Bisdemethoxycurcumin (BDMC) Alters Gene Expressionassociated Cell Cycle, Cell Migration and Invasion and Tumor Progression in Human Lung Cancer NCI-H460 Cells

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Abstract. Background/Aim: Lung cancer is one of the most common malignancies and a predominant cause of cancerrelated death. It can metastasize in almost all organs, and currently, while new cases are increasing, treatment is still insufficient. Bisdemethoxycurcumin (BDMC), one of the components of turmeric, has been known to possess biological activities. However, the effects of BDMC on the genetic level remain unclear. Materials and Methods: Human lung cancer NCI-H460 cells were treated with 35 µM BDMC for 24 h and cells were harvested for total RNA extraction. The purified RNA was used for cDNA synthesis, labeling, microarray hybridization, and flour-labeled cDNA on-chip hybridization. The expression Console software (Affymetrix) with default RNA parameters was used to detect and

quantitate concentrations of fluorescent molecules. The key genes involved and their possible interaction pathways were analyzed by the GeneGo software. Results: Seven genes, such as CCNE2 (cyclin E), associated with cell cycle, were over 4-fold overexpressed, 22 genes, such as ERCC6L (excision repair cross-complementing rodent repair deficiency, complementation group 6-like) associated with DNA damage and repair, were from 3- to 4-fold overexpressed and 266, such as cell division cycle, S-phase associated kinase and associated with cell death, genes were from 2- to 3-fold overexpressed. Conclusion: BDMC induced changes in gene expression that may reveal cytotoxic information on the genetic level while presenting novel biomarkers or targets for treatment of human lung cancer in the future.

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Key Words: Bisdemethoxycurcumin, cDNA, human lung cancer NCI-H460 cells.

Lung cancer is one of the major causes of death due to cancer worldwide and early metastatic dissemination and resistance to therapy are causal factors giving a median survival of less than 12 months (1). Recently, in lung cancer patients, new improvements have focused on chemotherapy and molecular-targeted therapy, however, the outcome of patients remains unsatisfactory (2). Non small-cell lung cancer (NSCLC) accounts for 85% of lung cancer cases (3, 4) and is also the main cause of cancer-related death worldwide (5) and the overall 5-year survival rate of NSCLC patients remains lower than 15% (6). Thus, new therapeutic strategies for NSCLC are urgently needed. Many current investigations are focusing on anticancer drugs

0258-851X/2015 \$2.00+.40

originating from natural products. Furthermore, in psychiatric medicine, herbals and phytochemicals have been recognized to be of great interest, as complementary and/or alternative therapies (7).

Turmeric, a ground rhizome of Curcuma longa, is widely used in Asian traditional medicine for wound healing, inflammatory conditions, and blood purification (8, 9). In traditional Chinese medicine, turmeric has a long history of treating diseases associated with abdominal pain (10). Bisdemethoxycurcumin (BDMC), one of the components of turmeric, has been shown to possess anti-inflammatory and anti-proliferative activities (11), anti-metastasis potency via the differentially down-regulation of ECM degradation enzymes (12) and to induce rapid DNA double-strand breaks in the human colon cancer HCT116 cells (13). BDMC inhibited cell invasion and motility and modulated MMP-3 expression in human invasive breast carcinoma cells (14). BDMC decreased inducible nitric oxide synthase (iNOS) expression and nitric oxide (NO) production (15) and inhibited the Wnt/beta-catenin pathway (16). Furthermore, the dual activity of BDMC on topoisomerases-IIa (TOP2A) has been used in novel therapeutic strategies in order to induce apoptosis in cancer cells (17).

Existing studies included patients treated with targetedagents based on specific molecular alterations of neoplastic cells (18, 19). Thus, molecular alterations in cancer cells are called to attention for aiding the development of specific targeted drugs. Moreover, in clinical practice, predictive biomarkers have been assessed and increased the availability of anticancer drugs corresponding to specific molecular alterations (20). In the past decade, the identification of specific predictive and/or prognostic molecular alterations for NSCLC and colorectal carcinoma (CRC) have been the center of attention (21, 22). Currently, the development of targeted-therapeutics for NSCLC treatment aim at defining genetic abnormalities in NSCLC such as mutations in EGFR or a fusion of the EML4, while the ALK gene has been recognized as a target for first-line therapies in NSCLC (23-25).

Although BDMC has been shown to induce cell death in NSCLC cells, the exact genes affected by BDMC remain unknown. Thus, we investigated altered gene expression in NCI-H460 cells after exposure of cells to BDMC and results indicated that affected genes are involved in apoptosis pathways.

Materials and Methods

Chemicals and reagents. DMSO was obtained from Sigma Chemical Co. (St. Louis, MO, USA). Culture medium RPMI-1640, fetal bovine serum (FBS), 1% L-glutamine, 100 U/ml penicillin and 100 µg/ml streptomycin were obtained from Gibco BRL (Grand Island, NY, USA). BDMC was dissolved in DMSO and stored at -20°C before use in experiments.

Human lung cancer cells. NCI-H460 human non-small cell lung cancer cells were obtained from the Food Industry Research and Development Institute (Hsinchu, Taiwan). The cells were cultured at 37°C in a humidified atmosphere of 95% air and 5% CO₂ air in RPMI-1640 medium supplemented with 10% (FBS), 1% L-glutamine and 1% penicillin/streptomycin (Invitrogen, Carlsbad, CA, USA). Cultured NCI-H460 cells were split every 4 days for maintaining exponential growth and were harvested with 0.025% trypsin and 0.52 mM EDTA in phosphate buffer saline (PBS), plated at required cell numbers and allowed to adhere for 24 h before BDMC treatment.

cDNA Microarray assay for gene expression in NCI-H460 cells after exposure to BDMC. NCI-H460 cells at a density of 5×10⁵ cells/ml were maintained on 12-well plates with RPMI-1640 medium for 24 h and then incubated with or without 35 µM of BDMC for 24 h. At the end of incubation, cells were collected from each treatment and total RNA was extracted by using the Qiagen RNeasy Mini Kit (Qiagen, Inc, Valencia, CA, USA), as described previously (26) and individually quantitated and used for cDNA synthesis, labeling and microarray hybridization, followed by flour-labeled cDNA hybridizing their complements on the chip (Affymetrix GeneChip Human Gene 1.0 ST array, Affymetrix, Santa Clara, CA, USA) (26). The resulting localized concentrations of fluorescent molecules on the chip were detected and quantified (Asia BioInnovations Corporation, Taipei, Taiwan) and data were further analyzed by the Expression Console software (Affymetrix) with default RMA parameters (26, 27). Data are representative of three separate assays.

Statistical analysis. All results were presented as mean±SD of three independent experiments. Significant differences between BDMC-treated and -untreated groups were considered if at least a 2-fold change was recorded. +, Up-regulation; –, down-regulation.

Results

BDMC induced both up-regulation and down-regulation of gene expression in NCI-H460 cells. After NCI-H460 cells were treated with or without 35 µM of BDMC for 48 h, they were extracted for total RNA from each treatment. Isolated total RNA was quantified, followed by cDNA microarray analysis and results are shown in Tables I and II. Table I shows that 7 genes were over 4-fold up-regulated, such as CCNE2 (cyclin E), 22 genes were from over 3- to 4-fold such as ERCC6L (excision repair cross-complementing rodent repair deficiency, complementation group 6-like), and 266 genes were over from 2- to 3-fold up-regulated, such as cell division cycle 6 homolog (CDC6), cell division cycle associated 5 (CDCA5), cell division cycle 25 homolog A (CDC25A) and cell division cycle associated 7-like (CDCA7L) associated with cell division, S-phase kinaseassociated protein 2, E3 ubiquitin protein ligase (SKP2), Cdk5 and Abl enzyme substrate 2 (CABLES2), Cdk5 and Abl enzyme substrate 1 (CABLES1) and cyclin E1 (CCNE1).

Table II indicates that 41 genes were over 4-fold down-regulated such as *DDIT3* associated with DNA damage; 57 genes were down-regulated from 3- to 4 -fold such as *DDIT4* associated with DNA damage, *CCPG1* associated with cell

Table I. H460 BDMC-treated vs. H460-BDMC-control.

