

# Differential microRNA expression and identification of putative miRNA targets and pathways in head and neck cancers

NURUL-SYAKIMA AB MUTALIB<sup>1</sup>, CHEAH YOKE-KQUEEN<sup>1,2</sup>, SABARIAH ABDUL RAHMAN<sup>2,3</sup>,  
SHIRAN MOHD SIDIK<sup>3</sup>, AVATAR SINGH MOHAN SINGH<sup>4</sup> and LEE LEARN-HAN<sup>1</sup>

<sup>1</sup>Department of Biomedical Science, Faculty of Medicine and Health Sciences, <sup>2</sup>UPM-MAKNA Cancer Research Laboratory, Institute of Bioscience, <sup>3</sup>Department of Pathology, Faculty of Medicine and Health Sciences, University Putra Malaysia, 43400 UPM Serdang, Selangor Darul Ehsan; <sup>4</sup>Department of Otorhinolaryngology, Taiping General Hospital, Jalan Taming Sari, 34000 Taiping, Perak, Malaysia

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**Abstract.** MicroRNAs (miRNAs) are small noncoding RNAs that involved in various cancer-related cellular processes. Diverse studies on expression profiling of miRNAs have been performed and the data showed that some miRNAs are up-regulated or down-regulated in cancer. Until now, there are no data published on the miRNA expression in head and neck cancers from Malaysia. Hence, this study aimed to investigate potentially crucial miRNAs in head and neck cancer patients from Malaysian populations. A global miRNA profiling was performed on 12 samples of head and neck cancer tissue using microarray analysis followed by validation using real-time RT-PCR. Microarray analysis identified 10 miRNAs that could distinguish malignant head and neck cancer lesions from normal tissues; 7 miRNAs (hsa-miR-181a-2\*, hsa-miR-29b-1\*, hsa-miR-181a, hsa-miR-181b, hsa-miR-744, hsa-miR-1271 and hsa-miR-221\*) were up-regulated while 3 miRNAs (hsa-miR-141, hsa-miR-95 and hsa-miR-101) were down-regulated. These miRNAs may contribute in a simple profiling strategy to identify individuals at higher risk of developing head and neck cancers, thus helping in the elucidation of the molecular mechanisms involved in head and neck cancer pathogenesis.

## Introduction

Head and neck cancers include cancers that originate from the nasopharynx, oral cavity, lips, nose and paranasal sinus, larynx,

oropharynx, hypopharynx, thyroid and the salivary glands. In 2006, 2884 cases of head and neck cancers were reported in Peninsular Malaysia with nasopharyngeal cancer ranked first constituting 34% of the cases. Malaysia is a multiracial country that consist of three major ethnicities; Malay, Chinese and Indian. Distribution of head and neck cancers are varied according to ethnicity; laryngeal, oral and pharyngeal cancers are more common in Indians, followed by Malay and Chinese. On the other hand, nasopharyngeal cancer is most common among the Chinese followed by the Malays, the indigenous East Malaysians and Indians.

The worldwide five-year relative survival rate for head and neck cancers is generally <50% and has remained unchanged for more than 30 years (1). Furthermore, the survival rate for patients diagnosed with regional or distant disease (with spread to nearby or distant organs and lymph nodes) was even lower (2). One of the factors that contribute to this poor survival rate is the late-stage diagnosis (1). Due to this fact, a molecular approach that enables the early stage detection of these cancers may be very helpful in improving prognosis.

MicroRNAs (miRNA) are endogenous, small, noncoding RNAs of 17-25 nucleotides that are thought to regulate almost 30% of human genes (3,4). The first miRNA discovery was in *Caenorhabditis elegans* and later studies elucidated that miRNAs are also widely conserved in animals and plants (4-6). Its crucial involvement in various cancer-related cellular processes, such as cell differentiation, cell division (7), apoptosis (8) and cell cycle regulation (9) have made it important to investigate the role of this molecule in head and neck cancers especially in a multiracial country like Malaysia.

Microarray is a global expression analysis that permits interrogation of the expression of thousands of genes simultaneously in a high-throughput manner, offering precious information on disease pathology and progression, thus leading to an improved diagnostic approach for cancer (10). This analysis method has been widely used for miRNA expression profiling and altered miRNA expression in head and neck cancer has been investigated by several groups (11) internationally but none has been conducted in Malaysia. Thus this study is the first to provide information about differences in

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*Correspondence to:* Dr Cheah Yoke-Kqueen, Department of Biomedical Science, Faculty of Medicine and Health Sciences, and UPM-MAKNA Cancer Research Laboratory, Institute of Bioscience, University Putra Malaysia, 43400 UPM Serdang, Selangor Darul Ehsan, Malaysia  
E-mail: ykcheah@medic.upm.edu.my

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Table I. Demographics and clinical characteristics of patients with head and neck cancer.

