

Research Paper

Potential Urinary miRNA Biomarker Candidates for the Accurate Detection of Prostate Cancer among Benign Prostatic Hyperplasia Patients

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Abstract

MicroRNAs (miRNAs) are a class of short (~22nt), single stranded RNA molecules that function as post-transcriptional regulators of gene expression. MiRNAs can regulate a variety of important biological pathways, including: cellular proliferation, differentiation and apoptosis. Profiling of miRNA expression patterns was shown to be more useful than the equivalent mRNA profiles for characterizing poorly differentiated tumours. As such, miRNA expression “signatures” are expected to offer serious potential for diagnosing and prognosing cancers of any provenance. The aim of this study was to investigate the potential of using deregulation of urinary miRNAs in order to detect Prostate Cancer (PCa) among Benign Prostatic Hyperplasia (BPH). To identify the miRNA signatures specific for PCa, miRNA expression profiling of 8 PCa patients, 12 BPH patients and 10 healthy males was carried out using whole genome expression profiling. Differential expression of two individual miRNAs between healthy males and BPH patients was detected and found to possibly target genes related to PCa development and progression. The sensitivity and specificity of miR-1825 for detecting PCa among BPH individuals was found to be 60% and 69%, respectively. Whereas, the sensitivity and specificity of miR-484 were 80% and 19%, respectively. Additionally, the sensitivity and specificity for miR-1825/484 in tandem were 45% and 75%, respectively. The proposed PCa miRNA signatures may therefore be of great value for the accurate diagnosis of PCa and BPH. This exploratory study has identified several possible targets that merit further investigation towards the development and validation of diagnostically useful, non-invasive, urine-based tests that might not only help diagnose PCa but also possibly help differentiate it from BPH.

Key words: Prostate Cancer, Benign Prostatic Hyperplasia, Urine, MicroRNA, Biomarkers.

Introduction

Globally, PCa is the second most common, non-cutaneous cancer among men (1, 2). Various studies have observed a positive correlation between age and incidence of PCa (2, 3, 4). Environmental factors, particularly diet, have also been reported to potentially play a role in a man’s risk for PCa (5). A number of techniques are currently available for as-

sessing prostate health, the most common of which include digital rectal exam (DRE), trans-rectal ultrasonography (TRUS), and prostate specific antigen-based testing (PSA) (6, 7).

Since PSA is primarily produced in the prostate and is therefore relatively prostate specific, it was proposed that a sharp but consistent increase in serum

PSA was indicative of exponential prostate cell growth, as might occur in PCa (8). PSA test had been validated for detecting PCa using any serum level above the empirically determined cutoff of 4ng/mL (9). PSA's use as a marker for PCa was soon limited by the many instances of BPH that were also associated with an increased serum PSA above the 4ng/mL cutoff, and especially in the range of 4-10ng/mL (10).

Serum PSA testing for the early detection of PCa has a sensitivity of around 86% and a specificity of approximately 33%, depending on patient's age and local prevalence of disease (11, 12). These problems have formed the basis for much controversy surrounding PSA's use as a regular blood-based screening tool for PCa (13, 14). As such, the search for novel PCa biomarkers, especially from non-invasive fluids, has been ongoing (15, 16).

MicroRNAs (miRNAs) are a class of short (~22nt), single stranded RNA molecules that function as post-transcriptional regulators of gene expression (17). MiRNAs have been found to play a significant role in carcinogenesis (18). They do so via their ability to function as regulators of tumour suppressors and oncogenes (19). MiRNA expression "signatures" are expected to offer serious potential for diagnosing and prognosing cancers of any provenance (20, 21).

Oncomirs, as they are known, function to down-regulate traditional tumour suppressor genes. An over-abundance of such microRNAs has been implicated in the progression of certain cancers (22). For example, PTEN, a tumour-suppressor gene that is responsible for promoting apoptosis via the PI3K-AKT-mTOR pathway, is a predicted target of microRNAs belonging to the miR-17-92 cluster (23, 24). While miRNAs such as those in the 17-92 cluster have been observed functioning as oncogenes, other miRNAs such as let-7 have been observed acting like tumour suppressors (22). MiRNAs that function as tumour suppressors do so by targeting traditional oncogenes, i.e. genes that promote cellular proliferation or that block apoptosis. For example, a number of human cancers have been shown to harbour a deletion in the chromosomal region encoding the let-7 microRNA (25). In particular, down-regulation of let-7 has been observed in several human lung cancers (26).

While differences in methodology and sampling likely form the basis for such inconsistencies, it's through studies like these that an association between PCa and several miRNAs has already been tentatively established (27). To date however, no miRNA-based diagnostic test has yet been approved for the early detection of PCa. In this study, the miRNA expression profiles in urine samples collected from PCa patients, BPH patients as well as from healthy male group were

analyzed. MiRNA profiling was firstly carried out on miRNA isolated from urine samples pooled from each group using LC Sciences patented microarray technology. Furthermore, we evaluated the deregulation of 2 miRNAs, revealed from our data, as potential non-invasive urinary biomarkers for the accurate diagnosis of prostate cancer among Benign Prostatic Hyperplasia Patients.

