

Research article

Open Access

The gene expression profiles of primary and metastatic melanoma yields a transition point of tumor progression and metastasis

Adam I Riker*¹, Steven A Enkemann², Oystein Fodstad¹, Suhu Liu¹, Suping Ren¹, Christopher Morris¹, Yaguang Xi¹, Paul Howell¹, Brandon Metge¹, Rajeev S Samant¹, Lalita A Shevde¹, Wenbin Li³, Steven Eschrich², Adil Daud², Jingfang Ju¹ and Jaime Matta⁴

Address: ¹Mitchell Cancer Institute-University of South Alabama, 315 North University Boulevard, MSB 2015, Mobile, Alabama 36688, USA, ²H. Lee Moffitt Cancer Center and Research Institute, 12902 Magnolia Drive, Tampa, Florida 33612, USA, ³Xuanwu Hospital, 45 Changchu St., Xuanwu District, Beijing, 10053, P.R.C and ⁴Department of Pharmacology, Physiology and Toxicology, Ponce School of Medicine. P.O. Box 7004, Ponce, 00732, Puerto Rico

Email: Adam I Riker* - ariker@usouthal.edu; Steven A Enkemann - steven.enkemann@moffitt.org; Oystein Fodstad - ofodstad@usouthal.edu; Suhu Liu - sliu@usouthal.edu; Suping Ren - sren@usouthal.edu; Christopher Morris - cmorris@usouthal.edu; Yaguang Xi - xi@usouthal.edu; Paul Howell - paulhowell@usouthal.edu; Brandon Metge - bmtege@usouthal.edu; Rajeev S Samant - rsamant@usouthal.edu; Lalita A Shevde - lsamant@usouthal.edu; Wenbin Li - neure55@hotmail.com; Steven Eschrich - steven.eschrich@moffitt.org; Adil Daud - adil.daud@moffitt.org; Jingfang Ju - jjju@usouthal.edu; Jaime Matta - jmatta@psm.edu

* Corresponding author

Published: 28 April 2008

Received: 15 December 2007

BMC Medical Genomics 2008, 1:13 doi:10.1186/1755-8794-1-13

Accepted: 28 April 2008

This article is available from: <http://www.biomedcentral.com/1755-8794/1/13>

© 2008 Riker et al; licensee BioMed Central Ltd.

This is an Open Access article distributed under the terms of the Creative Commons Attribution License (<http://creativecommons.org/licenses/by/2.0>), which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly cited.

Abstract

Background: The process of malignant transformation, progression and metastasis of melanoma is poorly understood. Gene expression profiling of human cancer has allowed for a unique insight into the genes that are involved in these processes. Thus, we have attempted to utilize this approach through the analysis of a series of primary, non-metastatic cutaneous tumors and metastatic melanoma samples.

Methods: We have utilized gene microarray analysis and a variety of molecular techniques to compare 40 metastatic melanoma (MM) samples, composed of 22 bulky, macroscopic (replaced) lymph node metastases, 16 subcutaneous and 2 distant metastases (adrenal and brain), to 42 primary cutaneous cancers, comprised of 16 melanoma, 11 squamous cell, 15 basal cell skin cancers. A Human Genome U133 Plus 2.0 array from Affymetrix, Inc. was utilized for each sample. A variety of statistical software, including the Affymetrix MAS 5.0 analysis software, was utilized to compare primary cancers to metastatic melanomas. Separate analyses were performed to directly compare only primary melanoma to metastatic melanoma samples. The expression levels of putative oncogenes and tumor suppressor genes were analyzed by semi- and real-time quantitative RT-PCR (qPCR) and Western blot analysis was performed on select genes.

Results: We find that primary basal cell carcinomas, squamous cell carcinomas and thin melanomas express dramatically higher levels of many genes, including *SPRR1A/B*, *KRT16/17*, *CD24*, *LOR*, *GATA3*, *MUC15*, and *TMPRSS4*, than metastatic melanoma. In contrast, the metastatic melanomas express higher levels of genes such as *MAGE*, *GPR19*, *BCL2A1*, *MMPI4*, *SOX5*, *BUB1*, *RGS20*, and more. The transition from non-metastatic expression levels to metastatic expression levels occurs as melanoma tumors thicken. We further evaluated primary melanomas of varying Breslow's tumor

thickness to determine that the transition in expression occurs at different thicknesses for different genes suggesting that the "transition zone" represents a critical time for the emergence of the metastatic phenotype. Several putative tumor oncogenes (*SPP-1*, *MITF*, *CITED-1*, *GDF-15*, *c-Met*, *HOX* loci) and suppressor genes (*PITX-1*, *CST-6*, *PDGFRL*, *DSC-3*, *POU2F3*, *CLCA2*, *ST7L*), were identified and validated by quantitative PCR as changing expression during this transition period. These are strong candidates for genes involved in the progression or suppression of the metastatic phenotype.

Conclusion: The gene expression profiling of primary, non-metastatic cutaneous tumors and metastatic melanoma has resulted in the identification of several genes that may be centrally involved in the progression and metastatic potential of melanoma. This has very important implications as we continue to develop an improved understanding of the metastatic process, allowing us to identify specific genes for prognostic markers and possibly for targeted therapeutic approaches.

Background

In the United States, the overall incidence of melanoma is increasing at a rate faster than any other cancer, with recent estimates for the lifetime risk of developing invasive melanoma at 1/49 [1]. Patients diagnosed with metastatic melanoma (AJCC stage IV) have an overall poor prognosis, and 6 out of every 7 deaths associated with all types of skin cancer are caused by metastatic melanoma [1-3]. Whereas patients with thin primary tumors are cured after appropriate surgical intervention, most patients with advanced disease do not respond to available therapies, currently limited to only 2 FDA approved agents to treat stage III and IV melanoma. For most cancers, earlier detection methods have the greatest impact on survival and further contributing to better cure rates, with targeted treatment regimens depending upon an understanding of the molecular profile of a tumor. Similarly, we must improve our understanding of the mechanisms that lead to progressive and metastatic disease, as this will most likely provide the greatest hope for patients in reducing the number of deaths due to metastatic melanoma (MM).

The development of melanoma begins with the malignant transformation of normal human epithelial melanocytes (NHEM) located within the basement membrane of the skin. However, the exact cellular mechanisms that occur and the genes and molecular pathways involved in this process remain obscure. Early work has shown a direct correlation between the thickness of the primary cutaneous melanoma (PCM) and its metastatic capacity, either via the lymphatic system or hematogenously [4,5]. Once melanoma has metastasized by either route, the overall survival for patients greatly diminishes [6,7].

Many investigators have examined the differences between primary melanoma and the late stage metastatic disease. In the last few years, the power of microarray technology has been utilized to survey transcriptional differ-

ences that might provide insight into the metastatic process [8-15]. The earliest reports used a handful of samples to first suggest that MM was characterized by a large number of gene expression differences from cultured melanocytes or benign nevi [8,16]. Since then, many reports have corroborated this notion in principle, although each study has focused on a different aspect of the disease process. Some studies have used cell lines to compare melanocytes to melanoma [8,12], while others have utilized tissue samples to look at melanoma samples relative to benign nevi or normal skin [10,11,15,16]. Several studies have focused on the clinical aspects of the disease, attempting to identify gene expression signatures that correlate with metastasis or survival [9,13,17-19]. Together, these studies suggest that many genes contribute to the signature that is detectable in patients with MM. We wished to further examine the exact timing of when these gene expression patterns arise.

A few studies have begun to suggest that the metastatic expression pattern emerges during the vertical growth phase of primary melanomas. Initially, Smith *et al.* (2005) [10] showed a distinct transition point of gene expression change compared to normal skin, benign nevi, melanoma in situ (MIS), PCM, and MM samples. However, they did not specifically describe the Breslow's tumor thickness of their primary melanoma samples. More recently, Soikkeli *et al.* [19] in studying the draining lymph nodes for melanoma, concluded that many of the "molecular traits" of micrometastases were already present in primary melanoma. The Melanoma Group of the European Organization for Research and Treatment of Cancer (EORTC) has recently reported that melanoma patients with an average tumor thickness of 2.0 mm had a favorable prognosis whereas patients with thicker tumors (5.3 mm) fared worse [13]. Although no comparisons were made to patients with metastatic disease, this study reported a 254-gene classifier that significantly correlated with the metastatic dissemination of cutaneous

melanoma. Similarly, Jaeger *et al.* [14] showed that different primary tumors could be grouped based upon gene expression.

In the current study, we have attempted to identify gene expression patterns that segregate non-metastatic tumors from MM, further examining thin PCM tumors to document the emergence of the metastatic signature. Additionally, we attempt to identify key genes involved in this process, as opposed to those genes that simply document the metastatic profile of a melanoma cell.

Methods

Tissue specimens

Over a 3 year period we surgically procured tumor samples from patients with primary cutaneous melanoma (PCM), squamous cell carcinoma (SCC), basal cell carcinoma (BCC) and metastatic melanoma (MM). All samples were obtained under an Investigational Review Board (IRB) approved tissue procurement protocol (MCC#13448, IRB#101751; PSM# 990914-JM, 020318-JM). Upon surgical removal of the primary melanoma, a single surgical oncologist (A.I.R.) utilized a scalpel to macrodissect and procure a portion of each tumor, careful to avoid central areas of necrosis and surrounding stroma. With bulky melanoma tumors, macrodissection usually results in the isolation of 90–95% pure tumor cells, with little interdigitating stroma or other contaminating tissues.

All samples for this study were confirmed to contain > 95% melanoma cells by a dermatopathologist. All samples were cryopreserved in liquid nitrogen and securely de-identified through a centralized database. We analyzed 40 MM samples, composed of 22 bulky, macroscopic (replaced) lymph node metastases, 16 subcutaneous and 2 solid organ metastases (adrenal and brain), and compared them with 42 primary cutaneous cancers [16 PCM, 11 SCC, 15 BCC]. PCM consisted of 2 melanoma in situ (MIS), 2 thin melanomas (< 1 mm), 3 intermediate-thickness melanomas (1–4 mm), and 9 thick melanomas (> 4 mm). Additionally, we included 4 samples of normal human skin and 1 sample of cultured, intermediate-pigmented, human epithelial melanocytes (NHEM).

