

Identification of large-scale characteristic genes of Müllerian inhibiting substance in human ovarian cancer cells

SUK WOO NAM¹, YUN SUNG JO², JUNG WOO EUN¹, JAE YEN SONG², KI SUNG RYU²,
JUNG YOUNG LEE¹, JOON MO LEE², DAVID T. MacLAUGHLIN³ and JANG HEUB KIM²

Departments of ¹Pathology, Microdissection Genomics Research Institute, and ²Obstetrics and Gynecology, College of Medicine, The Catholic University of Korea, Seoul, Korea; ³Pediatric Surgical Research Laboratories, Massachusetts General Hospital and Harvard Medical School, Boston, MA 02114, USA

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Abstract. The purpose of this study was to investigate the large-scale characteristic molecular signature of Müllerian inhibiting substance (MIS) in human ovarian cancer cells through expression genomics. To understand the comprehensive molecular mechanisms by which MIS inhibits ovarian cancer cell growth, we identified the large-scale characteristic molecular changes elicited by MIS in the human ovarian cancer cell line OVCAR-8, using DNA microarray analysis. Combined serial gene expression analysis from 0 to 96 h after MIS treatment of OVCAR-8 cells resulted in 759 genes which showed at least a 2-fold change in overexpression or underexpression compared to non-treatment groups. Of the 759 outlier genes, 498 genes were mapped to known biological cellular processes, and the resultant major pathways included metabolism, signal transduction, cell growth and apoptosis. Among these pathways, 68 genetic elements were dissected as cell cycle-related genes induced by MIS. Although cellular phenotypic changes by MIS were observed after 24 h of treatment, the characteristic large-scale molecular changes were observed from 48 to 96 h of exposure to MIS. This finding may imply that the suppressive role of MIS on ovarian cancer cells could be cumulative in that the metabolic disturbance of MIS is followed by arrest at the G1/S cell cycle checkpoint. We suggest 759 outlier genes comprise the characteristic molecular signature of MIS, which may be responsible for the suppressive effect on OVCAR-8 cells. Although the precise biological mechanisms underlying these outlier genes should be validated, the genetic elements described herein provide promising therapeutic interventions for ovarian cancer.

Introduction

Müllerian inhibiting substance (MIS), also known as anti-Müllerian hormone, is a member of the transforming growth factor- β (TGF- β) superfamily of growth and differentiation response modifiers. Members of this family play an important role in mesenchymal-epithelial interaction, cell growth and proliferation, extracellular matrix production and tissue remodeling (1). In contrast to other members of this family, which exert a broad range of functions in multiple tissues, the principal function of MIS is to induce regression of the Müllerian ducts during sex differentiation in mammalian, avian and reptilian embryos (2,3). The Müllerian duct, which forms from the coelomic epithelium, develops into the fallopian tubes, uterus, cervix, proximal vagina and surface epithelium of the ovary in females. MIS is produced in fetal and postnatal Sertoli cells in the testes at high levels which persist until puberty, after which, the concentration of MIS drops to basal levels, where it persists throughout life. In females, MIS is undetectable in the fetal and early postnatal ovary or in serum but becomes detectable in the granulosa cells of the developing ovarian follicles and in the serum of the adolescent and adult, and is never detected after menopause in healthy women (4-7). Mutations in MIS or the MIS type II (MISRII) receptor in humans cause persistent Müllerian duct syndrome, in which males retain Müllerian ducts in the presence of testes and the Y chromosome (8,9). Transgenic female mice that overexpress MIS, demonstrate complete ablation of the ovary, along with undeveloped uterus and oviducts. In male mice overexpression of MIS leads to feminized genitalia, undescended testes, and poorly differentiated Wolffian ducts (10). In addition to its significant role in sexual development, MIS as a hormone probably plays an important role in the adult gonads of either sex (11); in fact recent evidence suggests other non-Müllerian sites of synthesis and responsiveness including motor neurons (12).

As a member of the TGF- β family, MIS is thought to exert its effects through two distinct single membrane spanning serine-threonine receptors, types I and II. The binding of MIS to the type II receptor (MISRII) leads to heterodimerization with an MISRI, initiating a signaling cascade. There has been

Correspondence to: Dr Jang Heub Kim, Department of Obstetrics and Gynecology, College of Medicine, The Catholic University of Korea, #62 Yeouido-dong, Yeongdeungpo-gu, Seoul 150-713, Korea
E-mail: janghkim@catholic.ac.kr

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only one gene or the MIS type II receptor found in all animal species examined thus far. There are, however, several candidates for the MIS type I receptor (13). Since the type I receptors are nearly ubiquitously expressed, which type I receptor partners the type II receptor could be tissue and/or species specific. It is interesting to note that loss-of-function mutations created in mouse MISRII produce Leydig cell tumors in males and early follicular depletion in females (11,14). Thus, lack of proper MIS signaling seems to predispose mice to uncontrolled division of MIS-responsive cells. This also implies that MIS may play a similar role in humans.