Materials and Methods

Urine sample collection.

After approval from the Research Ethics Board at Brock University (St. Catharines, ON, Canada), urine samples were collected from 29 Egyptian males present at Alexandria University's General Hospital (Alexandria, Egypt). Urine samples were collected into 50mL Corning tubes containing Norgen's Urine Preservative Solution (Cat# 18126) (Norgen Biotek, Thorold, ON, Canada). The samples were acquired from among three groups of individuals, each described by a questionnaire completed by the patients' attending physician. Group I urine samples were collected from eight individuals diagnosed with Prostate Cancer (PCa). Group II urine samples were collected from twelve individuals with Benign Prostatic Hyperplasia (BPH). Urine samples from Group III were collected from ten healthy males (Table 1). Urine samples collected from men with other ailments, including Diabetes mellitus, kidney disease, and/or infections such as chronic HBV, HBV/HCV co-infection or Schistosomiasis, were excluded from participation. Each man included in the study was also given a PSA test as part of sample collection. Furthermore, urine samples were collected from the same patients two years from their initial participation in the study; a PSA test was also performed for this second collection.

Table 1. Clinical parameters regarding the males involved with this study.

	Healthy Males	Prostate Cancer Patients	Benign Prostatic Hyperplasia Patients
Number of males	10	8	12
PSA Level (ng/mL)	(0.8 ± 0.6)	(7.9 ± 3.0)	(4.9 ± 1.4)
HCV or HBV Status	-Ve	-Ve	-Ve
Diabetes mellitus	-Ve	-Ve	-Ve
Schistosomiasis	-Ve	-Ve	-Ve
Kidney diseases	-Ve	-Ve	-Ve

Isolation of total RNA from urine.

Total RNA was isolated from 2.5mL of urine using the Urine Total RNA Purification Maxi Kit (Slurry Format) (Cat# 29600) (Norgen Biotek,

Thorold, ON, Canada). The isolation was performed according to the manufacturer's instructions.

MicroArray-based MicroRNA Expression Profiling in Pooled Samples.

Differential expression analysis of miRNAs between the pooled samples was performed using LC Science's μ Paraflo™ MiRNA Microarray technology.

Relative expression profiling for candidate miRNAs using RT-qPCR.

Relative miRNA expression levels for the 5 candidate miRNAs (miR-1234, miR-1238, miR-1913, miR-486-5p, miR-1825, miR-484 and miR-483-5p) were analyzed among the PCa group, BPH group and the healthy male group. Candidate miRNAs were reversibly-transcribed using 3 μ l of RNA template and 0.5 μ l gene-specific stem-loop RT primer (Table 2). The reaction was then made up with RNase-free water to 10 μ l and incubated for 5 mins at 70°C and was then held at 4°C. During the 4°C incubation, a mix of 4 μ l 5X first strand Buffer, 2 μ l of 0.1mM DTT, 1 μ l of 10mM dNTPs, 0.5 μ l SuperScript III (Invitrogen) and 2.5 μ l nuclease-free water were added to the initial reaction for a final reaction volume of 20 μ l. The final reaction was then incubated at 25°C for 5 min, 50°C for 30 min, 70°C for 15 min with a final hold at 4°C. For controls, minus All samples were analyzed using the iCycler iQ real time PCR detection system with iQ SYBR Green Supermix (BioRad Laboratories). For each sample, 3 μ l from its corresponding RT reaction were mixed in a 10 μ l iQ SYBR Green Supermix, 6.76 μ l nuclease-free water and 0.12 μ l each specific forward and reverse miRNA primer (Table 2) in a total of 20 μ l. The expression levels were normalized to 5S rRNA levels. Relative gene expression was expressed as the LOG₂ of fold change. Fold changes were calculated using the $\Delta\Delta$ Ct method (28).

Results

MicroArray-based MicroRNA Expression Profiling in Pooled Samples

Total RNA was isolated from 2.5mL of each urine sample using the Urine Total RNA Purification Maxi Kit (Slurry Format, Cat# 29600; Norgen Biotek, Thorold, ON, Canada). Purified RNA from each sample within a group was then pooled, resulting in three samples: PCa, BPH and Healthy Control. Differential expression analysis of miRNAs between the pooled samples was performed using LC Science's μ Paraflo™ MiRNA Microarray technology. To determine fold change, the signal intensity of each miRNA (in pooled PCa or BPH) was divided against the corresponding miRNA level for the healthy con-

trol group and the LOG₂ of the quotient was then obtained.

Table 2. Synthetic oligonucleotides used in this study.