RNA isolation, purification and hybridization

A portion of each cryopreserved tissue sample was dissolved in TRIzol® (Invitrogen, Carlsbad, CA), purified according to manufacturer's recommendations, and further purified on RNeasy columns (Qiagen Inc., Valencia, CA). RNA integrity was verified by both gel electrophoresis and the Agilent 2100 Bioanalyzer (Agilent Technologies, Palo Alto, CA). A total of 5 ug of RNA was processed using established Affymetrix protocols for the generation of biotin-labeled cRNA and the hybridization, staining,

and scanning of arrays as outlined in the Affymetrix technical manuals [43,44]. The processed RNA was hybridized to Human Genome U133 Plus 2.0 arrays from Affymetrix, Inc. (Santa Clara, CA), and scanned on an Affymetrix GeneChip® scanner 3000 at 2.5 µm resolution. A more complete description of this process is available in Dobbin *et al.*, 2005 [45]. The tissue samples were processed in three independent groups.

Cell lines and tissue culture

Freshly excised melanoma samples were placed into culture media (RPMI 1640 and 5% FCS) and tissue procurement and expansion of daughter cell lines was established utilizing previously published techniques [46,47]. All cell lines were passaged and expanded in vitro, utilizing RPMI 1640 and 5% FCS. All cell lines were split and passaged < 10 times and characterized by flow cytometry and cytopspin preparations for cellular confirmation of melanoma cell purity (data not shown). The cell lines, MCC77 and MCC80a were derived from primary melanoma samples with TC80b derived from a metastatic lymph node (from the same patient). The cell lines, MCC12A and MCC12F, were derived from 2 different subcutaneous melanoma nodules from the same patient. There were 3 cell lines examined from metastatic samples, MCC66C, MCC72 and MCC89. The NHEM were cultured according to the manufacturer directions (Cambrex BioScience, Walkersville, MD). Several cell lines were obtained from the National Cancer Institute, Surgery Branch (624-Mel, 624.38-Mel and A375). These cell lines are derived from patients with stage IV melanoma and are considered highly aggressive cell lines by all standard measures of analysis.

Real-time and semi-quantitative RT-PCR

First-strand cDNA synthesis was performed using Superscript III RT (Invitrogen). Subsequently, the cDNA was utilized for semi-quantitative PCR utilizing intron-spanning primers and optimized reaction conditions. We normalized each sample with β-Actin as an internal control, comparing each sample with AlphaEase®FC image analysis software (Alpha Innotech, San Leandro, CA), followed by densitometric analysis of the integrated values for each sample. The expression levels of putative oncogenes and tumor suppressor genes were analyzed by real-time quantitative RT-PCR (qPCR) using Assays-on-Demand Gene Expression Assays (Applied Biosystems, Foster City, CA): *SPP1* (*osteopontin*, assay ID Hs00167093_m1), *GDF15* (*growth differentiation factor 15*, assay ID Hs00171132_m1), *PITX1* (*paired-like homeodomain transcription factor 1*, assay ID Hs00267528_m1), *DSC3* (*desmocollin 3*, assay ID Hs00170032_m1), *CST6* (*cystatin E/M*, assay ID Hs00154599), *POU2F3* (*POU domain, class 2, transcription factor 3*, assay ID Hs00205009) and *GAPDH* (assay IDHs99999905_m1) as the internal stand-

ard. Utilizing normal skin as the calibrator, the relative quantitation values of a target template for each sample were expressed as $2^{-\Delta\Delta Ct}$. Briefly, qPCR analysis was performed utilizing 40 ng of total cDNA in a 25 μ l reaction volume (Applied Biosystems). We performed qPCR utilizing established techniques, with all samples performed in triplicate and run on an ABI/PRISM 7500 Sequence Detector System (Applied Biosystems).

Gene microarray analysis and bioinformatics

MAS 5.0 analysis software was used to generate signal values for all probe sets based upon a mean intensity of 500, subsequently exported and iteratively normalized as a whole group to create the final normalization based upon the most stable gene expression measurements across all samples [48,49]. Genes highly expressed in metastatic melanomas but not in PCM, BCC, and SCC, or the converse, were specifically identified using visual inspection, t-tests and Pearson's correlation [50] [see Additional file 1]. An initial set of 2,014 Affymetrix probe sets were identified to discriminate between metastatic tumor samples and non-metastatic tumor samples and these were used for clustering. The normalized probe set values were \log_2 transformed and "mean-centered" across all clustered samples. Hierarchical clustering was then performed using absolute correlation and complete linkage in Eisen's cluster [51]. The complete microarray data is available from the Gene Expression Omnibus website [52].

Significance Analysis of Microarrays (SAM) was performed in order to identify a more extensive list of differentially expressed genes expressed between PCM and MM [53]. The first SAM analysis utilized all of the arrayed samples, with the MM samples opposed to all non-metastatic samples, inclusive of BCC, SCC and normal skin. A second SAM comparison utilized 6 thin PCM samples opposed to 6 MM samples. The median false discovery rate threshold was set at 5% for this comparison, with a final gene list generated by intersecting the two resulting gene lists. Following all microarray analyses, the identified probe sets were annotated based on the sequence of the probes used on the arrays [54].

Western blot analysis

Whole cell extracts from PCM and MM cell lines were prepared by directly lysing cells in SDS sample buffer. Expression of SPP-1 protein was assessed in cell lysate and serum-free conditioned medium. Briefly, 4×10^6 cells were plated in 5% FBS containing medium; 24 hours later, the growth medium was replaced with serum-free medium. The conditioned media and cell lysates were harvested 24 hours later and resolved using a 12.5% SDS-PAGE. Proteins were transferred to a PVDF membrane and probed with the anti-human SPP-1 mouse monoclonal antibody (Sigma, St. Louis, MO) (1:1000) followed by a

secondary antibody conjugated to horseradish peroxidase (Amersham Biosciences, Piscataway, NJ) and detected using chemiluminescence (Santacruz Biotechnology, Santa Cruz, CA). The osteopontin band (SPP-1) was visualized at ~55–65 kDa. Daughter melanoma cell lines derived from the freshly procured melanoma samples (with the exception of A375) were lysed by M-PER[®] Mammalian Protein Extraction Reagent (Pierce, Rockford, IL) and processed according to manufacturer instructions. A total of 15 μ g of protein from each experimental condition were electrophoresed on 10% SDS-PAGE and transferred to nitrocellulose membranes (Bio-Rad, Hercules, CA). Immunostaining was performed with the following primary antibodies: DSC3 (Santa Cruz) 1:200; CLCA2 (Novus Biologicals, Littleton, CO) 1:500; PDGFRL (Novus Biologicals) 1:500; α -tubulin (Cell signaling, Danvers, MA), 1:1000. Immunocomplexes were visualized using an enhanced chemiluminescence (ECL) Western Blotting Substrate (Pierce). The intensity of the bands were scanned with a Fujifilm intelligent dark box II and analyzed with Fujifilm Las-1000 Lite V1.3 software.

Results

Gene expression differences between primary cutaneous cancer and metastatic melanoma

An initial training set of 23 tumors revealed 2,014 Affymetrix probe sets with a greater than 2-fold difference in the average gene expression level between the MM and primary cutaneous cancers. This preliminary list, consisting of 1,141 well characterized and 471 poorly characterized human genes, was quite large, indicating a substantial difference between metastatic tumors and non-metastatic tumors. The expression differences allowed for a relatively robust gene classification of tissue samples into groups of metastatic samples and non-metastatic primary tumors. All tumor samples were clustered based on these gene expression values and individually identified as metastatic or non-metastatic based upon the characteristics of tumor samples in the same cluster.

The initial set of samples comprised a training set for which 22 of 23 samples were correctly partitioned into a cluster containing primary melanoma or the cluster containing MM samples. A single primary melanoma with a Breslow's tumor thickness of 90 mm was misclassified as a MM sample. We then analyzed and classified 2 independent test sets comprised of primary skin cancer (BCC, SCC and PCM) and MM samples, utilizing the same genes. Co-clustering led to the correct identification and classification of 56 of 60 melanoma samples. In general, the misidentified samples were thick primary melanomas classified as MM. Of note, several normal human skin samples were analyzed and found to classify as non-metastatic by their gene expression profiles.

We utilized all of the arrayed samples, exclusive of samples that were misclassified, in order to generate a more comprehensive list of genes that appeared to be differentially expressed between MM and PCM using SAM. This analysis identified 4,343 significantly different probe sets again indicating that malignant melanoma tumors are substantially different from the non-malignant tumors. All defined probe sets had an average difference greater than 2-fold between MM and PCM [see Additional file 2]. This list consisted of 1,667 Affymetrix probe sets that detect 279 poorly defined transcripts, 114 minimally defined genes, and 907 well characterized human genes. From this list, 303 genes were highly expressed in MM compared to 997 genes that were more highly expressed in the non-metastatic primary skin cancers and normal skin.

A subset of the full gene list is shown in Table 1, illustrating the trends in gene expression when MM samples are compared to *all* primary cutaneous cancers (BCC/SCC/PCM), or to only *very early* PCM (MIS/Thin), considered ≤ 1.5 mm in Breslow's thickness for this analysis. We found higher expression levels in MM for several genes implicated or known as melanoma-associated tumor antigens (*MAGE*, *CSAG2*) and some genes previously implicated in melanoma progression (*GDF15*, *MMP14*, *SPP-1*), cell cycle progression (*CDK2*, *TYMS*, *BUB1*) and the prevention of apoptosis (*BIRC5*, *BCL2A1*). Among the 997 genes with reduced expression in MM samples, several are implicated in keratinocyte differentiation and epidermal development, such as *loricrin* (*LOR*), *involucrin* (*IVL*), *keratin-5* (*KRT5*) and *plakophilin* (*PKP1*), suggesting a loss of epidermal characteristics. The expression changes suggest

Table 1: Differential gene expression between metastatic melanoma and non-metastatic cutaneous tumors