Characteristics	No.
Gender	
Man	6
Woman	3
Age	
Year, median (range)	60 (20-83-year old)
Race	
Malay	5
Chinese	2
Indian	2
TNM staging	
T1	0
T2	2
T3	2
T4	5
Pathological lymph node status	
N0	3
N <sup>+</sup>	6
Anatomical site	
Buccal	2
Supraglottic	2
Nasopharynx	2
Retromolar	1
External ear	1
Nasal cavity	1
TNM, tumor node metastasis.	

miRNA expression signatures between head and neck cancer and normal tissues among Malaysians.

## Materials and methods

**Sample collection.** This study was comprised of 9 head and neck cancer patients that underwent surgical resection at Taiping General Hospital, Perak, Malaysia. All diagnoses were histologically verified by the pathologists. Ethics approvals were obtained from the local research Ethics Committees (National Institutes of Health and Medical Research Ethics Committee, Faculty of Medicine and Health Sciences, University Putra Malaysia, Malaysia). All patients gave written informed consent. The tissues were cut <0.5 cm in at least one dimension and the specimens were submerged in five volumes of RNAlater (Ambion, Austin, TX, USA) at room temperature for 2 h (to allow the solution to thoroughly penetrate the tissue) and kept at -80°C until they were subjected to total-RNA extraction. RNAlater is an aqueous, non-toxic tissue storage reagent that rapidly permeates tissues to stabilize and protect cellular RNA. This solution eliminates the need to instantly process tissue samples or to freeze samples in liquid nitrogen for later processing.

Table II. Designation of the total-RNA samples.

Samples	Type
RL1	Cancer
RL3	Cancer
RL7	Cancer
RL8	Cancer
RL9	Cancer
RL19	Cancer
RL17Ca	Cancer
RL25Ca	Cancer
RL29Ca	Cancer
RL17N	Normal
RL25N	Normal
RL29N	Normal

There were 12 samples of total-RNA from 9 patients; 9 cancer tissues and 3 normal tissues. RL17, RL25 and RL29 are paired tissues (cancer and normal), whereas only cancer tissues are available for the rest.

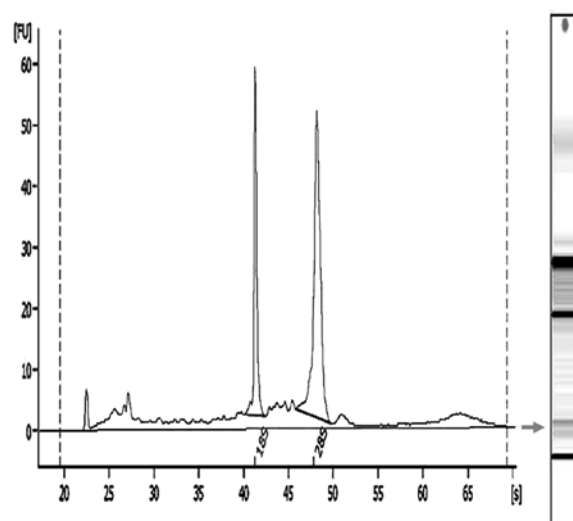


Figure 1. Electropherogram obtained from bioanalyzer analysis of total-RNA extracted from head and neck tissue. The arrow represents the presence of small RNA species including miRNAs. Highly intact RNA (RIN<sub>≥</sub>7) is indicated by sharp peaks of 18S and 28S.

Histologically-verified cancer and paired normal tissues were obtained from 3 patients (2 females and 1 male; 63, 59 and 46-year old, respectively) while for the rest only cancerous tissues were available. Normal tissues were >2 cm away from the tumor. Full patient demographics are provided in Table I and the designation of the total-RNA samples is summarized in Table II.

**Total-RNA extraction.** Total-RNA was extracted using the Total-RNA Purification kit (Norgen Biotek, Canada). Prior to extraction, sterile, RNase-free screw-cap 2 ml tubes were filled with ~700 mg of 0.1 mm glass beads (Biospec Products, Bartlesville, OK). Then 600  $\mu$ l of lysis solution and 6  $\mu$ l of  $\beta$ -mercaptoethanol were added to each tube. Tissue (10 mg)

Table III. Differentially expressed miRNAs in head and neck cancer samples compared to normal samples.