Name	Sequence
5S rRNA (Forward)	5'GCCATACCACCTGAACG3'
5S rRNA (Reverse)	5'AGCCTACAGCACCCGGTATT3'
SL-miR-486-5p	5'GTCGTATCCAGTGCAGGGTCCGAGG TATTCGCACTGGATACGACCTCCG3'
miR-486-5p (Forward)	5'TCCTGTACTGAGCTGCC3'
SL-miR-1825	5'GTCGTATCCAGTGCAGGGTCCGAGG TATTCGCACTGGATACGACGGAGA3'
miR-1825 (Forward)	5'TCCAGTGCCTCCTCT3'
SL-miR-484	5'GTCGTATCCAGTGCAGGGTCCGAGG TATTCGCACTGGATACGACATCGG3'
miR-484 (Forward)	5'TCAGGCTCAGTCCCCTC3'
SL-miR-1238	5'GTCGTATCCAGTGCAGGGTCCGAGG TATTCGCACTGGATACGACGGGGC3'
miR-1238 (Forward)	5'CTTCCTCGTCTGTCT3'
SL-miR-1913	5'GTCGTATCCAGTGCAGGGTCCGAGG TATTCGCACTGGATACGACTGGCA3'
miR-1913 (Forward)	5'TCTGCCCCCTCCGCTGC3'
SL-miR-1234	5'GTCGTATCCAGTGCAGGGTCCGAGG TATTCGCACTGGATACGACGTGGG3'
miR-1234 (Forward)	5'TCGGCCTGACCACCCAC3'

Of the 894 miRNAs assayed, seventeen were found to differ significantly in their levels between the three groups (ANOVA p-value < 0.01). These miRNAs included: miR-1826, miR-1234, miR-1238, miR-191*, miR-484, miR-191, miR-451, miR-486-5p, miR-1913, miR-625*, miR-1825, miR-92a, miR-92b, miR-483-5p, miR-486-3p, miR-563, miR-940 (Figure 1). Of these seventeen, all but miR-483-5p was down-regulated in PCa with respect to the healthy control group. Similarly in BPH, miR-1826 were both found to be up-regulated, and the rest down-regulated with respect to the healthy control group. From the microarray results: each of the seventeen miRNAs except for miR-1826, were deregulated in the same direction in both PCa and BPH (when each pool was compared to the healthy control pool).

From these seventeen, only certain miRNAs were selected for further analysis. Based on published literature suggesting that a miRNA's mRNA target was potentially involved in cancer (targets determined using TargetScan and MiRanda databases as outlined by Griffiths-Jones *et al.* in 2006) (29), seven of miRNAs were selected for further expression analysis among individual urine samples. The selected miRNAs included miRs: 1234, 1238, 1913, 486-5p, 1825, 484 and 483-5p. Their putative mRNA targets and the literature proposing each target's potential role in cancer are summarized in Table 3.

Validation of MicroRNA Deregulation among Individual Prostate Cancer Samples using RT-qPCR

Seven of the selected miRNAs were, according to microarray data on pooled samples, deregulated in PCa with respect to the healthy control group. In order to evaluate those miRNA's potential as candidate biomarkers for PCa, the relative expression of each was examined in all individual PCa samples by using RT-qPCR. To allow for a comparison of expression levels between samples, the expression of each miRNA was normalized against each sample's own content of 5S ribosomal RNA. Relative expression between each PCa sample and the average expression of the healthy control group was then calculated as LOG₂ fold change values using the following equation: LOG₂ (2^{-ΔCt} (target miRNA) PCa / 2^{-ΔCt} (target miRNA) Healthy Control). For PCR data, an absolute LOG₂ value of one or more was considered deregulated, while a value of less than one was considered minimally deregulated / unchanged. Table 4 lists the relative expression of each of the seven miRNAs, for each of the initially collected PCa samples and with respect to the average expression of the healthy control group.

Interestingly a substantial majority of PCa samples expressed the same type of deregulation for a miRNA: miR-1825 and miR-484. MiR-1825 was

up-regulated in seven out of eight PCa samples (88%); furthermore in the eighth PCa sample miR-1825 expression was unchanged with respect to healthy. PCR results showed that miR-484 was down-regulated in six out of the eight PCa samples (75%), up-regulated in one and unchanged in another. Figure 2 illustrates these percentages visually.

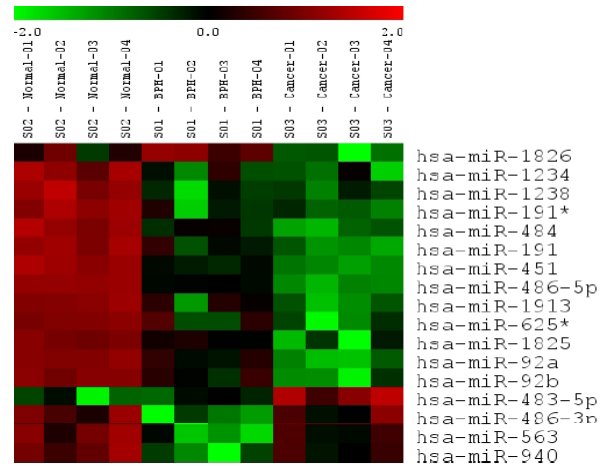


Figure 1. Expression of Eighteen Significantly Deregulated MiRNAs between Pooled PCa, BPH and Healthy by Microarray.

Table 3. Selected microRNAs and their Putative Targets.