Gene symbol	Increase in metastatic melanoma relative		Gene symbol	Decrease in metastatic melanoma relative	
	MIS/Thin	to BCC/SCC/MIS/Thin		MIS/Thin	to BCC/SCC/MIS/Thin
MAGEA3/A6/A12	125/119/57	27/25/29	CALML5	193	228
MAGEA1/A2/A5	24/57/10	13/31/6	DSC1/3	186/64	198/71
CSAG2	76	36	PKP1	166	240
TRIM51	51	35	CLCA2	162	177
GDF15	30	47	DSG1/3	160/73	178/119
GYPC	18	14	LY6D	143	147
SPP1	15	7	SERPINB3/B5/B7	111/199/106	184/227/144
KIFCI	15	3	C19orf33	122	135
RGS20	14	14	FLG	112	112
C1orf90	13	15	KRT5/16/17	49/111/105	62/196/274
BCL2A1	12	12	KLK7/8/10/11	99/23/27/65	112/32/81/83
SOX5	15	8	LOR	95	98
SLC16A4	12	29	LGALS7	84	89
AKT3	11	9	CST6	82	56
PEG10	11	10	TRIM29	79	119
BUB1	14	3	SFN	77	125
RASGRF1	8	12	ASAH3	69	56
MMP14	8	6	GATA3	63	54
SPRED1	6	4	CBLC	60	64
GPR19	6	5	RAB25	59	78
CDK2	6	7	SI00A14/A7/A7LI	27/57/21	44/60/82
HOXA10	3	4	ICEBERG	52	48
HOXB6/7/9	4/5/3	6/7/3	IVL	50	76
HEY1	7	16	ELOVL4	38	34
DUSP4	8	10	CXCL14	36	37
DUSP6	8	6	FOXNI	33	34
CDC45L	7	8	AQP3	29	31
CDC6	9	4	TP73L	29	48
RRM2	6	4	MUC15	25	21
TYMS	4	3	RORA	24	22
BIRC5	4	2	CD24	19	38

The initial statistical analysis addressed the question of whether there were genes differentially expressed (increased or decreased) between primary cancers (BCC/SCC/PCM) and metastatic cancers (metastatic melanoma). Column 1 (MIS/thin) are comprised of only such lesions, while column 2 (BCC/SCC/MIS/Thin) is comprised of all 4 histologic subtypes. All of the annotated genes listed above had a > 2-fold up- or down-regulation in gene expression with the full names of each gene, accession number and gene identification provided in supplementary table 1. Abbreviations: MIS, melanoma-in-situ, BCC, basal cell carcinoma, SCC, squamous cell carcinoma, Thin, thin melanomas < 1.5 mm in Breslow's thickness.

important comparative differences between non-metastatic and metastatic tumors, with genes putatively defining cellular function as it relates to the metastatic process.

A further analysis of the functional classes of genes changed using the gene ontology revealed that 15 genes associated with keratinocyte differentiation and 32 genes involved in epidermis development were down-regulated in the metastatic samples [see Additional file 3]. These losses were complemented by the increased expression of genes involved in several cellular processes, such as DNA repair, protein transport, melanocyte differentiation, muscle development, nervous system development and carbohydrate metabolism. Table 1 further illustrates that the magnitude of change in those genes under-expressed was much greater on average than the level of change in over-expressed genes. Overall, the losses in gene expression are both greater in number and magnitude compared to the gains in gene expression in MM samples.

Identification of a transition point in gene expression between non-metastatic and metastatic tumor samples

Figure 1B displays the comparative gene expression levels of 177 genes from our list across the spectrum of tissue samples examined, revealing a consistent level of expression through all of the presumed "non-metastatic" samples (normal skin, BCC, SCC, MIS, Thin). A marked change in the gene expression levels is seen, beginning with the I.M. thickness PCM (average Breslow's tumor thickness of 2.1 mm), progressively increasing or decreasing to the expression level representative of MM lesions and daughter MM cell lines. All of the thick PCM [average Breslow's thickness 19 mm] exhibited gene expression patterns similar to those of MM samples.

Comparative analysis of gene expression patterns in primary melanomas of different Breslow's thickness

The apparent transition zone for gene expression could represent a critical time period where many tumorigenic events occur simultaneously or may simply reflect the outgrowth of an aggressive and/or metastatic cell phenotype. To address this issue, we evaluated the gene expression levels in PCM of increasing Breslow's thickness and that of MM samples. Table 2 (left columns) reveals the relative change in gene expression for a subset of genes throughout the spectrum of increasingly thick PCM and MM samples.

Several genes, such as the *MAGE* genes, exhibited a steady and consistent increase in gene expression over the entire range of tumor thicknesses. However, we found a single major shift in expression for most genes when thinner primary tumors were directly compared to thicker ones. This was most apparent when comparing I.M. thickness to thick PCM, with the majority of genes showing the great-

est comparative increase in gene expression during this period. Notable exceptions were genes such as *SPP1*, *HOXA10* and *MMP14*, for which the greatest differential increase in expression was at the comparative interface between thin and I.M. thickness tumor samples. Other genes, such as *MMP19*, *CTH*, *PDGFRL*, *C16orf34* and *GPR19*, showed the greatest comparative increase in expression when comparing MIS to thin PCM lesions.

We also found a similar phenomenon for genes with decreased expression in primary tumors relative to more advanced lesions (Table 2, right columns). Here, however, the largest proportion of the gene expression change occurred between thick PCM and MM samples. We found very little expression of *keratin* (6B, 16, 17) and *SPRR1* (A, B) in MM compared to all primary melanomas, including thick lesions. Several genes, such as *TMPRSS4*, *STAR*, *ST7L*, *HAS3*, *FGFR3*, *CASZ1* and *HR*, were found to have gene expression changes at the very earliest stages of tumor thickening. Together, the gene expression patterns do not shift in a coordinated fashion as would be expected as the result of the outgrowth of a clonal aggressive or metastatic cell type. Rather, the data suggests that a series of events occur as PCM tumors thicken that may ultimately influence the expression of different groups of genes. One limitation of this analysis is that only a few tumor samples were evaluated at the intermediate thicknesses, which allows the possibility that some of the observed differences might be tumor specific rather than stage specific.

Comparative transcriptomic analysis of normal human epithelial melanocytes to primary and metastatic melanoma samples

We next compared the gene expression profiles of cultured NHEM to PCM and MM samples (Table 3), acknowledging the inherent limitations associated with the comparisons of cultured cells and freshly procured tumor samples [20]. We found large differences in gene expression when comparing NHEM to early, non-metastatic PCM (MIS/thin lesions only) and to MM samples suggesting that PCMs are already quite different from melanocytes. Genes such as *KRT14*, *GJA1*, *S100A7*, *S100A9*, and *EHF* were higher in most melanomas while genes such as *CITED-1*, *GDF15*, *QPRT*, *OCA2*, *c-MET* and *MME* were more highly expressed in NHEM. Many of the genes that increased (or decreased) in MM relative to thin PCM could be attributed to a shift toward expression levels similar to NHEM. For example, we found a decreased expression of keratinocyte proteins such as loricrin and involucrin. However, most of the melanoma associated antigens, including the *MAGE* genes, *PRAME*, *S100A8*, *TRAG3* and *MMP19*, were more highly expressed in the MM samples than in NHEM.

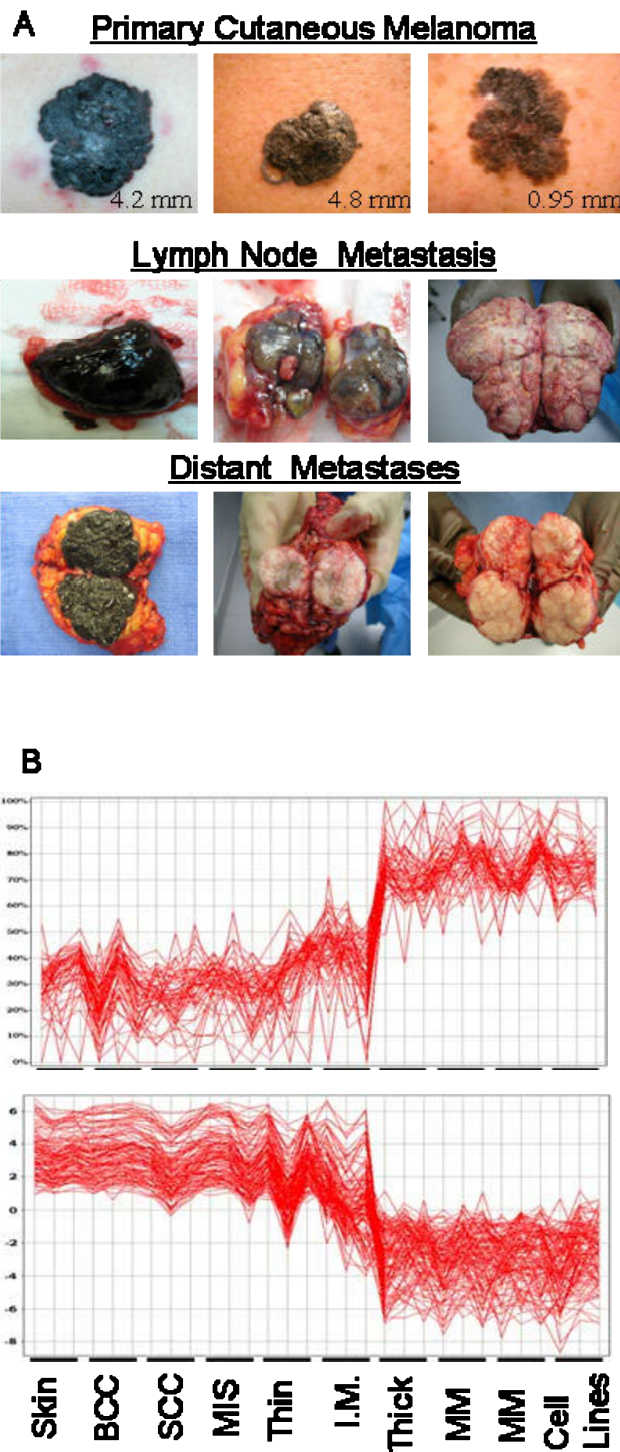


Figure 1

Clinical primary and metastatic melanoma samples procured and cryopreserved at the time of surgery. A: Intraoperative illustrations of the spectrum of PCM and MM samples procured. The PCM represents varying tumor thicknesses, measured utilizing Breslow's depth of invasion. All procured lymph node metastases were macroscopically involved, often completely replacing the entire lymph node parenchyma. Distant metastatic (subcutaneous and solid organ) melanoma often exhibited varying degrees of pigmentation, however, surrounding stroma was avoided in procurement of melanoma samples. B: A distinct change in the gene expression patterns is apparent within the comparative groups of thin/I.M. to thick PCM samples. Gene over-expression (upper graph) is evident at the I.M. thickness sample set, with an average Breslow's tumor thickness of 2.1 mm and 19 mm for thick melanomas. Contrary, there is a decrease in gene expression (lower graph) of the same set of genes, with a comparative difference in gene down-regulation evident at the same interphase of I.M. to thick PCM. Proceeding from left to right: normal skin, BCC, SCC, MIS, I.M., thick primary, metastatic melanoma (subcutaneous, lymph node and distant) and melanoma cell lines derived from patients with stage IV melanoma.

Table 2: Comparative analysis of gene expression changes in primary and metastatic melanoma.

