tissue sample. We found complete concordance between protein and mRNA expression. Twelve normal human uterine cervical specimens were positive for the MIS receptor. We also observed that MISRII protein and mRNA were expressed by all cervical cancers (100%). There was little difference in the intensity of expression for MISRII protein and mRNA between normal cervix and cervical cancer. Intensity of expression for MISRII protein and mRNA is nearly the same when non-invasive and invasive cervical squamous cell carcinoma and adenocarcinoma are compared. In Figs. 2 and 3, the cancer cell nests of squamous cell carcinoma diffusely and moderately express MISR II protein and mRNA in the cell membrane, but the cancer cells of adenocarcinoma express somewhat strong intensity. There is tendency for increased receptor protein levels in adenocarcinoma compared to squamous cell carcinoma, but it does not reach statistical significance. There were slight differences in the intensity of expression for...
MISR II protein and mRNA according to the differentiation of cervical cancer. Our data on cervical cancer is comparable with previously published studies on ovarian cancer, however, the results on the frequency of expression of MISRII in our study on cervical cancer are higher (100%) than those previously reported (69-89%) in ovarian cancer (22,33,34). This apparent high rate of receptor positivity in cervical cancers makes the idea plausible that using MIS as a treatment may be offered to the majority of the cases. We emphasize that all human cervical carcinoma tissues express MISRII and intensities of MISRII II protein and mRNA expression show only slight differences according to cell type or differentiation of cervical cancer. Expression of MISRII protein and mRNA is shown in all of normal cervical tissues and cervical cancer tissues in this study, therefore MIS may also be an effective targeted therapy for cervical cancer.

Cervical cancer cell lines used in the investigation include CaSkI, SiHa cell lines, which are HPV 16-positive, HeLa, HeLa S-3, ME-180 cell lines, which are HPV 18-positive, C33A and HT3 cell lines, which are HPV-negative. C33A and HT3 cell lines are Rb protein inactivated caused by exon mutation. Early studies on response assessment in cervical cancer to MIS showed suppressive effect on proliferation through MIS binding to cervical cancer HeLa S-3 cell surface (35). MIS treatment on HT-3 cell line represented 47% of suppressive effect on proliferation through colony inhibition assay (36). MISRII (63 kDa) was expressed in all CaSkI, SiHa, C33A cell lines, and MTT assay after MIS treatment showed that suppressive effect on proliferation was the greatest in C33A with 70-80% suppression, followed by 30-40% in CaSkI, and least in SiHa cell line (26). In our previous study, MISRII was expressed in CaSkI, SiHa, and C33A cell lines, and MTT assay after MIS treatment showed that suppressive effect on proliferation was 28-43, 15-24 and 7-11% in C33A, CaSkI and SiHa cell line, respectively (28). Furthermore, when transfected C33A was treated with inactive MIS, there was no significant decrease in cell line colony, however, when treated with active MIS, C33A colony was markedly decreased (90-95% decrease) (26). Similarly, in treatment with TGF-β (5 µg/ml) on cervical cell line, HPV-negative HT-3 cells showed 70% suppression, HPV 18-positive HeLa showed 25-50% suppression in cell proliferation, whereas, little decrease was observed in HPV 16-positive CaSkI, and SiHa cell lines. No suppression was seen in HPV-negative C33A and HPV 18-positive HeLa S-3, and ME-180 (37,38). Likewise, suppressive effects of MIS on ovarian cancer cell line also differ with HPV infection. The suppression effect of MIS is more significant in OVCAR5, ovarian epithelial cell carcinoma line which is not associated with HPV, compared to human ovarian cancer cell line HOSE 6-3, which is immortalized by HPV tumor proteins E5, and E7 (23). Overall, the anti-proliferative effect of MIS is more effective in HPV-negative cells whereas, it is limited in HPV-positive cells. Also the suppressive effect differs between various HPV types. Since HPV is established as a strong risk factor in cervical carcinoma, and most of cervical cancer patients are associated with HPV infection, the presence of HPV infection has an important role in the treatment effect of MIS and its TGF-β superfamily. In our experiments, MIS-treated human carcinoma cells achieved in vivo showed negative correlation between the MIS exposure time and the viability of cervical cells which is consistent with other experiments. When HPV 16, 18, 31, 33, 52, 58-positive cervical cancer cells were treated with 71 nM MIS for 24 and 48 h, anti-proliferative effect was seen in 15.5 and 28.4%, respectively. This finding corresponds with results (suppression up to 30-40%) in HPV 16-positive CaSkI cell line by Babie et al (26), and those (suppressed up to 15-24%) of our previous study (28). Furthermore, the infected rate and distribution of HPV type in this study were similar with meta-analysis in Korea women (39). HPV goes through various stages to change Rb family related mechanisms, which decrease MIS effect on HPV infected cervical cancer (26,28,40). Thus, MIS will have a more potent anti-proliferative effect in cervical cancer cells which are non-HPV infected or low risk HPV-related.

According to our experiment in cell cycle distribution data after exposure to MIS in cervical cancer cell, S and G2/M phases were shortened and G0/G1 and sub G0/G1 phases were lengthened. Therefore, MIS plays an important role in G0/G1 inducing cellular apoptosis by causing arrest at G1 phase and increasing cells at sub-G0/G1 phase. This finding corresponds with results of our previous study. In C33A cells, the sub-G0/G1 and G1 phases were increased 6 and 3% at 24 h and 10 and 8% at 48 h compared to controls, respectively. In CaSkI cells, the sub-G0/G1 and G1 phase was increased 3 and 5% at 24 h and 6 and 8% at 48 h compared to controls, respectively, but SiHa cells demonstrated minimal response in our previous study (28). Furthermore, annexin-V-FITC staining methods of MIS exposure in cervical carcinoma cells showed negative correlation between cellular survival and MIS exposure time.

In conclusion, MISRII is present in both normal cervix and cervical carcinoma tissues, and MIS shows receptor-mediated anti-proliferative effect on cervical cells in vitro. Thus, these data suggest that MIS may be used as a biological modifier or therapeutic modulator in MISRII-expressed tumors.

References