An increasing number of reports suggest that MIS is a multi-functional hormone in addition to its major role in sexual development in the embryo. Since most ovarian tumors originate from the surface epithelium of the ovary, which is of Mullerian duct origin (15,16), the effect of MIS on the growth of ovarian cancer cells has been an area of intense study. Many previous studies have demonstrated that MIS inhibits the growth of Müllerian duct-derived tumors (17-19) and several cancer cell lines or primary cells *in vitro* (20-23). It was also demonstrated that OVCAR-8, an epithelial ovarian cancer cell line, expresses MISRII and responds to MIS by growth inhibition mediated through a retinoblastoma protein (p-RB)-independent mechanism involving the up-regulation of p16 at the protein level. This suggests that MIS requires p16 for growth inhibition in ovarian cancer cells, which in turn arrests cell growth at the G1/S phase and subsequently augments apoptosis (19). Although several reports have described the anti-neoplastic properties of MIS, little insight into the molecular nature of anti-tumor growth or apoptosis has been gained. Recently, DNA microarray technology has enabled the genome-wide analysis of gene transcript levels, and, as such, has yielded great insight into the molecular nature of cancer. A number of studies have demonstrated the usefulness of this technique for identifying novel cancer markers and for classifying human cancers at the molecular level for improved diagnosis and prediction of outcome (24-27).

Thus, to better understand the molecular mechanisms by which MIS inhibits the growth of ovarian cancer cells, we analyzed global transcript levels in the series context of different MIS treatment time points in OVCAR-8 cells and identified characteristic genes which may reflect the molecular signature for the anti-tumor effect of MIS by utilizing large-scale gene expression analysis of DNA microarray.

Materials and methods

Recombinant human MIS. Recombinant human MIS was purified, and its biological activity was confirmed in the Pediatric Surgical Research Laboratories at the Massachusetts General Hospital from serum-free and serum-containing conditioned media as described earlier (28).

Cells and cell culture. The human ovarian cancer cell line OVCAR-8 was grown in Dulbecco's modified Eagle's medium and 10% female fetal bovine serum, 1% penicillin/streptomycin and 1% L-glutamine for no more than 5 passages, and subcultures were initiated at 80% confluency. The cultures

were maintained in a humidified atmosphere of 5% CO₂ at 37°C.

Microarray fabrication. The 60-mers of the Human Oligolibrary™ representing 18,664 LEADS™ clusters plus 197 controls (GAPDH) were purchased from Compugen/Sigma-Genosys and spotted onto a glass slide at the microarray core facility of the Microdissection Genomics Research Institute at the College of Medicine, The Catholic University of Korea as previously reported (29,30).

Large-scale analysis of gene expression profiling by using oligonucleotide arrays. In brief, total cellular RNA was extracted from the vehicle control and MIS-treated cells using a TRIzol® reagent (Gibco, Grand Island, NY) and quantified using NanoDrop (NanoDrop Technologies, Wilmington, DE). In addition, RNA quality was checked by Bioanalyzer 2100 (Agilent Technologies, Santa Clara, CA), and intact RNA samples (260/280 ratio >1.8) were used for hybridization. Human universal reference RNA (Stratagene, La Jolla, CA) was used as a reference RNA. From each total RNA sample, 20 µg was primed with oligo-dT and labeled with Cy3-dUTP or Cy5-dUTP (New England Nuclear, Wilmington, MA) by reverse transcription. Labeled cDNA targets were concentrated to ~15 µl using Microcon-YM30 (Amicon, Bedford, MA). Hybridization was carried out at 42°C for 16 h using the MAUI Hybridization System (BioMicro Systems, Salt Lake City, UT). After hybridization, the microarray was washed twice with buffers containing 2X standard sodium citrate (SSC) and 0.1% SDS for 2 min, 1X SSC and 0.1% SDS for 3 min, 0.2X SSC for 3 min, 0.05X SSC for 2 min, and finally rinsed with distilled water for 2 min. This protocol was repeated twice in independent experiments on the OVCAR-8 cells.