Gene Symbol	Relative increase in gene expression					Gene Symbol	Relative decrease in gene expression				
	MIS to Thin	Thin To IM	IM to Thick	Thick to Met	MIS to Met		MIS to Thin	Thin To IM	IM to Thick	Thick to Met	MIS to Met
MAGEA2	<2	8.6	<u>9.4</u>	3.1	30.7	SPRR1A	<2	<2	<2	<u>129.4</u>	239.3
MAGEA3	<2	8.1	<u>10.7</u>	3	83.8	SPRR1B	<2	<2	<2	<u>52.8</u>	100.8
MAGE A6	<2	7.1	<u>9.8</u>	2.7	103	KRT16	<2	<2	<2	<u>68.3</u>	195.5
MAGEA1,2	<2	2.5	<u>8.9</u>	2.6	81.5	KRT17	<2	2.6	<2	<u>27.4</u>	57.6
MAGE A1	<2	<2	<u>11.1</u>	<2	25.4	KRT6B	<2	<2	<2	<u>39.7</u>	100.9
MAGE A5	<2	2.2	<u>3.9</u>	<2	11.7	AQP3	<2	<2	3.2	<u>6.1</u>	43.2
MMP19	<u>5.4</u>	<2	<2	<2	7.4	CD24	<2	<2	2	<u>4.4</u>	18.7
PDGFRL	<u>22.3</u>	3.6	<2	<2	6.3	FLG	<2	<2	3	<u>24.1</u>	140
C16orf34	<u>5.4</u>	<2	<2	<2	18.5	IVL	<2	2.2	<2	<u>13.9</u>	84.1
CTH	<u>3.9</u>	<2	<2	<2	8.8	KLK7	<2	<2	6.5	<u>8.9</u>	128
GPR19	<u>4.2</u>	<2	3	<2	23.9	LGALS7	<2	<2	3.4	<u>17.3</u>	109.3
SPP1	<2	<u>13.1</u>	<2	<2	44.9	LOR	<2	<2	2.4	<u>22.6</u>	120.4
HOXA10	<2	<u>3.3</u>	<2	<2	3.8	RAB25	<2	<2	3.3	<u>10.3</u>	88.4
MMP14	<2	<u>3.6</u>	<2	2.1	9	SFN	<2	<2	<2	<u>10.6</u>	24.2
AKT3	<2	<2	<u>7.5</u>	<2	14.3	C19orf33	3.1	<2	4.6	<u>13.3</u>	220.3
BCL2A1	<2	2.2	<u>4.8</u>	<2	18.7	ASAH3	<2	<2	<u>25.9</u>	<2	60
BIRC5	<2	<2	<u>3.7</u>	<2	3.4	KRT15	<2	<2	<u>27.1</u>	2.2	104.9
BUB1	<2	<2	<u>9.4</u>	<2	10.8	ELOVL4	<2	2.3	<u>14.7</u>	<2	41.9
CDC45L	<2	<2	<u>13</u>	<2	9.1	GATA3	2.5	<2	<u>14.3</u>	<2	23.5
CDK2	<2	<2	<u>4.8</u>	<2	8.4	MUC15	<2	2.2	<u>11.9</u>	<2	25
CSAG2	<2	<2	<u>19.6</u>	2.6	54.5	SCEL	<2	<2	<u>28.8</u>	<2	71.8
DUSP4	2	<2	<u>5.7</u>	<2	12.5	TP73L	<2	<2	<u>5.5</u>	3.4	41.2
DUSP6	<2	<2	<u>3.7</u>	<2	10.5	RORA	<2	<2	<u>6.2</u>	2.2	26.6
GYPC	<2	<2	<u>12.2</u>	<2	14.2	POU2F3	<2	<2	<u>13.9</u>	2.2	73.9
HEY1	<2	<2	<u>5.7</u>	<2	9.3	ICEBERG	<2	2.7	<u>6.7</u>	4.5	32.3
KIFC1	<2	<2	<u>10.4</u>	<2	16	CASZ1	<2	<u>4.8</u>	2.8	<2	12.3
PEG10	<2	2.4	<u>4.1</u>	<2	11.3	HR	<2	<u>3.7</u>	<2	<2	7.7
RASGRF1	<2	<2	<u>5.1</u>	<2	9.9	TMPRSS4	<u>8.7</u>	<2	<2	3.1	42
RGS20	3.4	<2	<u>9.2</u>	<2	34.3	STAR	<u>4.9</u>	<2	<2	<2	11.1
SLC16A4	<2	<2	<u>4.4</u>	<2	26.4	ST7L	<u>4.3</u>	<2	3.6	<2	8.4
SOX5	<2	<2	<u>12.3</u>	<2	19.8	LTB4R	<u>4</u>	<2	2.3	<2	9.9
TRIM51	<2	3.1	<u>15.9</u>	<2	63.6	HAS3	<u>4.9</u>	<2	<2	2.5	16.6
TYMS	<2	<2	<u>3.6</u>	<2	4.7	FGFR3	<u>3.9</u>	<2	3.2	<2	7.6

NOTE: All annotated genes listed above with a < 2 indicates that any difference between tumors for each comparative analysis was less than 2-fold. Underlined numbers indicate the greatest change in gene expression across varying PCM tumor thickness for each gene. Abbreviations: MIS, melanoma-in-situ, Thin, thin melanomas < 1.0 mm in Breslow's thickness, I.M., intermediate thickness between 1–4 mm, with thick melanomas > 4 mm. PCM, primary cutaneous melanoma.

Identification of putative oncogenes and tumor suppressor genes in melanoma

A perusal of the gene expression differences between PCM and MM samples identifies numerous putative oncogenes and tumor suppressor genes (TSG). Table 4 lists several of the known oncogenes and TSG previously implicated in tumor types. The gene with the largest increase in expression (13.2 fold) was *osteopontin* (*SPP1*). Although not previously identified as an oncogene, *osteopontin* expression has been shown to strongly correlate with melanoma invasion and tumor progression [21]. The lineage-specific oncogene, *MITF*, previously shown to act as a master regulator of melanocyte development and a critical survival oncogene amplified in melanoma, showed a 3.7 fold increase [22-24]. Of the other listed genes, *GDF15*, *c-Met* and the *HOX* loci have been shown to act as possible oncogenes in breast cancer, squamous cell lung cancer, prostate and pancreatic cancer. Several of the putative

melanoma TSGs have also been previously shown to contribute to the development and progression of cancer in other tumor histologies.

The shifts in gene expression occur at different stages of the thickening process for each of the oncogenes and TSGs listed in table 4. Some of the genes show a progressive and steady increase (or decrease) in gene expression as tumors of greater thickness are compared. For other genes, such as *SPP-1*, *GDF15* (putative oncogenes), *PITX-1* and *CST6* (putative TSG), the major shifts in gene expression appear to occur at distinct but different times during the thickening of the primary melanoma tumors. This observation suggests that these changes may occur spontaneously but eventually may accumulate to contribute to the final metastatic phenotype.

Table 3: Comparative transcriptomic analysis of normal human epithelial melanocytes to thin primary cutaneous and metastatic melanoma samples

Fold Higher Expression in Melanoma				Fold Lower Expression in Melanoma			
Gene Symbol	NHEM c/t MIS/Thin	Gene Symbol	NHEM c/t Metastatic Melanoma	Gene Symbol	NHEM c/t MIS/Thin	Gene Symbol	NHEM c/t Metastatic Melanoma
KRT14	6787	GJA1	759	MME	106	MAP4	20
GJA1	5929	SEPP1	338	CITED1	77	OCA2	10
EHF	5487	KRT14	306	GDF15	64	TRIM7	7
SCEL	3931	MAGEA2	301	PAEP	47	CITED1	6
CLCA2	3689	TRAG3	242	RPEL1	45	TRPM4	6
SI00A7	3609	EHF	193	HES6	43	MME	5
KRTDAP	3416	SI00A9	160	ESDN	37	TRAP150	5
DSCI	2782	SI00A7	129	QPRT	35	FER1L3	4
GJB6	2576	SCEL	126	OCA2	19	QPRT	4
CXCL14	2484	SLC22A3	125	RENBP	17	KLF8	4
LOR	2308	EPHA3	124	NR4A3	16	RPEL1	4
KRT6A	1989	KRTDAP	121	Siat7c	16	PACE4	4
PKPI	1835	SI00A8	120	C6orf168	15	HPCAL1	4
SERPINB3	1778	ZIC1	119	BCL2A1	14	ACTRIA	4
SI00A9	1768	CXCL14	118	NTT73	14	MET	3
KRT15	1545	IL18	108	PSCD3	14	RAB32	3
GATA3	1347	PRAME	102	HPCAL1	13	TYR	3
PPL	1339	MAGEA6	94.7	MET	12	IRF6	3
IMUP	1250	PLCB4	88	ALS2CR3	12	GDF15	3
ICEBERG	1123	CLCA2	87	PTPLA	12	HINT3	3
KRT6B	1114	GJB6	85	TBCID7	11	SLC30A1	3
CSTA	693	MAGEA3	84	OAI	11	TRPV2	3
CST6	687	MMP19	59	TYR	11	RSN	3

NOTE: Abbreviations: NHEM: Normal Human Epithelial Melanocytes, c/t: Compared to, MIS, melanoma-in-situ, Thin, thin melanomas < 1.0 mm in Breslow's thickness

Validation of select candidate genes by semi- and quantitative RT-qPCR analysis

To further validate the expression of putative TSG and oncogenes in our melanoma panel, RT-qPCR was performed on 20 previously arrayed samples, comprised of 7 PCM and 13 MM samples. Figure 2A shows an overall decreased level of mRNA expression of TSG and increased mRNA expression of oncogenes compared to normal skin. This was consistent for all PCM samples compared to MM, although not statistically significant when comparing across Breslow's thicknesses. We then re-examined and compared the RT-qPCR data to the microarray data in order to validate the overall degree of agreement, finding a significant correlation for all TSG and oncogenes examined (Figure 2B).

Utilizing semi-quantitative PCR analysis, we also examined several primary and MM daughter cell lines derived from the freshly procured melanoma samples, normal skin and NHEM for oncogene and TSG mRNA expression (Figure 3A). Consistent with the data obtained from microarray analysis, several of the melanoma derived cell lines exhibited high levels of expression for several of the

reported oncogenes and a higher percentage of loss of TSG. Overall, we found a favorable correlation between the microarray results and both quantitative and semi-quantitative PCR analysis for daughter and non-daughter primary and MM cell lines.