miRNA	Unpaired t-test P-value	Fold-change log <sub>2</sub> (cancer/normal)	Cancer vs. normal	Chromosomal location
hsa-miR-29b-1*	0.00315	1.18	up-regulated	7:130212758-130212838 (-)
hsa-miR-181a-2*	0.000613	1.11	up-regulated	9:126494542-126494651 (+)
hsa-miR-221*	0.0273	0.80	up-regulated	X:45490529-45490638 (-)
hsa-miR-1271	0.0439	0.65	up-regulated	5:175794949-175795034 (+)
hsa-miR-744	0.00264	0.61	up-regulated	17:11925941-11926038 (+)
hsa-miR-181b	0.0163	0.60	up-regulated	1:197094625-197094734 (-)
hsa-miR-181a	0.0321	0.43	up-regulated	1:197094796-197094905 (-)
hsa-miR-141	0.00181	-0.46	down-regulated	12:6943521-6943615 (+)
hsa-miR-95	0.00577	-0.46	down-regulated	4:8057928-8058008 (-)
hsa-miR-101	0.0202	-0.51	down-regulated	1:65296705-65296779 (-)

was placed into the tube and subjected to lysis using a Biospec mini-bead beater (BioSpec Products) at the maximum speed until the tissue became small fragments (~40-80 sec, at 20 sec intervals). Subsequent RNA extraction steps followed the manufacturer's instruction. The absorbance at 260/280 nm and concentration of the RNA were determined by using the Biophotometer 6131 (Eppendorf, USA) and the RNA integrity was confirmed with the Agilent 2100 Bioanalyzer (Agilent Technologies, Santa Clara, CA, USA). The RNA integrity number (RIN) was calculated for every case. Total-RNA of 12 samples were chosen for microarray based on the RIN number ( $\geq 7$ ) and an example of electropherogram is shown in Fig. 1.

**miRNA microarray.** miRNA microarray was performed using the Human v2 MicroRNA Expression Profiling kit (Illumina, San Diego, CA) on BeadChip. All RNA samples were first normalized to approximately 90 ng/ $\mu$ l. For each sample, 5  $\mu$ l of normalized RNA was used for the array. All reactions were performed in duplicate. The BeadArray Reader was used to measure the fluorescence intensity. Images were imported into Illumina GenomeStudio and data were analyzed using Illumina's GenomeStudio Gene Expression Module.

The signal values of spike controls were averaged and used as a cut-off value, and sample with signal intensity below the cut-off value were excluded, followed by quantile normalization of the signals intensity on cancer tissues and normal tissues. The signal values were averaged from both replicates. Signals significantly detected were defined as those with a detection  $P < 0.01$ . After filtration with detection P-value, 493 miRNAs were found to be accurately detected from the array and were further analyzed in Excel 2007 (Microsoft, Seattle, WA). Microsoft Excel has been widely used in micro-array statistical analysis including in miRNA microarray (12-15). The unpaired t-test (t-test with unequal variance) was performed between cancer and normal sample groups and the P-values were calculated. Differentially expressed miRNAs were those with  $P < 0.05$ . The signal ratio or the expression ratio was logarithmically transformed to the base 2 to obtain the fold-change ( $> 0$ , up-regulated;  $< 0$ , down-regulated) and the significance was calculated by using  $-\log_{10}$  of the P-value obtained from the unpaired t-tests (16). A heat map was

generated for the differentially expressed miRNAs. All of the microarray data presented in this study are in accordance with the Minimum Information about a Microarray Experiment (MIAME) guidelines.

**Validation of miRNA microarray data by quantitative RT-PCR (qRT-PCR).** Six miRNAs were chosen for validation (hsa-miR-221\*, hsa-miR-744, hsa-miR-181b, hsa-miR-181a, hsa-miR-141 and hsa-miR-95) and qRT-PCR was performed using the NCode™ EXPRESS SYBR®-GreenER™ miRNA qRT-PCR kit Universal (Invitrogen, Carlsbad, CA, USA). Total-RNA input without additional enrichment was used to conserve precious samples. RNU-48 was used as normalization control or reference gene. Total-RNA (1  $\mu$ g) was subjected to a reverse transcription reaction according to the manufacturer's protocol.

For miRNA qPCR, the Universal qPCR Primer provided in the kit was used as the reverse primer. The forward qPCR primer was specific for the miRNA sequence of interest and was obtained from the NCode™ miRNA database (<http://escience.invitrogen.com/ncode>). However, in some miRNAs, qPCR using primers mentioned above were unsuccessful, thus miRNA-specific primer pairs were designed using Primer-BLAST (<http://www.ncbi.nlm.nih.gov/tools/primer-blast>). Primer-BLAST was developed at NCBI to help users design specific primers to the input PCR template. It utilizes Primer3 to design PCR primers followed by submission to BLAST search and then automatically analyzed to avoid primer pairs that can cause unspecific amplification of targets.

The qPCR reaction was run in triplicate using the Rotorgene 6000 (Qiagen, Valencia, CA, USA) in a 20  $\mu$ l volume according to manufacturer's protocol with adjustment of annealing time to 15 sec. The qRT-PCR amplification products were subsequently analyzed by melting curve analysis and confirmed by 1.8% agarose gel electrophoresis. A negative control without cDNA template was included to assess the specificity of the qPCR reaction. The fold-change of the miRNAs expression was calculated using the  $2^{-\Delta\Delta Ct}$  method (17).

