Onco-miR-155 targets SHIP1 to promote TNFα-dependent growth of B cell lymphomas

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INTRODUCTION

B cell survival and fate determination are strongly dependent upon phosphatidylinositol 3-kinase (PI3K) signalling (Alizadeh et al, 2000; Fruman, 2004). PI3K catalyses the conversion of membrane phosphatidylinositol-(4,5)-bisphosphate to phosphatidylinositol-(3,4,5)-trisphosphate (PIP3), which acts as a second messenger to recruit pleckstrin-homology domain containing adapters and kinases such as 3'-phosphoinositide-dependent kinase (PDK), AKT, phospholipase C γ2 (PLCγ2), Bruton’s tyrosine kinase (BTK), downstream of kinase (DOK) and others. Subsequent activation/inactivation of additional effectors including serum and glucocorticoid-inducible kinase (SGK), target of rapamycin (TOR), protein phosphatase 2A (PP2A), forkhead box O (FoxO) and Cyclin D/E mediates diverse biological responses such as survival, proliferation, migration, adhesion and differentiation. PI3K signals are antagonized by two lipid phosphatases: the 3'-inositol phosphatase (PTEN) and the 5'-inositol phosphatase (SHIP). It was recently discovered that in contrast to T cells, B cells do not undergo malignant transformation upon PTEN deletion (AV Miletic, A Mills, D Mills, IM Pedersen, H Morse, J Ravetch, S Bolland, RC Rickert, unpublished work). Similarly, deletion of SHIP1 in B cells is insufficient to generate B cell lymphoma. However, concomitant ablation of both SHIP1 and PTEN in murine B cells induces lethal lymphoma resembling DLBCL with 100% penetrance, revealing a novel role for SHIP1 as a tumour-suppressor (Miletic et al, unpublished work).

Disease in B cell non-Hodgkin lymphoma (B-NHL) patients is graded according to the International Prognostic Index (IPI), which (among other things) assigns low scores for the involvement of single lymph nodes, intermediate scores for progression to multiple lymph node involvement and high scores for systemic nodal and non-lymphoid metastases.
Significant effort towards understanding disease progression in human patients has suggested that B-NHL, in addition to multiple cancer-promoting genetic ‘hits’, is fuelled by non-specific stimuli such as B cell antigen receptor (BCR)-mediated recognition of undefined self-antigens or soluble survival factors. In support of this latter idea, B-NHL tumours often display several hallmarks of antigen-mediated clonal selection or cytokine dependence. DLBCL is clinically, morphologically and genetically a heterogeneous group of malignant proliferation of large lymphoid B cells that accounts for approximately 40% (25,000 cases/year) of adult NHLs (Coiffier, 2001).

Standard chemotherapy has been recently expanded from CHOP (Cyclophosphamide/Doxorubicin/Vincristine/Prednisolone) to R-CHOP with the inclusion of Rituximab, an anti-CD20 monoclonal antibody, which improved treatment success to an overall 3-year relapse free survival of DLBCL patients at 53–63.1% (Habermann et al, 2006; Pfreundschuh et al, 2008). Two prognostically different subgroups of DLBCL have been identified with distinct gene expression profiles either characteristic of normal germinal centre (GC) B cells or of activated memory B cells. The GC B-cell-like subgroup was correlated with a significantly better prognosis (5-year survival: 76%) in comparison to the activated B cell-like (ABC or non-GC) subgroup (5-year survival: 16%) (Alizadeh et al, 2000; Shipp et al, 2002). Furthermore, previous work by Rai et al and a very recent report published during the preparation of this manuscript revealed unique miR signatures for the two classes of DLBCL (Malumbres et al, 2009; Rai et al, 2008).

In the present study we show elevated miR-155 levels and concomitant suppression of SHIP1 expression in non-GC compared to the GC-DLBCL cell lines or primary tumour samples. Furthermore, we demonstrate that augmented miR-155 expression in DLBCL is a consequence of increased TNFα production rather than genomic mutation, and can therefore be reversed in vitro and in vivo through neutralization of TNFα, with subsequent inhibition of cell proliferation and tumour growth.

RESULTS

To determine whether a correlation exists between survival of patients with B cell malignancies and expression levels of PTEN and SHIP, we utilized ONCOMINE to query published cDNA array results. Analysis of data originating from gene expression studies by Alizadeh and Rosenwald (Alizadeh et al, 2000; Rosenwald et al, 2002) revealed that SHIP1 levels are significantly decreased in DLBCL compared to more indolent chronic lymphocytic leukaemias (CLL) or follicular lymphoma (FL) (Fig 1A). Strikingly, SHIP1 levels within the non-GC subset displayed strong correlation with overall survival (Fig 1B) suggesting that SHIP1 expression levels are useful prognostic indicators of survival among DLBCL patients.