MicroRNA	Type of Deregulation	Putative Target	Reference
miR-1234	Down in PCa & BPH vs Healthy	DAB2-Interacting-Protein	30, 32, 33
miR-1238	Down in PCa & BPH vs Healthy	SASH1 (SAM & SH3 domain containing protein 1)	33
miR-1913	Down in PCa vs BPH vs Healthy	HSPG2 (heparan sulphate proteoglycan 2)	34
miR-486-5p	Down in PCa vs BPH vs Healthy	CD40 (tumour necrosis factor receptor)	35
miR-1825	Down in PCa vs BPH vs Healthy	DDR1 (discoidin domain receptor tyrosine kinase 1)	36, 37, 38
miR-484	Down in PCa vs BPH vs Healthy	UBR5 (ubiquitin protein ligase E3 component n-recognin 5)	39
miR-483-5p	Up in PCa vs BPH & Healthy	Selenoprotein O	40

Table 4. LOG₂ fold change in the expression of the 7 miRNA among PCa patients relative to average healthy expression. (Red – Two fold or more up-regulated, Green – Two fold or more down-regulated, and Black – Minimally Deregulated / Unchanged).

Sample ID	LOG ₂ Fold Difference (PCa / Average Healthy Control)						
	miR-1234	miR-1238	miR-1913	miR-486-5p	miR-1825	miR-484	miR-483-5p
PCa 1	0.46	4.50	2.46	-0.31	5.54	-10.48	0.04
PCa 2	2.26	-6.60	3.06	2.89	10.54	0.16	0.34
PCa 3	-5.84	-4.10	-8.12	-5.81	5.74	-7.73	-8.83
PCa 4	-0.14	-7.41	1.06	1.69	6.64	-1.64	-6.86
PCa 5	-3.54	-3.10	-2.64	-3.71	1.04	-6.24	-6.06
PCa 6	0.96	2.40	-1.04	0.89	5.04	1.06	-2.66
PCa 7	2.46	1.80	0.86	1.49	5.84	-1.64	5.04
PCa 8	-3.94	-13.29	-2.04	-4.31	0.44	-5.04	-4.86

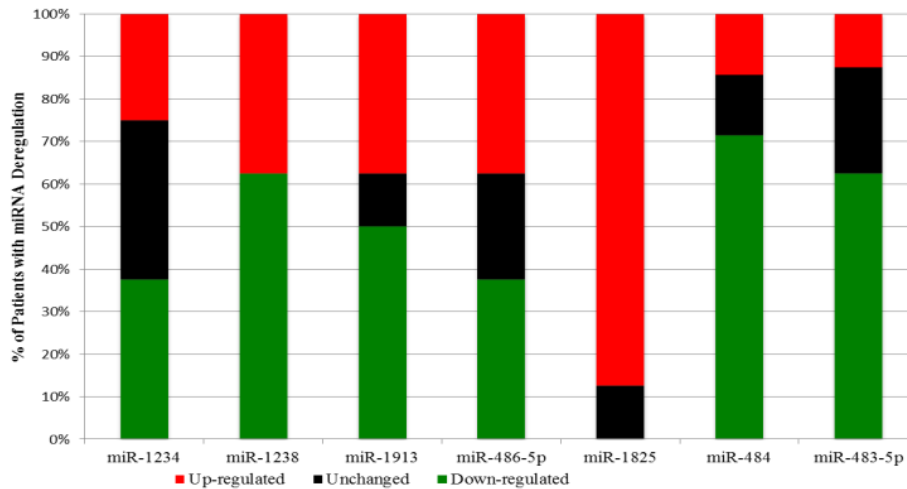


Figure 2. Percentage of the Initial PCa Samples with two fold or more deregulation of a microRNA.

Screening for Deregulation of miR-1825 and miR-484 in BPH Samples by RT-qPCR

Since only miR-1825 and miR-484 showed consistent up- and down-regulation across a majority of PCa samples by PCR (88% and 75%, respectively), they were selected for further evaluation among BPH samples; a group of patients who are usually falsely identified as having PCa by PSA testing. Table 5 displays the LOG₂ fold changes for the expression of miR-1825 and miR-484 among the initially collected BPH samples with respect to the average expression in the healthy control group. MiR-1825 was found to have variable expression between BPH samples. In particular miR-1825 was found to be up-regulated in 4 of the 12 samples or 33% of them, minimally deregulated or unchanged in 6 (50%) and down-regulated in 2 (17%). MiR-484 on the other hand was found to be two or more fold down-regulated in all of the initially collected BPH samples (100%). Figure 3 illustrates the percentage of initially collected BPH samples with up-regulation or miR-1825 and down-regulation of miR-484.

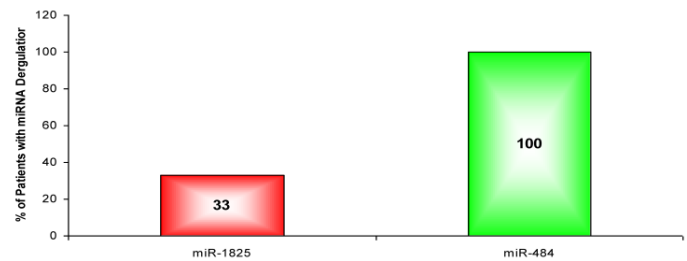


Figure 3. Percentage of BPH patients with two or more fold deregulation of miR-1825 or miR-484. (Red – Two fold or more up-regulated, Green – Two fold or more down-regulated).

