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Epigenetic Inactivation of the miR-124-1 in Haematological Malignancies

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Abstract

miR-124-1 is a tumour suppressor microRNA (miR). Epigenetic deregulation of miRs is implicated in carcinogenesis. Promoter DNA methylation and histone modification of miR-124-1 was studied in 5 normal marrow controls, 4 lymphoma, 8 multiple myeloma (MM) cell lines, 230 diagnostic primary samples of acute myeloid leukaemia (AML), acute lymphoblastic leukaemia (ALL), chronic myeloid leukaemia (CML), chronic lymphocytic leukaemia (CLL), MM, and non-Hodgkin’s lymphoma (NHL), and 53 MM samples at stable disease or relapse. Promoter of miR-124-1 was unmethylated in normal controls but homozgyously methylated in 4 of 4 lymphoma and 4 of 8 myeloma cell lines. Treatment of 5-Aza-2’-deoxycytidine led to miR-124-1 demethylation and re-expression of mature miR-124, which also associated with emergence of euchromatic trimethyl H3K4 and consequent downregulation of CDK6 in myeloma cells harboring homozygous miR-124-1 methylation. In primary samples at diagnosis, miR-124-1 methylation was absent in CML but detected in 2% each of MM at diagnosis and relapse/progresison, 5% ALL, 15% AML, 14% CLL and 58.1% of NHL (p<0.001). Amongst lymphoid malignancies, miR-124-1 was preferentially methylated in NHL than MM, CLL or ALL. In primary lymphoma samples, miR-124-1 was preferentially hypermethylated in B- or NK/T-cell lymphomas and associated with reduced miR-124 expression. In conclusion, miR-124-1 was hypermethylated in a tumour-specific manner, with a heterochromatin histone configuration. Hypomethylation led to partial restoration of euchromatic histone code and miR re-expression. Infrequent miR-124-1 methylation detected in diagnostic and relapse MM samples showed an unimportant role in MM pathogenesis, despite frequent methylation found in cell lines. Amongst haematological cancers, miR-124-1 was more frequently hypermethylated in NHL, and hence warrants further study.

Introduction

DNA methylation involves the addition of a methyl group to the number 5 carbon of the cytosine ring in the CpG dinucleotide, by catalyzing the cytosine into methylcytosine through DNA methyltransferase [1,2]. Cancer cells are characterized by global DNA hypomethylation but gene-specific hypermethylation of promoter-associated CpG islands of tumour suppressor genes (TSGs), resulting in transcriptional repression, and hence serve as an alternative mechanism of gene inactivation. Based on a pathway-specific approach, multiple TSGs across pathways including cell cycle regulation, JAK/STAT signalling, Wnt1 signalling, and DAP kinase-associated intrinsic tumour suppression have been shown to be inactivated by gene hypermethylation in leukaemia, lymphoma and multiple myeloma [3,4,5,6,7,8,9,10,11].

MicroRNA (miR) is a single-stranded, non-coding RNA molecule of 22-25 nucleotides, which leads to downregulation of target protein expression [12]. miRs are involved in carcinogenesis. miRs can be either oncogenic (oncomir) when TSGs are targeted, or tumour suppressive (tumour suppressor miRs) when oncogenes are targeted [12,13]. Little is known about the role of hypermethylation of tumour suppressor miRs in haemic cancers.

Recently, miR-124-1 has been shown to be hypermethylated in multiple cancers [14,15]. By luciferase assay, miR-124-1 has been shown to downregulate CDK6 translation by binding on the 3’ untranslated region (3’ UTR) of the CDK6 mRNA, and also reduce the retinoblastoma protein phosphorylation, thereby demonstrating the tumour suppressor role of miR-124-1 [14].

In this study, we aimed to study the role of miR-124-1 methylation in a wide range of haematological malignancies including acute myeloid leukaemia (AML), chronic myeloid leukaemia (CML), acute lymphoblastic leukaemia (ALL), chronic lymphocytic leukaemia (CLL), multiple myeloma (MM) and non-Hodgkin’s lymphoma (NHL).