**Prediction of putative miRNA targets and pathways through in silico analysis.** Targets of miRNA differentially expressed

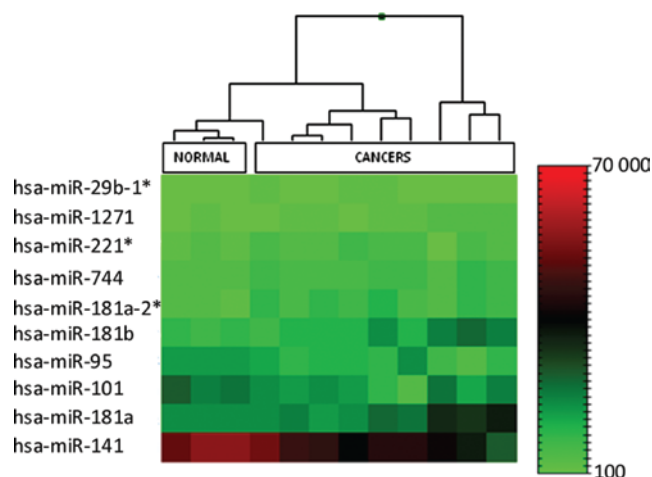


Figure 2. Microarray heat map showing the statistically significant ( $P<0.05$ ) differentially expressed miRNAs and the hierarchical clustering of the samples. The analyzed samples are in columns and the miRNAs are presented in rows. The sample-clustering tree appears at the top. The color scale shown represents the relative expression level of a miRNA; red illustrates a high expression level and green illustrates a low expression level.

in head and neck cancers were determined using the DIANA-microT version 4.0 (freely available at [http://diana.cslab.ece.ntua.gr/microT\\_v4](http://diana.cslab.ece.ntua.gr/microT_v4)). On the other hand, for identification of putative pathways targeted by miRNAs of interest, the DIANA-miRPath (<http://microrna.gr/mirpath>) was utilized. Both *in silico* tools mentioned above were connected to the KEGG (Kyoto Encyclopedia of Genes and Genomes), a database used for representation and analysis of molecular networks involving diseases and drugs (18).

## Results

Microarray analysis was performed to determine the miRNA expression profile in head and neck cancers. When the expression of miRNAs was compared between the normal head and neck samples and malignant samples, 10 miRNAs were found to be aberrantly expressed as summarized in Table III. Among the 10 miRNAs, 7 were up-regulated and 3 were down-regulated in cancer samples compared to normal samples.

A representative heat map of the statistically significant ( $P<0.05$ ) results is shown in Fig. 2. The hierarchical clustering based on an Euclidean algorithm generated a tree with a clear distinction of samples in two main groups, represented by normal tissues and cancer tissues. The Volcano plot in Fig. 3 displays the relationship between the significance of miRNAs detected and the fold-change between normal and cancer tissues.

In order to confirm the results obtained from the miRNA microarray, the expression of 6 miRNAs was analyzed by qRT-PCR in the same samples analyzed by the microarray. The fold change of the expression of miRNAs was calculated according to Livak and Schmittgen (19). Results of the qRT-PCR are consistent with those of the microarray analysis. miRNA hsa-miR-221\*, hsa-miR-744, hsa-miR-181b and hsa-miR-181a were up-regulated while hsa-miR-141 and hsa-miR-95 were down-regulated (Fig. 4). These results validate the results obtained by the microarray analysis.

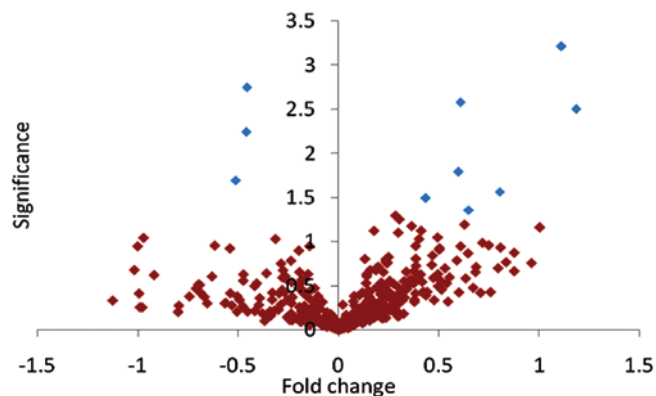


Figure 3. A Volcano plot representing the significance of miRNAs vs. the fold-change. The blue color diamonds represent miRNAs which are differentially expressed with  $P<0.05$ . The up-regulated miRNAs are signified by a positive fold-change value and *vice versa*.