Table 5. LOG₂ fold changes of the expression of miR-1825 & miR-484 among BPH patients relative to average healthy expression. (Red – Two fold or more up-regulated, Green – Two fold or more down-regulated, and Black – Minimally Deregulated).

Sample ID	LOG ₂ Fold Difference (BPH / Average Healthy Control)	
	miR-1825	miR-484
BPH1	2.44	-1.74
BPH2	4.04	-4.54
BPH3	0.34	-3.54
BPH4	0.64	-3.64
BPH5	-2.86	-3.84
BPH6	-0.36	-2.64
BPH7	-0.06	-4.04
BPH8	-2.95	-7.43
BPH9	1.04	-3.74
BPH10	0.94	-2.74
BPH11	0.44	-1.34
BPH12	2.64	-2.64

Screening for the Status of miR-1825 and miR-484 in PCa Patients: Two Years Later

In order to validate the initial findings, miR-1825 and miR-484 were assessed for up- and down-regulation in urine samples that were collected two years from the initial collection and from the same individual PCa patients. Again, relative expression was examined by RT-qPCR and both transcripts were normalized against the intrinsic 5S-rRNA present in each sample. Relative expression between samples was then calculated as LOG₂ fold change values according to the following equation: LOG₂ (2^{-ΔCt (target miRNA) PCa} / 2^{-ΔCt (target miRNA) Healthy Control}). Table 6 lists the LOG₂ fold change between each PCa sample that was collected after two years and the average for the original healthy control group, for both miR-1825 and miR-484. While miR-1825 was initially found to be two or more fold up-regulated in 88% of the PCa samples, after two years it was only up-regulated among 50% of them or in four of the eight. MiR-484 meanwhile, while initially found to be down-regulated in 75% of PCa samples, was after two years down-regulated in 88% of them or in seven out of eight. Figure 4 illustrates the percentage of PCa

patients with up- and down-regulation of miR-1825 and miR-484 at each time point.

Screening for the Status of miR-1825 and miR-484 in BPH Patients: Two Years Later

Since up- and down-regulation of miR-1825 and miR-484 was common to urine samples collected from individuals with PCa, including samples collected from the same cohort two years later, it was decided to assay their expression in BPH samples collected two years later, again by RT-qPCR. It should be noted that upon follow-up, only eight of the original twelve BPH patients participated. Table 7 lists the LOG₂ fold change values for miR-1825 and miR-484 in the BPH samples that were collected after two years; expression is relative to the average expression of the healthy control group. MiR-1825 was found to be two or more fold up-regulated in two of the eight BPH samples or 25% of them, down from 33% previously. MiR-484 was found to be two or more fold down-regulated in four of the eight BPH patients (50%), down from 100% observed two years prior. Figure 5 illustrates the percentage of BPH patients with up- and down-regulation of miR-1825 and miR-484 at both time points.

Table 6. LOG₂ fold changes of the expression of miR-1825 & miR-484 among PCa patients relative to average healthy expression after two years. (Red – Two fold or more up-regulated, Green – Two fold or more down-regulated, and Black – Minimally Deregulated).

Sample ID	LOG ₂ Fold Difference (PCa / Average Healthy Control)	
	miR-1825	miR-484
PCa 1	2.24	-1.34
PCa 2	2.54	-1.24
PCa 3	6.24	-1.04
PCa 4	7.44	2.46
PCa 5	-4.16	-2.84
PCa 6	-1.66	-3.14
PCa 7	-6.45	-6.24
PCa 8	0.14	-2.94

Table 7. LOG₂ fold changes of the expression of miR-1825 & miR-484 among BPH patients relative to average healthy expression after two years. (Red – Two fold or more up-regulated, Green – Two fold or more down-regulated, and Black – Minimally Deregulated).

Sample ID	LOG ₂ Fold Difference (BPH / Average Healthy Control)	
	miR-1825	miR-484
BPH1	-1.16	-3.84
BPH4	0.74	0.96
BPH6	-2.86	-1.34
BPH7	2.14	2.16
BPH9	0.24	-0.74
BPH10	0.14	-2.44
BPH11	1.94	-1.64
BPH 12	0.84	-0.54

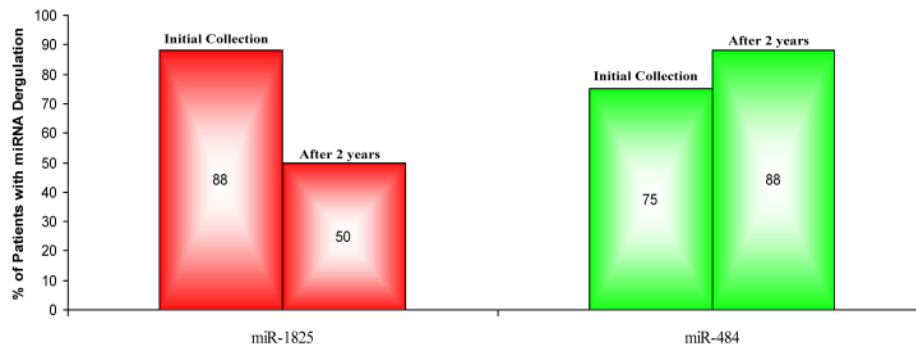


Figure 4. Percentage of PCa Patients with deregulation of miR-1825 or miR-484, initially and after two years. (Red – Two fold or more up-regulated, Green – Two fold or more down-regulated).

