Metabolic Regulation of Protein N-Alpha-Acetylation by Bcl-xL Promotes Cell Survival

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SUMMARY

Previous experiments suggest a connection between the N-alpha-acetylation of proteins and sensitivity of cells to apoptotic signals. Here, we describe a biochemical assay to detect the acetylation status of proteins and demonstrate that protein N-alpha-acetylation is regulated by the availability of acetyl-CoA. Because the antiapoptotic protein Bcl-xL is known to influence mitochondrial metabolism, we reasoned that Bcl-xL may provide a link between protein N-alpha-acetylation and apoptosis. Indeed, Bcl-xL overexpression leads to a reduction in levels of acetyl-CoA and N-alpha-acetylated proteins in the cell. This effect is independent of Bax and Bak, the known binding partners of Bcl-xL. Increasing cellular levels of acetyl-CoA by addition of acetate or citrate restores protein N-alpha-acetylation in Bcl-xL-expressing cells and confers sensitivity to apoptotic stimuli. We propose that acetyl-CoA serves as a signaling molecule that couples apoptotic sensitivity to metabolism by regulating protein N-alpha-acetylation.

INTRODUCTION

Increasing evidence suggests that specific metabolic alterations associated with cancer cells may not be ancillary to their transformation but are instrumental to their tumorigenic potential by mediating cell proliferation, growth, and survival (Vander Heiden et al., 2009). Many oncogenes and tumor suppressor genes known to promote excess cell proliferation also alter biosynthetic (or anabolic) processes. For example, Akt expression stimulates glucose uptake and glycolysis, the pentose phosphate pathway, and fatty acid synthesis. c-Myc expression promotes glutamine metabolism as well as purine and pyrimidine biosynthesis. Furthermore, mutations in genes encoding metabolic enzymes have been identified by cancer genetic association studies (Vander Heiden et al., 2009). How specific metabolites contribute to increased proliferation and apoptotic resistance in tumor cells remains a central unanswered question.

The proto-oncogene Bcl-xL has a prominent role in promoting cell survival and cancer development (Boise et al., 1993). It is well established that Bcl-xL protects against apoptosis by directly binding and inhibiting Bax/Bak oligomerization-mediated mitochondrial permeabilization. However, certain Bcl-xL mutants, such as F131V/D133A and G148E, that are unable to bind to Bax or Bak, nevertheless retain 70%–80% antiapoptotic activity of WT Bcl-xL (Cheng et al., 1996). Curiously, Bcl-xL has also been shown to regulate mitochondrial respiration and metabolism (Gottlieb et al., 2000; Vander Heiden et al., 1999). Whether the metabolic function of Bcl-xL contributes to its role in mediating apoptotic resistance is unclear.

Our unexpected identification of an N-terminal acetyltransferase, Arrest Defective 1 (dARD1), in a genome-wide RNA interference (RNAi) screen in Drosophila cells for apoptotic regulators (Yi et al., 2007) prompted us to posit that protein N-alpha-acetylation, a major N-terminal modification, links cell metabolism
to apoptotic induction in cancer cells. Since dARD1 is epistatic to diap1, which encodes for a direct inhibitor of caspases in Drosophila, and ARD1 is required for caspase activation in mammalian cells (Yi et al., 2007), the role for ARD1 in mediating caspase activation is evolutionarily conserved. How ARD1 regulates caspase activation has not yet been illustrated.

In mammalian cells, protein N-alpha-acetylation is mediated by the highly conserved N-acetyltransferase protein complexes (NatA, NatB, NatC, NatD, and NatE). The NatA complex consists of the catalytic subunit, Arrest Defective 1 (hNaa10p/ARD1), and the auxiliary subunit, N-acetyltransferase 1 (NAT1/hNaa15p/NATH), whereas NatB consists of N-terminal acetyltransferase 3 (hNaa20p/NAT3) and mitochondrial distribution and morphology 20 (hNaa25p/Mdm20). Although the Nat complexes are implicated in regulating cell-cycle progression, cell proliferation, and tumorigenesis, the mechanisms that connect N-alpha-acetylation to the cellular protein apparatus are unknown (Ametzazurra et al., 2008; Polevoda and Sherman, 2003; Starheim et al., 2008, 2009). Recent N-acetylome studies reveal incomplete acetylation status of proteins (Arnesen et al., 2008; Goetze et al., 2009). Although a commonly accepted view is that partial acetylation results from the degenerate nature of protein N-terminal sequences, we considered the possibility that protein N-alpha-acetylation might be regulated, an alternative hypothesis that had not been tested as a result of technical limitations.

