Nimboide targets BCL2 and induces apoptosis in preclinical models of Waldenström's macroglobulinemia

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Neem leaf extract (NLE) has medicinal properties, which have been attributed to its limonoid content. We identified the NLE tetranorterpenoid, nimboide, as being the key limonoid responsible for the cytotoxicity of NLE in various preclinical models of human B-lymphocyte cancer. Of the models tested, Waldenström's macroglobulinemia (WM) cells were most sensitive to nimboide, undergoing significant mitochondrial mediated apoptosis. Notably, nimboide toxicity was also observed in drug-resistant (bortezomib or ibrutinib) WM cells. To identify putative targets of nimboide, relevant in WM, we used chemoinformatics-based approaches comprised of virtual in silico screening, molecular modeling and target–ligand reverse docking. In silico analysis revealed the antiapoptotic protein BCL2 was the preferential binding partner of nimboide. The significance of this finding was further tested in vitro in RS4;11 (BCL2-dependent) tumor cells, in which nimboide induced significantly more apoptosis compared with BCL2 mutated (Jurkat BCL2Ser70-Ala) cells. Lastly, intraperitoneal administration of nimboide in WM tumor xenografted mice, significantly reduced tumor growth and IgM secretion in vivo, while modulating the expression of several proteins as seen on immunohistochemistry. Overall, our data demonstrate that nimboide is highly active in WM, as well as other B-cell cancers, and engages BCL2 to exert its cytotoxic activity.

Blood Cancer Journal (2014) 4, e260; doi:10.1038/bcj.2014.74; published online 7 November 2014

INTRODUCTION
Waldenström’s macroglobulinemia (WM) is an indolent Non-Hodgkin’s lymphoma characterized by clonal proliferation of lymphoplasmacytoid B-cells in the bone marrow, spleen and the lymph nodes. Patients can present with signs or symptoms associated with cytopenias, lymphadenopathy, splenomegaly and/or hepatomegaly. The malignant WM tumor clone characteristically produces and secretes excessive amounts of immunoglobulin M (IgM), which can cause hyperviscosity syndrome.1

The pathological behavior of WM cells relies on aberrant B-cell receptor and Toll-like receptor signaling and is intricately supported by malignancies in initiation of apoptosis.2,3 In this dynamic, a complex shift between functionally active proapoptotic and anti-apoptotic proteins from the BCL2 family occurs and bestows the malignant cells with a significant survival advantage.3–5 This defective apoptotic signaling is associated with aggressive clinical behavior, chemoresistance and a poor prognosis in many B-cell cancers, including WM.4–7 Thus, B-cell receptor- or Toll-like receptor-induced proliferation and abnormal apoptotic signaling act in concert for the assurance of the malignant WM cells survival.

There are currently no Food and Drug Administration approved treatments for WM, and the disease remains incurable. Management of symptomatic malignancy is currently conducted using therapies commonly employed for other lymphoid cancers, such as alkylating agents (cyclophosphamide), purine analogs (fludarabine), monoclonal antibodies (CD20 targeting rituximab) and the use of the proteasome inhibitor, bortezomib. However none of these strategies is curative and the median survival of WM patients is ~ 5 years.8

Over the last three decades ~ 70% of the Food and Drug Administration approved drugs have been developed from natural sources.9 Naturally occurring compounds remain a critical reservoir for drug development. Several are currently under investigation for cancer therapy either as whole compounds or their components.70,11 Neem is one such natural herb with demonstrable anti-cancer properties and is a source of several limonoids, which are a class of oxygenated triterpenes called tetranorterpenoids. These limonoids are responsible for the anti-tumor effects of neem leaf extract (NLE).12 Of the NLE tetranorterpenoids, it has been reported that nimboide is the most cytotoxic; however, its mechanism of action remains to be conclusively established. Being a primary constituent of NLE, nimboide is anticipated to trigger multiple cell death pathways through activation of apoptosis machinery (through BCL2 family proteins),13 induction of heat shock proteins and the tumor suppressor TP53.13–16

Unfortunately, because of their mechanistic ambiguity, many phytochemicals with observable anti-tumor efficacy are often precluded from being developed as viable therapeutic agents. The experimental deduction of a precise biological target for a particular ligand is a complicated and time-consuming process. Analogous to experimental target finding techniques (affinity chromatography, nuclear magnetic resonance), an emerging technology that capitalizes on computational approaches is increasingly being implemented to determine the biological