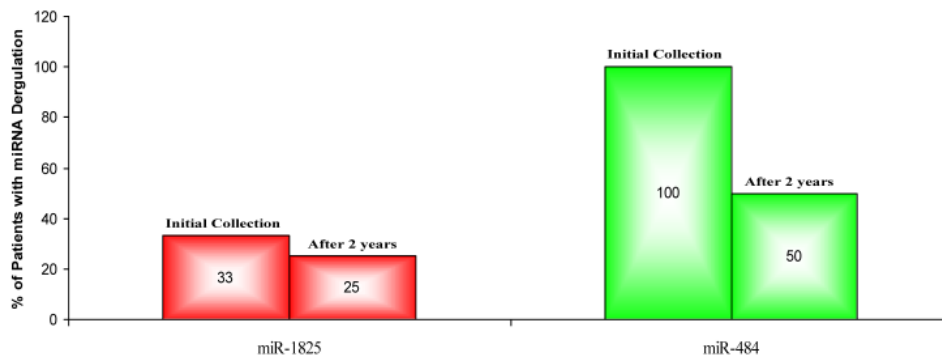


Figure 5. Percentage of BPH patients with two or more fold deregulation of miR-1825 or miR-484; both initially and after two years. (Red – Two fold or more up-regulated, Green – Two fold or more down-regulated).

Diagnostic Utility of miR-1825 and miR-484 for Diagnosing Prostate Cancer

To evaluate the potential of using aberrant expression of miR-1825 or miR-484 for diagnosing PCa the medical records, specifically PSA values and clinical diagnoses, were updated for all BPH patients as part of the two-year follow-up. It should be noted that all BPH patients were as a result diagnosed twice, both initially and after two years. Interestingly, during their second assessment it was discovered that four of the BPH patients had PCa (Table 8).

By comparing putative diagnoses (of PCa) made using miRNA deregulation against the actual clinical diagnoses, the sensitivity and specificity for using miR-1825 up-regulation and miR-484 down-regulation (either separately or in tandem) for diagnosing PCa were determined. The following criteria were applied: patients with a positive clinical diagnosis and exhibiting deregulation of either miR-1825 or miR-484 or deregulation of both were considered true positives; patients with a positive diagnosis and not exhibiting deregulation of any of the miRNAs (either separately or in tandem) were

considered false negatives; patients with a negative diagnosis and exhibiting deregulation of either miR-1825, miR-484 or deregulation of both were considered false positives; and finally, patients with a negative diagnosis and not exhibiting deregulation of either miR-1825 or miR-1825 or deregulation of both, were considered true negatives.

The above-mentioned criteria were applied to each sample that had a clinical diagnosis (including PCa samples), yielding the sensitivity and specificity values for diagnosing PCa. Using miR-1825 to diagnose PCa resulted in a sensitivity of 60% and a specificity of 69%, while using miR-484 to diagnose PCa resulted in a sensitivity of 80% and a specificity of 19%. Additionally, the sensitivity and specificity of using tandem deregulation of both microRNAs at the same time was 45% and 75%, respectively. Furthermore, the sensitivity and specificity of a panel, employing deregulation of both microRNAs as well as an abnormal PSA value (more than or equal to 4ng/mL) for detecting PCa was 40% and 81% respectively. Table 9 summarizes the sensitivity and specificity values for the various targets.

Table 8. Diagnosing PCa by Assessing Deregulation of miR-1825 or miR-484 among BPH Samples.

	Initial Urine Collection		Clinical Diagnosis for PCa	Serum PSA Levels (ng/mL)	Urine Collection After 2 Years			Serum PSA Levels (ng/mL)
	Test Result for PCa				Clinical Diagnosis for PCa	Test Result for PCa		
	miR-1825	miR-484				miR-1825	miR-484	
BPH1	PCa	PCa	Healthy	4.3	Healthy	PCa	PCa	3.9
BPH2	PCa	PCa	Healthy	5.6	N/A	N/A	N/A	N/A
BPH3	Healthy	PCa	Healthy	3.9	N/A	N/A	N/A	N/A
BPH4	Healthy	PCa	Healthy	5.8	Healthy	Healthy	Healthy	9.6
BPH5	Healthy	PCa	Healthy	2.4	N/A	N/A	N/A	N/A
BPH6	Healthy	PCa	Healthy	5.1	Healthy	PCa	Healthy	4.8
BPH7	Healthy	PCa	Healthy	6.9	PCa	Healthy	Healthy	4.2
BPH8	Healthy	PCa	Healthy	3.7	N/A	N/A	N/A	N/A
BPH9	PCa	PCa	Healthy	4.3	Healthy	Healthy	PCa	10.5
BPH10	Healthy	PCa	Healthy	5.1	Healthy	PCa	PCa	9.8
BPH11	Healthy	PCa	Healthy	7.4	PCa	PCa	PCa	4.2
BPH12	PCa	PCa	Healthy	3.8	Healthy	Healthy	Healthy	9.4

Table 9. Diagnostic Utility of miR-1825 and miR-484 for Diagnosing PCa.