Scanning and data analysis. The arrays with hybridized targets were scanned using an Axon scanner, and the images were analyzed using GenePix® Pro 4.1 software (Molecular Devices, Sunnyvale, CA). Spots of poor quality identified by visual inspection were removed from further analysis. The resulting data collected from each array were submitted to the BioArray Software Environment (BASE) database at the microarray core facility at the Department of Pathology, College of Medicine, The Catholic University of Korea, Seoul, (<http://genomics.catholic.ac.kr/>). Data were normalized using the method of Linear Models for Microarray Data (LIMMA) and R-package for Statistics for Microarray Analysis (SMA). Spots <50 µm were eliminated for analysis unless otherwise specified. Pearson's correlation coefficient was calculated using the S-Plus program. Cluster and TreeView programs were used for visualization of data (27,29,31). Pathway analysis was performed with ArrayXPath (<http://www.snubi.org/software/ArrayXPath/>).

Methylthiazolotetrazolium (MTT) assay. Cells (3,000/well) were seeded in 96-well plates in OVCAR-8 media. After 24 h, the cells were exposed to vehicle control or different concentrations (5, 10, or 15 µg/ml) of MIS for 24 and 48 h. 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%

SPANDIDOS was added to each well. Cells were incubated for 30 min at the end of which time 200 μ l DMSO (Sigma, St. Louis, MO) 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).

Cell cycle analysis. OVCAR-8 cells were exposed to MIS or vehicle control buffer for 24 h, and the cells were collected by trypsinization. The cells were fixed with 100% methanol and stored for 30 min at 20°C and washed with PBS. Following centrifugation, the cells were resuspended in 1 ml DNA staining solution (20 μ g/ml propidium iodide, 200 μ g/ml DNase-free RNase) and incubated in the dark at 37°C for 30 min. The cells were analyzed on a FACS Vantage SE Flow Cytometer (Becton Dickinson, San Jose, CA). The forward scatter and red fluorescence >600 nm were measured, and the results were analyzed using Cell Quest™ software and Modfit LT 3.0 program (Verity Software House, Topsham, ME).

Annexin V analysis. MIS-treated cells were stained with Annexin V and propidium iodide (PI) using the Annexin V-FITC Apoptosis Detection Kit I (BD Biosciences, San Diego, CA) according to the manufacturer's protocol. Briefly, following drug treatment, 1×10^5 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 $MgCl_2$, and 2 mM calcium chloride]. Subsequently, 5 μ l of Annexin V-FITC and PI were added to the cells which were then incubated for 15 min at room temperature in the dark. After this incubation, 400 μ l of binding buffer was added, and cells were analyzed using a FACS Vantage SE Flow Cytometer. Data analyses were conducted using Cell Quest software.

Results

MIS inhibits the growth of OVCAR-8 ovarian cancer cells and augments apoptosis. Before initiating experiments on MIS in ovarian cancer cells, we confirmed that the OVCAR-8 cells cultured in our laboratory expressed MISRII on their cell surface by immunohistochemical staining and by Western blot analysis using an anti-MISRII antibody (data not shown). Growth inhibition of OVCAR-8 cells by MIS was evident at both 24 and 48 h. As shown in Fig. 1, the cell viability was decreased to 65-85% of the control, and this inhibitory effect was dose-dependent. In order to determine whether this anti-growth activity was related to cell cycle regulation, the cell cycle phase distribution after MIS treatment was analyzed by flow cytometry (Fig. 2A). When cells were incubated with 10 μ g/ml of MIS, the G0/G1 phase was significantly increased 11% at 24 h and 13% at 48 h compared to controls. This finding could be partially explained by the fact that MIS may induce G1/S arrest at an early time point subsequently leading to cell death. In fact, induction of cell death by MIS in OVCAR-8 cells was detected but only at 48 h by FACS analysis of cells stained with Annexin V-FITC and PI (Fig. 2B).

Identification of large-scale molecular changes and characteristic molecular signature of MIS. To identify a