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Received 15 July 2014; revised 15 September 2014; accepted 18 September 2014
Institutional Animal Care and Use Committee of Mayo Clinic. A xenograft orientation.18 This in turn can be used to predict the binding of numerous biological proteins, a molecular model can be designed where the bioactive compound is ‘reverse docked’ to the protein to determine its binding site and preferred orientation.18 This in turn can be used to predict the binding affinity between the compound and the protein, which is quantitatively relayed by a scoring function.18 The utility of these rapid and cost-effective virtual methods has been applied to elucidate the ligand–target (including unperceived off-target) interactions for synthetic therapeutics such as PRIMA-1,20 4 H-tamoxifen,21 torcetrapib22 as well as organic compounds such as vitamin E21 catechin (a polyphenol)23 and baicalein (a flavonoid).24

Through a systematic approach combining in silico computational techniques and experimental methods, we investigated nimbolide to ascertain its exact protein targets and decipher its mechanism of anti-tumor activity in B-cancer cells. As such, this is the first mechanistic analysis of nimbolide demonstrating 1) its activity in WM cancer cells in vitro, 2) the use of virtual screening methods to classify its molecular targets and 3) its ability to be safely administered and produce anti-tumor effects in an in vivo model of hematologic malignancy.

MATERIALS AND METHODS

Cell lines, cell culture and animal experiments

Heparinized peripheral blood was obtained from healthy human donors (n = 3). Peripheral blood mononuclear cells from healthy human donors were isolated as previously described.13 WM (BCWM.1, MWCL-1 and RPCI-WM1) cell lines, multiple myeloma cell lines (U266, KMS11 and OPM2) and their unique corresponding bortezomib resistant (BR) or ibrutinib-resistant (IR) clones (BCWM.1/BR, BCWM.1/IR, MWCL-1/BR, U266/BR, OPM2/BR and KMS11/BR), which were developed in our laboratory, were used in experiments. In addition, BCL2-dependent leukemia cells (RS4;11), BCL2 KMS11/BR), which were developed in our laboratory, were used in experiments. (IR) clones (BCWM.1/BR, BCWM.1/IR, MWCL-1/BR, U266/BR, OPM2/BR and KMS11/BR), which were developed in our laboratory, were used in experiments. All cell lines were cultured in RPMI-1640 containing 10% FBS and penicillin (100 U ml⁻¹) and streptomycin (100 μg ml⁻¹). Culture medium was replaced every three days. Cell viability was maintained at ~90% and was measured by trypan blue exclusion assay using ViCell-XR viability counter.

All animal experiments were performed with the approval of the Institutional Animal Care and Use Committee of Mayo Clinic. A xenograft model of WM was established as previously described.25,26 Please see the Supplemental Materials & Methods for additional information.

RESULTS

NLE induces cell death in preclinical models of B-cell cancers and its cytotoxic effects are primarily mediated through nimbolide in a tumor-specific manner

We recently reported the proapoptotic effects of NLE in malignant CLL patient cells.13 To investigate if this effect was also evident in other B-cell cancers, we evaluated the activity of NLE and its individual limonoids in cancer cell lines representing human MM, Non-Hodgkin's lymphoma and WM in vitro. Tumor cell viability was determined by trypan blue exclusion assay after treating the cancer cell lines with NLE and its individual limonoids, nimbolide, azadirachtin or gedunin. In comparing the antineoplastic effects of NLE and its individual limonoids, nimbolide to ascertain its exact protein targets and decipher its mechanism of anti-tumor activity in B-cancer cells. As such, this is the first mechanistic analysis of nimbolide demonstrating 1) its activity in WM cancer cells in vitro, 2) the use of virtual screening methods to classify its molecular targets and 3) its ability to be safely administered and produce anti-tumor effects in an in vivo model of hematologic malignancy.

Figure 1. Nimbolide significantly reduces cell viability and induces apoptotic cell death in B-cell cancer models. MM (U266, U266/BR, KMS, OPM2, OPM2/BR), WM (BCWM.1, BCWM.1/BR) and Non-Hodgkin's lymphoma (Raji) cell lines were treated with NLE or its limonoid components for 24 h.

(a) Cell viability was determined by trypan blue exclusion assay. All cell lines showed significant sensitivity to nimbolide with maximum effect noted in the WM model. Cells from each representative model were treated with DMSO and used as a control. (b) MM, WM and Non-Hodgkin's lymphoma (Raji) cell lines were treated with indicated concentrations of NLE or its limonoid components and apoptosis was measured by ELISA (Roche, Indianapolis, IN, USA). All plasma cell cancer models demonstrated significant apoptosis in presence of nimbolide. Notably the proapoptotic effect of nimbolide was most significant in BR models. *p < 0.05; **p < 0.01; BR, bortezomib-resistant model.

