

Effect of UVB irradiation on microRNA expression in mouse epidermis

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Abstract. The aim of this study was to assess the effects of UVB irradiation on miRNA expression in the mouse epidermis. We determined miRNA expression profiles in the epidermis of UVB irradiated mice and untreated mice, and conducted TargetScan and Gene Ontology analyses to predict miRNA targets. Three miRNAs were downregulated and three were upregulated in the epidermis of UVB irradiated mice compared with untreated mice, and were predicted to be associated with photocarcinogenesis, hypomethylation and apoptosis. miRNAs are potentially involved in the pathogenesis of photodamage, and may aid in the treatment and prevention of UVB-induced dermatoses.

Introduction

Ultraviolet (UV) radiation from the sun, particularly its UVB component (290-320 nm), is the major cause of skin cancer. UV radiation is also known to elicit various other adverse effects, including erythema, sunburn, inflammation, hyperplasia, hyper-pigmentation, immunosuppression, premature skin aging and photocarcinogenesis (1,2). These cell events are mediated through gene activation or suppression (3). Despite intensive investigations into the regulation of gene expression in skin cells, in the majority of cases the precise molecular events remain to be elucidated. Novel methods should be investigated in this area.

MicroRNAs (miRNAs) are short, non-coding RNAs of approximately 22 nucleotides that are thought to regulate gene expression through sequence-specific base pairing with the 3'-untranslated region (3'-UTR) of target mRNAs. To elucidate the molecular mechanisms underlying photodamage, especially skin carcinogenesis by UVB, miRNA expression profiles in UVB irradiated cells have been investigated using miRNA microarrays (4-6). Guo *et al* investigated the differen-

tial expression profiles of miRNAs in NIH₃T₃ cells in response to UVB irradiation (4). Pothof *et al* found that miRNA expression changes and stress granule formation were most pronounced within the first hours following UVB irradiation, suggesting that miRNA-mediated gene regulation operates earlier than most transcriptional responses (5). The miRNA response may be related to the DNA damage response and cell proliferation (6). However, no studies are currently available on miRNA profiling in response to UVB irradiation *in vivo*. In this study, we compared the profiles of miRNA expression in 3 pairs of UVB irradiated and untreated mice epidermis, in order to reveal the specific underlying mechanisms associated with photodamage.

Materials and methods

Animals and UV light source. Animal care and handling complied with protocols approved by the Nanjing Medical University Institutional Animal Care and Use Committee and employed measures to minimize pain and discomfort. Female C57BL/6 mice (12 weeks old) were obtained from the Chinese Academy of Science, Shanghai SLAC Laboratory Animal Co. (Shanghai, China) and were maintained in a pathogen-free barrier facility at Nanjing Medical University. The source of UVB was a BLE-1T158 (Spectronics Corp., Westbury, NY, USA). A Kodacel filter (TA401/407; Kodak, Rochester, NY, USA) was used to block wavelengths of <290 nm (UVC). The UVB dose was quantified using a Waldmann UV meter (model no. 585100; Waldmann Co., VS-Schwenningen, Germany) and 180 mJ/cm² of UVB was delivered to the dorsal skin of each mouse.

Animal treatments. The C57BL/6 mice were divided into two groups of three animals. The mice in the control group did not receive any treatment. The mice in the second group received UVB (180 mJ/cm²). Following 24 h of UVB irradiation, the dorsal irradiated skin was collected. The epidermis was harvested for analysis by heat separation from the dermis.