Validation of gene expression and protein translation of select candidate genes by Western Blot analysis and immunohistochemistry

To independently verify and validate the gene expression changes at the protein level, we studied several suspected oncogenes and TSG for protein expression by Western Blot analysis. Osteopontin (SPP1) protein expression was examined from melanoma cell lysates and conditioned cell free media derived from 2 primary and 6 MM daughter cell lines (Figure 3B, a, b). Interestingly, the protein expression level of 2 subcutaneous MM nodules (MCC12A, 12F) procured from the same patient greatly differed. Similarly, we examined 2 paired cell lines (MCC80a from a primary melanoma from a synchronous metastatic lymph node, MCC80b), showing a slight increase in SPP1 protein expression in the latter. Several other melanoma cell lines exhibited minimal SPP1 pro-

Table 4: Differential expression of putative tumor oncogenes and suppressor genes in melanoma

Oncogenes				
Gene	Fold Increase	Interval of Increase	Activating Mechanisms in other Tumor Histologies	Affected Tumor Types
SPP-1	13.2	Thin to IM	C-Met activation via $\alpha v\beta 3$ receptor; Inhibition of apoptosis	Breast, HCC, Prostate, CRC, Head & Neck
MITF	3.7	Progressive increase	Somatic alteration via gene amplification (Chr.#3p13-3p14)	None, Lineage Specific for Melanoma
CITED-1 (cbp/p300 transactivator)	12.4	IM to Thick	Activation of Stat-3, Ras/MAPK kinase signaling via Ets1, Ets2	Thyroid
GDF15 (PLAB)	22.7	IM to Thick	Lineage specific activation or repression of ERK1/2; Integrator of AKT pathway	Breast, CRC, Gastric, Prostate, Pancreatic
c-Met	14.5	Thick to Met	Ras-Associated Protein (Rap1)/ERK/MAPK, rac1, Grb2, PI3K, src activation	CRC, Breast, Ovarian, Pancreatic, Liver
HOX Locus (A3, A10, B6, B7, B13)	2.1 – 5.0	Progressive increase	Downstream activation of WT-1, NFKB, NR4A3, BCL2, p53	AML, Breast, SCLC
Tumor Suppressor Genes				
Gene	Fold Decrease	Interval of Decrease	Suppressor Mechanisms in other Tumors Histologies	Affected Tumor Types
PITX-1	13.9	Thin to IM	Ras Pathway (RASAL1)	Barrett's [Esophagus] Prostate, Bladder
CST6 (CST E/M)	66.7	IM to Thick	Hypermethylation	Breast, Glioma
PDGFRL	7.3	IM to Thick	Gene Deletion from Chr.# 8p21.3-p22	HCC, CRC, NSCLC
DSC3	42.8	Progressive decrease	Hypermethylation	Breast
POU2F3	49	Thin to IM	Hypermethylation	Cervical
CLCA2	162	MIS/Thin to MM	Hypermethylation	Breast

Note: Tables represent a partial list of identified tumor oncogenes and tumor suppressor genes (TSG's) in PCM and MM samples. The fold increase represents the greatest fold change noted throughout all comparisons of each PCM tumor thickness to MM. The activating/suppressive mechanism and affected tumor type are also identified. Abbreviations: HCC, hepatocellular carcinoma, CRC, colorectal carcinoma, NSCLC, non-small cell lung cancer, SCLC, squamous cell lung cancer, AML, acute myelogenous leukemia.

tein expression (MCC12F, 66C, 80a and 89), with comparable findings noted between melanoma cell lysates and conditioned cell media.

Analysis of suspected TSG in 3 PCM and 3 MM cell lines revealed a very low level of protein expression of DSC3 in 6/6 cell lines (3c) with 4/6 (2/3 primary and 2/3 MM) cell lines expressing the protein for CLCA2 (3d). Interestingly, we found PDGFRL protein expression in a single primary cell line, with no evidence of expression in any of the metastatic cell lines (3e). Lastly, we chose to utilize existing databases that have previously determined the cellular staining patterns utilizing immunohistochemistry in skin and melanoma samples, available for viewing at the Human Protein Atlas website [55]. This website also highlights those available genes with available immunohistochemical staining patterns of both primary and MM samples [see Additional file 2].

Discussion

In this study, we have examined the gene expression profiles of a number of cutaneous tumors with the intention of highlighting the differences between metastatic tumors and non-metastatic tissues. We have profiled normal skin, melanocytes, BCC, SCC, and early stage melanomas to establish a gene expression pattern associated with the

non-metastatic state. It is well known that BCC and SCC have little if any metastatic potential, and thus adding such specimens substantially contributed to the overall validity of our "non-metastatic" gene signature. Likewise, similar findings are noted for MIS lesions and very early primary melanomas (< 0.75 mm in Breslow's depth). Opposed to this is the metastatic state, which exhibits vastly different expression levels in thousands of genes and is observed in thick PCM and MM tumors. The principal finding of this study is that normal tissues and non-metastatic cutaneous tumors can be distinguished from metastatic melanoma tumors on the basis of the expression level of these genes. We have reported 1,300 transcripts that differ in expression between thin PCM and MM, many of which differ more than 100-fold. This robust signature suggests that gene expression may be useful for identifying metastatic potential in PCM tumors.

A number of previous studies have attempted to use gene expression signatures to identify the genes associated with the establishment or identification of metastatic tumors. Using cell lines it has been shown that normal melanocytes differ greatly from tumor derived cells [8,12]. Benign nevi have been shown to differ from thick or metastatic melanoma tumors [10,11]. Also, several recent studies have shown that MM or thicker melanoma tumors con-

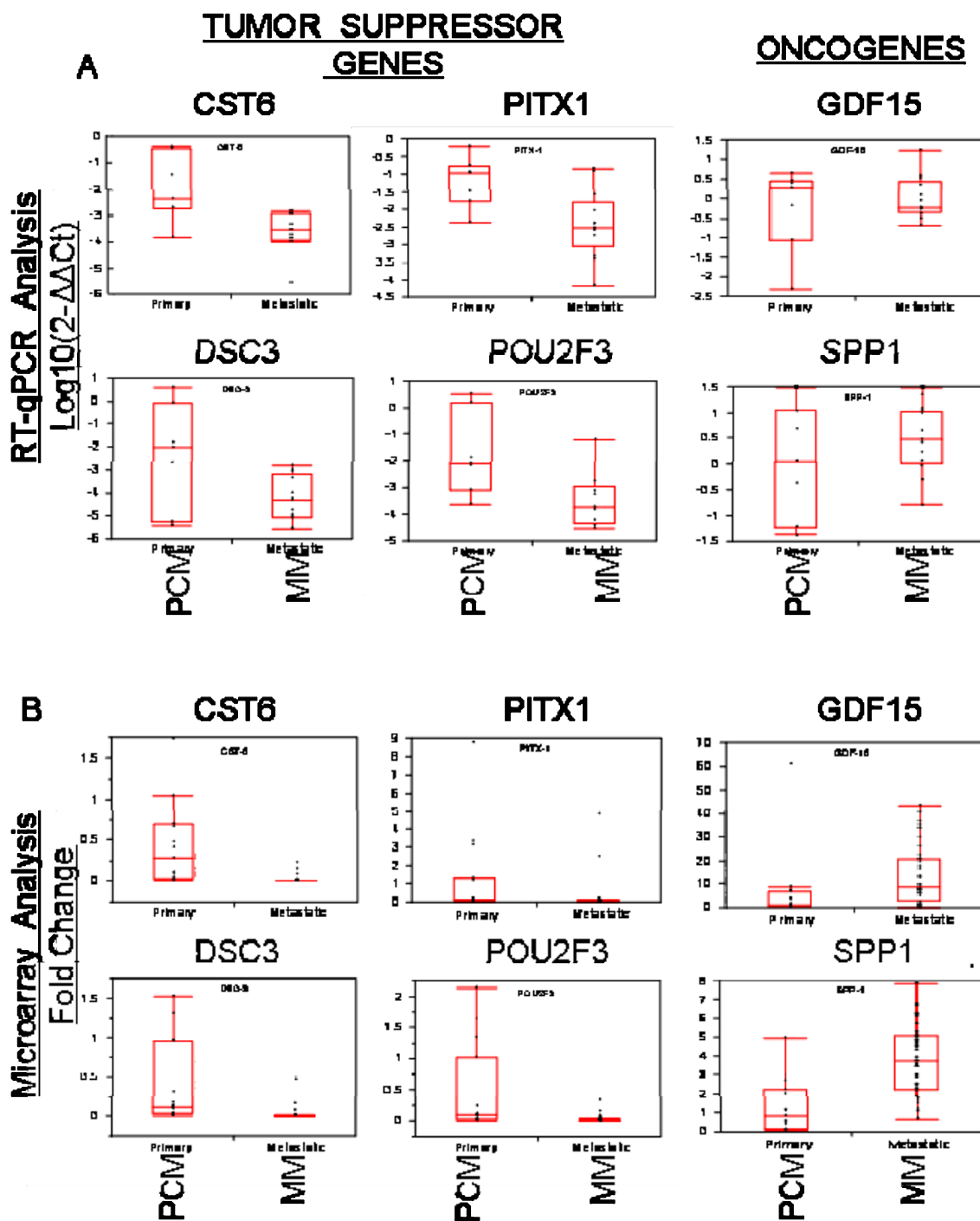


Figure 2

Comparative gene expression analysis of putative oncogenes and tumor suppressor genes. A: Comparative gene expression analysis utilizing qPCR data of putative tumor suppressor genes (CST6, DSC3, PITX1, POU2F3) and oncogenes (GDF15, SPP-1) in PCM (n = 7) and MM (n = 13) samples. Relative quantitation of target gene expression for each sample was determined using the equation $2^{-\Delta\Delta C_t}$, where GAPDH was used as the internal reference and normal skin as the calibrator. Values were Log base 10 transformed (y-axis) so that all values below zero represent a down-regulation in gene expression and values above zero represent an up-regulation in gene expression, compared to normal skin. B: Correlative microarray analysis of gene expression levels in primary and metastatic melanoma samples compared with normal skin. The statistical differences of gene expression between primary (PCM) and metastatic melanoma (MM) samples were analyzed by Wilcoxon's signed rank test; two-tailed significance level was set at $\alpha = 0.05$. Compared to PCM samples (n = 7), the expression levels of 4 putative tumor suppressor genes (CST6, $p < 0.0001$; DSC3, $p < 0.0001$; PITX1, $p = 0.0043$; POU2F3, $p < 0.0001$) were significantly decreased in MM samples (n = 40), while the expression of putative oncogenes (GDF15, $p = 0.0027$; SPP1, $p < 0.0001$) were significantly increased in MM samples.