Materials and Methods

Patient samples

Diagnostic bone marrow or tissue samples were obtained in 20 ALL, 20 AML, 11 CML in chronic phase, 50 CLL, 55 MM and...
74 NHL patients. Diagnosis of leukaemia and lymphoma were made according to the French-American-British Classification and WHO Classification of Tumours respectively [16,17,10,19]. Of the 20 ALL patients, there were eleven male and nine female patients with a median age of 33 years (range: 13–62). There were six common ALL, one early B precursor, ten precursor B ALL, and three pre-T ALL. Of the AML patients, there were nine male and eleven female with a median age of 41.5 years (range: 20–72). The AML cases comprised three M1, fourteen M2, two M4 and one M5 subtype. Of the 50 CLL patients, there were twenty three (46%) patients with limited stage (<stage II) and twenty seven (54%) with advanced stage (≥stage II) disease with a median age of 65.5 years (range: 37–91) [3]. Forty (80%) were male. The median presenting lymphocyte count was 17×10^9/L (range: 10–256×10^9/L). Of the 55 MM patients, the median age was 57 (25–87) years. The diagnosis of MM was based on standard criteria [20]. Apart from five patients with insufficient clinical data, there were seven (14%) Durie-Salmon stage I, thirteen (26%) stage II, twenty-one follicular: grade 1 to 2, eight nodal marginal zone, three pre-T ALL. Of the AML patients, there were nine male and ten female patients with a median age of 35 years (range: 13–62). There were 30 female patients including those at stable disease, refractory relapse, repeated chemotherapy, a total of 53 serial samples from 12 MM patients including those at stable disease, refractory relapse, relapse, primary refractory disease or refractory disease progression were included. Of the 74 patients with NHL, there were 17 patients with peripheral T cell lymphoma (two anaplastic large cell [ALCL], four angio-immunoblastic T-cell [AITL], eleven peripheral T-cell, not otherwise specified [PTCL, NOS]), 10 with natural killer (NK)/T-cell lymphoma, 47 patients with B-cell lymphoma (twenty-one follicular: grade 1 to 2, eight nodal marginal zone, three mantle cell lymphoma and fifteen diffuse large B-cell lymphoma). The study has been approved by Institutional Review Board of Queen Mary Hospital with informed consent.

Cell lines and culture
Lymphoma (SU-DHL-1, SUP-M2, SUP-T1, and KARPAS-299) and MM (KMS-12-PE, MOLP-3, OPM-2, and U-266) cell lines were purchased from Deutsche Sammlung von Mikroorganismen und Zellkulturen GmbH (DSMZ), Braunschweig, Germany). LP-1 and RPMI-8226 were kindly gifts from Dr Orlowski (Department of Hematology/Oncology, MD Anderson Cancer Center, USA). WL-2 was kindly provided by Dr Andrew Zannettino (Myeloma and Mesenchymal Research Laboratory, Division of Haematology, Institute of Medical and Veterinary Science, Adelaide, Australia). NC-IH929 was purchased from American Type Culture Collection (ATCC). Cell cultures were maintained in RPMI media 1640 (IMDM for LP-1) (Invitrogen, Carlsbad, CA, USA), supplemented with 10% fetal bovine serum (Invitrogen, Carlsbad, CA, USA), 50 U/ml penicillin, and 50 μg/ml streptomycin (Invitrogen, Carlsbad, CA, USA) in a humidified atmosphere of 5% CO₂ at 37°C.

Methylation-specific polymerase chain reaction (MSP)
DNA was extracted from bone marrow samples of ALL, AML, CML, CLL, and MM at diagnosis, diagnostic tissues (either lymph node or nasal biopsy in nasal NK-cell lymphoma) in patients with NHL, and lymphoma and MM cell lines by standard method. MSP for aberrant gene promoter methylation was performed as previously described [8]. Treatment of DNA with bisulfite for conversion of unmethylated cytosine to uracil (but unaffected methylated cytosine) was performed with a commercially available kit (EpìTect Bisulfite Kit, Qiagen, Hilden, Germany). Primers used for the methylated MSP (M-MSP) and unmethylated MSP (U-MSP) were published previously [14]. DNA from normal bone marrow donors (N = 5) was used as negative control, while enzymatically methylated control DNA (CpGenome Universal Methylated DNA, Chemicon, Temecula, CA, USA) was used as positive control in all the experiments.

Bisulfite genomic sequencing (BGS)
Bisulfite-treated DNA was used as template. Promoter region of miR-124-1 was amplified and cloned using TOPO TA Cloning Kit (Invitrogen, Carlsbad, CA, USA) according to the manufacturer’s instructions. Primers used were published previously [14].