RNA isolation and miRNA microarray. Total RNA isolation and the miRNA enrichment procedure were performed using a mirVana miRNA Isolation kit (Ambion, Austin, TX, USA) according to the manufacturer's instructions. RNA concentration was quantified using a NanoDrop spectrophotometer (Thermo Fisher, Waltham, MA, USA). The integrity of the RNA was evaluated using an Agilent 2100 Bioanalyzer

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(Agilent Technologies, Santa Clara, CA, USA). RNA labeling and hybridization on the Agilent miRNA microarray chips were performed using a miRNA labeling reagent and hybridization kit (Agilent Technologies) at 37°C for 30 min. Total RNA samples (100 ng) were treated with calf intestine alkaline phosphatase (Takara Bio Inc., Dalian, China), denatured using 100% DMSO (Sigma-Aldrich, St. Louis, MO, USA) at 100°C for 8 min in a thermal cycler, and then transferred to an ice-water bath to prevent reannealing of the RNA. The RNA samples were then labeled with pCp-Cy3 using T4 RNA ligase (Ambion) and incubated at 16°C for 2 h. The labeled samples were hybridized to Agilent mouse miRNA microarrays, which contained probes for 627 mouse miRNAs and 39 mouse viral miRNAs as catalogued in the Sanger Centre Database version 10.1 (<http://microrna.sanger.ac.uk>). Hybridization was performed in SureHyb chambers (Agilent Technologies) for 24 h at 55°C. The microarrays were then washed using Agilent prepared buffers. The microarray images were scanned using the Agilent microarray scanner, and gridded and analyzed using Agilent Feature Extraction Software, version 9.5.1 (Agilent Technologies). Normalization was performed using the per-chip median normalization method and the median array (7).

Quantitative real-time PCR analysis for miRNA expression. Expression levels of mmu-miR-233 and mmu-miR-141 were validated using quantitative real-time PCR (qRT-PCR). Primers for qRT-PCR were synthesized by Invitrogen (Shanghai, China). cDNA synthesis was performed using a miScript Reverse Transcription kit (Qiagen, Hilden, Germany) according to the manufacturer's instructions. qRT-PCR was performed using a miScript SYBR-Green PCR kit (Qiagen, Hilden, Germany). The reactions were incubated in a 96-well optical plate at 95°C for 15 min, followed by 40 cycles of 15 sec at 94°C, 30 sec at 55°C and 30 sec at 70°C. Expression analysis was performed in triplicate for each sample. mmu-Actin was used as the normalization control. miRNA expression levels were quantified using an ABI Prism 7300 Sequence Detection System (Applied Biosystems, Foster City, CA, USA).

Target prediction and function analysis. TargetScan software was used to predict miRNA targets. To evaluate the TargetScan target predictions for all single miRNAs, we searched for significantly over-represented Gene Ontology (GO) terms among all target genes for all differential miRNAs separately using Gostat software (<http://gostat.wehi.edu.au/cgi-bin/goStat.pl>) (8). In brief, the program determines all the annotated GO terms and all the GO terms that are associated (i.e., in the path) with the genes analyzed. It then counts the number of appearances of each GO term for the genes inside the group and for the reference genes. First, we pasted the mouse RefSeq ID of the target genes into the text area, we then chose 'mgi' (*Mus musculus*) from the available GO gene-association databases. For the remaining options we selected the default values. The majority of significant GO terms usually represent the same subset of genes, since the genes may have several GO annotations that are similar. Fisher's exact test was performed to determine whether the observed difference was significant. For each GO category, this resulted in a P-value whereby the observed counts were due to chance. In addition, pathway

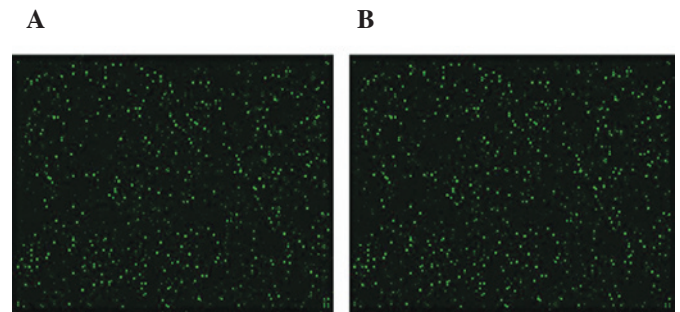


Figure 1. Expression of different miRNAs in mouse skin of two groups detected by microarrays. (A) Untreated mouse epidermis. (B) UVB irradiated epidermis. miRNA, microRNA; UVB, ultraviolet radiation B.

analysis of the targets was performed using DAVID Bioinformatics Resources 2008 (<http://david.abcc.ncifcrf.gov/>).