Nimbolide was considerably lower (200 nM) (Supplementary Figure 1). Apoptosis ELISA confirmed that nimbolide was the most potent inducer of apoptotic cell death, and this was independent of the disease model studied (Figure 1b). This effect was evident even in the drug-resistant models; bortezomib sensitive cell lines as well as BR derivatives were equally susceptible to nimbolide. Although cell viability (Figure 1a) was most reduced in presence of NLE, programmed cell death (apoptosis ELISA using cell culture supernatants) was more evident in nimbolide treated cells, with WM models showing the greatest sensitivity. Observing that apoptosis was the primary mode of nimbolide-induced WM cell death, we treated WM cells with different concentrations of nimbolide ranging from 0.1 to 1 μM for 6 h and conducted annexin-V and PI staining followed by flow cytometry to assess the degree of apoptosis across the different models. Nimbolide was able to produce apoptotic cell death in a dose-dependent manner (Figure 2a). Using our most drug-resistant model (RPCI-WM1 that was established from a therapy-resistant and terminal WM patient),26 we noted significant induction of apoptosis with nimbolide (26% apoptosis). Interestingly, the ibrutinib-resistant (IR) WM subclones were the most sensitive with ~53% of the cells undergoing apoptosis (Figure 2b). Under similar experimental conditions, nimbolide exerted minimal effect on the induction of apoptosis in peripheral blood mononuclear cells from healthy donors (Figure 2c). Lastly, tumor
cell apoptosis was also confirmed on immunoblot analysis by detection of PARP-1, initiator caspase-9 and executioner caspase-3 cleavage, signifying involvement of the intrinsic (death receptor independent) apoptotic system (Figure 2d). All together, these experiments confirmed that nimbolide is the most potent component of NLE and it induces cytotoxicity through apoptotic mechanisms.

The mitochondrial transmembrane potential of WM cells is altered in presence of nimbolide

To further understand the mechanism of apoptosis induced by nimbolide in WM cell lines in vitro, we focused our study on the effect of nimbolide on mitochondrial integrity. Disturbance of the mitochondrial transmembrane potential (Δψm), through an increase in mitochondrial outer membrane permeability (MOMP) is a hallmark of apoptosis that occurs autonomous of death receptor activation.27 Thus we investigated whether Δψm was altered in the presence of nimbolide. Wild-type, BR and IR WM cell lines (total n = 7) were treated with increasing concentrations of nimbolide for 6 h followed by incubation with tetramethylrhodamine methyl ester (TMRM) for 15 min. MOMP was measured by examining TMRM fluorescence in tumor cells versus control cells (% MOMP) using flow cytometry. Nimibolide treatment resulted in dose-dependent MOMP activation in all the WM cell lines tested (Figure 3a). Nimibolide-treated cells showed substantially less TMRM fluorescence, which specified disruption of the Δψm and suggested mitochondrial leakiness and caspase release from mitochondria into the cytoplasm. To further confirm the consequence of compromised mitochondrial integrity, we investigated the activation of caspases 9 and 3, which signify the complete activation of the intrinsic apoptotic pathway. Both initiator caspase-9 and executioner caspase-3 cleavage was significantly increased in cells treated with nimibolide as compared with control untreated cells (see Figure 2d).

Nimbolide restores the cytotoxic activity of bortezomib or ibrutinib in drug-resistant WM models

We previously demonstrated that the proapoptotic effects of nimibolide are maintained despite drug resistance. In our BR model system we investigated if nimibolide could restore bortezomib sensitivity. On a 48 h MTS assay, we noted that nimibolide significantly decreased the established EC50 of bortezomib (50-fold) in BCWM.1/BR cells (Figure 4a). Similar effects were noted in our IR model, wherein; we assessed the ability of nimibolide to restore the IR cells sensitivity toward ibrutinib. We examined cell death in BCWM.1/IR cells using escalating doses of ibrutinib (5–20 μM). Even at the highest dose tested (20 μM), ibrutinib could only achieve 10.37% cell death. We observed that when a sub-lethal concentration of nimibolide (0.1 μM) was added to ibrutinib-treated BCWM.1/IR cells, the combination activity