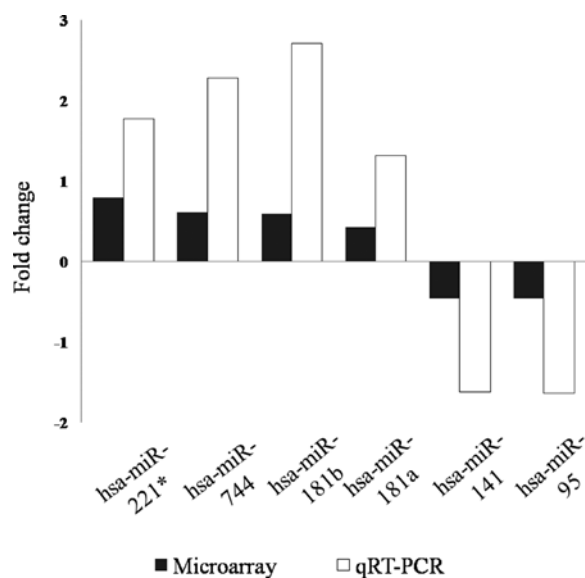


Figure 4. Validation of miRNA microarray results by qRT-PCR in the head and neck cancer samples. The black bars represent values from microarray analysis, whereas the white bars represent qRT-PCR data.

*In silico analysis of dysregulated miRNAs in head and neck cancers.* Using the medium or the strict Score Threshold setting in the DIANA-microT, genes targeted by particular miRNAs were revealed along with the KEGG pathway enrichment in the target genes. Results obtained through the DIANA-microT version 4.0 are summarized in Table IV. Only the results associated with cancer-related KEGG pathways are displayed.

Using the DIANA mirPath, the list of pathways was narrowed to choose only the pathways that are involved in human cancers according to the KEGG pathway database ([http://www.genome.jp/kegg-bin/show\\_pathway?hsa05200+1956](http://www.genome.jp/kegg-bin/show_pathway?hsa05200+1956)) (Table V).

## Discussion

*miRNAs differentially expressed in carcinoma and normal tissues of the head and neck.* In recent years, there have been a number of studies that have profiled the miRNA expression

in various cancers and these differential expressions may be applied to distinguishing tumors and normal tissues (20-30). Moreover, some miRNAs have even been identified as cancer markers or prognostic markers for cancer patients (21,26,30-32). However, data on a miRNA signature of head and neck cancers in Malaysians have not been reported. Hence, the aim of this study was to assess the profiles of miRNA in Malaysian patients diagnosed with head and neck cancers.

miRNAs are differentially expressed between normal and cancerous tissues in agreement with this study. A miRNA expression array was used to determine the miRNA profiles in head and neck cancer tissues and the results showed that the miRNA expression profile can definitely distinguish head and neck cancer tissues from normal tissues. Comparison of the expression of miRNAs resulted in 10 differentially expressed miRNAs; 7 were up-regulated and 3 were down-regulated. The most interesting finding was that, the number of up-regulated miRNAs in carcinoma samples was higher than the number of down-regulated miRNAs. This finding is in agreement with previously published miRNA profiling studies, most of which showed a higher degree of up-regulation than down-regulation of miRNAs in cancer (33,34). Defects in the miRNA processing machinery components could explain the overall increased expression of miRNAs (35,36).

The clustering analysis performed on all the samples based on miRNA expression produced a clustering tree that successfully separated normal from cancer samples. However we did not obtain any differentially expressed miRNAs associated with different stages of head and neck cancers. This could be due to the fact that samples in this study were mostly of advanced stages (stage IV) and there was no sample of stage I. Furthermore, the small size of our sample could represent a limit for the statistical analysis.

Owing to the tissue specificity of miRNAs (37-39), different miRNAs are up-regulated or down-regulated in different types of cancer. Among the interesting findings obtained from this study are the up-regulation of miRNAs of the miR-181 family; hsa-miR-181a-2\*, hsa-miR-181a and hsa-miR-181b. Considered as oncogenic, the miR-181 family has also been found to be up-regulated in carcinomas of the ovary (22) and breast (40). Other miRNA studies in head and neck cancers also revealed the up-regulation of the miR-181 family (34,41,42). Using the mouse as an animal model, Wang *et al* also demonstrated the oncogenic property of this miRNA family (43). On the contrary, hsa-miR-181a and hsa-miR-181b were found to be significantly down-regulated and to function as tumor suppressors in glioma (44). These remarkable discoveries underscore the fact that the same miRNA can act differently among various cancer types; it is thus important to study its function in head and neck carcinogenesis. So far no in-depth study has been conducted to determine the functionality of hsa-miR-181a-2\*. An extended study of this miRNA family is warranted to investigate its role in head and neck cancers. In this study, the miR-181 family could be considered as an oncomiR (a miRNA that acts as an oncogene) due to its up-regulation in carcinoma tissues. However, functional studies must be performed to validate its cancer-related properties in head and neck cancers.