5-Aza-2’-deoxycytidine (5-AzadC) treatment
For treatment with 5-AzadC (Sigma-Aldrich, St. Louis, MO, USA), cells were seeded in six-well plates at a density of 1×10^6 cells/ml, and cultured with 1 μM of 5-AzadC for 3 days. Cells on day 0 and day 3 of 5-AzadC treatment were harvested.

RNA isolation and stem-loop reverse transcription-polymerase chain reaction (RT-PCR)
Total RNA was isolated using mirVana™ miR Isolation Kit (Ambion, Austin, TX, USA), according to the manufacturer’s instructions. RT was performed using Taqman® MicroRNA RT Kit and Taqman® MicroRNA Assay Kit (Applied Biosystems, Foster City, CA, USA), according to the manufacturer’s instructions. Total RNA was reverse transcribed in 1 mM dNTPs, 50 U MultiScribe™ Reverse Transcriptase, 1× RT Buff, 3.8 U RNase Inhibitor, and 1× stem-loop RT primer at following thermal cycling condition: 16°C for 30 minutes, 42°C for 30 minutes, and 85°C for 5 minutes. Quantitative real-time PCR of miR-124 was performed using 1.35 μl of 1:15 diluted RT product in 1× Taqman® Universal PCR Master Mix, and 1× Taqman® Assay at 95°C for 10 minutes, followed by 40 cycles of 95°C for 15 seconds and 60°C for 1 minute. RNU48 was used as reference for data analysis using the 2^(-∆∆Ct) method [21].

Chromatin immunoprecipitation (ChiP)
ChiP assays were conducted according to manufacturer’s instructions (Upstate, Cat# 17-610). Cells of 2×10^6 were fixed in 1% formaldehyde for each ChiP. Fixed cells were washed by cold PBS, resuspended in lysis buffer, and sheared into fragments ranging between 200 and 800 bp in size on ice using 431A cup horn (Misonix, Farmingdale, NY, USA). ‘Input’ of 1% was reserved as control, immunoprecipitation was performed by 4°C overnight incubation with anti-H3K1me3 (Upstate, 04-745), anti-H3Kme3 (Upstate, 17-625), anti-H3K8ac (Upstate, 17-658), and normal rabbit IgG respectively. Immunoprecipitated complex was collected by magnetic protein A beads. The complex was washed, treated with protease K, and reverse cross-linked by heat. Primers used for ChiP-PCR of miR-124-1 were forward: 5’- CAA AGA GCC TTT GGA AGA CG -3’ and reverse: 5’- GGA AGA GGG GTG GGT AGA AG -3’. ChiP-PCR was also controlled by GAPDH promoter and Alu repeats [22,23].

Western blot for CDK6
Cells were harvested and lysed in RIPA buffer (50 mM Tris-HCl, pH 7.4, 150 mM NaCl, 0.2% SDS, 1% Triton X-100, 2 mM EDTA) supplemented with protease inhibitors including 4 μg/ml aprotinin, 2 μg/ml leupeptin, 1 μg/ml pepstatin A, 20 μg/ml PMSF, and 1 mM Na₂VO₃. Cell debris was removed by centrifugation at 10,000 g for 5 minutes at 4°C. Protein lysate was denatured in an equal volume of loading buffer (100 mM Tris-HCl, pH 6.8, 200 mM DTT, 4% SDS, 0.2% bromophenol blue, 20% glycerol), heated at 95°C for 5 minutes, and separated...
Figure 1. Methylation of \textit{miR-124-1}. (A) Schematic diagram showing the distribution of CpG dinucleotides (solid vertical lines) over the precursor (solid black box) and mature \textit{miR-124-1}. Sequence analysis of the M-MSP product from bisulfite-treated positive control DNA showed that the cytosine \([C]\) residues of CpG dinucleotides were methylated and remained unchanged, whereas all the other C residues were unmethylated and were converted to thymidine \([T]\), indicating complete bisulfite conversion and specificity of MSP. Grey bars indicated the amplification regions of the MSP, ChIP, and BGS primers. (B) U-MSP showed that the methylated positive control \([P]\) was totally methylated, and all five normal controls \([N1–N5]\) were unmethylated. In the M-MSP, the methylated control was positive (methylated) but all normal controls were negative (unmethylated). For the cell lines, SUP-T1, SUP-M2 (ALK\(^+\)), SU-DHL-1 (ALK\(^+\)), KARPAS-299 (ALK\(^+\)), KMS-12-PE, LP-1, OPM-2, and WL-2 were completely methylated of \textit{miR-124-1}. (C) Bisulfite genomic sequencing for the bisulfite-treated promoter region of \textit{miR-124-1} of normal controls \([N1–N5]\), lymphoma and myeloma cell lines of different methylation statuses (MM, UM, or UU), and the methylated positive control were depicted. Unmethylated (empty circle) and methylated (filled circle) CpG dinucleotides were shown by eight independent clones for each sample.