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**Figure 2.** Nimibolide induces dose-dependent apoptosis in WM cells. (a and b) WM cell lines were treated with nimibolide at different concentrations for 6 h. Cell death was analyzed by annexin V staining followed by flow cytometry. Nimibolide induced apoptosis was dose dependent with maximum cell death at 1 μM, most notably in ibrutinib-resistant (BC/IR) cells (51% cell death). (c) Peripheral blood mononuclear cells from healthy human donors (n = 3) were treated with indicated concentrations of nimibolide and stained with annexin V to assess apoptosis. Toxicity from nimibolide was minimal ranging between 0.15 and 4.7% of % control cell death, with the latter only seen at the highest concentration tested (1 μM). (d) Western blot analysis of WM cells treated with nimibolide at 0 and 1 μM confirms apoptosis by cleavage of caspases 3, 9 and PARP-1. All experiments were performed in triplicate. NS, Not statistically significant; **P < 0.01. WM tumor cell lines: BC, BCWM.1; BC/IR, BCWM.1/IR; BC/BR, BCWM.1/BR; MW, MWCL-1; RP, RPCI-WM1
subsequently incubated with 20 μM TMRM for 15 min followed by 92.04 Å², solvent accessible surface area of 651.41 Å² and 9,9a,10a,10b,10c-decahydro-2 H,5 H cyclopenta[b]furo[2',3',4':4,5]
10cR)-8-(3-furyl)-2a,5a,6a,7-tetramethyl-2,5 dioxo-2a,5a,6,6a,8,
calculated geometric volume of 417.32 Å². In addition, nimbolide
has excellent characteristics exemplified by zero violations of
either Lipinski Rule of 5 or Jorgensen Rule of 3.

Chemoinformatics and molecular modeling of nimbolide–target
interactions identifies BCL2 as a primary binding partner of
nimboide

The rapid and potent cytotoxic effect of nimbolide on all the B-cell
cancer cell lines tested was intriguing and indicated that the
limonoid targets important survival pathways biologically relevant
to lymphoid cells and independent of acquired resistance to
bortezomib or ibrutinib. To further understand the intracellular
targets of nimbolide, we performed a chemoinformatics analysis
to determine its biochemical and molecular properties
(Supplementary Table 1). Briefly, nimbolide is a 466.52 Dalton
small molecule with an atomic formula of C$_{27}$H$_{30}$O$_7$ (chemical
structure shown in Supplementary Figure 2), which has the
IUPAC nomenclature methyl-[l2-ar5-ar6,5,6-ar8-r,9-ar10-as1,10br,
10 cr]-8-(3-furyl)-2a,5a,6a,7-tetramethyl-2,5 dioxo-2a,5a,6a,8,
9,9a,10a,10b,10c-decahydro-2 H,5 H cyclopenta[b]furo[2,3,3',4,4,5]
naphtho[2,3-d]furan-6-yl]acetate. It has a polar surface area of
92.04 Å², solvent accessible surface area of 651.41 Å² and
calculated geometric volume of 417.32 Å³. In addition, nimbolide
has excellent characteristics exemplified by zero violations of
either Lipinski Rule of 5 or Jorgensen Rule of 3.

Next, we conducted virtual screening using in silico target
identification coupled with detailed atom–molecular docking
experiments (reverse docking) to identify targets of nimbolide
relevant in WM. Molecular optimization algorithms to determine
relative ranking of protein affinity with nimbolide were utilized.38–37
Using the pre-docking filter LASSO (Ligand Activity
by Surface Similarity Order) as a broad screen, we first identified a
pool of over 50 proteins with which nimbolide potentially interacted (top 3 proteins shown in Supplementary Table 2). Next
we conducted a docking analysis on all proteins of interest and
their different isoforms (total protein targets, n = 86) from the
pre-docking screen. Using comprehensive algorithms (see
Supplementary Methods section) that ensured statistical
mechanics output and thermodynamic validity, we identified the
BCL2 protein proper as displaying the highest affinity for
nimboide
with a binding energy of -84.86 (MM-GBSA score).

We were encouraged to find that our in silico analysis corroborated prior findings that BCL2 is an important and
preferential target of nimbolide.13–16 This finding was validated
in WM cells, where BCL2 protein levels were assessed by western
blot and appeared to decrease after cell exposure to the limonoid
(Figure 6). Recognizing that nimbolide is a promiscuous molecule
(as many phytochemicals tend to be), with additional targets,18
we sought to examine other potential binding partners identified in
the virtual screening campaign. Whereas BCL2 ranked highest in
terms of binding affinity, HSP90 and PI3K (p110) ranked second
and third most likely by MM-GBSA calculations to interact with
nimboide
respectively (see Supplementary Table 2). In nimbolide
treated WM cells, we did not observe a direct shift in HSP90;
however, a marked induction of HSP70 and a decrease in MAPK
signaling (p-ERK1/2) was observed, which was indicative of
HSP90 inhibition.9,40 Next we looked for evidence of PI3K (p110) modu
lation in nimbolide-treated cells and found its downstream
targets, protein kinase B (AKT) and p-AKT, were notably decreased.
In addition to its well-known effects on AKT, PI3K inhibition has
been noted to mitigate NFκB signaling.41 We found a marked