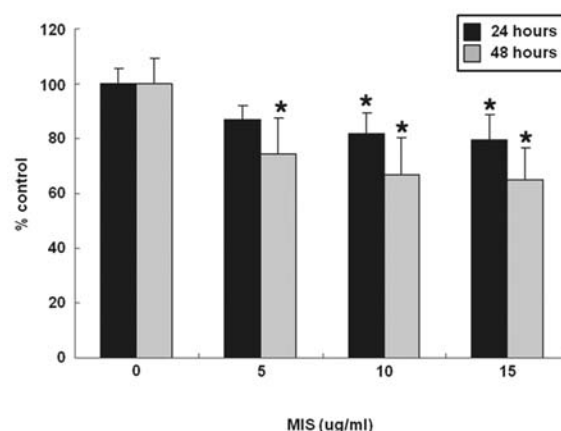


Figure 1. Inhibition of cell growth by MIS in OVCAR-8 ovarian cancer cells as assessed by the cell viability test using the MTT assay. OVCAR-8 cells were treated with MIS at each indicated concentration for 24 and 48 h. After incubation with or without MIS, cells were stained with MTT, and the absorbance was read at 550 nm. Results are presented as a percentage of the control which was calculated using the equation: (mean absorbance of treated cells/mean absorbance of control cells) \times 100. Data are expressed as the mean \pm standard deviation (SD) from four independent experiments. * $P < 0.05$ as compared to corresponding control cells.

molecular signature which is specifically induced by MIS, serial gene expression analysis using large-scale comprehensive DNA microarray experiments was conducted. Since MIS showed sustained growth inhibitory activity for at least 48 h, OVCAR-8 cells were continuously treated with MIS for up to 96 h, and the RNAs were harvested for microarray analysis at multiple time points as early as 6 h. As shown in Fig. 3A, unsupervised hierarchical clustering of 9,073 genetic elements, which passed the minimum selection, and filtering criteria (see Materials and methods) resulted in two separate subclusters on the dendrogram; a short-term exposure group (0 to 24 h of MIS treatment) and a long-term exposure group (48 to 96 h) group. To identify genes that were continuously changing due to MIS action, mean values of the entire pool of genetic elements in non-treated cells (0 h of MIS) was subtracted from that of the long-term exposure group. The 759 outlier genes from this analysis were then re-examined in the expression data of the short-term exposure group and then shown as a heat map (Fig. 3B). Of 759 outlier genes, 416 genes were up-regulated and 343 genes were under-expressed in the long-term MIS exposure group compared to the controls. As shown in this figure, most of the outlier genes showed gradual and continuous changes from early MIS exposure with a few exceptions (Fig. 3B).

Molecular dissection of cell growth regulation by MIS through pathway mining analyses. By an in depth examination of the genes identified as candidate MIS targets, it was possible to locate downstream signaling pathways under the control of long-term MIS treatment of OVCAR-8 cells. This process was accomplished by subjecting the 759 MIS-regulated genes to the pathway mining tool in the PANTHER Classification System (<http://www.pantherdb.org/>). Of the 759 outlier genes, 498 genes were mapped into known pathways, and the resultant biological processes were