Statistical analysis. To identify miRNA that was differentially expressed among the groups, a Student's t-test was performed using SPSS version 12.0 (SPSS, Chicago, IL, USA). $P < 0.05$ was considered to indicate a statistically significant difference.

The threshold cycle (Ct) value for the genes was determined using SDS software, version 1.2 (Applied Biosystems). The Ct is the cycle number at which fluorescence is generated as a reaction crosses the threshold. The expression levels of mmu-miR-188-5p, mmu-miR-22, mmu-miR-233, mmu-miR-125a-5p, mmu-miR-146a and mmu-miR-141 were normalized by subtracting their Ct values from that of the internal control mmu-actin, to obtain ΔCt . The $\Delta\Delta Ct$ method for relative quantitation of gene expression was used to determine the miRNA expression levels. ΔCt was calculated by subtracting the Ct of actin from the Ct of the miRNA of interest. $\Delta\Delta Ct$ was calculated by subtracting the ΔCt of the reference sample (control) from the ΔCt of each sample. Fold change was generated using the equation $2^{\Delta\Delta Ct}$. Experiments were repeated in triplicate. Statistical significance was measured using the Student's t-test; $p < 0.05$ was considered to indicate a statistically significant difference.

Results

Significantly differentially expressed miRNAs. From the miRNA microarray results, six miRNAs (mmu-miR-188-5p, mmu-miR-223, mmu-miR-22, mmu-miR-125a-5p, mmu-miR-146a and mmu-miR-141) were found to be differentially expressed in the UVB treatment group compared with the control group ($P < 0.05$). The microarray images are shown in Fig. 1.

Validation of microarray data by miRNA-specific qRT-PCR analysis. To confirm the microarray findings, we measured the expression levels of six differentially expressed miRNAs (mmu-miR-188-5p, mmu-miR-223, mmu-miR-22, mmu-miR-125a-5p, mmu-miR-146a and mmu-miR-141), using qRT-PCR. The expression of mmu-miR-188-5p, mmu-miR-223 and mmu-miR-22 was increased in the UVB-treated epidermis compared with the control. mmu-miR-125a-5p, mmu-miR-146a and mmu-miR-141 were downregulated following UVB irradiation. These results suggest that the expression levels of the

Table I. Differentially expressed miRNAs and putative targets.

	miRNA	Published targets	Putative targets
Three miRNAs expressed significantly more in UVB group than in control group	mmu-miR-188-5p	Mef2c	RSPO3, PHF3, RNF144A, ZFP91, RAP2C
	mmu-miR-223		PHF20L1, GPR155, RHOB, FBXW7, NFAT5
	mmu-miR-22		FUT9, CLIC4, SNX30, PTGS1, TET2
Three miRNAs expressed significantly less in UVB group than in control group	mmu-miR-125a-5p	ZEB1/TCF8	MSRB3, EIF1AD, IRF4, TMEM170B, ENPEP
	mmu-miR-146a		IRAK1, TRAF6, APPL1, FAM62B, MED1
	mmu-miR-141		AKAP2, PALM2-AKAP2, ZEB2, FAT3, TRHDE

miRNA, microRNA; UVB, ultraviolet radiation B.

Table II. Gene Ontology (GO) terms for the target genes of treatment responsive miRNAs.