Here, we developed a biochemical approach to assess the status of endogenous levels of protein N-alpha-acetylation. Using this assay, we show that protein N-alpha-acetylation levels are sensitive to alterations in metabolism and Bcl-xL expression. Bcl-xL overexpression leads to reduced levels of acetyl-CoA and hypoacetylation of protein N termini through a Bax/Bak-independent mechanism. Conversely, bcl-x−/− mouse embryonic fibroblasts show increased levels of acetyl-CoA as well as protein N-alpha-acetylation levels. Protein N-alpha-acetylation deficiency in Bcl-xL-overexpressing cells contributes to apoptotic resistance since increasing acetyl-CoA production can rescue this deficiency in protein N-alpha-acetylation and sensitize Bcl-xL cells to cell death. Our study suggests that regulation of acetyl-CoA availability and protein N-alpha-acetylation may provide a Bax/Bak-independent mechanism for Bcl-xL to regulate apoptotic sensitivity.

RESULTS

We confirmed that ARD1 is necessary for cell death induced by the DNA-damaging agent doxorubicin in multiple cell lines of different origins, including Drosophila Kc (Yi et al., 2007), HeLa, HT1080, and U2OS cells (Figures 1A–1D). In addition, HeLa and U2OS cells deficient for NATH were also resistant to doxorubicin treatment, recapitulating the apoptotic resistant phenotype of ARD1 knockout cells (Figures 1A–1D). Thus, the acetylation activity of the NatA complex serves to influence the sensitivity of these cells to apoptosis. Next, we tested whether NatA influences apoptotic sensitivity to other DNA damaging agents. We found that ARD1 knockout cells are also resistant to cisplatin and ultraviolet (UV) treatment (Figure 1E). However, these cells remained sensitive to tumor necrosis factor (TNFalpha) and cyclohexamide treatment, which specifically activates apoptosis through the death receptor pathway (Figure 1F). We conclude that protein N-alpha-acetylation regulates apoptotic sensitivity downstream of DNA damage.

Since N-alpha-acetylation has been suggested to affect protein stability (Polevoda and Sherman, 2003), we examined whether protein synthesis and/or protein turnover might be affected by acetylation status. We tested whether ARD1 substrates such as caspase-2 and Chk1 (see results below) are destabilized in ARD1 knockout cells using cyclohexamide, an inhibitor of protein synthesis. Deficiency in ARD1 did not lead to decreases in the cellular levels of these proteins compared to that of control (Figure S1A available online). The steady-state levels of total cellular proteins in ARD1 knockout cells were similar to the levels in control cells (Figure S1B). We also tested whether general protein stability is altered in ARD1 or NATH knockout cells (Figure S1C). By pulse-chase 35S-Met labeling experiments, we observed that neither general protein synthesis nor turnover was affected in ARD1 or NATH knockout cells. Thus, protein N-alpha-acetylation mediated by NatA complex is not required to maintain protein stability globally. In addition, we verified that cell-cycle progression is unaffected in cells deficient for ARD1/NATH (Figure S1D). Taken together, these data suggest that the NatA complex may influence apoptotic sensitivity by mediating protein N-alpha-acetylation of key apoptotic components.