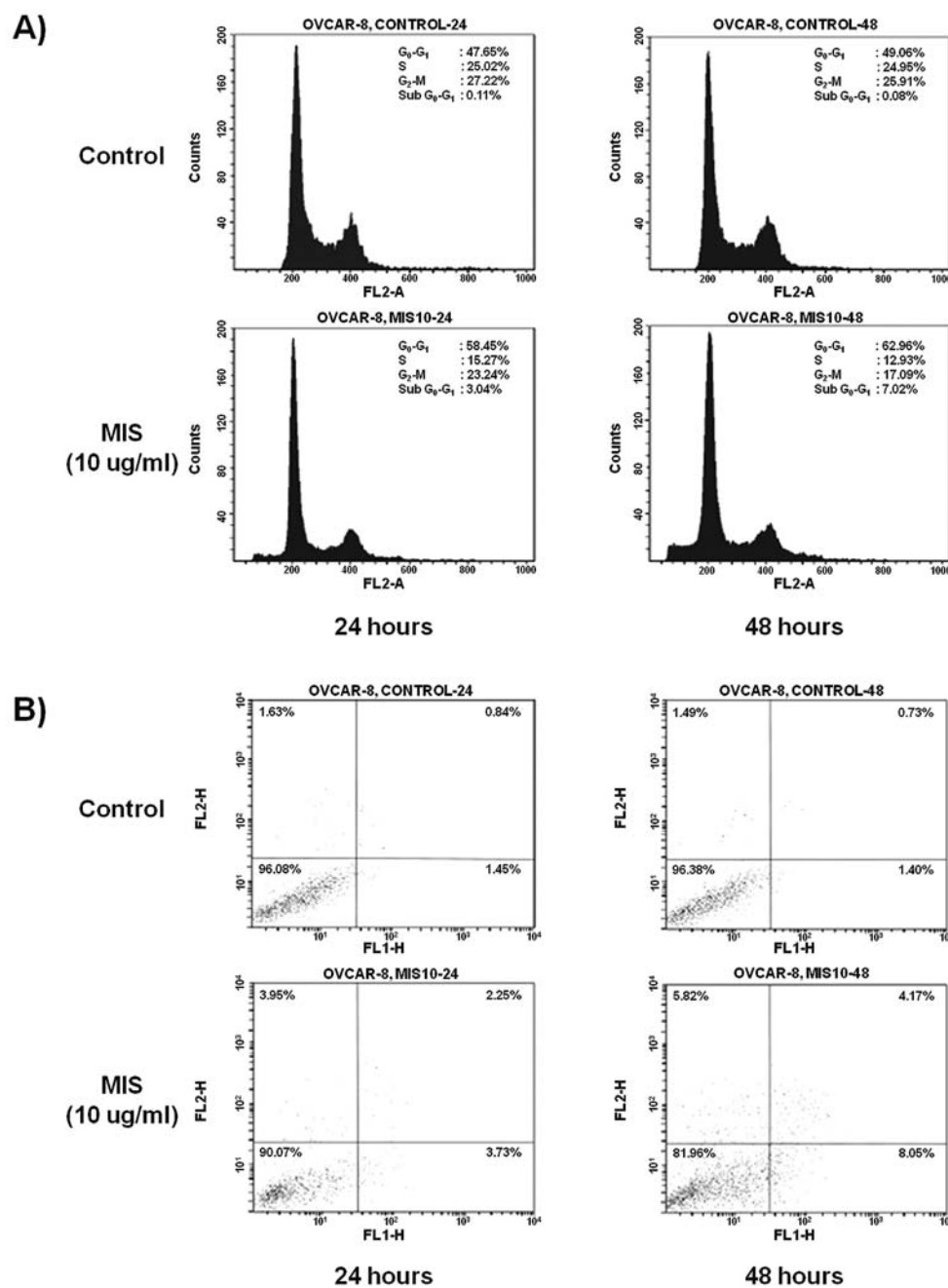


Figure 2. Cell cycle arrest and induction of apoptosis by MIS in OVACAR-8 ovarian cancer cells. Cells were treated with MIS (10 μ g/ml) for 24 and 48 h. (A) FACS analysis of the cell cycle. Cells were harvested with ice-cold PBS/EDTA, fixed in 95% ethanol after incubation with MIS and stained with propidium iodide in an RNase solution with sodium citrate buffer for 30 min after which they were immediately analyzed. (B) Quantification of cells undergoing apoptosis. OVACAR-8 cells were treated with 10 μ g/ml of MIS for 24 and 48 h, stained with Annexin V-FITC and PI, and analyzed by FACS. Quadrant rectangular dot grams from a representative of 3 independent experiments are shown.

categorized (Table I). As listed in Table I, the major processes in which the MIS characteristic genes are involved include signal transduction, nucleic acid, and protein metabolism and cell growth. In order to narrow these down to cell growth regulatory mechanisms which are mediated by MIS, we retrieved all genetic elements relating to cell growth processing from the 759 outlier genes (Fig. 4A). A total of 68 genetic elements were found as cell growth-related genes. Among these, 41 genes were underexpressed and 27 genes were overexpressed at long-term exposure time points. Furthermore, we dissected genetic elements, which are

involved in the G1/S cell cycle checkpoint (Fig. 4B). The transcript level of p16^{INK4A} did not change; however, the CDKs, particularly CDK2, 4 and 6 seemed to be down-regulated by MIS as were E2F2, 3 and 4, binding partner of DP1. The downstream molecules such as MCMs, PCNA, RANGAP seemed to be suppressed by the down-regulation of the E2F/DP1 complex (Fig. 4B) whereas, HDAC5 and 6 exhibited a very strongly increased expression level from 48 h of MIS treatment.

In addition, we also dissected the cell death-related molecular signature of MIS from the 759 outlier genes



SPANDIDOS PUBLICATIONS summary of the biological processes that are part of characteristic molecular signature of MIS in OVCAR-8 cells.

Biological process	No. of genes
Signal transduction	108
Nucleoside, nucleotide and nucleic acid metabolism	90
Protein metabolism and modification	78
Mesoderm and ectoderm development	42
Cell proliferation and differentiation	38
Cell cycle	23
Cell adhesion	23
Oncogenesis	19
Apoptosis	18

There were 759 genes which showed expression more than 2-fold above or below those of the untreated controls. Of the mapped genes, 498 were grouped by biological process.

(Fig. 5). Although this molecular signature should be validated and detailed regulatory cascade provided in apoptosis signal, it is obvious that caspase 1, TNF receptor and MAP3K13, well known death-related factors, were overexpressed by MIS.