	GO term	Function	P-value
Three miRNAs expressed significantly more in UVB group than in control group	0043283	Biopolymer metabolic process	3.64E-18
	0050794	Regulation of cellular process	3.64E-18
	0050789	Regulation of biological process	3.56E-17
	0065007	Biological regulation	3.56E-17
	0031323	Regulation of cellular metabolic process	4.37E-15
Three miRNAs expressed significantly less in UVB group than in control group	0043283	Biopolymer metabolic process	9.13E-16
	0050789	Regulation of biological process	3.06E-13
	0005515	Protein binding	3.06E-13
	0050794	Regulation of cellular process	3.06E-13
	0010468	Regulation of gene expression	3.06E-13

miRNA, microRNA; UVB, ultraviolet radiation B.

four miRNAs observed in the arrays were consistent with those observed using qRT-PCR (Fig. 2).

Target prediction and function analysis of differentially expressed miRNA. The prediction of miRNA-regulated gene targets is a crucial step in understanding the functions of miRNA. We used TargetScan to obtain predicted gene targets for all differentially expressed miRNAs. As expected, these miRNA genes could potentially regulate several hundred targets. We then examined the significant GO categories and Kyoto Encyclopedia of Genes and Genomes pathways (KEGG) (Tables I-III).

Discussion

Our present study revealed miRNAs that are sensitive to UVB and baicalin treatment. We analyzed skin tissues from mice in two groups (those irradiated with UVB and controls) 24-h post-irradiation, using a miRNA microarray platform that was able to assess the expression of 627 mouse miRNAs and 39 mouse viral miRNAs. To select differentially expressed miRNAs from the microarray data, we set a cut-off limit at $p < 0.05$. The differentially expressed miRNAs were mmu-miR-188-5p, mmu-miR-223, mmu-miR-22, mmu-miR-125a-5p, mmu-miR-146a and mmu-miR-141.

Among the UVB downregulated miRNAs, miR-141 has been described as a member of the miR-200 family.

Korpál *et al* (9) found that the miR-200 family miRNAs inhibit epithelial-mesenchymal transition and cancer cell migration by direct targeting of the E-cadherin transcriptional repressors ZEB1 and ZEB2. These findings suggested that the downregulated expression of miR-141 induced by UVB irradiation is involved in the repression of E-cadherin, thereby enhancing migration and invasion during cancer progression. The base excision repair protein MED1 is a predicted target of mmu-miR-146a. MED1 interacts with the mismatch repair protein MLH1 and has a key role in the maintenance of genomic stability, with dual functions in DNA damage response and repair (10,11). MED1 acts as a thymine and uracil DNA N-glycosylase on T:G and U:G mismatches that occur at cytosine-phosphate-guanine (CpG) methylation sites due to the spontaneous deamination of 5-methylcytosine and cytosine, respectively. This event indicates that MED1 is involved in the removal of methylated DNA (12). Abnormal DNA methylation (including hypermethylation and hypomethylation) is a hallmark of the majority of cancers, including colon, lung, prostate and breast cancers, and contributes to carcinogenesis by silencing tumor suppressor genes, upregulating oncogenes and/or reducing genomic stability (13,14). Mittal *et al* observed global DNA hypomethylation and reduced maintenance methylation in UV-exposed mouse skin (15). In view of these findings, whether or not the downregulated expression of miR-146a contributes to UV-induced DNA hypomethylation via MED1 requires further investigation.

Table III. Pathway analysis of target genes of treatment responsive miRNAs using DAVID Bioinformatics Resources 2008.

	KEGG pathway	P-value
Three miRNAs expressed significantly more in UVB group than in control group	ErbB signaling pathway	9.45
	MAPK signaling pathway	8.01
	Dorsoventral axis formation	6.67
	Prostate cancer	6.53
	Chronic myeloid leukemia	5.94
Three miRNAs expressed significantly less in UVB group than in control group	Chronic myeloid leukemia	13.54
	Notch signaling pathway	9.32
	Dorsoventral axis formation	7.83
	MAPK signaling pathway	6.29
	Pancreatic cancer	5.67

miRNA, microRNA; UVB, ultraviolet radiation B; KEGG, Kyoto Encyclopedia of Genes and Genomes.