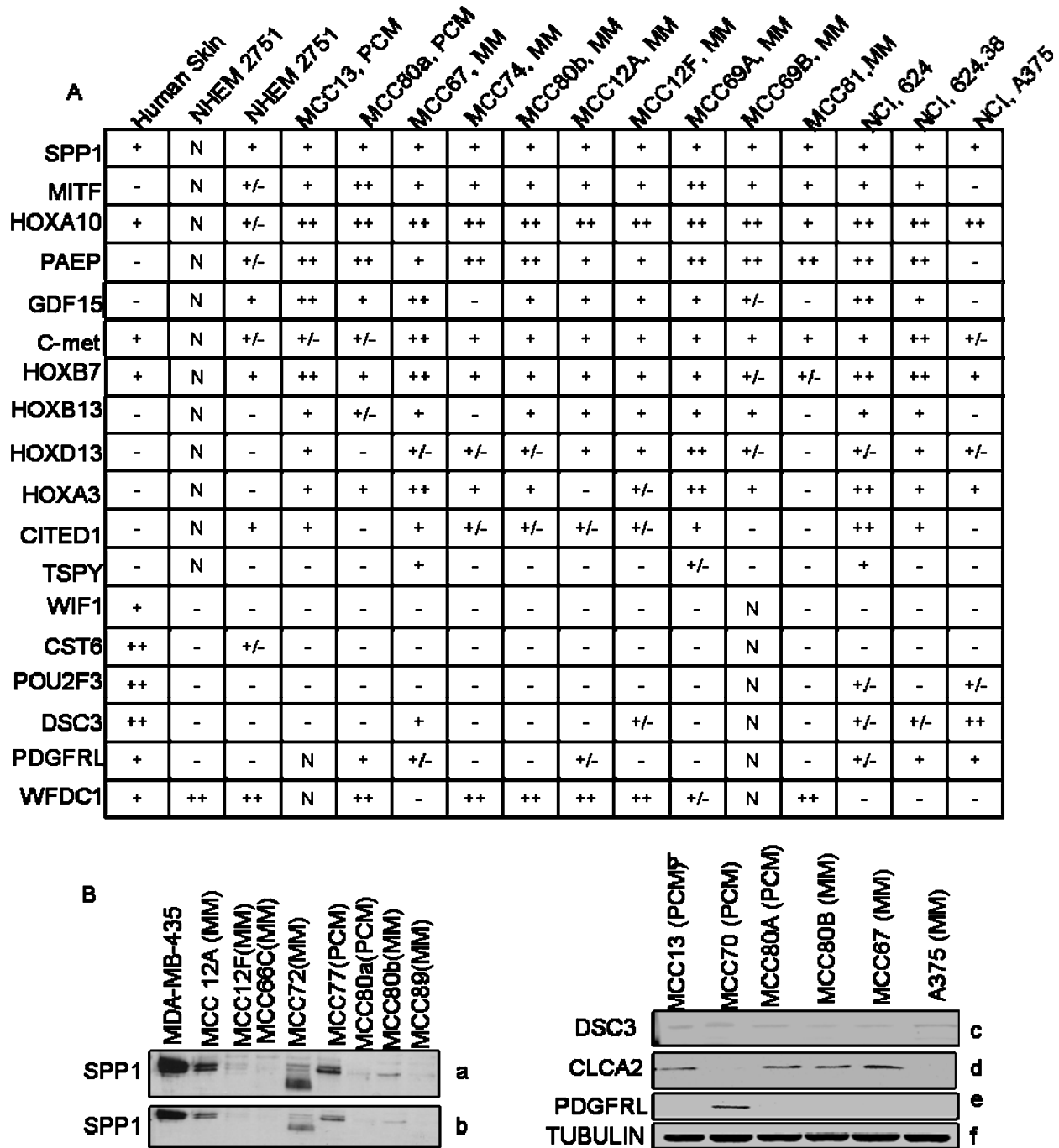


Figure 3
Semi-quantitative RT-PCR analysis of oncogene and tumor suppressor genes. A: A panel of 12 putative oncogenes and 6 TSG were analyzed, with the relative levels of mRNA expression as follows: negative band (-), faintly visible band (+/-), visible band (+), strongly visible band (++), N = Not Done. β -actin served as the internal comparative control. The grey values of PCR products of each gene are analyzed by the AlphaEase 3 software and standardized according to β -actin in every sample. B: a, b, Daughter melanoma cell lines secrete SPP1 (osteopontin). The melanoma cell lysates and conditioned cell-free media was resolved by SDS-PAGE and transferred to a PVDF membrane. The blot was probed with anti-SPP1 antibody. c-f, Antibodies for DSC-3 (c), CLCA2 (d), PDGFRL (e) and α -tubulin (f) as an internal control. Lanes 1–3 are PCM and lanes 4–6 are MM cell lines.

tain one or more distinctive signatures that may indicate the more aggressive phenotype with respect to clinical outcome [9,13,14,17-19]. Although each of these studies was unique in the samples they used and the gene expression signatures they report, we believe there is a very high degree of similarity between all of the "signatures".

The recent report by Jaeger *et al.* coincides very well with the current study [14]. Although they compared tumors to normal skin they utilized U133A arrays from Affymetrix, which contain a subset of the same probes found on the U133 Plus 2.0 arrays used in the present study. We compared the common probe sets of both arrays, finding a 65% overlap between the two gene lists. This is an exceedingly high level of concordance between different microarray studies utilizing tumor samples; probably reflecting the fact that the two studies used nearly identical microarray platforms.

There remains considerable debate in the literature as to why comparative microarray studies do not completely coincide with each other. We believe that the primary reason for such discrepancies is that strict p-values and other restrictive criteria are often utilized to generate gene lists. If we relax our criteria only slightly to generate a larger list than reported in supplementary table 1, we find a stunning 90% concordance with the study of Jaeger *et al.* The remaining differences between the two studies appear to be a few differences that exist between normal skin and metastatic melanoma rather than between primary melanoma and metastatic melanoma and a few genes that exist within a small subset of melanoma tumors and therefore most likely reflect subtle differences in the spectrum of tumors analyzed by the two studies (based on our data).

In all of the recent gene expression studies an attempt has been made to compare normal, benign, or non-metastatic cells or tissues to clinically aggressive tumors. In most of these studies, it is not clear where the shift in gene expression occurs. However, in the current study, we used a larger number and variety of non-metastatic tissues to illustrate the fact that the metastatic signature is robustly different from all the non-metastatic conditions that one might use for comparison. We have also included a documented gradation of tumor thicknesses to better resolve thin primary melanomas from thick primary melanomas which has helped to illustrate that the metastatic signature emerges as primary melanoma tumors thicken.

We have approximated what might occur in an individual tumor by grouping the melanoma samples into subgroups from thin through thick and proceeding on to metastatic tumors (Figure 1B). This illustration follows the analysis of Smith *et al.* [10] who compared normal skin,

benign nevi, melanoma in situ (MIS) and MM samples to show a distinct transition point of gene expression associated with the vertical growth phase of melanoma tumors. However, that study did not specifically describe the Breslow's tumor thickness of their primary melanoma samples leaving open the question of what was metastatic and what was a non-metastatic tumor sample. We believe that our data resolves this issue, showing that MIS and thin melanomas do not have much, if any, of the gene expression measures associated with metastasis. The metastatic signature emerges when primary melanomas begin to thicken. This data suggests that thickening tumors is the best arena for further evaluating the cellular changes leading to metastasis.

Thin melanomas are the most difficult samples to acquire in a research setting where RNA integrity can be preserved and thus our analysis is limited by the few samples of thin melanoma we were able to collect during this study. Thus, it is probable that we have not identified the optimal progression in the transition from non-metastatic gene expression to metastatic gene expression. However, by looking more closely at the various thicknesses of tumors, we observed that the gene expression changes do not occur synchronously. Certain changes occur early and others appear later in the progression from thin to thick tumors (Table 2). Each change brings the expression pattern of primary melanoma tumors closer to the expression pattern observed in metastatic tumors. This suggests that the phenotype of tumors becomes more metastatic-like as the tumor gets thicker and argues against the outgrowth of a cell with the full metastatic signature.

Our data also indicates that many phenotypic changes may be occurring during this thickening period. Our gene ontology analysis suggests a fundamental shift in the functional properties of the cells comprising a tumor. There is a reduced expression or loss of genes involved in the processes of keratinocyte differentiation, epidermal development, cell adhesion and cell-to-cell signaling. This loss of cell-stromal interactions may reflect the gain of migratory potential for the metastatic cell type. This is opposed by functional gains associated with the increased expression of genes involved in melanocyte differentiation, nervous system development, protein transport, carbohydrate metabolism and DNA repair. The unidirectional shift in these classes of genes, the number of genes involved, and the extent to which each gene changed (fold-changes of 30 or more) all suggest a developmental change, rather than a regulation of cellular metabolism. Further supporting such a fundamental change is the observation by Alonso *et al.* that an epithelial-mesenchymal transition occurs as the metastatic signature emerges [18].

The thickening of PCM is also when the increased expression of putative proto-oncogenes and decreased expression of putative TSGs occurs (Table 3). Although most of these oncogenes and TSGs have not been demonstrated to function in melanoma progression, they represent key factors to consider when understanding the emergence of the metastatic phenotype. *Growth differentiation factor-15* (*PLAB\MIC-1\GDF-15*) shows a striking correlation between expression level and metastatic phenotype in several tumor types [25,26]. Osteopontin (*SPP-1*) is a secreted phosphoglycoprotein that has been implicated in tumor progression and invasive behavior in many tumor types [27]. *SPP-1* expression has been strongly correlated with invasive melanoma but found highly expressed in only 72% of invasive primary melanomas [21]. Our data shows that *SPP-1* gene expression increases when thin primary melanomas thicken (Table 2) and that *SPP-1* is expressed in daughter melanoma cell lines and secreted as a soluble protein (Figure 2). Other identified genes, such as *Cbp/p300-interacting transactivator (CITED-1)*, *hepatocyte growth factor receptor (c-MET)*, and various homeobox genes have yet to be investigated in melanoma although some have been previously identified in metastatic melanoma [11].