Fold chan	ige Gene symbol	mRNA Description
7.03	CYP1A1	Cytochrome P450, family 1, subfamily A, polypeptide 1
5.67	FAM111B	Family with sequence similarity 111, member B
5.53	TIPARP	TCDD-inducible poly(ADP-ribose) polymerase
1.83	CCNE2	Cyclin E2
4.35	DSCC1	Defective in sister chromatid cohesion 1 homolog (S. cerevisiae)
1.04	NAP1L2	Nucleosome assembly protein 1-like 2
1.02	SNAI2	Snail homolog 2 (Drosophila)
3.80	CLSPN	Claspin
3.77	ZNF483	Zinc finger protein 483
3.70	ORC1	Origin recognition complex, subunit 1
3.66	FIGNL1	Fidgetin-like 1
3.65	LPCAT2	Lysophosphatidylcholine acyltransferase 2
3.58	ERCC6L	Excision repair cross-complementing rodent repair deficiency, complementation group 6-like
3.57	VDR	Vitamin D (1,25- dihydroxyvitamin D3) receptor
3.52	ID3	Inhibitor of DNA binding 3, dominant negative helix-loop-helix protein
3.45	IFIT2	Interferon-induced protein with tetratricopeptide repeats 2
3.44	CLSPN	Claspin
3.42	METTL7B	Methyltransferase like 7B
3.30	HECW1	HECT, C2 and WW domain containing E3 ubiquitin protein ligase 1
3.18	ID1	Inhibitor of DNA binding 1, dominant negative helix-loop-helix protein
3.17	SKA3	Spindle and kinetochore associated complex subunit 3
3.11	MBOAT2	Membrane bound O-acyltransferase domain containing 2
3.06	WNK3	WNK lysine deficient protein kinase 3
3.05	DLX2	Distal-less homeobox 2
3.04	NRG4	Neuregulin 4
3.01	TLR3	Toll-like receptor 3
3.00	FEN1	Flap structure-specific endonuclease 1
3.00	CRYZ	Crystallin, zeta (quinone reductase)
3.00	MCM10	Minichromosome maintenance complex component 10
2.99	DHFR	Dihydrofolate reductase
2.99	CROT	Carnitine O-octanoyltransferase
2.99	GINS3	GINS complex subunit 3 (Psf3 homolog)
2.98	CDC6	Cell division cycle 6 homolog (S. cerevisiae)
2.95	RALGPS2	Ral GEF with PH domain and SH3 binding motif 2
2.95	DHFR	Dihydrofolate reductase
2.93	KLHL13	Kelch-like 13 (Drosophila)
2.93	C10orf122	Chromosome 10 open reading frame 122
2.91	HSPA1A	Heat shock 70kDa protein 1A; heat shock 70kDa protein 1B
2.91	HSPA1B	Heat shock 70kDa protein 1B; heat shock 70kDa protein 1A
2.90	SLC1A1	Solute carrier family 1 (neuronal/epithelial high affinity glutamate transporter, system Xag), member 1
2.89	TDO2	Tryptophan 2,3-dioxygenase
2.89	TMEM14A	Transmembrane protein 14A
2.89	HSPA1A	Heat shock 70kDa protein 1A; heat shock 70kDa protein 1B
2.87	EML1	Echinoderm microtubule associated protein like 1
2.86	ASF1B	ASF1 anti-silencing function 1 homolog B (S. cerevisiae)
2.85	C3orf14	Chromosome 3 open reading frame 14
2.85	STOX1	storkhead box 1
2.84	HSPA1A	Heat shock 70kDa protein 1A; heat shock 70kDa protein 1B
2.83	STS	Steroid sulfatase (microsomal), isozyme S
2.83	HSPA1A	Heat shock 70kDa protein 1A; heat shock 70kDa protein 1B
2.82	SNORA3	Small nucleolar RNA, H/ACA box 3; ribosomal protein L27a
2.80	SFN	Stratifin
2.80	GTF2I	General transcription factor IIi
2.79	DHFR	Dihydrofolate reductase; dihydrofolate reductase pseudogene
2.78	PLEK2	Pleckstrin 2
2.78	RN5S494	RNA, 5S ribosomal 494
2.74	CDCA5	Cell division cycle associated 5
2.72	PKI55	DKFZp434H1419

Table I. Continued

Table I. Continued

Fold change	Gene symbol	mRNA Description	
2.70	TMOD2	Tropomodulin 2 (neuronal)	
2.66	PLCE1	Phospholipase C, epsilon 1	
2.66	GMNN	Geminin, DNA replication inhibitor	
2.65	CEP78	Centrosomal protein 78kDa	
2.65	CXorf57	Chromosome X open reading frame 57	
2.65	RNU5D-1	RNA, U5D small nuclear 1	
2.65	FOXRED2	FAD-dependent oxidoreductase domain containing 2	
2.63	POLR2A	Polymerase (RNA) II (DNA directed) polypeptide A, 220kDa	
2.63	IFIT1	Interferon-induced protein with tetratricopeptide repeats 1	
2.62	PIK3R3	Phosphoinositide-3-kinase, regulatory subunit 3 (gamma)	
2.62	CDC25A	Cell division cycle 25 homolog A (S. pombe)	
2.62	ADI1	Acireductone dioxygenase 1	
2.59	DTL	Denticleless E3 ubiquitin protein ligase homolog (Drosophila)	
2.57	BLM	Bloom syndrome, RecQ helicase-like	
2.57	FAM81A	Family with sequence similarity 81, member A	
2.55	BBS5	Bardet-Biedl syndrome 5	
2.54	DSN1	DSN1, MIND kinetochore complex component, homolog (S. cerevisiae)	
2.53	NT5DC2	5'-nucleotidase domain containing 2	
2.53	FRMD3	FERM domain containing 3	
2.52	KIAA1009	KIAA1009	
2.51	PRKAR2B	Protein kinase, cAMP-dependent, regulatory, type II, beta	
2.51	POLE2	Polymerase (DNA directed), epsilon 2, accessory subunit	
2.51	SKP2	S-phase kinase-associated protein 2, E3 ubiquitin protein ligase	
2.51	IQCG	IQ motif containing G	
2.49	SMAD9	SMAD family member 9	
2.49	SHCBP1	SHC SH2-domain binding protein 1	
2.49	HMGCS1	3-hydroxy-3-methylglutaryl-CoA synthase 1 (soluble)	
2.48	NEFL	Neurofilament, light polypeptide	
2.48	NAGK	N-acetylglucosamine kinase	
2.48	BAMBI	BMP and activin membrane-bound inhibitor homolog (Xenopus laevis)	
2.47	FAM40B	Family with sequence similarity 40, member B	
2.47	ZNF136	Zinc finger protein 136	
2.46	IFIT3	Interferon-induced protein with tetratricopeptide repeats 3	
2.45	CCDC138	Coiled-coil domain containing 138	
2.45	SEPW1	Selenoprotein W, 1	
2.45	OR2AP1	Olfactory receptor, family 2, subfamily AP, member 1	
2.44	HSPB11	Heat shock protein family B (small), member 11	
2.44	LINC00324	Long intergenic non-protein coding RNA 324	
2.43	ID2	Inhibitor of DNA binding 2, dominant negative helix-loop-helix protein	
2.43	MEPCE	Methylphosphate capping enzyme	
2.43	EHHADH	Enoyl-CoA, hydratase/3-hydroxyacyl CoA dehydrogenase	
2.41	SNRNP25	Small nuclear ribonucleoprotein 25kDa (U11/U12)	
2.41	SDAD1	SDA1 domain containing 1	
		Zinc finger protein 701	
2.41	ZNF701		
2.40	FGD6	FYVE, RhoGEF and PH domain containing 6	
2.40	TRIP13	Thyroid hormone receptor interactor 13	
2.40	BRMS1L	Breast cancer metastasis-suppressor 1-like	
2.40	DPF1	D4, zinc and double PHD fingers family 1	
2.39	MCM4	Minichromosome maintenance complex component 4	
2.39	CABLES2	Cdk5 and Abl enzyme substrate 2	
2.39	GPX3	Glutathione peroxidase 3 (plasma)	
2.39	NPTX2	Neuronal pentraxin II	
2.39	TMEM171	Transmembrane protein 171	
2.37	ESCO2	Establishment of cohesion 1 homolog 2 (S. cerevisiae)	
2.37	ZNF280A	Zinc finger protein 280A	
2.37	BFSP1	Beaded filament structural protein 1, filensin	
2.37	GUCY1B3	Guanylate cyclase 1, soluble, beta 3	
2.36	JAKMIP2	Janus kinase and microtubule interacting protein 2	
2.36	C17orf53	Chromosome 17 open reading frame 53	

Table I. Continued

Fold change	Gene symbol	mRNA Description
2.35	ZNF117	Zinc finger protein 117
2.35	APOBEC3B	Apolipoprotein B mRNA editing enzyme, catalytic polypeptide-like 3B
2.35	GDAP1	Ganglioside induced differentiation associated protein 1
2.35	ZNF611	Zinc finger protein 611
2.34	HSPH1	Heat shock 105kDa/110kDa protein 1
2.34	MCM2	Minichromosome maintenance complex component 2
2.34	CCDC56	Coiled-coil domain containing 56
2.33	DHCR24	24-dehydrocholesterol reductase
2.32	CDCA2	Cell division cycle associated 2
2.32	TSPAN12	Tetraspanin 12
2.32	SSX1	Synovial sarcoma, X breakpoint 1
2.32	DYX1C1	Dyslexia susceptibility 1 candidate 1; DYX1C1-CCPG1 readthrough (non-protein coding)
2.32	WDR52	WD repeat domain 52
2.31	WDHD1	WD repeat and HMG-box DNA binding protein 1
2.31	CERK	Ceramide kinase
2.31	ALDH3A1	Aldehyde dehydrogenase 3 family, member A1
2.31	THNSL1	Threonine synthase-like 1 (S. cerevisiae)
2.31	TMEM229B	Transmembrane protein 229B
2.30	PRKAR1B	Protein kinase, cAMP-dependent, regulatory, type I, beta
2.30	RPA2	Replication protein A2, 32kDa
2.30	CENPW	Centromere protein W
2.30	PCNA	Proliferating cell nuclear antigen
2.30	DDX46	DEAD (Asp-Glu-Ala-Asp) box polypeptide 46
2.30	RNY4P10	RNA, Ro-associated Y4 pseudogene 10
2.29	EIF4EBP2	Eukaryotic translation initiation factor 4E binding protein 2
2.29	FOXM1	Forkhead box M1
2.29	FAR2	Fatty acyl CoA reductase 2
2.28	C15orf42	Chromosome 15 open reading frame 42
2.28	GINS1	GINS complex subunit 1 (Psf1 homolog)
2.28	C2orf44	Chromosome 2 open reading frame 44
2.28	CYP4Z2P	Cytochrome P450, family 4, subfamily Z, polypeptide 2 pseudogene
2.28	LYZL2	Lysozyme-like 2
2.27	SCRN1	Secernin 1
2.26	ACAT2	Acetyl-CoA acetyltransferase 2
2.26	NSF	N-ethylmaleimide-sensitive factor; vesicle-fusing ATPase-like
2.25	DENND2C	DENN/MADD domain containing 2C
2.25	ABRACL	ABRA C-terminal like
2.25	SYNJ2	Synaptojanin 2
2.25	POP4	Processing of precursor 4, ribonuclease P/MRP subunit (S. cerevisiae)
2.25	TUBA4A	Tubulin, alpha 4a
2.25	TCTN2	Tectonic family member 2
2.24	CKAP2	Cytoskeleton associated protein 2
2.24	ZWINT	ZW10 interactor
2.24	RN5S180	RNA, 5S ribosomal 180
2.23	CARD6	Caspase recruitment domain family, member 6
2.23	C11orf82	Chromosome 11 open reading frame 82
2.23	RAB27A	RAB27A, member RAS oncogene family
2.23	IDI1	Isopentenyl-diphosphate delta isomerase 1
2.23	ACACB	Acetyl-CoA carboxylase beta
2.23	SAMD9	Sterile alpha motif domain containing 9
2.23	DYDC2	DPY30 domain containing 2
2.22	IQGAP2	IQ motif containing GTPase activating protein 2
2.22	CDCA7L	Cell division cycle associated 7-like
2.21	EFCAB10	EF-hand calcium binding domain 10
2.21	MYBL1	V-myb myeloblastosis viral oncogene homolog (avian)-like 1
2.21	UMPS	Uridine monophosphate synthetase
2.21	MYL6	Myosin, light chain 6, alkali, smooth muscle and non-muscle
2.21	COMMD4	COMM domain containing 4
2.20	SGK3	Serum/glucocorticoid regulated kinase family, member 3; C8orf44-SGK3 readthrough