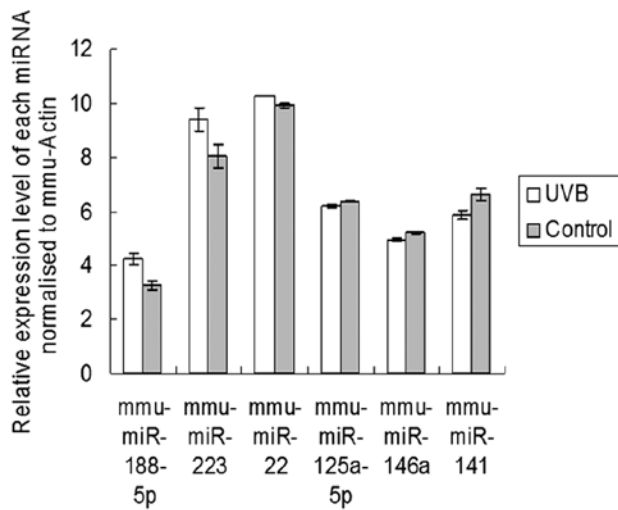


Figure 2. qRT-PCR analysis of dysregulated miRNA expression. UVB, ultraviolet radiation B; qRT-PCR, quantitative real-time PCR; miRNA, microRNA.

In the present study, it is clear that mmu-miR-188-5p, mmu-miR-223 and mmu-miR-22 were upregulated following UVB irradiation. Johnnidis *et al* found that miR-223 mutant mice have an expanded granulocytic compartment resulting from a cell-autonomous increase in the number of granulocyte progenitors (16). These authors also revealed that Mef2c, a transcription factor that promotes myeloid progenitor proliferation, is a target of miR-223 (16-18). Their data support a model in which miR-223 acts as a fine-tuner of granulocyte production and the inflammatory response. We demonstrated that miR-223 is expressed in the skin of untreated mice, and markedly increases following UVB irradiation. These findings suggest that mmu-miR-223 is sensitive to UVB and elicits a direct or indirect effect of UVB-induced inflammation. RhoB, a predicted target of miR-223, has been reported to protect human keratinocytes from UVB-induced apoptosis through epidermal growth factor receptor signaling (19), suggesting that miR-223 plays a role in regulating keratinocyte survival

following UVB exposure. Little is known about the function of miR-188-5p and miR-22.

To further analyze the correlation between patterns of target gene expression and their functional implications, we classified the target genes of all differentially expressed miRNAs into several function categories using Gostat software (<http://gostat.wehi.edu.au/cgi-bin/goStat.pl>) (8). GO terms concerning the regulation of cellular and biological processes are listed in Table II. The targets of these nine miRNAs were selected for pathway analysis using DAVID Bioinformatics Resources 2008 (<http://david.abcc.ncifcrf.gov/>) (Table III).

It should be noted that certain miRNAs reported to be involved in the response to UVB irradiation were not observed in this study. For example, miR-21 is known to be involved in the progression of cancer and has been described as an oncogenic miRNA (20), but when it appears together with miR-24, it inhibits growth. Guo *et al* found that miR-21 and miR-24 appeared together following 12-h exposure to 50 J/m² UVB (4) and their observations of the cell cycle and apoptosis appeared to reveal a sub-G1 DNA content fraction and apoptotic cells 12-h post-irradiation, suggesting that miR-21 and miR-24 together inhibited growth. However, the present study results did not show any changes in miR-21 or miR-24 in the UVB group. This finding may be due to differences in animal selection, UVB dose, chip fabrication or a weakness in the microarray technology.

In conclusion, the focus of this study was to investigate the differential expression profiles of miRNAs in the skin of mice following UVB irradiation. Although our investigation is at a preliminary stage, we believe this study provides a basis for further investigation of the function of expression profiles in miRNAs in signal transduction pathways induced by UVB treatment. miRNAs may therefore be new research hotspots for the prevention or treatment of skin cancer caused by UVB.

Acknowledgements

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