The TSGs listed in table 3 have been implicated in the evolution of the metastatic state via down-regulation and inactivation. *PITX-1* has been identified as a TSG in several tumor types, including lung cancer and Barrett's esophagus leading to esophageal adenocarcinoma [28-30]. Loss of *PITX-1* expression is seen in thin primary melanoma samples, with a 14-fold decrease in gene expression in thin primary compared to I.M. samples. Many of the other TSGs listed in table 3 have been down-regulated by epigenetic silencing in other tumor histologies [31-38]. Recently, Muthusamy *et al.* described the epigenetic silencing of novel tumor suppressors in melanoma samples, identifying 17 genes not previously known to be hypermethylated. [39]. We have found 2 genes that are in common with this list. We are actively examining the identified TSGs for evidence of epigenetic silencing via hypermethylation of promoter CpG islands.

We also found several known melanoma tumor antigens (*MAGE*, *TRAG3*, *PRAME*) highly expressed in thicker tumors. This is consistent with studies that have shown an increase in tumor antigen expression with advanced disease [8,40-42]. Some of these antigens were seen in nearly all tumor samples while others were only seen in a subset of the melanoma tumors. Nonetheless, these surface antigens provide a mechanism for identifying the metastatic cell arising in early melanoma tumors.

Based upon their distinct gene expression profiles, the four classes of tumors could be correctly classified greater

than 90% of the time. The major difficulty in classification was for thick primary melanoma tumors that appeared to have the gene expression signature of MM tumors. The difficulty in classifying these tumors suggested that they might represent a transitional state between primary tumors and truly metastatic tumors. If we accept this assumption, then 81 of our 82 tumor samples were correctly classified based upon their gene expression signature. With the ability to correctly and accurately classify tumors into distinct classes, we can begin to investigate the mechanisms responsible for malignant tumor formation, invasion, progression and metastasis.

Conclusion

Based upon our molecular analysis of primary cutaneous (BCC, SCC, PCM) and MM samples, we conclude that PCM and MM have distinct gene expression profiles that may be useful for tumor tissue classification. Additionally, squamous cell and basal cell carcinomas share certain gene expression patterns with primary melanomas that are distinct from MM. This is important to try and develop prognostic markers for patients based upon the distinct molecular profile of their melanoma, further able to distinguish between a metastatic versus non-metastatic gene profile. This will allow for the physician and patient to have a more informed discussion as to how to proceed with clinical treatment options based upon the gene expression profile.

Authors' contributions

AIR is responsible for the initial conception, overall hypothesis and design of this manuscript, including the original draft and all subsequent revisions. AIR is also responsible for the procurement and cryopreservation of melanoma tissue specimens. SAE performed the molecular studies with gene microarray analysis as well as the bioinformatics data derived from all samples. SAE also assisted with the draft of this manuscript. JFJ and OF assisted with the draft of this manuscript and the numerous revisions and formatting for the final version. SL, SR, PH and BM all assisted with outstanding technical assistance, including the PCR analysis, western blotting experiments and performing all other confirmatory and functional data associated with this manuscript. All were also involved in the drafting and revisions for this manuscript. CM assisted with the statistical interpretations of the functional data. RSS and LAS both assisted with the Western blotting and cell analysis of *SPP* expression and were involved with the draft and revisions of this manuscript. WL was instrumental in the *in vitro* expansion of the melanoma cell lines and associated experiments with all daughter cell lines. JFJ and YX assisted with the semi- and quantitative PCR experiments and provided excellent guidance as to quality control for many of the experiments. SE provided the SAM analysis. AD provided some

of the initial concepts and design for all subsequent experiments and was involved in the draft and revisions for this manuscript. JM assisted with the initial concept and design of the experiments and was also involved with the draft and revisions for this manuscript. All authors read and approved the final manuscript.

Additional material

Additional file 1

Detailed Gene Microarray Analysis and Bioinformatics. This document provides a more detailed and in-depth description of the statistical and biostatistical process involved in the supervised analysis of the generated data. We have also added a description of patient demographics.

Click here for file

[<http://www.biomedcentral.com/content/supplementary/1755-8794-1-13-S1.doc>]

Additional file 2

Final Supervised Gene List. The data in this file describes the following: Supervised Gene list, symbol, GenBank reference ID #, Entrez Med ID #, UniGene ID #, Human Protein Atlas Tissue Staining Link for each gene.

Click here for file

[<http://www.biomedcentral.com/content/supplementary/1755-8794-1-13-S2.xls>]

Additional file 3

Gene Ontology and Biological Processes. This file shows the ontological comparisons of various numbers of genes associated with each biological process between primary and metastatic melanoma.

Click here for file

[<http://www.biomedcentral.com/content/supplementary/1755-8794-1-13-S3.pdf>]

Acknowledgements

Grant support: National Functional Genomics Center at the H. Lee Moffitt Cancer and Research Institute [NCRR RCMI-NIH Program Grant 2-G12-RR03050, AIR] and from the Abraham Mitchell Clinical Research Scholarship at the University of South Alabama, Mitchell Cancer Institute [AIR]. Jaime Villa M.D. is acknowledged for the procurement of the non-melanoma skin cancer samples. Richard Jove Ph.D., and Ralf Beuttner Ph.D. for their expert technical expertise and assistance with this manuscript.

References

- Jemal A, Siegel R, Ward E, Murray T, Xu J, Thun MJ: **Cancer Statistics, 2007**. *CA Cancer J Clin* 2007, **57(1)**:43-66.
- Balch CM, Buzaid AC, Atkins MB: **A new American Joint Committee on Cancer staging system for cutaneous melanoma**. *Cancer* 2000, **88(6)**:1484-1491.
- Eton O, Legha SS, Moon TE, Buzaid AC, Papadopoulos NE, Plager C, Burgess AM, Beidikian AY, Ring S, Dong O, Glassman AB, Balch CM, Benjamin RS: **Prognostic factors for survival of patients treated systemically for disseminated melanoma**. *J Clin Oncol* 1998, **16(3)**:1103-1111.
- Haddad FF, Stall A, Messina J, Brobeil A, Ramnath E, Glass LF, Cruse CW, Berman CG, Reintgen DS: **The progression of melanoma nodal metastasis is dependent on the tumor thickness of the primary lesion**. *Ann Surg Oncol* 1999, **6(2)**:144-149.
- Cascinelli N, Belli F, Santinami M, Fait V, Testori A, Ruka W, Cavaliere R, Mozzillo N, Rossi CR, MacKie RM, Nieweg O, Pace M, Kirov K: **Sentinel lymph node biopsy in cutaneous melanoma: The WHO Melanoma Program experience**. *Ann Surg Oncol* 2000, **7(6)**:469-474.
- Balch CM, Buzaid AC, Soong SJ, Atkins MB, Cascinelli N, Coit DG, Fleming ID, Gershenwald JE, Houghton A, Kirkwood JM, McMasters KM, Mihm MF, Morton DL, Reintgen DS, Ross MI, Sober A, Thompson JA, Thompson JF: **Final version of the American Joint Committee on Cancer staging system for cutaneous melanoma**. *J Clin Oncol* 2001, **19(16)**:3635-3648.
- Balch CM, Soong SJ, Gershenwald JE, Thompson JF, Reintgen DS, Cascinelli N, Urist M, McMasters KM, Ross MI, Kirkwood JM, Atkins MB, Thompson JA, Coit DG, Byrd D, Desmond R, Zhang Y, Liu PY, Lyman GH, Morabito A: **Prognostic factors analysis of 17,600 melanoma patients: Validation of the American Joint Committee on Cancer melanoma staging system**. *J Clin Oncol* 2001, **19(16)**:3622-3634.
- Hoek K, Rimm DL, Williams KR, Zhao H, Ariyan S, Lin A, Kluger HM, Berger AJ, Cheng E, Trombetta ES, Wu T, Niinobe M, Yoshikawa K, Hannigan GE, Halaban R: **Expression profiling reveals novel pathways in the transformation of melanocytes to melanomas**. *Cancer Res* 2004, **64(15)**:5270-5282.
- Haqq C, Nosrati M, Sudilovsky D, Crothers J, Khodabakhsh D, Pulliam BL, Federman S, Miller JR, Allen RE, Singer MI, Leong SP, Ljung BM, Sagebiel RW, Kashani-Sabet M: **The gene expression signatures of melanoma progression**. *Proc Natl Acad Sci USA* 2005, **102(17)**:6092-6097.
- Smith AP, Hoek K, Becker D: **Whole-genome expression profiling of the melanoma progression pathway reveals marked molecular differences between nevi/melanoma in situ and advanced-stage melanomas**. *Cancer Biol Ther* 2005, **4(9)**:1018-1029.
- Talantov D, Mazumder A, Yu JX, Briggs T, Jiang Y, Backus J, Atkins D, Wang Y: **Novel genes associated with malignant melanoma but not benign melanocytic lesions**. *Clin Can Res* 2005, **11(20)**:7234-7242.
- Mischiati C, Natali PG, Sereni A, Sibilio L, Giorda E, Cappellacci S, Nicotra MR, Mariani G, Di Filippo F, Catricala C, Gambari R, Grammatico P, Giacomini P: **cDNA-array profiling of melanomas and paired melanocyte cultures**. *J Cell Physiol* 2006, **207(3)**:697-705.
- Winnepenninckx V, Lazar V, Michiels S, Dessen P, Stas M, Alonso SR, Avril M, Romero PLO, Robert T, Balacescu O, Eggemont AMM, Lenoir G, Sarasin A, Tursz T, Oord JJ van den, Spatz A: **Melanoma Group of the European Organization for Research and Treatment of Cancer: Gene expression profiling of primary cutaneous melanoma and clinical outcome**. *J Natl Cancer Inst* 2006, **98(7)**:472-482.
- Jaeger J, Koczan D, Thiesen HJ, Ibrahim SM, Gross G, Spang R, Kunz M: **Gene expression signatures for tumor progression, tumor subtype, and tumor thickness in laser-microdissected melanoma tumors**. *Clin Can Res* 2007, **13(3)**:806-815.
- Jensen E, Lewis J, McLoughlin J, Alvarado M, Daud A, Messina J, Enkemann SA, Yeatman TJ, Sondak V, Riker AI: **Down-regulation of pro-apoptotic genes is an early event in the progression of malignant melanoma**. *Ann Surg Oncol* 2007, **14(4)**:1416-1423.
- Becker B, Roesch A, Hafner C, Stolz W, Dugas M, Landthaler M, Vogt T: **Discrimination of melanocytic tumors by cDNA array hybridization of tissues prepared by laser pressure catapulting**. *J Invest Dermatol* 2004, **122(2)**:361-368.
- Mandruzzato S, Callegaro A, Turcatel G, Francescato S, Montesco MC, Chiarion-Sileni V, Mocellin S, Rossi CR, Biccato S, Wang E, Marincola FM, Zanovello P: **A gene expression signature associated with survival in metastatic melanoma**. *J Transl Med* 2006, **4**:50.
- Alonso SR, Tracey L, Ortiz P, Perez-Gomez B, Palacios J, Pollan M, Linares J, Serrano S, Saez-Castillo AI, Sanchez L, Pajares R, Sanchez-Aguilera A, Artiga MJ, Piris MA, Rodriguez-Peralto JL: **A high-throughput study in melanoma identifies epithelial-mesenchymal transition as a major determinant of metastasis**. *Cancer Res* 2007, **67(7)**:3450-3460.
- Soikkeli J, Lukk M, Nummela P, Virolainen S, Jahkola T, Katainen R, Harju L, Ukkonen E, Saksela O, Holttä E: **Systematic search for the best gene expression markers for melanoma micrometastasis detection**. *J Pathol* 2007, **213(2)**:180-189.
- Vogl A, Sartorius U, Vogt T, Roesch A, Landthaler M, Stolz W, Becker B: **Gene expression profile changes and their daughter cell lines: Implication for vaccination trials**. *J Invest Dermatol* 2005, **124(2)**:401-404.