Table I. Continued

Fold change	e Gene symbol	mRNA Description
2.20	IFI44	Interferon-induced protein 44
2.20	SLC10A5	Solute carrier family 10 (sodium/bile acid cotransporter family), member 5
2.20	DIO2	Deiodinase, iodothyronine, type II
2.19	DBR1	Debranching enzyme homolog 1 (S. cerevisiae)
2.19	ZNF680	Zinc finger protein 680
2.19	MFSD5	Major facilitator superfamily domain containing 5
2.18	ZNF382	Zinc finger protein 382
2.18	RCBTB2	Regulator of chromosome condensation (RCC1) and BTB (POZ) domain containing protein 2
2.17	PSG3	Pregnancy specific beta-1-glycoprotein 3
2.17	ZNF480	Zinc finger protein 480
2.17	RFWD3	Ring finger and WD repeat domain 3
2.17	CENPQ	Centromere protein Q
2.17	PPP2R3A	Protein phosphatase 2, regulatory subunit B", alpha
2.17	RPL41	Ribosomal protein L41
2.17	TUBB4B	Tubulin, beta 4B class IVb
2.17	CABLES1	Cdk5 and Abl enzyme substrate 1
2.16	BRCA1	Breast cancer 1, early onset
2.16	TRAIP	TRAF interacting protein
2.16	SLC25A21	Solute carrier family 25 (mitochondrial oxoadipate carrier), member 21
2.16	RPS6KA6	Ribosomal protein S6 kinase, 90kDa, polypeptide 6
2.15	SAMHD1	SAM domain and HD domain 1
2.15	SHMT1	Serine hydroxymethyltransferase 1 (soluble)
2.15	DNAJA1	DnaJ (Hsp40) homolog, subfamily A, member 1
2.15	HMOX1	Heme oxygenase (decycling) 1
2.15	REEP1	Receptor accessory protein 1
2.14	GINS4	GINS complex subunit 4 (Sld5 homolog)
2.14	SEPT3	Septin 3
2.14	KIAA0101	KIAA0101
2.14	TIMELESS	Timeless homolog (Drosophila)
2.14	MEIG1	Meiosis expressed gene 1 homolog (mouse)
2.14	CORO2A	Coronin, actin binding protein, 2A
2.14	C12orf26	Chromosome 12 open reading frame 26
2.13	FAM171B	Family with sequence similarity 171, member B
2.13	ATP6V0D1	ATPase, H+ transporting, lysosomal 38kDa, V0 subunit d1
2.13	MSH6	MutS homolog 6 (E. coli)
2.13	FN3KRP	Fructosamine 3 kinase related protein
2.13	TUBB3	Tubulin, beta 3 class III
2.13		
	DNAJC6 HYAL1	DnaJ (Hsp40) homolog, subfamily C, member 6
2.12		Hyaluronoglucosaminidase 1
2.12	ZBTB9	Zinc finger and BTB domain containing 9
2.11	PCYT2	Phosphate cytidylyltransferase 2, ethanolamine
2.11	BEND6	BEN domain containing 6 Sorrete BNA effector melecule homeles (Archidensis)
2.11	SRRT	Serrate RNA effector molecule homolog (Arabidopsis)
2.11	SHQ1	SHQ1 homolog (S. cerevisiae)
2.11	PHTF1	Putative homeodomain transcription factor 1
2.10	NAV1	Neuron navigator 1
2.10	MAP2	Microtubule-associated protein 2
2.10	TCF19	Transcription factor 19
2.10	Clorf112	Chromosome 1 open reading frame 112
2.10	TIMM21	Translocase of inner mitochondrial membrane 21 homolog (yeast)
2.10	TACC3	Transforming, acidic coiled-coil containing protein 3
2.10	ZNF468	Zinc finger protein 468
2.10	C2CD3	C2 calcium-dependent domain containing 3
2.10	GCA	Grancalcin, EF-hand calcium binding protein
2.10	MVK	Mevalonate kinase
2.10	DDX23	DEAD (Asp-Glu-Ala-Asp) box polypeptide 23
2.09	COMMD9	COMM domain containing 9
2.09	RPGRIP1L	RPGRIP1-like
2.09	ZNF675	Zinc finger protein 675

Table I. Continued

Fold char	nge Gene symbol	mRNA Description
2.09	RFC5	Replication factor C (activator 1) 5, 36.5kDa
2.09	FTSJ3	FtsJ homolog 3 (E. coli)
2.08	PCYT2	Phosphate cytidylyltransferase 2, ethanolamine
2.08	DHX9	DEAH (Asp-Glu-Ala-His) box polypeptide 9
2.08	GEMIN5	Gem (nuclear organelle) associated protein 5
2.08	C6orf225	Chromosome 6 open reading frame 225
2.07	TCF19	Transcription factor 19
2.07	SMG8	Smg-8 homolog, nonsense mediated mRNA decay factor (C. elegans)
2.07	GIMAP2	GTPase, IMAP family member 2
2.07	UST	Uronyl-2-sulfotransferase
2.07	KBTBD8	Kelch repeat and BTB (POZ) domain containing 8
2.07	HRSP12	Heat-responsive protein 12
2.07	CASP8AP2	Caspase 8 associated protein 2
2.07	CENPP	Centromere protein P
2.07	ISG20L2	Interferon stimulated exonuclease gene 20kDa-like 2
2.06	TUBB4B	Tubulin, beta 4B class IVb
2.06	TUBB2A	Tubulin, beta 2A class IIa
2.06	DCLRE1B	DNA cross-link repair 1B
2.06	RAD51	RAD51 homolog (S. cerevisiae)
2.06	TSGA10	Testis specific, 10
2.06	VPS33A	Vacuolar protein sorting 33 homolog A (S. cerevisiae)
2.06	PRMT7	Protein arginine methyltransferase 7
2.05	MSX2	Msh homeobox 2
2.05	МСМ3	Minichromosome maintenance complex component 3
2.05	POP1	Processing of precursor 1, ribonuclease P/MRP subunit (S. cerevisiae)
2.05	KIF3C	Kinesin family member 3C
2.05	MTBP	Mdm2, transformed 3T3 cell double minute 2, p53 binding protein (mouse) binding protein, 104kDa
2.05	SLFN13	Schlafen family member 13
2.05	ZNF695	Zinc finger protein 695
2.05	CTR9	Ctr9, Paf1/RNA polymerase II complex component, homolog (S. cerevisiae)
2.05	GCNT3	Glucosaminyl (N-acetyl) transferase 3, mucin type
2.04	<i>FANCM</i>	Fanconi anemia, complementation group M
2.04	TSPYL2	TSPY-like 2
2.04	OIP5	Opa interacting protein 5
2.04	MTMR4	Myotubularin related protein 4
2.04	PLCG1	Phospholipase C, gamma 1
2.04	MYBL2	V-myb myeloblastosis viral oncogene homolog (avian)-like 2
2.04	WDR52	WD repeat domain 52
2.04	TEX9	Testis expressed 9
2.03	CCNE1	Cyclin E1
2.03	EFNB2	Ephrin-B2
2.03	CBX5	Chromobox homolog 5
2.03	HMGN2	High mobility group nucleosomal binding domain 2
2.03	TM2D2	TM2 domain containing 2
2.02	TCEB3	Transcription elongation factor B (SIII), polypeptide 3 (110kDa, elongin A)
2.02	CHAF1B	Chromatin assembly factor 1, subunit B (p60)
2.02	QDPR	Quinoid dihydropteridine reductase
2.02	PRAME	Preferentially expressed antigen in melanoma
2.02	TP53I3	Tumor protein p53 inducible protein 3
2.01	SGOL2	Shugoshin-like 2 (S. pombe)
2.01	PIM2	Pim-2 oncogene
2.01	CISD1	CDGSH iron sulfur domain 1
2.01	TP53INP1	Tumor protein p53 inducible nuclear protein 1
2.01	CXCL14	Chemokine (C-X-C motif) ligand 14
2.01	AHSA1	AHA1, activator of heat shock 90kDa protein ATPase homolog 1 (yeast)
2.01	FLJ25715	Uncharacterized protein FLJ25715
2.01	ZNF780B	Zinc finger protein 780B
2.01	FAM98A	Family with sequence similarity 98, member A
2.01	ZNF138	Zinc finger protein 138
2.01	SQRDL	Sulfide quinone reductase-like (yeast)

Table II. H460 BDMC-treated cells vs. H460-BDMC-control.