In order to determine the molecular mechanisms that mediate the observed decrease in SHIP1 expression in DLBCL, we determined the methylation status of the SHIP1 promoter in 44 non-GC and GC-DLBCL specimens, as well as screened for mutations in the coding regions and splice sequences, but did not detect any differences among the samples (not shown).

We next considered the possibility that post-transcriptional regulatory events mediated by small non-coding RNAs might alter the expression of SHIP1 in these haematopoietic malignancies. Indeed, in addition to differential expression of coding genes, non-GC-type lymphoma cells express elevated levels of several miRs. Scanning of the SHIP1 3’-untranslated region (3’-UTR) revealed perfect sequence complementarity with the seed sequence of miR-155 (Fig 1C), an onco-miR whose ectopic expression gives rise to B cell malignancies (Costinean et al, 2006), but whose cellular targets have remained elusive.

Using a prototypic cell line model of the GC- and a non-GC-subtypes of DLBCL, Lawrie et al showed that in addition to miR-155, miR-221 and miR-21 were also over-expressed in non-GC-type but not GC-type lymphoma cells (Lawrie et al, 2007). To determine miR-155 and SHIP1-mRNA levels in primary DLBCL patient samples without concern of potential contamination by non-malignant tissue, we isolated tumour cells by laser-capture-microdissection (LCM) from frozen lymph node biopsies. As shown in Fig 1D, miR-155 expression was significantly higher in non-GC-DLBCL compared to GC-DLBCL, whereas SHIP1 mRNA levels were lower in non-GC-DLBCL, consistent with the notion of attenuation of SHIP1 expression by miR-155.

To establish a link between miR-155 and SHIP1 expression beyond mere correlation, the non-GC-DLBCL cell line OCILY-3 was transfected either with a non-specific control-miR, miR-1 or miR-155. Neither miR-1 nor the control miR affected SHIP1 mRNA levels, whereas introduction of miR-155 resulted in a clear decrease in SHIP1 mRNA (Fig 2A). More importantly, neutralization of endogenous miR-155 by means of a synthetic anti-miR resulted in a dramatic increase in SHIP1 mRNA compared to a transfection of a non-specific anti-miR control (Fig 2B, left panel), and is not unique to OCILY-3 cells, but is also observed in the OCILY-10 and Toledo cells, a widely used cell line model representative of non-GC-DLBCL (Fig 2B, middle and right panel). In contrast, miR-155 did not alter the expression of hSHIP2 protein (Fig S1 of Supporting Information).

Lastly, analysing SHIP1 protein levels in B-cells isolated from WT and miR-155 transgenic (tg) mice, we found that SHIP1 is greatly reduced in miR-155 (tg) B-cells as compared to their WT counterparts (not shown), further supporting the notion that miR-155 post-transcriptionally regulates SHIP1 expression.

To investigate whether miR-155 targets and represses SHIP1 directly through 3’-UTR interaction, we inserted the 3’-UTR of hSHIP1 into a reporter plasmid at the 3’ end of luciferase mRNA driven by a CMV promoter (WT 3’-UTR-luc) and analysed the effect of miR-155 on luciferase expression. As anticipated, co-transfection of miR-155 attenuated expression of luciferase from the WT 3’-UTR-luc reporter (Fig 2C, bars 1 and 3), whereas no inhibition was observed when a control miR (miR-CTL) was used (Fig 2C, bar 2). Similarly, a mutated miR-155 (miR-155mut) whose seed sequence had been altered, failed to suppress the luciferase activity originating from WT 3’-UTR-luc
plasmid (Fig 2C, bar 4). To further demonstrate the specificity of this interaction, we also generated a luciferase reporter in which the hSHIP1 3'UTR was modified to be complementary to the seed sequence of miR-155mut (Mut 3'UTR-luc). This reporter system was resistant to the inhibitory effects of miR-155 (Fig 2C, bar 7), but was repressed by miR-155mut (Fig 2C, bar 8). These results demonstrate that miR-155 inhibits SHIP1 expression by directly interacting with the hSHIP1 3'UTR.

miR-155 is one of only a few miRs whose expression has been shown to be regulated by extra-cellular ligands. O’Connell et al demonstrated that miR-155 expression can be induced in macrophages by stimulation with lipopolysaccharide (LPS) or TNFα (O’Connell et al, 2007), and analysis of the activation pathway indicated that miR-155 is an AP1-responsive transcript involving the JNK pathway. B cells not only produce TNFα (Endres et al, 1999; Pasparakis et al, 1996), but they upregulate miR-155 in response to this cytokine in a JNK-dependent manner similar to macrophages (Fig S2 of Supporting Information). As previous studies showed that patients with non-GC-DLBCL display elevated serum TNFα levels compared to those suffering from GC-DLBCL or other B cell malignancies (Pedersen et al, 2005), the possibility that non-GC-DLBCL cells create an auto-stimulatory loop leading to elevated miR-155 levels by producing TNFα appeared intriguing.