Candidate Biomarker(s)	Sensitivity (%)	Specificity (%)
miR-1825	60	69
miR-484	80	19
miR-1825 and miR-484	45	75
miR-1825/miR-484/PSA	40	81
PSA (Experimental/Literature)	90/86	25/33

Discussion

MicroRNAs (miRNAs) are a class of short (~22nt), single stranded RNA molecules that function

as post-transcriptional regulators of gene expression (17). MiRNAs can regulate a variety of important biological pathways, including: cellular proliferation, differentiation and apoptosis (41). MiRNAs have also been found to play a significant role in carcinogenesis (18). They do so via their ability to function as regulators of tumour suppressors and oncogenes (19). As a result of their regulatory nature, patterns of miRNA expression appear to be tissue and even tumour specific (42). Moreover, profiling of miRNA expression patterns was shown to be more useful than the equivalent mRNA profiles for characterizing poorly differentiated tumours (21). As such, miRNA expres-

sion "signatures" are expected to offer serious potential for diagnosing and prognosing cancers of any provenance (20, 21).

Understanding the pathogenesis of viral-associated HCC is important for developing effective means of prevention and treatment of this highly malignant form of cancer (25). Currently, the identification of a non-invasive biological biomarker that can be used to screen high-risk patients is urgently needed to enhance HCC detection, resulting in earlier intervention and increasing the probability of successful treatment. Recent studies have already identified several miRNAs whose expression is deregulated in viral hepatitis infections and in HCC (29). Although important information was gathered about the hepatocarcinogenic effect of these deregulated miRNAs and the viral-host interactions that lead to the viral pathogenic effect, none of these studies evaluated the potential of using deregulated miRNA expression patterns to predict the development and progression of HCC among high-risk HCV infected patients.

The aim of this study was to investigate the potential for using deregulation of urinary miRNAs in order to accurately detect PCa among BPH patients. Profiling of pooled miRNA expression revealed that several miRNAs were differentially expressed between PCa, BPH and the healthy control group. Importantly, seventeen miRNAs were identified as being significantly deregulated among the three groups. In order to narrow the scope of the study, TargetScan and MiRanda databases were utilized for the identification of putative targets to the differentially expressed microRNAs, for the purpose of determining how each microRNA might be involved with cancer (23). This step revealed that microRNAs 1234, 1238, 1913, 489-5p, 1825, 484 and 483-5p were possibly playing a definable role in the development, progression or suppression of PCa.

Interestingly, miR-1234 (which was found to be down-regulated in PCa versus the pooled healthy control) was found to potentially regulate the Disabled homolog 2-interaction protein (DAB2IP). DAB2IP is thought to be a tumour suppressor gene; and is noticeably down-regulated in a number of PCa cell lines (30, 31, 32). A single nucleotide polymorphic variant of DAB2IP (rs1571801) is reportedly associated with an aggressive phenotype in PCa (43). Aberrant methylation of the DAB2IP promoter region has also been reported as a reason for its transcriptional silencing in PCa and hence progression of the disease (31).

Curiously, in this study's pooled data miR-1234 was observed to be down-regulated in PCa relative to the healthy control, as opposed to up-regulated as might have been expected if miR-1234 was acting to

silence the translation of DAB2IP, which appears to be common in some PCas. When expression levels were assayed by RT-qPCR among individual PCa samples, 25% actually showed up-regulation relative to the average healthy level. This is of course possible since the previously pooled samples represented an average signal for all patients. The up-regulation of miR-1234 that was observed among individual cancer samples might have been responsible for the down-regulation of DAB2IP. In 2003 Chen *et al.* (31) reported that down-regulation of DAB2IP in PCa was the result of aberrant methylation of the DAB2IP promoter region. Here it is proposed that the up-regulation of miR-1234 might be an alternative mechanism for the down-regulation DAB2IP.

Of some of the other deregulated microRNAs, miRs 1238 and 1913 might possibly regulate: SAM/SH3 domain-containing protein 1 (SASH1) and Heparan Sulfate Proteoglycan 2 (HSPG2), respectively, whereas miR-483-5p is predicted to target Selenoprotein-O and miR-486-5p, tumour necrosis factor receptor (CD40). Of these targets, SASH1 and HSPG2 are commonly affected in various cancers.

SASH1 has previously been reported as a candidate tumour suppressor gene, common to breast cancer (33). Down-regulation of *SASH1* has also been reported in aggressive and metastatic cancers, such that its down-regulation was prognostic of poor survival among breast cancer patients (33). Heparan Sulphate Proteoglycan-2 (*HSPG2*) on the other hand has been implicated in a variety of processes, including: maintenance of epithelial basement membrane integrity, growth factor binding and signaling, cellular differentiation, angiogenesis, neuro-muscular function and bone development. *HSPG2* can bind to several other basement membrane proteins including: laminin-1, fibronectin, nidogen, PRELP, and collagen IV via its core protein or heparin sulphate side-chains (44, 45). The loss of *HSPG2* and associated basement membrane structure is sometimes indicative of cancer, as can be the case in some invasive breast cancers (34).