Fold change	Gene symbol	mRNA Description
-2.01	MIR32	MicroRNA 32
-2.01	TOM1L1	Target of myb1 (chicken)-like 1
-2.01	MAML3	Mastermind-like 3 (Drosophila)
-2.02	KCNN4	Potassium intermediate/small conductance calcium-activated channel, subfamily N, member 4
-2.02	NCAM2	Neural cell adhesion molecule 2
-2.02	CPS1	Carbamoyl-phosphate synthase 1, mitochondrial
-2.02	SPRY3	Sprouty homolog 3 (Drosophila)
-2.02 -2.02	ICAM1	Intercellular adhesion molecule 1
-2.02 -2.03	AXL PRSS23	AXL receptor tyrosine kinase Protease, serine, 23
-2.03 -2.03	NR2F1	Nuclear receptor subfamily 2, group F, member 1
-2.03 -2.03	GADD45B	Growth arrest and DNA-damage-inducible, beta
-2.03	CDK17	Cyclin-dependent kinase 17
-2.03 -2.03	PDE4DIP	Phosphodiesterase 4D interacting protein
-2.03	NEDD1	Neural precursor cell expressed, developmentally down-regulated 1
-2.03	SCCPDH	Saccharopine dehydrogenase (putative)
-2.04	FN1	Fibronectin 1
-2.04	PDE1A	Phosphodiesterase 1A, calmodulin-dependent
-2.04	GPR126	G protein-coupled receptor 126
-2.04	GADD45A	Growth arrest and DNA-damage-inducible, alpha
-2.04	ST6GAL1	ST6 beta-galactosamide alpha-2,6-sialyltranferase 1
-2.04	SLPI	Secretory leukocyte peptidase inhibitor
-2.05	HIST1H2AC	Histone cluster 1, H2ac
-2.05	COL4A6	Collagen, type IV, alpha 6
-2.05	ACSS3	Acyl-CoA synthetase short-chain family member 3
-2.05	TRNAL45P	Transfer RNA leucine 45 (anticodon UAA) pseudogene
-2.05	RHBDD1	Rhomboid domain containing 1
-2.05	HIST1H2AL	Histone cluster 1, H2al; histone cluster 1, H2ah; histone cluster 1, H2ag; histone cluster 1,
		H2am; histone cluster 1, H2ak; histone cluster 1, H2ai
-2.05	SEPP1	Selenoprotein P, plasma, 1
-2.05	PTPN14	Protein tyrosine phosphatase, non-receptor type 14
-2.06	PTPRD	Protein tyrosine phosphatase, receptor type, D
-2.06	C6orf48	Chromosome 6 open reading frame 48
-2.06	PSAT1	Phosphoserine aminotransferase 1
-2.07	NT5C2	5'-nucleotidase, cytosolic II
-2.07	PTPRB	Protein tyrosine phosphatase, receptor type, B
-2.07	SLC2A14	Solute carrier family 2 (facilitated glucose transporter), member 14
-2.07 -2.07	ANP32A-IT1	ANP32A intronic transcript 1 (non-protein coding)
-2.07 -2.08	RASSF8 IFRD1	Ras association (RalGDS/AF-6) domain family (N-terminal) member 8 Interferon-related developmental regulator 1
-2.08 -2.08	SESN2	Sestrin 2
-2.08 -2.08	ITGB5	Integrin, beta 5
-2.08 -2.08	RORA	RAR-related orphan receptor A
-2.08	LPAR6	Lysophosphatidic acid receptor 6
-2.08 -2.08	KLRC4	Killer cell lectin-like receptor subfamily C, member 4
-2.08	CTAGE5	CTAGE family, member 5
-2.09	SLC39A8	Solute carrier family 39 (zinc transporter), member 8
-2.09	LAMB1	Laminin, beta 1
-2.09	TNFRSF19	Tumor necrosis factor receptor superfamily, member 19
-2.10	ITGA6	Integrin, alpha 6
-2.10	RCN3	Reticulocalbin 3, EF-hand calcium binding domain
-2.10	GREM2	Gremlin 2
-2.10	LOXL2	Lysyl oxidase-like 2
-2.10	NRN1	Neuritin 1
-2.11	PDGFA	Platelet-derived growth factor alpha polypeptide
-2.12	PMAIP1	Phorbol-12-myristate-13-acetate-induced protein 1
-2.12	ZC3H6	Zinc finger CCCH-type containing 6
-2.12	ALPK2	Alpha-kinase 2
-2.13	ST3GAL1	ST3 beta-galactoside alpha-2,3-sialyltransferase 1
-2.13	CEBPG	CCAAT/enhancer binding protein (C/EBP), gamma

Table II. Continued

Fold change	Gene symbol	mRNA Description
-2.13	RAB31	RAB31, member RAS oncogene family
-2.13	UNC13A	Unc-13 homolog A (C. elegans)
-2.13	<i>HIST1H2BF</i>	Histone cluster 1, H2bf; histone cluster 1, H2bc; histone cluster 1, H2bi; histone cluster 1,
		H2be; histone cluster 1, H2bg
-2.13	MME	Membrane metallo-endopeptidase
-2.13	TERF1P2	Telomeric repeat binding factor (NIMA-interacting) 1 pseudogene 2
-2.14	ANTXR1	Anthrax toxin receptor 1
-2.14	EML2	Echinoderm microtubule associated protein like 2
-2.14	SREBF2	Sterol regulatory element binding transcription factor 2
-2.14	TF	Transferrin
-2.15	ETV4	Ets variant 4
-2.15	F2RL1	Coagulation factor II (thrombin) receptor-like 1
-2.16	SOCS3	Suppressor of cytokine signaling 3
-2.16	LAMA1	Laminin, alpha 1
-2.16	ARRDC4	Arrestin domain containing 4
-2.17	RAB3B	RAB3B, member RAS oncogene family
-2.17	AGA	Aspartylglucosaminidase
-2.17	COL5A2	Collagen, type V, alpha 2
-2.17	IFITM1	Interferon induced transmembrane protein 1
-2.18	DGKD	Diacylglycerol kinase, delta 130kDa
-2.18	NRP2	Neuropilin 2
-2.18	ALS2CR8	Amyotrophic lateral sclerosis 2 (juvenile) chromosome region, candidate 8
-2.19	CCDC68	Coiled-coil domain containing 68
-2.19	PPFIBP1	PTPRF interacting protein, binding protein 1 (liprin beta 1)
-2.19	MARS	Methionyl-tRNA synthetase
-2.19	RPS6KA2	Ribosomal protein S6 kinase, 90kDa, polypeptide 2
-2.19	COL3A1	Collagen, type III, alpha 1
-2.20	DEPDC1	DEP domain containing 1
-2.20	AKR1B10	Aldo-keto reductase family 1, member B10 (aldose reductase)
-2.20	LONP1	Lon peptidase 1, mitochondrial
-2.20	ARHGEF10	Rho guanine nucleotide exchange factor (GEF) 10
-2.20	SCARNA5	Small Cajal body-specific RNA 5
-2.21	BACE2	Beta-site APP-cleaving enzyme 2
-2.22	RFK	Riboflavin kinase
-2.22	HIST1H3D	Histone cluster 1, H3d; histone cluster 1, H2ad; histone cluster 1, H3f; histone cluster 1, H3b; histone cluster 1, H3h; histone cluster 1, H3j; histone cluster 1, H3g; histone cluster 1, H3c; histone cluster 1, H3a; histone cluster 1, H3c; histone cluster 1, H3a
-2.22	DOCK4	Dedicator of cytokinesis 4
-2.22	RNY3P2	RNA, Ro-associated Y3 pseudogene 2
-2.23	CEBPB	CCAAT/enhancer binding protein (C/EBP), beta
-2.23	LIF	Leukemia inhibitory factor
-2.23	PDLIM5	PDZ and LIM domain 5
-2.24	LRRFIP1	Leucine rich repeat (in FLII) interacting protein 1
-2.24	PCSK1	Proprotein convertase subtilisin/kexin type 1
-2.25	PYCR1	Pyrroline-5-carboxylate reductase 1
-2.25	CRY1	Cryptochrome 1 (photolyase-like)
-2.26	PDE1C	Phosphodiesterase 1C, calmodulin-dependent 70kDa
-2.26	SOX9	SRY (sex determining region Y)-box 9
-2.26	SPDYE8P	Speedy homolog E8 (Xenopus laevis), pseudogene
-2.27	HIST1H3F	Histone cluster 1, H3f; histone cluster 1, H3b; histone cluster 1, H3h; histone cluster 1,
		H3j; histone cluster 1, H3g; histone cluster 1, H3i; histone cluster 1, H3e; histone cluster 1, H3c; histone cluster 1, H3d; histone cluster 1, H3a
-2.27	DUSP5	Dual specificity phosphatase 5
-2.27	PDK1	Pyruvate dehydrogenase kinase, isozyme 1
-2.27	PP7080	Uncharacterized LOC25845
-2.28	GALNTL1	UDP-N-acetyl-alpha-D-galactosamine:polypeptide N-acetylgalactosaminyltransferase-like 1
-2.28	FAM129A	Family with sequence similarity 129, member A
-2.28	ZBTB38	Zinc finger and BTB domain containing 38
-2.28	EPST11	Epithelial stromal interaction 1 (breast)
-2.28	CARS	Cysteinyl-tRNA synthetase