**P < 0.01. WM tumor cell lines: BC, BCWM.1; BC/IR, BCWM.1/IR; BC/BR, BCWM.1/BR; MW, MWCL-1; RP, RPCI-WM1.

Figure 3. Nimboide alters MOMP in WM cells. (a and b) WM cell lines were treated with nimboide at different concentrations for 6 h and
subsequently incubated with 20 nm TMRM for 15 min followed by flow cytometry for analysis of MOMP. MOMP was most activated at 1 μM,
particularly bortezomib-resistant (BC/BR) WM cells (69% MOMP). All experiments were performed in triplicate. **P < 0.01. WM tumor cell lines:
BC, BCWM.1; BC/IR, BCWM.1/IR; BC/BR, BCWM.1/BR; MW, MWCL-1; RP, RPCI-WM1.
cells. To test this, we used RS4;11 tumor cells, which are BCL2-independent cells and show that BCL2 and other potential targets have not been assessed in in vivo models of WM (or any B-cell cancer). We hypothesized that systemically delivered nimbolide would engage its target(s) in vivo to induce WM cell death at pharmacologically achievable doses. Using our murine xenograft model of WM, we administered nimbolide via intraperitoneal injection in two separate cohorts of mice, testing two doses, 100 and 200 mg kg$^{-1}$ (see Supplementary section). Compared with the control group (DMSO-treated mice), we observed a 48% reduction in tumor volume in the nimbolide treated mice (100 mg kg$^{-1}$ cohort, $P < 0.05$; Figures 8a and b and Supplementary Figure 4) and a corresponding decrease in human IgM secreted by the xenografted WM cells ($P < 0.05$) (Figure 8c). A higher dose (200 mg kg$^{-1}$) had a more pronounced impact on IgM reduction but not on tumor volume (vs that observed in the 100 mg kg$^{-1}$ mice cohort). Immunohistochemistry (IHC) analysis was conducted on tumor tissues obtained from the animals treated with nimbolide (100 mg kg$^{-1}$) or DMSO alone. In vivo activity was validated by a notable decrease of IgM and BCL2 in nimbolide treated mice vs control. Likewise, an increase in tumor cell apoptosis (cleaved caspase-3 stain) and decrease in cell proliferation (Ki-67 stain) was noted in these tissues (Figure 8d). These data provide sufficient evidence that nimbolide can be safely administered in WM xenografted mice, elicit a significant anti-tumor effect and modulate important WM cell death and apoptotic pathways.

**DISCUSSION**

Nimbolide is a potent limonoid triterpene that is derived from the neem plant. It has been shown to produce anti-tumor responses in various preclinical solid tumor cancer models including melanoma, osteosarcoma, choriocarcinoma, cervical cancer, breast cancer, prostate adenocarcinoma, colorectal carcinoma and glioblastoma. However, the study of its activity in hematologic malignancies is limited; it has only been examined in MM, T-cell leukemia, myeloid leukemia and monocytic leukemia/lymphoma cell lines. Our report provides critical insight into the mechanisms of nimbolide’s cytotoxic potential in lymphoid cells and reveals that malignant WM (lymphoplasmacytic) cells are highly sensitive to the limonoid.

We have previously demonstrated that the anti-leukemic capabilities of NLE are mediated through caspase activation and engagement of the intrinsic apoptotic pathway. Being a powerful component of NLE, we anticipated that nimbolide would catalyze a similar cascade of death receptor-independent apoptotic signaling events in WM cells in vitro, as evidenced by cleavage of caspase-9. For activation of initiator caspase-9, destabilization of mitochondrial membrane permeability must occur. As such, we detected a dose-dependent increase in MOMP (indicating loss of Δψm) in nimbolide treated WM cells and this correlated with nimbolide-mediated cytotoxicity. Although we identified caspase-9 induction and cleavage in nimbolide treated cells, nimbolide has been shown to also engage the death receptor-dependent apoptotic pathway. These observations

**Antitumor effects of nimbolide in WM cells**

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BCL2-dependent cells are highly sensitive to the cytotoxic effects of nimbolide

With virtual screening and immunoblot analysis pointing toward BCL2 as a primary target of nimbolide, we hypothesized that the effects of the limonoid would be catastrophic in tumor cells that are completely reliant on BCL2. Conversely, we postulated that activity of the limonoid would be diminished in BCL2-independent cells. To test this, we used RS4;11 tumor cells, which are BCL2-dependent43 and Jurkat BCL2$^{Ser70-Ala}$ cells, which served as a BCL2-independent model. Neo Jurkat cells were used as a neutral control. As anticipated, RS4;11 cells were sensitive to nimbolide, undergoing significant apoptosis ($P < 0.00008$, 82% apoptosis at 1 μM) and activation of MOMP in a dose-dependent manner as compared with Jurkat BCL2$^{Ser70-Ala}$ cells (28% apoptosis at 1 μM) (Figures 7a and b). These corroborated our in silico observations that BCL2 is a central mediator of nimbolide cytotoxicity.