To test this hypothesis, three non-GC-DLBCL cell lines (OCILY-3, OCILY-10, Toledo) were cultured in the presence of etnacept (Enbrel®), an antagonistic soluble TNFα receptor that finds widespread clinical use in the treatment of inflammatory diseases such as rheumatoid arthritis and Crohn’s disease. Strikingly, this anti-TNFα regimen led to a substantial
reduction in miR-155 expression in all three cell lines, but did not occur in the GC-DLBCL cell line SUDHL-4 (Fig 3A). Similar results were obtained when infliximab (Remicade\textsuperscript{1}), a neutralizing humanized monoclonal antibody against TNF\textalpha, was used (not shown). Importantly, reduced miR-155 levels were accompanied by a concomitant increase in SHIP1 protein levels in the non-GC-DLBCL cells (Fig 3B).

The biological significance of these observations is evidenced by the finding that eternacept imposes significant antiproliferative effects upon the three non-GC-DLBCL cell lines, but did not produce any growth modulation in the GC-DLBCL cells, or in Daudi Burkitt lymphoma (Fig 3C).

Our results thus far demonstrated that in non-GC-DLBCL, elevated levels of miR-155, and consequent abrogation of SHIP1 expression, are mediated through autocrine stimulation of cells by TNF\textalpha, a proinflammatory cytokine whose serum levels are known to be elevated in DLBCL patients (Pedersen et al, 2005). To explore the potential efficacy of anti-TNF\textalpha regimen as a treatment for non-GC-DLBCL patients, we employed xenograft models in which non-obese diabetic/severe combined immunodeficiency (Nod/SCID) mice were subcutaneously inoculated with non-GC-DLBCL Toledo cells. Upon establishment of palpable tumours, the animals received either 100\(\mu\)g eternacept or solvent intravenously every three days, and tumour size was measured after 2, 4 or 6 days. As shown in Fig 4A, eternacept treatment resulted in a slight, but detectable inhibition in tumour growth at day 4, and produced a substantial reduction in tumour burden after 6 days. In concurrence, the analysis of the excised xenograft tumours revealed increased SHIP1 protein levels in tumours from eternacept-treated mice compared to tumours isolated from animals that received phosphate buffered saline (PBS) (Fig 4B).

**DISCUSSION**

Soon after the discovery of the first mammalian miR some 12 years ago, it became evident that this class of molecules plays a critical role in global gene regulation, and a likely impact on cellular survival and death pathways. It is not surprising that tremendous strides have been made in defining miRNAs expression profile in human cancers. High throughput analyses,
utilizing various analytical methods demonstrated that miR expression was commonly dysregulated in a multitude of human cancers (Calin & Croce, 2006; Lu et al, 2005; Roldo et al, 2006). miR expression profiling has shown promise in defining malignant status in retrospective studies. It has even been suggested that microRNA expression profiling can distinguish cancers according to diagnosis and developmental stage of the tumour to a greater degree of accuracy than traditional gene expression analysis.

More than 50% of annotated human miR genes are located in fragile chromosomal regions that are susceptible to amplification, deletion or translocation during the course of tumour development (Calin et al, 2004). Moreover, recent evidence indicates that some miRs function either as oncogenes or tumour suppressors (Esquela-Kerscher & Slack, 2006; He et al, 2005; Johnson et al, 2005). The first study documenting abnormalities in miR expression in tumour samples focused on CLL. Deletion of chromosome 13q14 is the most frequent chromosomal abnormality in this disorder. Croce and coworkers demonstrated that tumour suppressor activity is likely provided by two miRNAs, miR-15a and miR-16-1 (Calin et al, 2002). Cimmino et al identified a conserved site for miR-15a and miR-16-1 in the 3'-UTR of the bcl-2 mRNA, which encodes the anti-apoptotic protein Bcl-2 (Cimmino et al, 2005).

miR-155 was the first onco-miR described, whose expression in B cells alone is sufficient to trigger malignant transformation,
The paper explained

PROBLEM:
Diffuse large B-cell lymphoma (DLBCL) is clinically, morphologically and genetically a heterogeneous group of lymphomas involving malignant proliferation of large lymphoid B cells. DLBCL accounts for approximately 40% of adult non-Hodgkin lymphomas (NHL). Two prognostically different subgroups have been identified, with distinct gene expression profiles characteristic of either normal germinal centre (GC) B cells, or activated memory B cells with the GC B cell-like subgroup being correlated with a significantly better prognosis. Recent work has revealed unique miR signatures in each sub-group, but the relation of these to tumour growth and malignancy and their value as prognostic indicators has not been fully clarified.