The down-regulation of hsa-miR-95 in carcinoma tissues observed in this study is in agreement with a similar study by Kozaki *et al* (41). In addition, hsa-miR-95 was also found to

be down-regulated in lung cancer tissues vs. noncancerous lung tissues (21). This finding signifies that hsa-miR-95 could be a tumor suppressor miRNA that leads to carcinoma of the head and neck as well as to lung cancer when its expression is low. On the other hand, Cheng *et al* discovered that inhibition of hsa-miR-95 in HeLa cells resulted in decreased cell growth (45). This contrast finding supports the notion that hsa-miR-95 is an oncomiR in that cervical cancer cell line.

#### *New miRNAs differentially expressed in head and neck cancers.*

Some of the differentially expressed miRNAs identified in this study have not been reported to be differentially expressed in other similar studies pertaining to head and neck cancers. Noteworthy is the expression level of hsa-miR-744, which is significantly up-regulated, signifying that it could be a miRNA signature specific for head and neck cancers. However, since samples used in this study were mostly of advanced stage tumors, perhaps hsa-miR-744 is the late diagnosis marker of head and neck cancers. This notion may similarly be applied to hsa-miR-141 and hsa-miR-101, which are significantly down-regulated. Even though no miRNA studies of head and neck cancers have identified hsa-miR-141, this miRNA has been found to be down-regulated in other cancers, such as gastric and renal cell carcinoma (46,47). Meanwhile hsa-miR-101 is one of the best studied miRNA in cancer, and its down-regulation has been reported in ovarian (22) and hepatocellular cancer (48). hsa-miR-101 has been proven to promote apoptosis and to suppress tumorigenicity in hepatocellular cancer (48). The tumor suppressing ability of hsa-miR-101 is further enhanced by a more recent finding that it plays a regulatory role in prostate tumorigenesis and that an increase of its levels may be a novel valuable strategy for prostate cancer treatment (49).

It is well known that hsa-miR-21 is highly expressed in the majority of human malignancies (27); however, in this study the observed difference of this oncomiR between cancer and normal tissues was not significant. This result is in agreement with other studies which found that hsa-miR-21 is not differentially expressed between cancer and normal tissues in prostate (23), gastric (26) and male breast cancer (50). On the contrary, almost all miRNA studies in head and neck cancers reported that hsa-miR-21 is up-regulated in cancers. This varied finding could be due to different approaches used for miRNA profiling as well as due to the method chosen for data analysis. Furthermore, the site of tissues chosen for expression profiling could contribute to this result. This study consists of tissues obtained from various anatomical sites in the head and neck; buccal, supraglottic, nasopharynx, retromolar, external ear and nasal cavity. The uniqueness of studied subjects is further enhanced by inclusion of three major races in Malaysia; Malay, Chinese and Indian.

The miRNA biogenesis pathway involves the stepwise processing of miRNA precursor transcripts containing hairpin structures in the nucleus as well as in the cytoplasm (51). After Drosha processing in the nucleus, miRNA containing hairpins are exported into the cytoplasm and subsequently cleaved by Dicer, resulting in the miRNA duplex (52). Both strands of miRNA duplexes are produced in equivalent amounts by transcription; however, their accumulation is mostly asymmetric at steady state. The most abundant strand of a processed

Table IV. Summary of genes and cancer-related pathways targeted by miRNAs differentially expressed in head and neck cancers.