Nimbolide significantly inhibits tumor growth in a xenograft model of WM

Although nimbolide has been examined in murine models of various cancers, its activity and effects on specific protein targets have not been assessed in in vivo models of WM (or any B-cell cancer). We hypothesized that systemically delivered nimbolide would engage its target(s) in vivo to induce WM cell death at pharmacologically achievable doses. Using our murine xenograft model of WM, we administered nimbolide via intraperitoneal injection in two separate cohorts of mice, testing two doses, 100 and 200 mg kg$^{-1}$ (see Supplementary section). Compared with the control group (DMSO-treated mice), we observed a 48% reduction in tumor volume in the nimbolide treated mice (100 mg kg$^{-1}$ cohort, $P < 0.05$; Figures 8a and b and Supplementary Figure 4) and a corresponding decrease in human IgM secreted by the xenografted WM cells ($P < 0.05$) (Figure 8c). A higher dose (200 mg kg$^{-1}$) had a more pronounced impact on IgM reduction but not on tumor volume (vs that observed in the 100 mg kg$^{-1}$ mice cohort). Immunohistochemistry (IHC) analysis was conducted on tumor tissues obtained from the animals treated with nimbolide (100 mg kg$^{-1}$) or DMSO alone. In vivo activity was validated by a notable decrease of IgM and BCL2 in nimbolide treated mice vs control. Likewise, an increase in tumor cell apoptosis (cleaved caspase-3 stain) and decrease in cell proliferation (Ki-67 stain) was noted in these tissues (Figure 8d). These data provide sufficient evidence that nimbolide can be safely administered in WM xenografted mice, elicit a significant anti-tumor effect and modulate important WM cell death and apoptotic pathways.

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were made in preclinical models of breast, colon and buccal pouch carcinoma and attest to the unique properties of this limonoid in inducing (perhaps simultaneously) both intrinsic and extrinsic apoptotic signaling cascades. However, the degree to which nimbolide distinctly activates the respective apoptotic systems and their overall contribution toward imminent cell death maybe tumor-cell type dependent and remains to be investigated in cells of lymphocytic lineage.

Currently, management of WM entails the use of bortezomib-based regimens with reports of promising activity of the BTK inhibitor, ibrutinib, in relapsed/refractory WM patients. However, as in most B-cell cancers, drug resistance eventually develops and remains a daunting yet inevitable clinical challenge. As such, we were encouraged to find that nimbolide’s cytotoxic effects were maintained in bortezomib or ibrutinib-resistant cell lines (BR and IR models, respectively) underscoring its potential role in these clinical settings. More intriguingly, BCWM.1/IR cells appeared most susceptible to nimbolide therapy exhibiting notably more apoptosis compared to ibrutinib sensitive (BCWM.1, MWCL-1) models. Using our IR model, we also observed that ibrutinib resistance could be ‘reversed’ by treating BCWM.1/IR cells with ibrutinib combined with a suboptimal dose of nimbolide. Together, the two agents induced significantly more apoptosis (48%; \( P < 0.05 \)), which indicates that nimbolide distinctively disrupts key oncogenic cellular networks engaged in ibrutinib resistance.

The capability of nimbolide to concurrently modulate several biological pathways has been previously described in solid tumor models where NFκB, TP53 and BCL2 associated proteins have been proposed to be the primary mediators of nimbolide’s cytotoxicity. It is likely that the anti-tumor effects of nimbolide are based on increased dependence of the tumor cell toward one (or more) of these pro-survival molecular pathways that are cell type-specific and are more vulnerable to modification by the limonoid. With compounds such as nimbolide, where the exact biological target is not known and the possibility of multiple oncotargets exists, the in silico technique of virtual screening is being increasingly utilized for its rapid and accurate prediction of compound–ligand interactions. Thus, to identify potential endogenous binding partners of nimbolide, we used a