Discussion

In previous studies, it has been demonstrated that recombinant human MIS inhibits the growth of ovarian cancer cells and various primary tumors *in vitro* and *in vivo* (19,32-34). These findings are significant since the MIS type II receptor has been detected in an increasing number of human gynecological tumors (35). In addition, it also suggests that MIS inhibits growth inhibition through the activation of p16, a negative cell cycle regulator (19), and consequently disrupts cell cycle progression and leads to apoptosis. In order to confirm the anti-neoplastic activity of MIS, we assessed the growth inhibition and apoptosis of MIS on OVCAR-8 cells, which was previously demonstrated to express MISRII and to respond to MIS by growth inhibition. Our current results are in concordance with previous reports and confirmed that MIS induces anti-growth activity on OVACR-8 cells. It appears that, when compared to the cell cycle arrest data presented here, apoptosis seems to play a minor role in the effects of MIS on these cancer cells.

Accumulating evidence suggests that MIS augments the protein levels of the p16^{INK4A}, a cyclin-dependent kinase (CDK) inhibitor and consequently down-regulates the CDK, particularly the cyclin D/CDK4 or 6 complex and inhibits G1/S cell cycle transition. Although similar cell cycle regulation by MIS has been investigated in other cancers (20,36), the systemic molecular circuit of this cell cycle regulation in ovarian cancer still remains unknown. Furthermore, as several lines of evidence suggest that MIS is a multi-functional hormone, the comprehensive molecular signature which is mediated by MIS in cells needs to be elucidated.

In this study, we present the results of an initial microarray study designed to further expand upon the existing

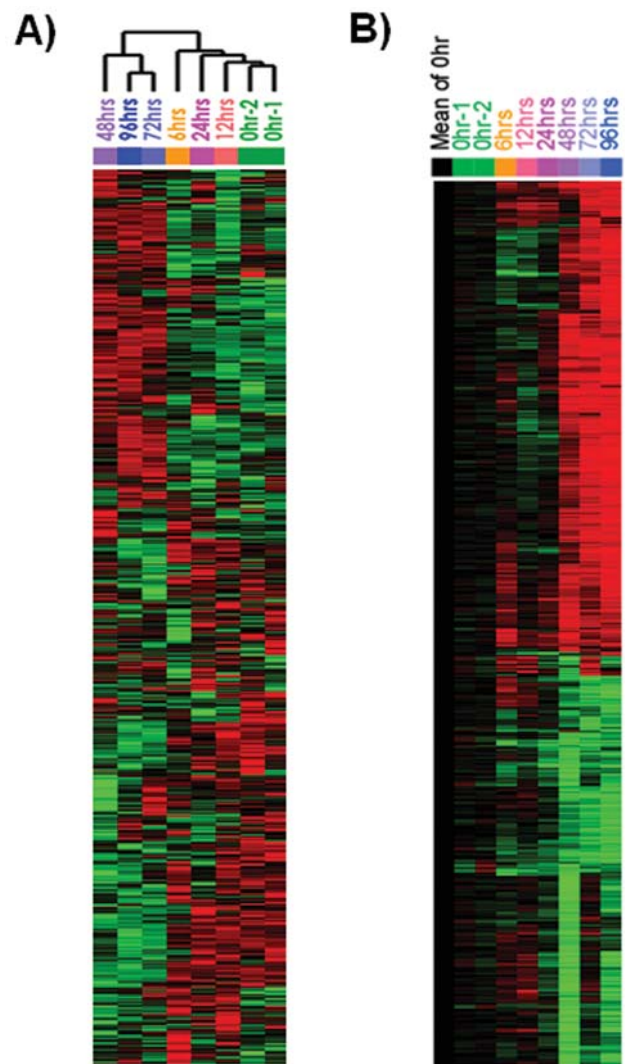


Figure 3. Differential gene expression profiling and identification of large-scale molecular changes in MIS-treated OVCAR-8 cells. (A) Unsupervised hierarchical clustering of 9,073 genetic elements with minimum selection and filtering criteria (see Material and methods) resulted in two separate subclusters on dendrogram, short-term exposure group (0 to 24 h of MIS-treatment) and long-term exposure (48 to 96 h) group. (B) A significant subset of outlier genes were further narrowed by filtering genes showing expression changes due to MIS treatment of cells. Briefly, the mathematical comparisons of genes between 0 h and the 48- to 96-h time points were performed by selection of genes showing at least 2-fold changes induced by MIS treatment compared to non-treated control (0 h). This analysis resulted in the identification of 759 outlier genes that were significantly differentially expressed in MIS-treated cells. These elements were then visualized as a heat map where the red color indicates that expression levels of genetic elements are higher than the mean value of non-treatment, and green color indicates that expression levels of genetic elements are lower than the mean value of non-treatment.

hypotheses related to the MIS mechanism of action and to discover the role played by MIS in mediating other signaling, metabolic and cell cycle pathways in cancer cells. Although biological validation for the candidate MIS-regulated pathways reported here should be followed by Western blot analysis and real-time PCR, the genetic elements that we suggest as cell growth-related genes may further our understanding of the molecular insights into the cross-talk of intracellular signaling on cell growth regulations by MIS.