RESULTS:
The authors identify the inositol phosphatase SHIP1 as a bona fide target of the oncogenic miR-155. They demonstrate that DLBCL cells display elevated levels of miR-155 and consequently diminished expression of SHIP1. They establish that both features are the result of autocrine stimulation by the pro-inflammatory cytokine TNF and go on to show that an anti-TNF regimen involving treatment with etanercept or infliximab is sufficient to reduce miR-155 levels and restore SHIP1 expression. These changes are accompanied by a reduction in cell proliferation and a substantial decrease in tumour burden in DLBCL xenografts.

IMPACT:
These findings strongly support the concept that cytokine-regulated miRs can function as a crucial link between inflammation and cancer, and illustrate the feasibility of anti-TNF therapy as a novel and immediately accessible (co)treatment for DLBCL.

MATERIALS AND METHODS

Cell culture
OCIly-3, OLLY-10 and Toledo cells were maintained in complete Iscove's DMEM (Dulbecco's modified Eagle's medium) + 20% human serum + 100 mg/ml penicillin/streptomycin + 2 mM L-glutamine (Invitrogen, Carlsbad, CA). SU1H-L-4 and Daudi cells were maintained in complete Roswell Park Memorial Institute (RPMI) medium + 10% foetal calf serum (FCS) + 100 mg/ml penicillin/streptomycin + 2 mM L-glutamine. Where indicated, cells were treated with 400 nM c-Jun N-terminal kinase (JNK) Inhibitor II (Calbiochem, San Diego, CA), and/or 20 ng/ml of human TNFα (Peprotech Inc., Rocky Hill, NJ).

Patient samples
Sections of frozen biopsies were stained for CD10 (clone: 56C6, Novocstra), Bcl-6 (clone: PCLBgp, DAKO) and multiple myeloma oncogene 1 (MUM1) (MUM1p, DAKO) in a Techmate 500 Immunostainer using DAKO Envision X5007 as a secondary antibody. CD10 positive or Bcl-6 positive, MUM1 negative samples were classified as GC B-LCL as described by Hans et al (Hans et al, 2004). For LCD, lymph node sections were stained with haematoxylin and eosin (H&E) staining kit (Molecular Machines Industries, Glattbrugg, Switzerland), and lymphoma cells (1,000 cells/patient specimen) were isolated using the MMi CellCut (MMI). All human material was handled in full compliance with NIH guidelines and IRB approval.

Luciferase Assays
The putative target site of the miR-155 seed sequence in the 3'-UTR of hSHIP1 (WT; 5'-AGCTTGGGGCTTCTTAATGCTTTCACCCCTCA-3') and 5'-CTAGTGAGGGGTGAAAGCATTAAGAAGCCCA-3' or a mutated variant (Mut; 5'-AGCTTGGGGCTTCTTAATGCTTTCACCCCTCA-3') and 5'-
293 T cells at 50% confluency were transfected in triplicate using lipofectamin with the pmiReport luciferase vectors and either miR-

CTATGGCGGCTGAAGCCAGACGCCAGCCCA-3\') were cloned into the pmirReport fire-destin near vector (Applied Biosystems, Foster City, CA). The miR

Transfection of miRs

RNA isolation and quantitative polymerase chain reaction (qPCR)

RNA was extracted using Trizol following the manufacturer’s protocol (Invitrogen), and miR kits (RT and TaqMan q-PCR primers, Applied Biosystems) were used for qPCR analysis of miR-155, U6 and U43 according to the manufacturer's instructions. qPCR analysis of SHIP1 and GAPDH mRNA was performed using specific primers for SHIP1 (SABiosciences, Frederick, MD) and GAPDH (Dharmacon). Relative expression was calculated using the comparative threshold cycle (Ct) method.

Western blot analysis

Cell lysates were subject to Western blot analysis using rabbit monoclonal antibodies against SHIP1, SHIP2 or GAPDH (Cell Signaling Tech, Danbars, MA), and blots were developed using ECL (GE Healthcare).

Xenografts

Nod/SCID mice were subcutaneously inoculated with 10^7 Toledo cells. Upon establishment of palpable tumours, the animals received either 100 μg etanercept or an equal volume of solvent intravenously every three days. Tumour size was measured after 2, 4 and 6 days. Tumours were excised at day 6, lysed and analysed by Western blot for SHIP1 expression. All experiments were approved by UCSD Institutional Animal Care Committee.

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The authors declare that they have no conflict of interest.

For more information

Lymphoma Research Foundation:  
http://www.lymphoma.org

Oncomine – Cancer profiling database:  
http://www.oncomine.org

References