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Role of miR-124-1 Methylation in Haemic Cancers

A

MW: DNA ladder;
B: blank;
AML: acute myeloid leukaemia;
CML: chronic myeloid leukaemia;
ALL: acute lymphoblastic leukaemia;
CLL: chronic lymphocytic leukaemia;
MM: multiple myeloma;
NHL: non-Hodgkin’s lymphoma;
P: positive control with methylated DNA

B

MW | B | NHL7 | NHL8 | NHL9 | NHL10 | NHL11 | NHL12
---|---|-----|-----|-----|-------|-------|-------
U | M | U | M | U | M | U | M

MW | N | NHL13 | NHL14 | NHL15 | NHL16 | NHL17 | NHL18
---|---|-------|-------|-------|-------|-------|-------
U | M | U | M | U | M | U | M

MW | P | NHL19 | NHL20 | NHL21 | NHL22 | NHL23 | NHL24
---|---|-------|-------|-------|-------|-------|-------
U | M | U | M | U | M | U | M

MW | NHL25 | NHL26 | NHL27 | NHL28 | NHL29 | NHL30 | NHL31
---|-------|-------|-------|-------|-------|-------|-------
U | M | U | M | U | M | U | M

C

ΔCt

Methylated Unmethylated

miR-124-1 methylation status

MW: DNA ladder; B: blank; N: normal donor;
NHL: non-Hodgkin’s lymphoma;
P: positive control with methylated DNA
on 0.2 μm nitrocellulose membrane (Bio-Rad, Hercules, CA). The membrane was blocked at room temperature for 1 hour in 5% skim milk diluted in PBS-Tween 20 (0.5% v/v). The membrane was then incubated with CDK6 primary antibody (1:1000) at 4°C overnight with shaking. After washing 3 times of 15 minutes each in PBS-Tween 20 (0.5% v/v), the membrane was incubated with anti-mouse horseradish peroxidase conjugate secondary antibody (1:1000) at room temperature for 1 hour. After washing 3 times of 15 minutes each in PBS-Tween 20 (0.5% v/v), signals were detected by ECL Western blotting detection reagents (Amersham Biosciences, Buckinghamshire, UK) and exposed to X-ray film.

Statistical analysis

The frequency of miR-124-1 methylation in different types of haematological cancers was computed by Chi-Square or Fisher Exact test. In CLL, correlation between miR-124-1 expression and categorical variables (gender, histological subtypes, lineage [B, T or NK/T] and nodal/extranodal presentation) were studied in 49 patients with complete clinical data by Student’s t-test and Chi-square test (or Fisher Exact test) respectively. Overall survival (OS) is measured from the date of diagnosis to the date of last follow-up or death. OS of patients with limited Rai stage (stages 0, I and II) were compared to those with advanced stage (stage III and IV). Survival is plotted by the Kaplan-Meier method and compared by the log-rank test. All p-values were two-sided. In NHL, correlation between miR-124-1 methylation with continuous (mean age) and categorical variables (gender and Rai staging) were studied by Student t-test and Chi-square test (or Fisher Exact test) respectively. Overall survival (OS) is measured from the date of diagnosis to the date of last follow-up or death. OS of patients with limited Rai stage (stages 0, I and II) were compared to those with advanced stage (stage III and IV). Survival is plotted by the Kaplan-Meier method and compared by the log-rank test. All p-values were two-sided. In NHL, correlation between miR-124-1 methylation with continuous (mean age) and categorical variables (gender, histological subtypes, lineage [B, T or NK/T] and nodal/extranodal presentation) were studied in 49 patients with complete clinical data by Student’s t-test and Chi-square test (or Fisher Exact test) respectively. Moreover, in 25 primary B-cell NHL samples in which both DNA and RNA were available, the mean expression of miR-124-1 in methylated and unmethylated lymphoma were compared by the Student’s t-test.