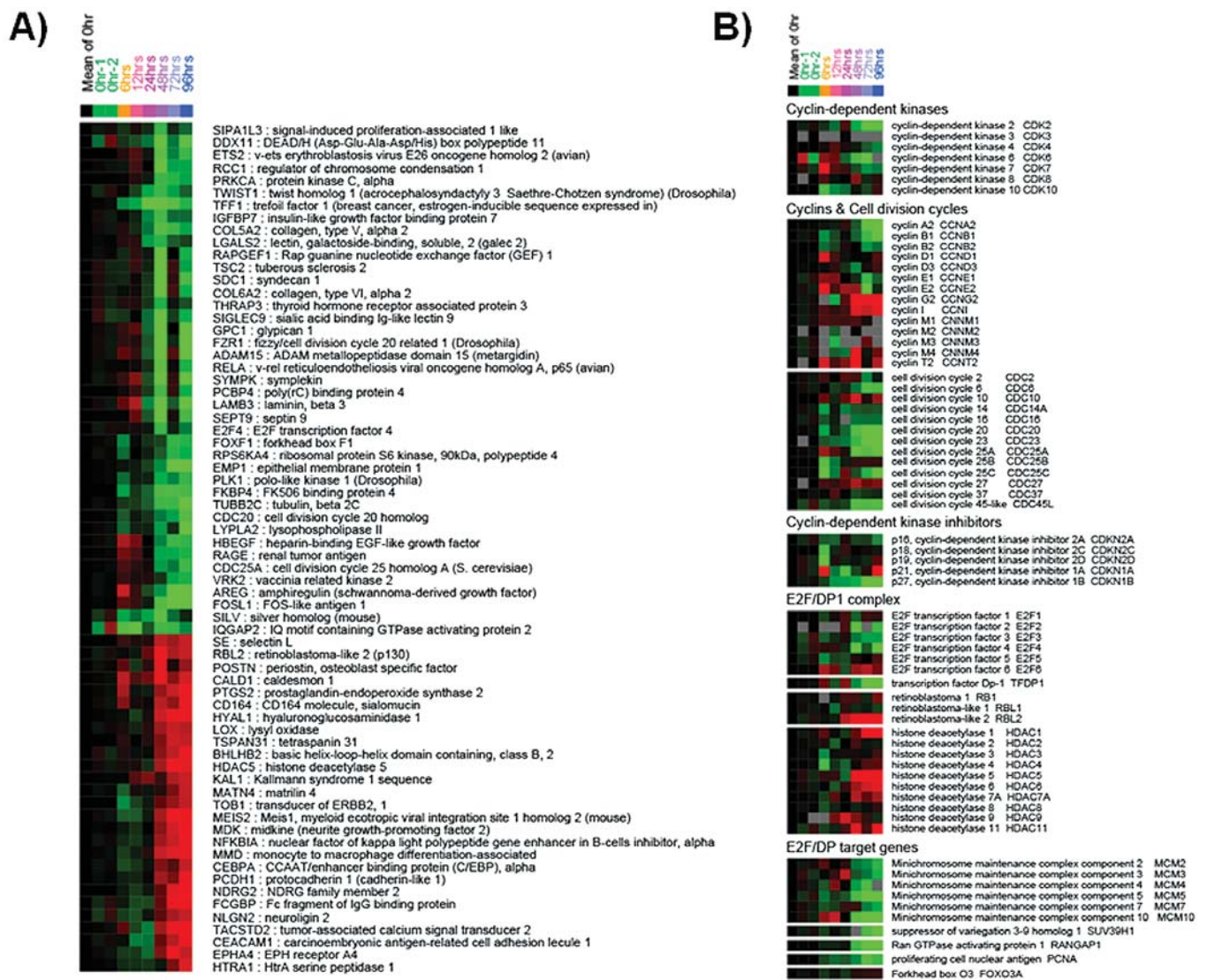


Figure 4. Selective expression profile of cell cycle-related genetic elements retrieved from 759 outliers. (A) A cell growth-associated molecular signature. Of 759 outlier genes, 68 genes were mapped into cell growth-related pathways such as cell growth and differentiation and cell cycle (Table I) through the PANTHER Classification System (<http://www.pantherdb.org/>). The expression patterns of genes were then visualized as a heat map. (B) Selective expression profile of genetic elements of G1/S transition of the cell cycle. The red color indicates that expression levels of genetic elements are higher than the mean value of non-treatment, and the green color indicates that expression levels of genetic elements are lower than mean value of non-treatment.