Table II. Continued

Fold char	nge Gene symbol	mRNA Description
-2.28	WDR69	WD repeat domain 69
-2.29	GPR65	G protein-coupled receptor 65
-2.30	XYLT1	Xylosyltransferase I
-2.30	TCEA1	Transcription elongation factor A (SII), 1
-2.30	F2R	Coagulation factor II (thrombin) receptor
-2.31	CXCL5	Chemokine (C-X-C motif) ligand 5
-2.31	LOC100128252	Uncharacterized LOC100128252
-2.31	EGR1	Early growth response 1
-2.32	FRY	Furry homolog (Drosophila)
-2.32	CEP70	Centrosomal protein 70kDa
-2.32 -2.33	SLC37A2	Solute carrier family 37 (glycerol-3-phosphate transporter), member 2
-2.33 -2.33	TRMT2B	TRNA methyltransferase 2 homolog B (S. cerevisiae)
-2.33 -2.33	CSGALNACT1	Chondroitin sulfate N-acetylgalactosaminyltransferase 1 Histone cluster 3, H2bb
-2.33 -2.34	HIST3H2BB ZNF585B	Zinc finger protein 585B
-2.34 -2.34	PCID2	PCI domain containing 2
-2.34 -2.34	XBP1	X-box binding protein 1
-2.34	CDH13	Cadherin 13, H-cadherin (heart)
-2.34	GPR37	G protein-coupled receptor 37 (endothelin receptor type B-like)
-2.35	SERPINB9	Serpin peptidase inhibitor, clade B (ovalbumin), member 9
-2.35	GFPT1	Glutaminefructose-6-phosphate transaminase 1
-2.35	SOCS2	Suppressor of cytokine signaling 2
-2.35	ZEB2	Zinc finger E-box binding homeobox 2
-2.35	IGFBP3	Insulin-like growth factor binding protein 3
-2.35	LCN1	Lipocalin 1
-2.36	TTC32	Tetratricopeptide repeat domain 32
-2.36	COL4A5	Collagen, type IV, alpha 5
-2.36	RN5S353	RNA, 5S ribosomal 353
-2.37	ZFP36	Zinc finger protein 36, C3H type, homolog (mouse)
-2.37	PLEKHG2	Pleckstrin homology domain containing, family G (with RhoGef domain) member 2
-2.37	DUSP5P	Dual specificity phosphatase 5 pseudogene
-2.38	CRLF2	Cytokine receptor-like factor 2
-2.38	LRRC63	Leucine rich repeat containing 63
-2.38	SLITRK4	SLIT and NTRK-like family, member 4
-2.39	ITPRIP	Inositol 1,4,5-trisphosphate receptor interacting protein
-2.40	OCLN	Occludin; occludin pseudogene
-2.40	PLAU	Plasminogen activator, urokinase
-2.40	KIAA0825	KIAA0825
-2.41	OSMR	Oncostatin M receptor
-2.41	LAT2	Linker for activation of T cells family, member 2
-2.41	SLC25A37	Solute carrier family 25 (mitochondrial iron transporter), member 37
-2.42	EDN1	Endothelin 1
-2.42	THSD4	Thrombospondin, type I, domain containing 4
-2.43	PLA2R1	Phospholipase A2 receptor 1, 180kDa
-2.43	CTIF	CBP80/20-dependent translation initiation factor
-2.43	CEP72	Centrosomal protein 72kDa
-2.43	BMP6	Bone morphogenetic protein 6 Pharmharmata and avvilling 2 2 (mitashandrial)
-2.45	PCK2	Phosphoenolpyruvate carboxykinase 2 (mitochondrial)
-2.45 2.45	DTWD1	DTW domain containing 1 Thrombospondin 1
-2.45 2.45	THBS1	1
-2.45 -2.45	PABPC1L	Poly(A) binding protein, cytoplasmic 1-like Alanine-glyoxylate aminotransferase 2-like 2
-2.45 -2.45	AGXT2L2 CPED1	Cadherin-like and PC-esterase domain containing 1
-2.45 -2.45	PCDHB17	· · · · · · · · · · · · · · · · · · ·
-2.45 -2.46	TCEA1	Protocadherin beta 17 pseudogene Transcription elongation factor A (SII), 1
-2.46 -2.46	ASNS	Asparagine synthetase (glutamine-hydrolyzing)
-2.46 -2.47	ASNS FLNB	Asparagine synthetase (glutamine-nydrotyzing) Filamin B, beta
-2.47 -2.47	MCL1	Myeloid cell leukemia sequence 1 (BCL2-related)
-2.47 -2.47	SPDYE8P	Speedy homolog E8 (Xenopus laevis), pseudogene
4.7/	SI DI EGI	opecay nomorog to (Achopus racers), pseudogene

Table II. Continued

	Gene symbol	mRNA Description
-2.48	RSPO3	R-spondin 3
-2.48	SLC2A13	Solute carrier family 2 (facilitated glucose transporter), member 13
-2.48	SLC7A1	Solute carrier family 7 (cationic amino acid transporter, y+ system), member 1
-2.48	DUSP4	Dual specificity phosphatase 4
-2.48	CLK1	CDC-like kinase 1
-2.48	CRELD1	Cysteine-rich with EGF-like domains 1
-2.48	TRNAQ52P	Transfer RNA glutamine 52 (anticodon UUG) pseudogene
-2.49 -2.49	RN5S506 SEMA3C	RNA, 5S ribosomal 506 Sema domain, immunoglobulin domain (Ig), short basic domain, secreted, (semaphorin) 3C
-2.49 -2.49	SSFA2	sperm specific antigen 2
-2. 4 9 -2.49	LOC344887	NmrA-like family domain containing 1 pseudogene
-2.50	SLC4A7	Solute carrier family 4, sodium bicarbonate cotransporter, member 7
-2.50	PDCD1LG2	Programmed cell death 1 ligand 2
-2.50	P4HA1	Prolyl 4-hydroxylase, alpha polypeptide I
-2.51	FAM27E3	Family with sequence similarity 27, member E3; family with sequence similarity 27, member E2
-2.53	KIAA0825	KIAA0825
-2.53	EFEMP1	EGF containing fibulin-like extracellular matrix protein 1
-2.53	FAM27E3	Family with sequence similarity 27, member E3; family with sequence similarity 27, member E2
-2.53	MIR21	MicroRNA 21
-2.53	GLRXP3	Glutaredoxin (thioltransferase) pseudogene 3
-2.54	GDF15	Growth differentiation factor 15
-2.54	LURAP1L	Leucine rich adaptor protein 1-like
-2.55	PRDM1	PR domain containing 1, with ZNF domain
-2.55	RNF19A	Ring finger protein 19A, E3 ubiquitin protein ligase
-2.55 -2.56	SLC4A11	Solute carrier family 4, sodium borate transporter, member 11 Nuclear receptor subfamily 4, group A, member 1
-2.56 -2.56	NR4A1	Sema domain, immunoglobulin domain (Ig), short basic domain, secreted, (semaphorin) 3A
-2.56 -2.56	<i>SEMA3A</i> <i>MIF4GD</i>	MIF4G domain containing
-2.56 -2.56	ANGPTL4	Angiopoietin-like 4
-2.57	FGG	Fibrinogen gamma chain
-2.57	ADAM19	ADAM metallopeptidase domain 19
-2.57	ITGA3	Integrin, alpha 3 (antigen CD49C, alpha 3 subunit of VLA-3 receptor)
-2.59	SNRPA1	Small nuclear ribonucleoprotein polypeptide A'
-2.59	FBXL13	F-box and leucine-rich repeat protein 13
-2.59	ADAMTS6	ADAM metallopeptidase with thrombospondin type 1 motif, 6
-2.59	GPRIN3	GPRIN family member 3
-2.59	STC2	Stanniocalcin 2
-2.60	FAM27E3	Family with sequence similarity 27, member E3; family with sequence similarity 27, member E2
-2.60	IL18R1	Interleukin 18 receptor 1
-2.60	SUCNR1	Succinate receptor 1
-2.61	BTG1	B-cell translocation gene 1, anti-proliferative
-2.61 2.62	PLAT	Plasminogen activator, tissue
-2.62 2.62	CCPG1 GDPD1	Cell cycle progression 1; DYX1C1-CCPG1 readthrough (non-protein coding) Glycerophosphodiester phosphodiesterase domain containing 1
-2.62 -2.63	HOXB9	Homeobox B9
-2.63 -2.64	CD55	CD55 molecule, decay accelerating factor for complement (Cromer blood group)
-2.64	LAMB3	Laminin, beta 3
-2.65	PSD3	Pleckstrin and Sec7 domain containing 3
-2.65	SLC35E3	Solute carrier family 35, member E3
-2.67	ODZ2	Odz, odd Oz/ten-m homolog 2 (Drosophila)
-2.68	FLJ44124	Uncharacterized LOC641737
-2.71	SLCO5A1	Solute carrier organic anion transporter family, member 5A1
-2.72	FOSL2	FOS-like antigen 2
-2.72	CTBS	Chitobiase, di-N-acetyl-
-2.73	FBN2	Fibrillin 2
-2.73	SLC9A3	Solute carrier family 9, subfamily A (NHE3, cation proton antiporter 3), member 3
-2.74	THSD7A	Thrombospondin, type I, domain containing 7A
-2.76	FSTL1	Follistatin-like 1
-2.76	PPAP2B	Phosphatidic acid phosphatase type 2B
-2.77	IL8	Interleukin 8