Results

MSP

Controls. Direct sequencing of the M-MSP products from the methylated positive control showed the expected nucleotide changes after bisulfite treatment, therefore confirming complete bisulfite conversion and specificity of MSP (Figure 1A). None of the five normal control bowels showed aberrant methylation of miR-124-1 (Figure 1B). The positive and negative controls showed the expected MSP results (normal DNA: U-MSP positive/M-MSP negative; methylated DNA: U-MSP negative/M-MSP positive).

Lymphoma cell lines. The profile of methylation of miR-124-1 of lymphoma cell lines was shown in Figure 1B. SUP-T1, SUP-M2 (ALK+), SU-DHL-1 (ALK+) and KARPAS-299 (ALK+) were homozygously methylated for miR-124-1.

Myeloma cell lines. The profile of methylation of miR-124-1 of myeloma cell lines was shown in Figure 1B. Apart from MOLP-8 and RPMI-8226, which were completely unmethylated (UU) of miR-124-1, KMS-12-PE, LP-1, OPM-2, and WL-2 were homozygously methylated (MM) for miR-124-1, whereas NCI-H929 and U-266 were hemizygously methylated (MU) for miR-124-1.

Bisulfite genomic sequencing confirmed miR-124-1 hypomethylation in five normal controls, hypermethylation in the methylated positive control, and the corresponding methylation statuses (MM, MU, and UU) as detected by MSP (Figure 1C).

Primary samples at diagnosis. miR-124-1 hypermethylation was not detected in any of the CML. On the other hand, miR-124-1 methylation was found in one (2%) MM samples at diagnosis, one (2%) MM samples at relapse/progression, 1 (5%) ALL, 3 (15%) AML, 7 (14%) CLL and 43 (58.1%) NHL samples (p<0.001) (Figure 2A). Amongst the lymphoid malignancies, there was significantly more frequent miR-124-1 methylation in NHL than MM, CLL or ALL (p<0.001). In CLL, there was no correlation between miR-124-1 methylation and age (p = 0.79), gender (p = 0.99), diagnostic lymphocyte count (p = 0.09); Hb (p = 0.98), platelet count (p = 0.42), advanced Rai stage (≥stage 2) (p = 0.69) and death (p = 0.41). The projected OS in CLL patients with and without miR-124-1 methylation were 86% and 62% (p = 0.36). Amongst lymphoma samples, miR-124-1 was methylated in seven NK/T (70.0%), thirty-one B-cell NHL (66.0%), and five T-cell NHL (29.4%) (p = 0.023). However, miR-124-1 methylation did not correlate with age (p = 0.457), gender (p = 0.99) or Ann Arbor stage (p = 0.105) of the lymphoma patients.

In order to determine the role of miR-124-1 methylation on the expression of miR-124-1 in primary samples, we analyzed the methylation status and expression level in 25 primary B-cell NHL samples in which both DNA and RNA were available. By MSP and stem-loop qRT-PCR, 22 samples displayed methylated MSP signals and three were completely unmethylated (Figure 2B). Moreover, methylation of miR-124-1 was associated with a lower level of miR-124-1 expression, and hence a higher ΔCt (Ct miR-124-1 - Ct RNU48) (p = 0.01) (Figure 2C).

5-AzadC treatment of lymphoma and myeloma cells

SU-DHL-1, KARPAS-299, KMS-12-PE, and WL-2 cells were completely methylated for miR-124-1. Upon 5-AzadC demethylation treatment, miR-124-1 U-MSP signal emerged on day 3, with re-expression of mature miR-124-1 as shown by Taqman stem-loop qRT-PCR (Figure 3A). 5-AzadC treatment led to augmentation of euchromatic histone code with abundance of trimethyl H3K4 at miR-124-1 promoter region (Figure 3B); GAPDH promoter and Alu repeat element, with the inherent hypo- and hypermethylated DNA, were used as biological controls for euchromatin and heterochromatin configurations (Figure 3B). Finally, demethylation of miR-124-1 by 5-AzadC with miR-124-1 re-expression led to downregulation of CDK6 (Figure 3C).

Discussion

There are several observations.