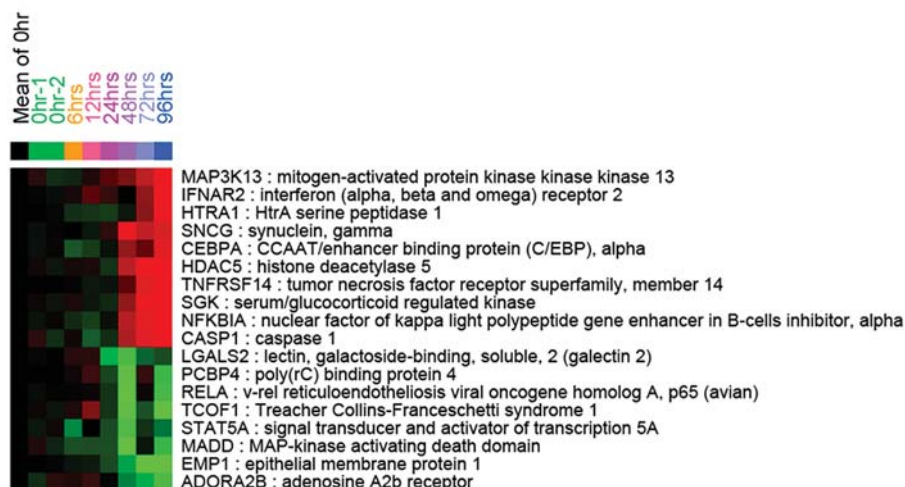


Figure 5. Selective expression profile of cell death-related genetic elements retrieved from 759 outliers. Of 759 outlier genes, 18 genes were mapped into apoptosis pathway (Table I) using the PANTHER Classification System (<http://www.pantherdb.org/>). The expression pattern of genes was then visualized as a heat map with the color indicators as described for the previous two figures.



SPANDIDOS PUBLICATIONS G1/S cell cycle regulation by MIS, we found that not induce most of the CDK inhibitors, but p19^{INK4D}

and p21^{Cip1} were slightly induced at 96 h. It is interesting to know that the transcript level of p16^{INK4A} was not changed by MIS in a previous study (19) and the same result was obtained from our analysis (Fig. 4B). Of the CDKs, particularly CDK2, 4 and 6 seemed to be down-regulated by MIS. The E2F/DP1 complex is an important transcription factor regulated by the G1/S checkpoint. Our results indicated that E2F2, 3 and 4 were down-regulated by MIS. Consequently, many of the downstream target molecules of the E2F/DP1 transcription factor such as MCMs, PCNA, RANGAP seemed to be suppressed by this down-regulation of the E2F/DP1 complex (Fig. 4B). Histone deacetylases (HDACs), which are known to play a crucial role in chromatin remodeling, are also important co-suppressors of the E2F/DP1 complex such as Rb-family molecules. In our results, HDAC5 and 6 exhibited a very strongly increased expression level from 48 h of MIS treatment. This also indicates that the enhanced expression level of HDAC5 or 6 is cooperatively suppressive on the E2F/DP1 transcription factor. In summary, our results suggest that the regulatory mechanism of MIS on the G1/S cell cycle checkpoint had a suppressive effect on CDKs and CDCs and the E2F/DP1 complex with transcriptional enhancement of certain HDAC family of E2F/DP1 co-suppressors, but did not seem to elevate transcript level of CDK inhibitors, a cell cycle negative regulator. The continuous changes in gene expression along with sequential time points of exposure to MIS suggest that gene expression changes may occur with early exposure to MIS, but transcriptomic expression characterized by MIS was induced after at least 48 h of long-term exposure. This indicated that the comprehensive molecular changes of the long-term exposure group were more unique than the short-term group as this short-term group clustered with non-treatment control.

These novel findings clarify, at a molecular level, the manner in which MIS regulates the proliferation of cancer cells. It remains to be determined whether these pathways are MIS targets in normal fetal or adult tissues, but these data may facilitate the design of new drug targets for MIS in combination with other drugs for the treatment of ovarian cancer.

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