21. Zhou Y, Dai DL, Martinka M, Mingwan S, Zhang Y, Campos EI, Dorocicz I, Tang L, Huntsman D, Nelson C, Ho V, Li G: **Osteopontin expression correlates with melanoma invasion.** *J Invest Dermatol* 2005, **124(5)**:1044-1052.
22. Garraway LA, Widlund HR, Rubin MA, Getz G, Berger AJ, Ramaswamy S, Beroukhi R, Milner DA, Granter SR, Du J, Lee C, Wagner SN, Li C, Golub TR, Rimm DL, Meyerson ML, Fisher DE, Sellers WR: **Integrative genomic analyses identify MITF as a lineage survival oncogene amplified in malignant melanoma.** *Nature* 2005, **436(7047)**:117-122.
23. Levy C, Khaled M, Fisher DE: **MITF: master regulator of melanocyte development and melanoma oncogene.** *Trends Mol Med* 2006, **12(9)**:406-414.
24. McGill GG, Haq R, Nishimura EK, Fisher DE: **C-Met expression is regulated by Mitf in the melanocyte lineage.** *J Biol Chem* 2006, **281(15)**:10365-10373.
25. Bauskin AR, Brown DA, Kuffner T, Johnen H, Luo XW, Hunter M, Breit SN: **Role of macrophage inhibitory cytokine-1 in tumorigenesis and diagnosis of cancer.** *Cancer Res* 2006, **66(10)**:4983-4986.
26. Wollmann W, Goodman ML, Bhat-Nakshatri P, Koshimoto H, Goulet RJ, Meharotra S, Morimiya A, Badve S, Nakshatri H: **The macrophage inhibitory cytokine integrates AKT/PKB and MAP kinase signaling pathways in breast cancer cells.** *Carcinogenesis* 2005, **26(5)**:900-907.
27. Coppola D, Szabo M, Boulware D, Muraca P, Alsarraj M, Chambers AF, Yeatman TJ: **Correlation of osteopontin protein expression and pathological stage across a wide variety of tumor histologies.** *Clin Can Res* 2004, **10(1)**:184-190.
28. Chen Y, Knosel T, Ye F, Pacyna-Gengelbach M, Deutschmann N, Petersen I: **Decreased PITX1 homeobox gene expression in human lung cancer.** *Lung Cancer* 2007, **55(3)**:287-294.
29. Kolschoten IG, van Leeuwen B, Berns K, Mullenders J, Beijersbergen RL, Bernards R, Voorhoeve PM, Agami R: **A genetic screen identifies PITX1 as a suppressor of RAS activity and tumorigenicity.** *Cell* 2005, **121(6)**:849-858.
30. Lord RV, Brabender J, Wickramasinghe K, DeMeester SR, Holscher A, Schneider PM, Danenberg PV, DeMeester TR: **Increased CDX2 and decreased PITX1 homeobox gene expression in Barrett's esophagus and Barrett's-associated adenocarcinoma.** *Surgery* 2005, **138(5)**:924-931.
31. Ai L, Kim WJ, Kim TY, Fields CR, Massoll NA, Robertson KD, Brown KD: **Epigenetic silencing of the tumor suppressor cystatin M occurs during breast cancer progression.** *Cancer Res* 2006, **66(16)**:7899-7909.
32. An Q, Liu Y, Gao Y, Huang J, Fong X, Liu L, Zhang D, Zhang J, Cheng S: **Deletion of tumor suppressor genes in Chinese non-small cell lung cancer.** *Cancer Lett* 2002, **184(2)**:189-195.
33. Fujiwara Y, Ohata H, Kuroki T, Koyama K, Tsuchiya E, Monden M, Nakamura Y: **Isolation of a candidate tumor suppressor gene on chromosome 8p21.3-p22 that is homologous to an extracellular domain of the PDGF receptor beta gene.** *Oncogene* 1995, **10(5)**:891-895.
34. Kim TY, Zhong S, Fields CR, Kim JH, Robertson KD: **Epigenomic profiling reveals novel and frequent targets of aberrant DNA methylation-mediated silencing in malignant glioma.** *Cancer Res* 2006, **66(15)**:7490-7501.
35. Lerebours F, Olschwang S, Thuille B, Schmitz A, Fouchet P, Buecher B, Martinet N, Galateau F, Thomas G: **Fine deletion mapping of chromosome 8p in non-small-cell lung carcinoma.** *Int J Cancer* 1999, **81(6)**:854-858.
36. Rivenbark AG, Jones WD, Coleman WB: **DNA methylation-dependent silencing of CST6 in human breast cancer cell lines.** *Lab Invest* 2006, **86(12)**:1233-1242.
37. Seitz S, Korsching E, Weimer J, Jacobsen A, Arnold N, Meindl A, Arnold W, Gustavus D, Klebig C, Petersen I, Scherneck S: **Genetic background of different cancer cell lines influences the gene set involved in chromosome 8 mediated breast tumor suppression.** *Genes Chromosomes Cancer* 2006, **45(6)**:612-627.
38. Zhang Z, Huettner PC, Nguyen L, Bidder M, Funk MC, Li J, Rader JS: **Aberrant promoter methylation and silencing of the POU2F3 gene in cervical cancer.** *Oncogene* 2006, **25(39)**:5436-5445.
39. Muthusamy V, Duraisamy S, Bradbury CM, Hobbs C, Curley DP, Nelson B, Bosenberg M: **Epigenetic silencing of novel tumor suppressors in malignant melanoma.** *Cancer Res* 2006, **66(23)**:11187-11193.
40. Barrow C, Browning J, MacGregor D, Davis ID, Sturrock S, Jungbluth AA, Cebon J: **Tumor antigen expression in melanoma varies according to antigen and stage.** *Clin Can Res* 2006, **12(3)**:764-771.
41. Brasseur F, Rimoldi D, Liénard D, Lethé B, Carrel S, Arienti F, Suter L, Vanwijck R, Bourlond A, Humblet Y: **Expression of MAGE genes in primary and metastatic cutaneous melanoma.** *Int J Cancer* 1995, **63(3)**:375-380.
42. Hodi FS: **Well-defined melanoma antigens as progression markers for melanoma: Insights into differential expression and host response based on stage.** *Clin Can Res* 2006, **12(3)**:673-678.
43. Van Gelder RN, von Zastrow ME, Yool A, Dement WC, Barchas JD, Eberwine JH: **Amplified RNA synthesized from limited quantities of heterogeneous cDNA.** *Proc Natl Acad Sci USA* 1990, **87(5)**:1663-1667.
44. Warrington JA, Nair A, Mahadevappa M, Tsyganskaya M: **Comparison of human adult and fetal expression and identification of 535 housekeeping/maintenance genes.** *Physiol Genomics* 2000, **2(3)**:143-147.
45. Dobbin KK, Beer DG, Meyerson M, Yeatman TJ, Gerald WL, Jacobson JW, Conley B, Buetow KH, Heiskanen M, Simon RM, Minna JD, Girard L, Misek DE, Taylor JM, Hanash S, Naoki K, Hayes DN, Ladd-Acosta C, Enkemann SA, Viale A, Giordano TJ: **Interlaboratory comparability study of cancer gene expression analysis using oligonucleotide microarrays.** *Clin Can Res* 2005, **11(2)**:565-572.
46. Riker AI: **The isolation and culture of melanoma cell lines.** In *Cancer cell culture: Methods and protocols* Edited by: Langdon S. Totowa: Human Press; 2004:93-100.
47. Riker AI, Panelli MC, Kammula US, Wang E, Wunderlich J, Abati A, Fetsch P, Rosenberg SA, Marincola FM: **Development and characterization of melanoma cell lines established by fine-needle aspiration biopsy: advances in the monitoring of patients with metastatic melanoma.** *Cancer Detect Prev* 1999, **23(5)**:387-396.
48. Li C, Wong WH: **Model-based analysis of oligonucleotide arrays: Expression index computation and outlier detection.** *Proc Natl Acad Sci USA* 2001, **98(1)**:31-36.
49. Wang Y, Lu J, Lee R, Gu Z, Clarke R: **Iterative normalization of cDNA microarray data.** *IEEE Trans Inf Technol Biomed* 2002, **6(1)**:29-37.
50. Hever L, Kruglyak S, Yooshep S: **Exploring expression data: Identification and analysis of co-expressed genes.** *Genome Research* 1999, **9(11)**:1106-1115.
51. Eisen MB, Spellman PT, Brown PO, Botstein D: **Cluster analysis and display of genome-wide expression patterns.** *Proc Natl Acad Sci USA* 1998, **95(25)**:14863-14868.
52. **Gene Expression Omnibus, accession number GSE7553** [<http://www.ncbi.nlm.nih.gov/geo/>]
53. Tusher VG, Tibshirani R, Chu G: **Significance analysis of microarrays applied to the ionizing radiation response.** *Proc Natl Acad Sci USA* 2001, **98(9)**:5116-5121.
54. Harbig J, Sprinkle R, Enkemann SA: **A sequence-based identification of the genes detected by probesets on the Affymetrix U133 plus 2.0 array.** *Nucleic Acids Res* 2005, **33(3)**:e31.
55. **Human Protein Atlas Website** [<http://www.proteinatlas.org/>]

Pre-publication history

The pre-publication history for this paper can be accessed here:

<http://www.biomedcentral.com/1755-8794/1/13/prepub>