miRNA	Summary	Putative target gene	Cancer-associated KEGG pathway
hsa-miR-29b-1*	88 target sites in 27 genes	BHLHB3, KLF12, TSGA14, CCDC117, NEUROD1	None
hsa-miR-181a-2*	58 target sites in 22 genes	SLC39A10, ENSG00000214728, FBN1, ENSG00000183604, PLCXD1	None
hsa-miR-221*	32 target sites in 11 genes	ADIPOQ, ERC2, MBNL1, ENSG00000127388, C16orf72	None
hsa-miR-1271	234 target sites in 82 genes	ATG9A, KIAA2022, SOX5, ERC2, NOVA1	None
hsa-miR-744	61 target sites in 33 genes; genes in pathways: 5	ILK ILK GNAS ENSG00000100181 ILK	Endometrial cancer PPAR signaling pathway Gap junction Wnt signaling pathway Focal adhesion
hsa-miR-181b	676 target sites in 181 genes; genes in pathways: 30	SOS1, MAPK1, RAP1B, PAK7  SOS1, MAPK1, PAK7, RPS6KB1 ACVR2A, MAPK1, ACVR2B, RPS6KB1 SOS1, MAPK1, RPS6KB1 NFATC2, IL1A, SOS1, MAP3K3, MAPK1, RAP1B NFATC2, MAPK1, NFAT5 MAPK1, RPS6KB1 SOS1, MAPK1, RAP1B, PAK7 SOS1, MAPK1 SOS1, MAPK1 WASL, SOS1, MAPK1, PAK7 SOS1, MAPK1 NFATC2, SENP2, NFAT5 LIFR, SOS1, SPRY4 WASL, MAPK1 LIFR, IL1A, ACVR2A, ACVR2B SOS1, MAPK1 SOS1, MAPK1 SOS1, MAPK1 SOS1, MAPK1 MAPK1 PRKCE MAPK1 MAPK1 ZIC2 IL1A MAPK1 MAPK1	Renal cell carcinoma  ErbB signaling pathway TGF- $\beta$ signaling pathway Acute myeloid leukemia MAPK signaling pathway  VEGF signaling pathway mTOR signaling pathway Focal adhesion Endometrial cancer Non-small cell lung cancer Regulation of actin cytoskeleton Glioma Wnt signaling pathway Jak-STAT signaling pathway Adherens junction Cytokine-cytokine receptor interaction Chronic myeloid leukemia Colorectal cancer Prostate cancer Gap junction Thyroid cancer Tight junction Bladder cancer Toll-like receptor signaling pathway Hedgehog signaling pathway Apoptosis Pancreatic cancer Melanoma
hsa-miR-181a	772 target sites in 206 genes; genes in pathways: 32	SOS1, MAPK1, RAP1B, PAK7, AKT3 SOS1, MAPK1, RPS6KB1, AKT3 SOS1, MAPK1, PAK7, RPS6KB1, AKT3 ID4, ACVR2A, MAPK1, ACVR2B, RPS6KB1 PDGFRA, SOS1, MAPK1, AKT3 NFATC2, PDGFRA, ATF2, SOS1, MAPK1, RAP1B, NLK, AKT3 WASL, MAPK1, SSX2IP, NLK PDGFRA, SOS1, MAPK1, AKT3 MAPK1, RPS6KB1, AKT3 PDGFRA, SOS1, MAPK1, AKT3 PDGFRA, SOS1, MAPK1, RAP1B, PAK7, AKT3 SOS1, MAPK1, AKT3 SOS1, MAPK1, AKT3	Renal cell carcinoma Acute myeloid leukemia ErbB signaling pathway TGF-beta signaling pathway Glioma MAPK signaling pathway  Adherens junction Colorectal cancer mTOR signaling pathway Prostate cancer Focal adhesion Endometrial cancer Non-small cell lung cancer

Table IV. Continued.

miRNA	Summary	Putative target gene	Cancer-associated KEGG pathway
		NFATC2, MAPK1, AKT3 PDGFRA, MAPK1, AKT3 SOS1, MAPK1, AKT3 PDGFRA, WASL, SOS1, MAPK1, PAK7 PDGFRA, SOS1, MAPK1 PRKCE, MPP5, AKT3 NFATC2, NLK, SENP2 SOS1, SPRY4, AKT3 MAPK1, AKT3 MAPK1, AKT3 MAPK1 MAPK1 AKT3 AKT3 ZIC2	VEGF signaling pathway Melanoma Chronic myeloid leukemia Regulation of actin cytoskeleton Gap junction Tight junction Wnt signaling pathway Jak-STAT signaling pathway Pancreatic cancer Toll-like receptor signaling pathway Thyroid cancer Bladder cancer Small cell lung cancer Apoptosis Hedgehog signaling pathway
hsa-miR-141	306 target sites in 97 genes; genes in pathways: 18	E2F3, CDC14A, YWHAG PRKCE, PTEN, PTENP1, MYH10 E2F3, PTEN, PTENP1 E2F3, PTEN, PTENP1 E2F3, PTEN, PTENP1 E2F3, PTEN, PTENP1 E2F3 PTEN, PTENP1 E2F3 PRKACB MAP2K4, PRKACB PTEN, PTENP1 PTEN, PTENP1 E2F3 E2F3 PRKACB MAP2K4 PRKACB PRKACB MAP2K4	Cell cycle Tight junction Glioma Melanoma Small cell lung cancer Prostate cancer Bladder cancer Endometrial cancer Non-small cell lung cancer Hedgehog signaling pathway MAPK signaling pathway Focal adhesion p53 signaling pathway Pancreatic cancer Chronic myeloid leukemia Apoptosis ErbB signaling pathway Wnt signaling pathway Gap junction Toll-like receptor signaling pathway
hsa-miR-95	15 target sites in 14 genes; genes in pathways: 3	GNAI2 GNAI2	Gap junction Tight junction
hsa-miR-101	335 target sites in 109 genes; genes in pathways: 24	PTCH1, FZD4 FBXW11, PTCH1 MAGI2, PRKCE, MAGI1 TGFB1, FZD4 TGFB1, ACVR2B TGFB1, MAP3K4, RAP1B FBXW11, FZD4 SOCS5 RAP1B GJA1 CADM1 RAP1B MITF TGFB1 TGFB1 GJA1 TGFB1 TGFB1, ACVR2B	Basal cell carcinoma Hedgehog signaling pathway Tight junction Colorectal cancer TGF- $\beta$ signaling pathway MAPK signaling pathway Wnt signaling pathway Jak-STAT signaling pathway Focal adhesion Cell communication Cell adhesion molecules (CAMs) Renal cell carcinoma Melanoma Adherens junction Pancreatic cancer Gap junction Chronic myeloid leukemia Cytokine-cytokine receptor interaction

Table V. Pathway analysis of multiple miRNAs performed with DIANA-mirPath.