Table II. Continued

Fold change	Gene symbol	mRNA Description
-2.77	JHDM1D	Jumonji C domain containing histone demethylase 1 homolog D (S. cerevisiae)
-2.77	PSD3	Pleckstrin and Sec7 domain containing 3
-2.78	TFPI2	Tissue factor pathway inhibitor 2
-2.79	PTPRM	Protein tyrosine phosphatase, receptor type, M
-2.80	SERPINB1	Serpin peptidase inhibitor, clade B (ovalbumin), member 1
-2.82	PAPPA2	Pappalysin 2
-2.85	TUBE1	Tubulin, epsilon 1
-2.86	PHLDA1	Pleckstrin homology-like domain, family A, member 1
-2.86	RGS2	Regulator of G-protein signaling 2, 24kDa
-2.86	OLFML2A	olfactomedin-like 2A
-2.87	SLC7A2	Solute carrier family 7 (cationic amino acid transporter, y+ system), member 2
-2.89	LAMA5	Laminin, alpha 5
-2.91	EDIL3	EGF-like repeats and discoidin I-like domains 3
-2.91	TRPC6	Transient receptor potential cation channel, subfamily C, member 6
-2.91	ULBP1	UL16 binding protein 1
-2.92	VEGFA	Vascular endothelial growth factor A
-2.92	LOC284861	Uncharacterized LOC284861
-2.94	BDNF	Brain-derived neurotrophic factor
-2.95	ABCC3	ATP-binding cassette, sub-family C (CFTR/MRP), member 3
-2.97	FHAD1	Forkhead-associated (FHA) phosphopeptide binding domain 1
-2.99	DDIT4	DNA-damage-inducible transcript 4
-3.02	ANTXR2	Anthrax toxin receptor 2
-3.03	HERPUD1	Homocysteine-inducible, endoplasmic reticulum stress-inducible, ubiquitin-like domain member 1
-3.04	DPP10	Dipeptidyl-peptidase 10 (non-functional)
-3.06	BEST1	Bestrophin 1
-3.06	TMEM144	Transmembrane protein 144
-3.06	OGT	O-linked N-acetylglucosamine (GlcNAc) transferase
-3.07	UNC5B	Unc-5 homolog B (C. elegans)
-3.07	GLRX	Glutaredoxin (thioltransferase)
-3.07	SPOCK1	Sparc/osteonectin, cwcv and kazal-like domains proteoglycan (testican) 1
-3.08	SLC6A9	Solute carrier family 6 (neurotransmitter transporter, glycine), member 9
-3.10	HSPA13	Heat shock protein 70kDa family, member 13
-3.11	BHLHE40	Basic helix-loop-helix family, member e40
-3.13	CYR61	Cysteine-rich, angiogenic inducer, 61
-3.15	DUSP1	Dual specificity phosphatase 1
-3.16	LSAMP	Limbic system-associated membrane protein
-3.18	CDH19	Cadherin 19, type 2
-3.19	GTPBP2	GTP binding protein 2
-3.20	CD36	CD36 molecule (thrombospondin receptor)
-3.20	HPGDS	Hematopoietic prostaglandin D synthase
-3.20	CLDN1	Claudin 1
-3.20	NRCAM	Neuronal cell adhesion molecule
-3.21	PITPNC1	Phosphatidylinositol transfer protein, cytoplasmic 1
-3.27	DLG2	Discs, large homolog 2 (Drosophila)
-3.28	PRKAA2	Protein kinase, AMP-activated, alpha 2 catalytic subunit
-3.29	ALDH1L2	Aldehyde dehydrogenase 1 family, member L2
-3.30	PDE3A	Phosphodiesterase 3A, cGMP-inhibited
-3.30	STARD4	StAR-related lipid transfer (START) domain containing 4
-3.31	NR1D1	Nuclear receptor subfamily 1, group D, member 1
-3.33	SLC25A37	Solute carrier family 25 (mitochondrial iron transporter), member 37
-3.35	H1F0	H1 histone family, member 0
-3.37	RAB39B	RAB39B, member RAS oncogene family
-3.39	CLGN	Calmegin
-3.40	IL6	Interleukin 6 (interferon, beta 2)
-3.40	VLDLR	Very low density lipoprotein receptor
-3.43	SMOX	Spermine oxidase
-3.43	SCARA5	Scavenger receptor class A, member 5 (putative)
-3.45	PLAUR	Plasminogen activator, urokinase receptor
-3.47	B4GALT1	UDP-Gal:betaGlcNAc beta 1,4- galactosyltransferase, polypeptide 1
-3.48	TSPAN8	Tetraspanin 8

Table II. Continued

roid chan	ge Gene symbol	mRNA Description
-3.49	APLN	Apelin
-3.51	DUSP6	Dual specificity phosphatase 6
3.53	PRICKLE1	Prickle homolog 1 (Drosophila)
3.53	NR4A2	Nuclear receptor subfamily 4, group A, member 2
3.59	CBS	Cystathionine-beta-synthase
3.59	LINC00173	Long intergenic non-protein coding RNA 173
3.61	TNFAIP3	Tumor necrosis factor, alpha-induced protein 3
3.62	SHISA3	Shisa homolog 3 (Xenopus laevis)
3.62	EPHA4	EPH receptor A4
3.70	TSC22D3	TSC22 domain family, member 3
3.70	KLRC3	Killer cell lectin-like receptor subfamily C, member 3; killer cell lectin-like receptor subfamily C, member 2
3.71	TFPI	Tissue factor pathway inhibitor (lipoprotein-associated coagulation inhibitor)
3.75	CXCL2	Chemokine (C-X-C motif) ligand 2
3.82	ANXA3	Annexin A3
3.84	SLC16A4	Solute carrier family 16, member 4 (monocarboxylic acid transporter 5)
3.84	<i>KLRC4-KLRK1</i>	KLRC4-KLRK1 readthrough; killer cell lectin-like receptor subfamily K, member 1
3.90	PLAC4	Placenta-specific 4
3.97	ABCC9	ATP-binding cassette, sub-family C (CFTR/MRP), member 9
4.03	TMEM27	Transmembrane protein 27
4.03	STC1	Stanniocalcin 1
4.05	SEMA3D	Sema domain, immunoglobulin domain (Ig), short basic domain, secreted, (semaphorin) 3D
4.05	TRIB3	Tribbles homolog 3 (Drosophila)
4.06	CTGF	Connective tissue growth factor
4.09	NUPR1	Nuclear protein, transcriptional regulator, 1
4.10	ABI3BP	ABI family, member 3 (NESH) binding protein
4.21	FBXO32	F-box protein 32
4.25	RPE65	Retinal pigment epithelium-specific protein 65kDa
4.35	PARK2	Parkinson protein 2, E3 ubiquitin protein ligase (parkin)
4.38	C3orf49	Chromosome 3 open reading frame 49
4.48	WARS	Tryptophanyl-tRNA synthetase
4.50	SAT1	Spermidine/spermine N1-acetyltransferase 1
4.51	DDIT3	DNA-damage-inducible transcript 3
4.52	TCP11L2	T-complex 11 (mouse)-like 2
4.65	KCNE4	Potassium voltage-gated channel, Isk-related family, member 4
4.65	MIR29A	MicroRNA 29a
4.66	ERRF11	ERBB receptor feedback inhibitor 1
4.67	AREG	Amphiregulin
4.70	AREG	Amphiregulin
4.80	CTH	Cystathionase (cystathionine gamma-lyase)
4.89	PAPPA	Pregnancy-associated plasma protein A, pappalysin 1
4.89	ESM1	Endothelial cell-specific molecule 1
4.97	C12orf39	Chromosome 12 open reading frame 39
5.03	TXNIP	Thioredoxin interacting protein
5.06	SMOC1	SPARC related modular calcium binding 1
5.10	ASS1P11	Argininosuccinate synthetase 1 pseudogene 11
5.23	LCN1	Lipocalin 1
5.39	ASS1	Argininosuccinate synthase 1
5.44	CHAC1	ChaC, cation transport regulator homolog 1 (E. coli)
5.47	CHL1	Cell adhesion molecule with homology to L1CAM (close homolog of L1)
6.06	GABRE	Gamma-aminobutyric acid (GABA) A receptor, epsilon; microRNA 452; microRNA 224
6.39	FSTL4	Follistatin-like 4
6.44	CD274	CD274 molecule
7.68	TGFB2	Transforming growth factor, beta 2
7.75	GABRE	Gamma-aminobutyric acid (GABA) A receptor, epsilon; microRNA 452; microRNA 224
8.71	SERPINB11	Serpin peptidase inhibitor, clade B (ovalbumin), member 11 (gene/pseudogene)
-9.43	FST	Follistatin
-9.43 -9.94	SLC2A3	Solute carrier family 2 (facilitated glucose transporter), member 3
10.77	GPR133	G protein-coupled receptor 133
10.//	UI NIJJ	O protein-coupled receptor 133

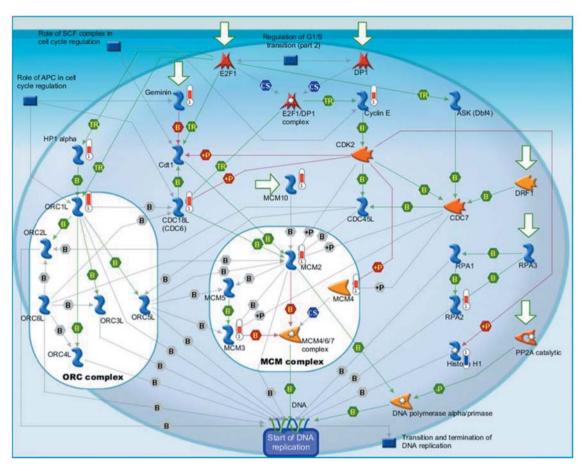


Figure 1. Cell cycle: Start of DNA replication. The top scored map (map with the lowest p-value) based on the enrichment distribution sorted by 'Statistically significant Maps' set. Experimental data from all files are linked to and visualized on the maps as thermometer like figures. Up ward thermometers have red color and indicate up regulated signals and down ward (blue) ones indicate down regulated expression levels of the genes.

cycle and 255 genes were down-regulated from 2 to 3 folds such as growth arrest and DNA-damage-inducible, alpha (GADD45A), DNA-damage-inducible transcript 4 (DDIT4), DNA-damage-inducible transcript 3 (DDIT3), cyclin-dependent kinase 17 (CDK17), CDC-like kinase 1 (CLK1), and cell cycle progression 1, DYX1C1-CCPG1 read through (CCPG1) associated with cell cycle, tumor necrosis factor receptor superfamily, member 19 (TNFRSF19), early growth response 1 (EGR1), programmed cell death 1 ligand 2 (PDCD1LG2), and ATP-binding cassette, sub-family C (CFTR/MRP), member 3 (ABCC3) and member 9 (ABCC9), intercellular adhesion molecule 1 (ICAM1), Ras association (RalGDS/AF-6) domain family (N-terminal) member 8 (RASSF8), Rho guanine nucleotide exchange factor 10 (ARHGEF10) and cell adhesion molecule with homology to L1CAM (close homolog of L1) (CHLI) associated with cell migration.