In Vitro Detection of Unmodified Protein N Termini

The lack of an immunological method to detect the acetylation status of protein N termini has limited our understanding of the mechanisms that regulate protein N-alpha-acetylation. To this end, we developed a selective biotin labeling method using an engineered protein ligase, termed subtiligase (Abrahmsén et al., 1991; Tan et al., 2007) that detects nonacetylated N termini of endogenous proteins. This approach was used to capture unmodified protein N termini resulting from caspase-mediated cleavage during apoptotic cell death (Mahrus et al., 2008). Unblocked N termini can be labeled using subtiligase, which preferentially biotinylates N-terminal amine groups consistent with the specificity of NatA or NatB (Abrahmsén et al., 1991; Mahrus et al., 2008). As the N termini of up to 80%–90% of cellular proteins may be blocked by a number of different modifications (Martinez et al., 2008), very few proteins will be biotin labeled by subtiligase as previously demonstrated (Mahrus et al., 2008). Thus, any protein that is biotin labeled by subtiligase in our assays most likely results from a specific loss in N-alpha-acetylation.

We utilized subtiligase to biotinylate free N termini of proteins in whole-cell lysates followed by avidin affinity purification and western blot analysis. Decreased levels of protein N-alpha-acetylation are expected to increase subtiligase-mediated protein biotinylation and conversely, increased levels of protein N-alpha-acetylation are expected to decrease subtiligase-mediated protein biotinylation (Figure 2A). First, we asked whether the subtiligase assay could be used to distinguish the N-alpha-acetylation status of protein N termini when the expression of the NatA complex is diminished by RNAi-mediated knockdown.
Figure 1. NatA Knockdown Suppresses Cell Death Induced by DNA Damage in HeLa, HT1080, and U2OS Cells

(A and B) HeLa cells were treated with doxorubicin (1.25 μg/ml, 20 hr for cell viability; 5 μg/ml, 8 hr for caspase activity).

(C) HT1080 cells were treated with doxorubicin (1.25 μg/ml, 20 hr).

(D) U2OS cells were treated with doxorubicin (1.25 μg/ml, 20 hr).

(E) HeLa cells were treated with cisplatin (40 μM) or UV (50 J/m² or 100 J/m²) for 24 hr.

(F) HeLa cells were treated with TNFalpha (10 ng/ml, 24 hr) and cyclohexamide (1 μg/ml, 24 hr) to induce death receptor mediated cell death.

Immunoblots were conducted in parallel to show extent of target knockdown. Data are represented as mean ± standard deviation (SD; n = 3). (Student’s t test: *p < 0.05; **p < 0.01; ***p < 0.001.) See also Figure S1.
Figure 2. A Subtiligase-Based Assay to Label Unacetylated Protein N Termini

(A) A schematic of the subtiligase assay. In brief, cells were gently lysed in NP-40 lysis buffer on ice. Cell debris was pelleted, and the resulting supernatant was subjected to a subtiligase reaction (1 mM subtiligase, 1 mM biotin-peptide, 2 mM DTT; 1 hr at room temperature). Biotinylated proteins were affinity purified with Neutravidin beads (Thermo Scientific) and analyzed by SDS-PAGE. Protein N-alpha-acetylation levels were determined by immunoblot against proteins in which the N-terminal residue corresponds to specificity of a Nat complex. Chk2 was included as a negative control, which should not be recognized by subtiligase based on sequence.

(B and C) Subtiligase reaction was conducted on HeLa cells transfected with siRNA against NATH (B) or ARD1 (C) as described in (A). Enrichment of biotin-labeled NatA substrates was observed in ARD1 or NATH-deficient cells compared to that of control. Enrichment in the amount of pulldown suggests a decrease in protein N-alpha-acetylation levels of NatA substrates in NatA-deficient cells. Blot quantification for all subtiligase assays was calculated relative to control and normalized to the corresponding lysate sample with ImageJ software.
ARD1 acetylates a subclass of proteins with Ser, Ala, or Thr as the newly exposed N-terminal residue after initiator Met (iMet) cleavage (Polevoda and Sherman, 2003). We tested 14–3–3ζ, which is known to be N-alpha-acetylated (Arnesen et al., 2009; Martin et al., 1993), and proteins that we predict to be N-alpha-acetylated based on their sequences, Chk1 and Msh2. Caspase-2, which is responsive to both DNA damage (Tinel and Tschopp, 2004) and metabolic stress (Nutt et al., 2005, 2009), is also a good candidate for acetylation by ARD1 as the second amino acid in the caspase-2 polypeptide is Ala. We observed that these proteins as well as caspase-2 were biotinylated to a higher extent by subtiligase in NATH or ARD1 knockdown cells than in control cells (Figures 2B and 2C). These data suggest that subtiligase can distinguish N-alpha-acetylation of multiple proteins that is dependent on NatA expression.