Firstly, in this study, we showed that miR-124-1 is not methylated in normal blood cells but is hypermethylated in lymphoma and myeloma cell lines, which can be re-expressed upon demethylating treatment. In cancer, miRs may be hypermethylated by two patterns. First, tumour suppressor miRs are expected to be hypomethylated in normal cells but hypermethylated in cancer cells [24]. On the other hand, some miRs may be hypermethylated in both normal and tumour cells, and therefore, hypermethylation of these miRs is tissue-specific but not tumour-specific. For example, miR-127 and miR-373 are hypermethylated in both the normal and cancer cells [24,25]. Therefore, our data and those from Lujambio et al. showed that miR-124-1 is differentially methylated in cancer
Role of miR-124-1 Methylation in Haemic Cancers

A

MW: DNA ladder;
B: blank;
N: normal donor;
D0: day 0 in 1μM 5-AzadC;
D3: day 3 in 1μM 5-AzadC;
P: positive control with methylated DNA

B

C

KMS-12-PE

5-AzadC

0 μM
0.5 μM
1 μM

CDK6

actin

1.00
0.53
0.38

WL-2

5-AzadC

0 μM
0.5 μM
1 μM

CDK6

actin

1.00
0.46
0.15
cells but not normal cells, consistent with its tumour suppressor role [14]. In addition to miR silencing associated with miR-124-1 methylation, miR expression could be restored by miR-124-1 demethylation, which was associated with restoration of the euchromatin code trimethyl H3K4. Moreover, miR-124-1 re-expression after hypomethylating treatment was associated with downregulation of CDK6 expression, consistent with data that CDK6 is a target of translation repression by miR-124-1 [14]. Furthermore, to ensure ChIP specificity, we showed two trimethyl H3K9 antibodies of different preparations generated comparable results, which further controlled the ChIP technically together with input and IgG controls; in addition to GAPDH and β-actin repeat element, with the inherent hypo- and hypermethylated DNA, which served as biological controls for euchromatin and heterochromatin configurations.

Secondly, as miR-124-1 is localized to chromosome 8p, where loss of heterozygosity (LOH) is frequently found in various solid cancers [26,27,28,29], and certain subtypes of NHL including mantle cell [30], and small B cell lymphoma [31]. Therefore, miR-124-1 hypermethylation may collaborate with LOH to result in biallelic miR-124-1 inactivation in NHL, thereby fulfilling the Knudson’s hypothesis [32]. This was supported by the finding that miR-124-1 was preferentially methylated in lymphoma, in particular, in B- and NK/T-cell lymphomas, which was associated with a lower expression of miR-124. Therefore, methylation of miR-124-1 might be important in lymphomagenesis.

Fourthly, miR-124-1 was preferentially hypermethylated in NK/T-cell lymphoma, which is an Epstein-Barr virus–associated aggressive extranodal lymphoma more frequently encountered in Asia, and Central and South America [33]. Various tumour suppressor genes have been shown to be frequently hypermethylated in NK/T-cell lymphoma including p73, CDKN2A, CDKN2B, hMLH1 and RARB [34], but hypermethylation of CDK6 is associated with disease progression in multiple myeloma. Therefore, miR-124-1 might be important in lymphomagenesis.

In summary, miR-124-1 hypermethylation is tumour-specific, associated with gene silencing, which can be reversed by hypomethylating treatment. Re-expression of miR-124 by 5-AzaC treatment was associated with emergence of a partial euchromatin histone code and consequent downregulation of CDK6. Amongst haematological malignancies, miR-124-1 is preferentially hypermethylated in NHL (in particular NK/T-cell lymphoma), in which methylation of miR-124-1 was associated with a lower expression of miR-124, and hence warrant further study in lymphoma. Finally, in MM, despite frequent miR-124-1 methylation in myeloma cell lines, miR-124-1 methylation was infrequent in primary samples including relapse samples, and hence unimportant in myeloma pathogenesis.

Author Contributions

Approval of the manuscript: CSC KYW CCS FL LPC WWLL RI GKHL DJY. Sample acquisition: CSC CCS FL LPC WWLL GKHL. Conceived and designed the experiments: CSC KYW DJY. Performed the experiments: KYW. Analyzed the data: CSC KYW RI. DJY. Contributed reagents/materials/analysis tools: CSC RL. Wrote the paper: CSC KYW.

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