BDMC affected the gene expression score measured by GeneGo analysis program in NCI-H460 cells by the number of

pathway networks. After total mRNA was isolated from BDMC-treated or -untreated cells and used for cDNA microarrays, the analysis was followed with further processing by GeneGo and the results are shown in Figures 1, 2 and 3. Figure 1 indicates that BDMC affected associated gene expression with cell cycle that starts from DNA replication in the early S phase. For example, BDMC up-regulates cyclin E and CDC6 but down-regulates CDK2 and CDC7, leading to affect the start of DNA replication in NCI-H460 cells. Figure 2 indicates that BDMC affected associated gene expression in development with TGF-beta-dependent induction of EMT via SMADs. For example, TGF-beta receptor type 1 and II were up-regulated by BDMC but SNAIL, SIP1 and MMP-2 were down-regulated affecting cell motility, adhesion and epithelialto-mesenchymal transition in NCI-H460 cells. Figure 3 indicates that BDMC affected gene expression, which associates with targets of tissue factor signaling in cancer. BDMC affected angiogenesis, inhibition of apoptosis, cell migration and invasion, leading to tumor progression.

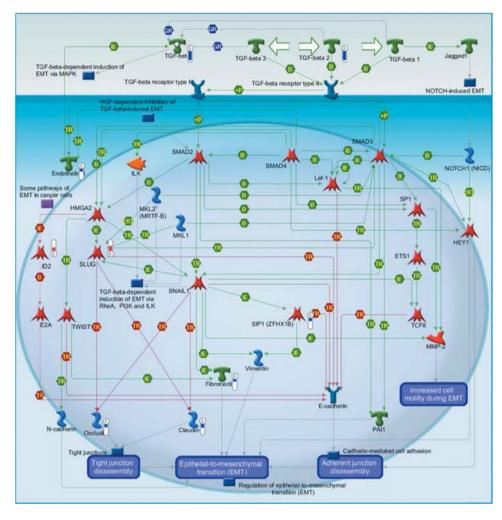


Figure 2. Development: TGF beta dependent induction of EMT via SMADs. The second top-scored map (map with the second lowest p-value) based on enrichment distribution, sorted by 'Statistically significant Maps' set. Experimental data from all files are linked to and visualized on the maps as thermometer like figures. Up ward thermometers have red color and indicate up regulated signals and down ward (blue) ones indicate down regulated expression levels of the genes.

Discussion

Several reports have shown that BDMC induces cytotoxic effects in human cancer cells *in vitro*, however, there is no available information on how BDMC affects gene expression and associated signaling pathways in human lung cancer cells. In the present study, we examined whether BDMC induced up- or down-regulation of genes associated with cell cycle, DNA replication, cell survival, cell migration and invasion and tumor progression in NCI H460 cells.

Table I indicates that *ERCC6L*, a gene associated with DNA damage and repair, was increased by 3.58-fold, four genes *CDC6*, *CDCA5*, *CDC25A* and *CDCA7L* associated with cell division, were increased by 2.98-, 2.74-, 2.62- and 2.22-fold, respectively, three genes *SKP2*, *CABLES2* and

CABLES1 associated with cell cycle, were increased by 2.51, 2.39, and 2.17 times respectively, three genes *CARD6*, *ATP6V0D1* and *CASP8AP2*, associated with cell death, were found increased by 2.23-, 2.13- and 2.07-times, respectively.

Table II demonstrates that BDMC suppressed expression of numerous genes associated with DNA damage, cell cycle, cell survival and cell migration and invasion. In particular, three genes *DDIT3*, *DDIT4* and *GADD45A* associated with DNA damage, were decreased by 4.51-, 2.99- and 2.04-times, respectively, three genes *CDK17*, *CLK1* and *CCPG1*, associated with cell cycle, were found decreased by 2.03-, 2.48- and 2.62-times, respectively, five genes *TNFRSF19*, *EGR1*, *PDCD1LG2*, *ABCC3* and *ABCC9* associated with cell death, were decreased 2.09-, 2.31-, 2.50-, 2.95- and 3.97-fold, respectively, four genes *ICAM1*, *RASSF8*, *ARHGEF10*

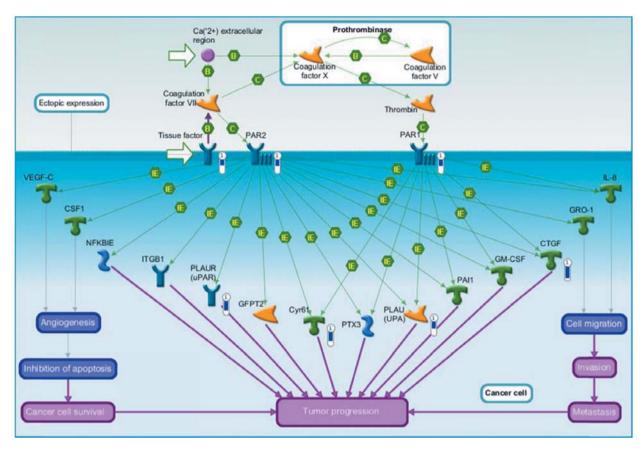


Figure 3. Expression targets of Tissue factor signaling in cancer. The third top-scored map (map with the third lowest p value) based on the enrichment distribution sorted by 'Statistically significant Maps' set. Experimental data from all files are linked to and visualized on the maps as thermometer like figures. Up ward thermometers have red color and indicate up regulated signals and down ward (blue) ones indicate down regulated expression levels of the genes.

and *CHL1*, associated with cell migration, were decreased 2.02-, 2.07-, 2.20-, and 5.47-fold, respectively.

It is well-documented that cell-cycle regulation is the key mechanism of cancer cell growth (28, 29). Results from Figure 1 show that BDMC up-regulated cyclin E and CDC6, however, down-regulated CDK2 and CDC7 affecting the start of DNA replication in NCI-H460 cells, and in turn affecting S phase. It has been reported that cyclin E/Cdk2 complex is involved in G1-S transition and is also associated with initiation of DNA synthesis. Furthermore, the cyclin A/Cdk2 complex is associated with the initiation of DNA synthesis and is also associated with the progression to S phase (30). It has also been reported that increase in Cdc7 and/or Dbf4 can arrest cells in G₁ phase, or slow down Sphase progression when cells are already in S phase (31). Furthermore, it has also been reported that Cdc6 and Mcm proteins are required to establish pre-replicative complex and the activities of Cdks and of Cdc7 kinase for triggering the G_1 -S transition (32).

Results from Figure 2 demonstrate that BDMC upregulated the transforming growth factor-beta (TGF- β) receptor type 1 and II, however down-regulated SNAIL, SIP1 and MMP-2 that were associated with cell motility, adhesion and epithelial-to-mesenchymal transition in NCI-H460 cells. It is well-known that TGF- β ligands play an important role in cell proliferation, extracellular matrix production, cell motility and apoptosis (33). It has also been reported that in the mammary gland of transgenic mice, if there was overexpression of active TGF- β 1 or an activated type I TGF- β 1 receptor (T β RI), it can lead to accelerated metastases derived from Neu-induced mammary tumors (34, 35).

The zinc-finger transcription factors Snail and SIP1 (Smad interacting protein 1) have been demonstrated to repress transcription of the E-cadherin (E-cad) gene by binding to E-boxes (CACCTG sequence) on the E-cad promoter (36, 37). E-cad plays a major role in the establishment and maintenance of intercellular adhesion, cell polarity, and tissue architecture (38, 39). Furthermore, matrix metalloproteinase

(MMP-1, MMP-2, MMP-7), and MT1-MMP expressions are strongly upregulated by Snail (40). Results from Figure 3 indicate that BDMC affects angiogenesis, inhibition of apoptosis, cell migration and invasion, leading to tumor progression. Thus, further studies are required to expand or append our current findings and possible understanding of them. In the present study, the genes affected by BDMC *in vitro* may offer certain insight on the cytotoxic mechanism of BDMC in the genetic level, which in turn may provide potentially useful biomarkers or targets for diagnosis and treatment of human lung cancer.

Acknowledgements

This work was supported by grant DMR-101-051 from China Medical University Hospital, Taichung, Taiwan.

References

- 1 Powell HA, Tata LJ, Baldwin DR, Potter VA, Stanley RA, Khakwani A and Hubbard RB: Treatment decisions and survival for people with small-cell lung cancer. Br J Cancer 110: 908-915, 2014.
- 2 McCulloch M, See C, Shu XJ, Broffman M, Kramer A, Fan WY, Gao J, Lieb W, Shieh K and Colford JM Jr.: Astragalus-based Chinese herbs and platinum-based chemotherapy for advanced non-small-cell lung cancer: meta-analysis of randomized trials. J Clin Oncol 24: 419-430, 2006.
- 3 Shaw AT and Engelman JA: ALK in lung cancer: past, present, and future. J Clin Oncol 31: 1105-1111, 2013.
- 4 Siegel R, Ma J, Zou Z and Jemal A: Cancer statistics, 2014. CA Cancer J Clin 64: 9-29, 2014.
- 5 Morra F, Luise C, Visconti R, Staibano S, Merolla F, Ilardi G, Guggino G, Paladino S, Sarnataro D, Franco R, Monaco R, Zitomarino F, Pacelli R, Monaco G, Rocco G, Cerrato A, Linardopoulos S, Muller MT and Celetti A: New therapeutic perspectives in CCDC6 deficient lung cancer cells. Int J Cancer 136: 2146-2157, 2015.
- 6 Tsai MF, Wang CC and Chen JJ: Tumour suppressor HLJ1: A potential diagnostic, preventive and therapeutic target in non-small cell lung cancer. World J Clin Oncol 5: 865-873, 2014.
- 7 Chiappedi M and Bejor M: Herbals and natural dietary supplements in psychiatric practice. Recent Pat CNS Drug Discov 5: 164-171, 2010.
- 8 Aggarwal BB, Sundaram C, Malani N and Ichikawa H: Curcumin: the Indian solid gold. Adv Exp Med Biol 595: 1-75, 2007.
- 9 Krishnaswamy K: Traditional Indian spices and their health significance. Asia Pac J Clin Nutr 17(Suppl 1): 265-268, 2008.
- 10 Aggarwal BB, Takada Y and Oommen OV: From chemoprevention to chemotherapy: common targets and common goals. Expert Opin Investig Drugs 13: 1327-1338, 2004.
- 11 Sandur SK, Pandey MK, Sung B, Ahn KS, Murakami A, Sethi G, Limtrakul P, Badmaev V and Aggarwal BB: Curcumin, demethoxycurcumin, bisdemethoxycurcumin, tetrahydrocurcumin and turmerones differentially regulate anti-inflammatory and antiproliferative responses through a ROS-independent mechanism. Carcinogenesis 28: 1765-1773, 2007.