To determine the validity of subtiligase assay, we measured the extent of protein N-alpha-acetylation by quantitative mass spectrometry using differential isotope labeling (Figure 2D). First, we tested whether we could detect the basal levels of N-alpha-acetylation of caspase-2 by mass spectrometry. We observed that the mass to charge ratio (m/z) of the N-terminal peptide of caspase-2 is shifted as expected with an acetyl modification (Figures S2A and S2B). Furthermore, we found a 30% reduction in the amount of N-alpha-acetylated caspase-2 in NATH-deficient cells relative to control by subtiligase assay as well as mass spectrometry (Figure 2B and Figure S2D). These results support the conclusion that caspase-2 is N-alpha-acetylated by ARD1.

Protein N-Alpha-Acetylation Promotes the Assembly of Caspase-2 Complex

As caspase-2 is a substrate of ARD1 (Figure 2 and Figure S2) and the activation of caspase-2 is inhibited by ARD1 or NATH knockdown (Figure 3A), we asked how N-alpha-acetylation of caspase-2 might influence caspase activation.

First we conducted mutagenesis analysis of caspase-2 to disrupt protein N-alpha-acetylation. We replaced the third residue of caspase-2 with Pro (A3P) as the presence of Pro in this position inhibits protein N-alpha-acetylation. The 3P mutation has been previously demonstrated to inhibit N-alpha-acetylation of other substrates, known as the XPX rule (Goetze et al., 2009). We also replaced the second Ala for Ser as a control to maintain N-alpha-acetylation (A2S) as well as iMet removal (Arnesen et al., 2009; Goetze et al., 2009). Generation of these targeted substitutions allows us to definitively test whether subtiligase can differentiate between acetylated and unacetylated forms of caspase-2. An increase in subtiligase-mediated biotinylation of A3P was detected, while very little A2S or wild-type caspase-2 was detected after biotin pull down, consistent with acetylation as the explanation for the lower biotinylation levels (Figure 3B). A defect in N-alpha-acetylation of A3P caspase-2, but not WT and A2S caspase-2 was confirmed by mass spectrometry (Figures S2A–S2C). Thus, subtiligase is an effective tool for detecting unmodified protein N termini.

The caspase-2 scaffolding complex, which promotes caspase-2 activation, includes the adaptor protein, receptor-interacting protein (RIP)-associated ICH-1/CEP-3 homologous protein with a death domain (RAIDD) (Duan and Dixit, 1997; Tinel and Tschopp, 2004). The ability of the N-terminal caspase-2 mutants to interact with RAIDD was assessed by coimmunoprecipitation. We found that RAIDD efficiently coimmunoprecipitated with WT and A2S but not with A3P caspase-2 (Figure 3C). This suggests that N-alpha-acetylation of caspase-2 facilitates its interaction with RAIDD.

Since acetyl-CoA is a key cofactor in N-alpha-acetylation, we speculated that the levels of N-alpha-acetylated caspase-2 might be dependent on expression of key metabolic enzymes that are responsible for production of cytoplasmic acetyl-CoA. To explore this question, we tested whether knockdown of ATP citrate lyase or acetyl-CoA synthetase (the two metabolic enzymes that utilize citrate and acetate, respectively) to generate acetyl-CoA, results in decreased levels of N-alpha-acetylated caspase-2. Indeed, we observed increased biotin labeling of caspase-2 in knockdown cells compared to control cells following subtiligase assay (Figure 3D). This suggests that caspase-2 is hypoacetylated when acetyl-CoA generation is reduced and therefore, protein N-alpha-acetylation is subject to metabolic regulation.