Figure 5. Detailed interaction of nimbolide with BCL2 as identified by in silico docking analysis. (a) Nimbolide binds with BCL2 in reported P4-binding pocket. Hydrophobic binding pocket termed P4 is shown with nimbolide bound. The protein is shown with solvent-accessible surface threshold attenuated to reveal crevices and side chain details, and carbon atoms are colored gray, oxygen atoms red and nitrogen atoms blue. A box denoting the P4 pocket is shown at higher resolution in panel (b) where nimbolide is shown in detail with nearby residues, in particular Asn134 as part of a loop segment at the end of the helix. (c) Nimbolide 2D structure is shown with chirality. The residues Ala100, Asp103, Phe104, Arg107, Tyr108, Phe112, Leu137, Trp144, Gly145, Arg146, Val148, Ala149, Phe198, Leu201 and Tyr202 all form close proximity with nimbolide. The interactions are given as hydrophobic in green, charge in blue or blue and dashed arrow indicate H-bond.
chemoinformatics-based approach to reverse dock all known human biological proteins that were similar in shape, structure and biochemical properties (descriptors) to nimbolide and with which it could potentially associate with. Others and we have previously reported that NLE modulates the expression of BCL2, an anti-apoptotic protein that plays a major role in preventing malignant B-cell (including WM) death.\textsuperscript{13–16} We hypothesized that the anti-tumor activity of nimbolide might also be mediated through similar interactions. Our analysis suggests the BCL2–nimbolide interaction is critical for the antitumor activity of nimbolide and for the first time predicts the exact amino acid residues along with the thermodynamic and molecular mechanical exchanges that occur between these two molecules.

BCL2 functional biology is complex and tightly regulated by a number of post-translational modifications, which include phosphorylation, proteolytic cleavage, ubiquitination and proteosomal degradation.\textsuperscript{55,56} Phosphorylation of BCL2 in tumor cells has been shown to prolong the stability of BCL2, enhance its antiapoptotic activity and delay G1/S transition of the cell cycle.\textsuperscript{55,59} Thus a post-translational modification, such as a phosphorylation-disrupting event, could destabilize BCL2 and render it vulnerable to proteolytic destruction, potentially explaining its partial down-regulation in nimbolide treated cells. BCL2 phosphorylation can occur at multiple sites in the flexible loop domain, but it occurs notably at Ser70, Ser87, Thr56 and Thr74. However, its phosphorylation at Ser70 is widely accepted to support malignant cell survival and resistance to therapy. Because \textit{in silico} screening identified BCL2 as the prime target we sought to validate the results \textit{in vitro} by testing the cytotoxicity of nimbolide in a tumor cell model where the BCL2 protein is absolutely critical for cell survival, comparing against a model where the function of BCL2 has been attenuated by mutation and is not detrimental to tumor subsistence. We used RS4;11 tumor cells, which have been shown to express increased BCL2.\textsuperscript{43} Using a BH3 profiling technique, it was demonstrated that RS4;11 cells functionally rely on BCL2 (versus other BCL2 antiapoptotic members) for survival and were most sensitive to the anti-tumor effects of the BH3 mimic ABT-737.\textsuperscript{43} For comparison, we also utilized Jurkat BCL2\textsubscript{Ser70-Ala} cells in which the Ser70 residue of the BCL2 flexible domain has been mutated to alanine and prevents full phosphorylation of BCL2, in effect reducing the antiapoptotic potential of the protein. Our finding that BCL2 mutated cells tolerated nimbolide treatment significantly (\(P < 0.00008\)) more so than RS4;11 cells provides further support to the notion that nimbolide affects BCL2 and that the phosphorylational capacity of the protein indeed impacts the ability of the limonoid to produce an apoptotic reaction.

Lastly, we resolved to determine whether nimbolide could mitigate WM cell tumor expansion \textit{in vivo} and how this effect was carried out. \textit{In vivo} kinetics and toxicity profiling of nimbolide has

![Figure 6. Nimbolide shifts expression of BCL2 and proteins associated with other potential targets, identified on in silico screen. WM cells (BCWM-1, MWCL-1 and RPCI-WM1) were treated with different concentrations of nimbolide for 6 h and western blot analysis was performed. Among the concentrations tested, nimbolide effects were most evident at 1 μM (nimboide 0.1 and 0.5 μM data not shown).](image)