- 12 Yodkeeree S, Chaiwangyen W, Garbisa S and Limtrakul P: Curcumin, demethoxycurcumin and bisdemethoxycurcumin differentially inhibit cancer cell invasion through the downregulation of MMPs and uPA. J Nutr Biochem 20: 87-95, 2009.
- 13 Basile V, Ferrari E, Lazzari S, Belluti S, Pignedoli F and Imbriano C: Curcumin derivatives: molecular basis of their anticancer activity. Biochem Pharmacol 78: 1305-1315, 2009.
- 14 Boonrao M, Yodkeeree S, Ampasavate C, Anuchapreeda S and Limtrakul P: The inhibitory effect of turmeric curcuminoids on matrix metalloproteinase-3 secretion in human invasive breast carcinoma cells. Arch Pharm Res *33*: 989-998, 2010.
- 15 Kim AN, Jeon WK, Lee JJ and Kim BC: Up-regulation of heme oxygenase-1 expression through CaMKII-ERK1/2-Nrf2 signaling mediates the anti-inflammatory effect of bisdemethoxycurcumin in LPS-stimulated macrophages. Free Radic Biol Med 49: 323-331, 2010.
- 16 Ryu MJ, Cho M, Song JY, Yun YS, Choi IW, Kim DE, Park BS and Oh S: Natural derivatives of curcumin attenuate the Wnt/beta-catenin pathway through down-regulation of the transcriptional coactivator p300. Biochem Biophys Res Commun 377: 1304-1308, 2008.
- 17 Belluti S, Basile V, Benatti P, Ferrari E, Marverti G and Imbriano C: Concurrent inhibition of enzymatic activity and NF-Y-mediated transcription of Topoisomerase-IIalpha by bis-Demethoxy Curcumin in cancer cells. Cell Death Dis 4: e756, 2013.
- 18 Douillard JY, Oliner KS, Siena S, Tabernero J, Burkes R, Barugel M, Humblet Y, Bodoky G, Cunningham D, Jassem J, Rivera F, Kocakova I, Ruff P, Blasinska-Morawiec M, Smakal M, Canon JL, Rother M, Williams R, Rong A, Wiezorek J, Sidhu R and Patterson SD: Panitumumab-FOLFOX4 treatment and RAS mutations in colorectal cancer. N Engl J Med 369: 1023-1034, 2013.
- 19 Mok TS, Wu YL, Thongprasert S, Yang CH, Chu DT, Saijo N, Sunpaweravong P, Han B, Margono B, Ichinose Y, Nishiwaki Y, Ohe Y, Yang JJ, Chewaskulyong B, Jiang H, Duffield EL, Watkins CL, Armour AA and Fukuoka M: Gefitinib or carboplatin-paclitaxel in pulmonary adenocarcinoma. N Engl J Med 361: 947-957, 2009.
- 20 Normanno N, Rachiglio AM, Roma C, Fenizia F, Esposito C, Pasquale R, La Porta ML, Iannaccone A, Micheli F, Santangelo M, Bergantino F, Costantini S and De Luca A: Molecular diagnostics and personalized medicine in oncology: challenges and opportunities. J Cell Biochem 114: 514-524, 2013.
- 21 Garraway LA: Genomics-driven oncology: framework for an emerging paradigm. J Clin Oncol 31: 1806-1814, 2013.
- 22 Li T, Kung HJ, Mack PC and Gandara DR: Genotyping and genomic profiling of non-small-cell lung cancer: implications for current and future therapies. J Clin Oncol 31: 1039-1049, 2013.
- 23 Bunn PA, Jr.: Worldwide overview of the current status of lung cancer diagnosis and treatment. Arch Pathol Lab Med 136: 1478-1481, 2012.
- 24 Eberhard DA, Johnson BE, Amler LC, Goddard AD, Heldens SL, Herbst RS, Ince WL, Janne PA, Januario T, Johnson DH, Klein P, Miller VA, Ostland MA, Ramies DA, Sebisanovic D, Stinson JA, Zhang YR, Seshagiri S and Hillan KJ: Mutations in the epidermal growth factor receptor and in KRAS are predictive and prognostic indicators in patients with non-small-cell lung cancer treated with chemotherapy alone and in combination with erlotinib. J Clin Oncol 23: 5900-5909, 2005.

- 25 Kwak EL, Bang YJ, Camidge DR, Shaw AT, Solomon B, Maki RG, Ou SH, Dezube BJ, Janne PA, Costa DB, Varella-Garcia M, Kim WH, Lynch TJ, Fidias P, Stubbs H, Engelman JA, Sequist LV, Tan W, Gandhi L, Mino-Kenudson M, Wei GC, Shreeve SM, Ratain MJ, Settleman J, Christensen JG, Haber DA, Wilner K, Salgia R, Shapiro GI, Clark JW and Iafrate AJ: Anaplastic lymphoma kinase inhibition in non-small-cell lung cancer. N Engl J Med 363: 1693-1703, 2010.
- 26 Wu SH, Hsiao YT, Chen JC, Lin JH, Hsu SC, Hsia TC, Yang ST, Hsu WH and Chung JG: Bufalin alters gene expressions associated DNA damage, cell cycle, and apoptosis in human lung cancer NCI-H460 cells in vitro. Molecules 19: 6047-6057, 2014.
- 27 Gardina PJ, Clark TA, Shimada B, Staples MK, Yang Q, Veitch J, Schweitzer A, Awad T, Sugnet C, Dee S, Davies C, Williams A and Turpaz Y: Alternative splicing and differential gene expression in colon cancer detected by a whole genome exon array. BMC Genomics 7: 325, 2006.
- 28 Li YG, Ji DF, Zhong S, Lin TB, Lv ZQ, Hu GY and Wang X: 1-deoxynojirimycin inhibits glucose absorption and accelerates glucose metabolism in streptozotocin-induced diabetic mice. Sci Rep 3: 1377, 2013.
- 29 Murray AW: Recycling the cell cycle: cyclins revisited. Cell 116: 221-234, 2004.
- 30 Moon J, Yu SJ, Kim HS and Sohn J: Induction of G(1) cell cycle arrest and p27(KIP1) increase by panaxydol isolated from Panax ginseng. Biochem Pharmacol 59: 1109-1116, 2000.
- 31 Guo B, Romero J, Kim BJ and Lee H: High levels of Cdc7 and Dbf4 proteins can arrest cell-cycle progression. Eur J Cell Biol 84: 927-938, 2005.
- 32 Hengstschlager M, Braun K, Soucek T, Miloloza A and Hengstschlager-Ottnad E: Cyclin-dependent kinases at the G1-S transition of the mammalian cell cycle. Mutat Res 436: 1-9, 1999.
- 33 Massague J: TGF-beta signal transduction. Annu Rev Biochem 67: 753-791, 1998.

- 34 Muraoka RS, Koh Y, Roebuck LR, Sanders ME, Brantley-Sieders D, Gorska AE, Moses HL and Arteaga CL: Increased malignancy of Neu-induced mammary tumors overexpressing active transforming growth factor beta1. Mol Cell Biol 23: 8691-8703, 2003.
- 35 Siegel PM, Shu W, Cardiff RD, Muller WJ and Massague J: Transforming growth factor beta signaling impairs Neu-induced mammary tumorigenesis while promoting pulmonary metastasis. Proc Natl Acad Sci USA 100: 8430-8435, 2003.
- 36 Batlle E, Sancho E, Franci C, Dominguez D, Monfar M, Baulida J and Garcia De Herreros A: The transcription factor snail is a repressor of E-cadherin gene expression in epithelial tumour cells. Nat Cell Biol 2: 84-89, 2000.
- 37 Comijn J, Berx G, Vermassen P, Verschueren K, van Grunsven L, Bruyneel E, Mareel M, Huylebroeck D and van Roy F: The two-handed E box binding zinc finger protein SIP1 downregulates E-cadherin and induces invasion. Mol Cell 7: 1267-1278, 2001.
- 38 Gumbiner BM: Cell adhesion: the molecular basis of tissue architecture and morphogenesis. Cell 84: 345-357, 1996.
- 39 Takeichi M: Cadherin cell adhesion receptors as a morphogenetic regulator. Science 251: 1451-1455, 1991.
- 40 Miyoshi A, Kitajima Y, Sumi K, Sato K, Hagiwara A, Koga Y and Miyazaki K: Snail and SIP1 increase cancer invasion by upregulating MMP family in hepatocellular carcinoma cells. Br J Cancer 90: 1265-1273, 2004.

Received July 22, 2015 Revised September 14, 2015 Accepted September 17, 2015