In this study both miR-1238 and miR-1913 were found by PCR, to be over-expressed in 38% of the initial PCa samples. Their up-regulation therefore might potentially have been causing the down-regulation of their putative targets. This mechanism might help explain the observations of lower than normal levels of *SASH1* and *HSPG2*, as can be observed in some cancers (33, 34).

Another microRNA observed as deregulated by microarray, miR-1825, putatively targets member-1 of the Discoidin Domain family of Receptors (*DDR1*). This miRNA was identified as being up-regulated in 88% of the initial PCa samples, a seemingly common

event. Over-expression of DDR1 has been reported in many epithelial-cell cancers; DDR1 therefore likely plays an important role in tumourigenesis (36, 37). DDR1 over-expression was reported to be prognostic of poor survival in several cohorts of patients with brain, breast, or lung cancers (38). Since DDR1 is a putative oncogene, it is proposed that miR-1825 might function as a tumor-suppressor by normally or in response to cancer, inhibiting DDR1's translation. Since a majority of PCa samples (88%) showed up-regulation of miR-1825, it is possible that these cancers might have developed independently of DDR1. It is therefore proposed that deregulation of miR-1825 might be a potential mechanism for the deregulation of DDR1, an event common to some cancers.

MiR-484, another miRNA identified by microarray as being deregulated in PCa may regulate the expression of the E3 ubiquitin-protein ligase (UBR5). UBR5 is thought to be involved with DNA repair (46) and has even reportedly been associated with cancer progression (39). Clancy *et al.* reported on the over-expression of a potentially mutated version of UBR5 found in breast and ovarian cancers and whose deregulation was cited as a possible cause of malignant progression (39). Here RT-qPCR data show a down-regulation in miR-484 among 75% of PCa patients. This down-regulation might even cause the over-expression of miR-484's putative target, UBR5, possibly resulting in cancer progression. It is therefore being proposed that miR-484 might function as a tumor-suppressor whereby its expression controls progression via the regulation of genes, such as UBR5, that have already been potentially associated with cancer progression (39).

Based on the aforementioned trends regarding the observed deregulation of miRNAs among individual PCa samples, two of the seven miRNAs were selected for further evaluation among BPH samples. Because miR-1825 was observed to be up-regulated in 88% of the initial PCa samples and because miR-484 was observed to be down-regulated in 75%, it was thought that their deregulation might have been indicative of PCa. In support of this supposition, the putative target for miR-1825 is DDR1, a potential oncogene, while miR-484 potentially regulates UBR5, another putative oncogene. Since BPH is commonly misdiagnosed as PCa (12) and leads to many unnecessary biopsies, and because first time prostate biopsies can miss cancer approximately 20% of the time (47, 48), it became important to evaluate any potential biomarker in samples collected from BPH patients.

Using RT-qPCR, the levels of both miR-1825 and miR-484 were evaluated in all twelve urine samples that were initially collected from the BPH cohort;

none of whom had at that time yet been diagnosed with PCa. Data showed that (similar to the PCa samples) miR-484 was down-regulated in the majority of the initially collected BPH samples. Interestingly however, miR-1825 was only up-regulated in 25% of them (four BPH samples). These numbers coupled with the fact that prostate biopsy can occasionally miss PCa (47, 48) suggest that some of the BPH patients might have possibly had or been in the process of developing PCa; alternatively it may have been indicative of a lack of specificity. In order to distinguish between these possibilities, the medical records for many of the BPH patients were updated two years after the initial collection; as well additional urine samples were also collected.

Upon assessment of miRNA expression among those newly collected BPH samples, miR-1845 was only observed to be up-regulated in 25% of the cohort. In addition; miR-484 was also observed to be down-regulated, though this time in only 50% of BPH samples as opposed to 100% of them. Importantly, coupled to these new expression levels, the updated medical records indicated that four of the eight BPH patients had been diagnosed with PCa within the last two years.

By combining the original diagnoses and expression levels with the updated levels and medical records, the ability of miR-1845 and/or miR-484 deregulation to predict PCa was investigated. Towards this, the sensitivity and specificity of using miR-1825 up-regulation and/or miR-484 down-regulation to detect PCa was calculated. This calculation revealed that deregulation of miR-1825 was 60% sensitive and 69% specific for detecting PCa, whereas miR-484 deregulation was 80% sensitive but only 19% specific. When both were combined, the sensitivity and specificity was 45% and 75%, respectively. And finally, the sensitivity and specificity for a panel including: miR-1825, miR-484 and PSA (at a 4.0ng/mL cutoff) were 40% and 81%, respectively.

Despite the marginal diagnostic qualities of these microRNAs, the sensitivity of miR-484 was on par with - and the specificity of all except miR-484 were superior to - the literature reported values for PSA testing (sensitivity of 86% and specificity of 33% (12)). It is therefore proposed that these signatures might (with further validation) ultimately be useful for detecting PCa.

Competing Interests

The authors have declared that no competing interest exists.

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