Figure 6. BCL2-dependent tumor cells are highly sensitive to nimbolide whereas BCL2-independent tumor cells are not. (a) RS4;11, Neo Jurkat and Jurkat BCL2\textsubscript{Ser70-Ala} tumor cells were treated with different concentrations of nimbolide for 6 h and apoptosis was measured by annexin V staining and flow cytometry. BCL2-dependent cells (RS4;11) were significantly sensitive to 1 μM nimbolide (82% cell death) as compared with BCL2-independent Jurkat BCL2\textsubscript{Ser70-Ala} tumor cells (28% cell death, \(P < 0.00008\)) or Neo Jurkat cells (40% cell death, \(P < 0.0001\)). (b) MOMP activation was also more significant in RS4;11 cells as compared with Jurkat BCL2\textsubscript{Ser70-Ala} or Neo Jurkat cells. \(P < 0.0001, \star \star P < 0.00008\).
been described previously, but no data exists in tumor models of lymphoid malignancy. The LD_{50} of nimbolide administered via intraperitoneal route in adult female mice has been reported as \(280 \text{ mg kg}^{-1} \text{ body weight}\). As such, for our initial study we opted to use two doses of nimbolide: a lower dose that was \(\sim 65\%\) of the LD_{50} (100 mg kg\(^{-1}\)) and a higher dose 200 mg kg\(^{-1}\), which was \(\sim 35\%\) of the LD_{50} reported. Using these doses, we anticipated minimal toxicity with preliminary indication of anti-tumor efficacy, which was more prominent in the higher dose cohort. However, we did not observe any additional tumor volume reduction in the 200 mg kg\(^{-1}\) arm; rather this dose was less tolerated resulting in the death of two mice. In mice treated with the 100 mg kg\(^{-1}\) dose, we noted an approximate 50\% decrease in tumor volume and IgM. This was an impressive finding and indicated that nimbolide had the potency to significantly (\(P < 0.005\)) kill WM cells in vivo, particularly in a xenograft model that contains tumor cells developed from a terminally refractory WM patient. Additionally, IHC analysis provided valuable insight into the putative in vivo targets of nimbolide. A notable decrease in BCL2 was evident but we did not find any change in TP53 in nimbolide treated mice. This is largely consistent with our in vitro findings that indicated a negligible decrease in TP53 protein, which was observed on immunoblot (data not shown). The ability of nimbolide to act through TP53-independent and TP53-dependent mechanisms\(^{14,46,56}\) is not entirely surprising as it is a derivative of NLE. We previously demonstrated that NLE is equally active in TP53 intact and TP53\(^{-}\) cells.

Figure 8. Nimbolide exerts anti-tumor activity in a human WM xenograft model. Female SCID mice (\(n = 20\)) were subcutaneously implanted with \(1 \times 10^6\) RPCI-WM1 cells (as described in the Supplementary methods). Progressive increase in IgM was measured in the serum and this corresponded with tumor growth in a time-dependent fashion. On average, WM tumor tissues grew to a median of 3 cm and IgM levels reached a median of 823 ng ml\(^{-1}\) by Day 30 post-implantation. (a) Average tumor reduction in the nimbolide 100 mg kg\(^{-1}\) cohort was 48\% with no additional anti-WM benefit noted in the 200 mg kg\(^{-1}\) cohort. (b) Tumors from two representative mice (control and nimbolide 100 mg kg\(^{-1}\)) show a significant reduction in nimbolide-treated mice (~50\%). (c) Human IgM levels measured by ELISA were significantly lower in nimbolide treated mice vs control (DMSO). Data from 15 mice is shown as representatives of the 3 different treatment groups. \(* P < 0.005\). (d) IHC analysis of tumor samples taken from DMSO and nimbolide (100 mg kg\(^{-1}\)) treated mice shows changes in IgM, Ki-67 and BCL2. The TP53 protein did not appear changed. Cleaved caspase-3 (Casp-3) was notably increased in nimbolide-treated mice.
CD19+ CLL cells and TP53 protein expression is decreased by NLE treatment. With TP53 mutations and deletions being prevalent in the more aggressive stages of cancer, this finding warrants attention because an anti-cancer compound with activity in TP53 defective patients is highly desirable.

In summary, we have extensively examined the biochemical features of nimbolide in preclinical models of B-cell cancer. Our data unequivocally demonstrate that its anti-tumor effects, most notably in WM cancer cells in vitro and in vivo, are mediated through the intrinsic apoptotic pathway. Nimbolide directly activates BCL2 and this appears to be a central mechanism for its cytotoxic activity in lymphoid cancer cells. Other potential targets such as HSP90 and PI3K (p110α), identified through in silico reverse docking, may also be affected as indicated by modulation of their associated proteins. Ongoing studies will examine these significance and interplay through a more in depth manner. Our investigations also show that nimbolide is capable of being considered for further development as an anti-WM therapeutic.

CONFLICT OF INTEREST

The authors declare no conflict of interest.

ACKNOWLEDGEMENTS

The experiments and analysis carried out in this study were supported by funding from the Leukemia and Lymphoma Society (A-C-K is a Leukemia and Lymphoma Scholar in Clinical Research) and the Daniel Foundation of Alabama (A-C-K). We thank Mrs Kelly Viola for her editorial assistance.

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