	Cancer-associated KEGG pathways (in up-regulated miRNAs)	Cancer-associated KEGG pathways (in down-regulated miRNAs)
Cytokine-cytokine receptor interaction	+	-
ECM-receptor interaction	+	+
Focal adhesion	+	+
Renal cell carcinoma	+	+
Glioma	+	+
Prostate cancer	+	+
ErbB signaling pathway	+	+
Colorectal cancer	+	+
Small cell lung cancer	+	+
Endometrial cancer	+	+
TGF- $\beta$ signaling pathway	+	+
mTOR signaling pathway	+	+
MAPK signaling pathway	+	+
Non-small cell lung cancer	+	+
Melanoma	+	+
Pancreatic cancer	+	+
Acute myeloid leukemia	+	+
Chronic myeloid leukemia	+	+
Wnt signaling pathway	+	+
Adherens junction	+	+
Thyroid cancer	+	+
VEGF signaling pathway	+	+
p53 signaling pathway	+	+
Apoptosis	+	+
Jak-STAT signaling pathway	+	+
Bladder cancer	+	+
Cell communication	+	+
Cell adhesion molecules (CAMs)	+	+
Basal cell carcinoma	+	+
PPAR signaling pathway	+	+
Cell cycle	+	+

Only pathways mentioned in the KEGG database as being involved in human cancer are included. All of the pathways involved were found to be targeted by differentially expressed miRNAs, with the exception of the cytokine-cytokine receptor interaction pathway which was found to be influenced only by groups of up-regulated miRNAs (plus sign only for the left column of up-regulated genes).

pre-miRNA is referred as 'miRNA', whereas the less abundant strand is known as 'passenger strand' or miRNA\* (53,54). Three of the ten differentially expressed miRNAs in this study were miRNAs\*; hsa-miR-29b-1\*, hsa-miR-181a-2\* and hsa-miR-221\*. To date, there are no studies conducted to determine and validate the function of these miRNA\*.

This research enables the identification of miRNAs that have the potential to be used as cancer markers in head and neck cancers. However, in order to identify miRNAs that could be used as markers in head and neck cancers, more miRNA expression profiling studies with a larger number of tumor samples are warranted. In addition, the differential expression of miRNAs in carcinoma samples suggests that miRNAs may be involved in cancer development (51). Furthermore, to clarify the role of individual miRNAs in head and neck cancers, the

genes, functions and pathways that are regulated by the differentially expressed miRNAs should be studied. Unfortunately, due to partial complementarity of miRNAs and their target sequences, target prediction is a very challenging task and sometimes imprecise, though with the aid of special prediction algorithms. Hence, miRNA functions, target genes and regulated pathways are in need for validation before being further exploited in cancer research.

*Cancer-related genes and pathways are targeted by miRNAs differentially expressed in head and neck cancers.* Single and combinatorial miRNA *in silico* analysis revealed that miRNAs dysregulated in our study targeted genes and pathways that are involved in cancers. The data obtained from DIANA-microT and DIANA-mirPath are summarized in Tables IV

and V, respectively. From the list, worth mentioning is the VEGF signaling pathway, which is influenced by all miRNAs differentially expressed in this study. The VEGF signaling pathway is well known as an important pathway in head and neck cancers and has been exploited in search for a targeted therapy of these diseases (55). Another interesting pathway regulated by our miRNAs is the p53 signaling pathway, a renowned pathway involved in tumor suppression. The p53 network suppresses tumor development through the coordinated activation of multiple transcriptional targets, and miRNAs differentially expressed in our study may act jointly with other effectors to hinder improper cell proliferation. Involvement of dysregulated miRNAs in these pathways demands future validation to prove their influence in head and neck cancers management.

In conclusion, this study is the first to describe a miRNA expression profile in Malaysian head and neck cancer patients with a focus on the identification of miRNAs differentially expressed in carcinomas vs. normal samples. Differentially expressed miRNAs in head and neck cancers vs. normal samples were identified through microarray. Up-regulation of some miRNAs from miR-181 family were observed, however, there is yet no functional study that validates its function in head and neck cancers. Three up-regulated and down-regulated miRNAs in head and neck cancers which have not been reported in other similar studies were revealed, hsa-miR-744, hsa-miR-141 and hsa-miR-101. In addition, simultaneous hierarchical clustering performed on the samples was able to distinguish cancers from normal samples. *In silico* analysis revealed that differentially expressed miRNAs in this study are involved in cancer-related pathways. Functional analysis shall follow to validate the target and functions of these miRNAs.

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