Regulation of Protein N-Alpha-Acetylation by Bcl-xL

Since decreased levels of protein N-alpha-acetylation leads to apoptotic deficiency, we reasoned that regulation of protein N-alpha-acetylation of certain apoptotic regulators might provide a mechanism to control apoptotic sensitivity. Bcl-xL, an antiapoptotic Bcl-2 family member, is known to have an effect on metabolism (Shimizu et al., 1999; Vander Heiden et al., 2001). We asked whether protein N-alpha-acetylation levels are sensitive to Bcl-xL expression using subtiligase assay. An increase in biotin labeling of caspase-2, -9, -3, and Bax was observed by Bcl-xL expression in 293T, HeLa, and Jurkat cells compared to that of control (Figures 4A, 4B, and 4C, respectively). Conversely, a decrease in biotin labeling was apparent in bcl-xL−/− mouse embryonic fibroblasts (MEFs) compared to that of bcl-xL+/− MEFs (Figure 4D). Since Bcl-xL is known for maintaining mitochondrial integrity by blocking oligomerization of Bax/Bak, we measured the levels of protein N-alpha-acetylation in Bax/Bak-deficient cells. Surprisingly, the levels of protein N-alpha-acetylation were similar in bax−/−, bak−/−, or bax/bak−/− (double knockout [DKO]) MEFs compared to that of WT MEFs by subtiligase assay (Figure 4E). This suggests that
Bcl-xL-mediated regulation of protein N-alpha-acetylation is independent of Bax/Bak.

Recent studies show that histone lysine acetylation is dependent on acetyl-CoA production in yeast and mammalian cells (Takahashi et al., 2006; Wellen et al., 2009). However, we found that lysine acetylation of histone H3 and H4 were unaffected in Bcl-xL cells compared to control (Figure S3). This suggests that histone lysine acetylation is not sensitive to the changes in acetyl-CoA levels associated with Bcl-xL expression.

We next tested whether protein N-alpha-acetylation levels in Bcl-xL cells are affected by changes in acetyl-CoA metabolism.

Figure 3. N-Alpha-Acetylation of Caspase-2 Is Required for Binding to RAIDD

(A) HeLa cells were transfected with siRNAs as indicated and treated with doxorubicin (5 μg/ml, 8 hr). Cell lysates were analyzed by SDS-PAGE, and caspase cleavage was assessed by immunoblot. Caspase-2 and caspase-3 activation is suppressed in ARD1 and NATH knockdown cells compared to that of control. Arrow indicates cleaved product of caspase-2.

(B) Subtiligase reaction was conducted on HEK293T cells transfected with FLAG-tagged WT, A3P, and A2S caspase-2 (active cysteine mutant C320G of caspase-2) as described in Figure 1. Enrichment of biotin-labeled A3P caspase-2 was observed compared to WT and A2S caspase-2. This suggests that A3P caspase-2 is hypoacetylated.

(C) HEK293T cells were transfected with FLAG-tagged WT, A3P, and A2S caspase-2 as well as with VSV-RAIDD as indicated. Caspase-2 was affinity purified with anti-FLAG agarose and eluted with FLAG peptide. The resulting eluants were subjected to SDS-PAGE for immunoblot analysis. RAIDD efficiently coimmunoprecipitates with WT and A2S but not with A3P caspase-2. Blots are representative of at least three independent experiments.

(D) Subtiligase reaction was conducted on lysates from HeLa cells transfected with siRNAs against acetyl-CoA synthetase or ATP citrate lyase as indicated. Increased biotin labeling of caspase-2 was observed in knockdown cells compared to control cells.

See also Figure S2.
Figure 4. Metabolic Regulation of Protein N-Alpha-Acetylation by Bcl-xL
(A–C) HEK293T, HeLa, or Jurkat cells were generated to express empty GFP vector or Bcl-xL. Cells were treated with acetate (50 mM, 24 hr) or citrate (10 mM, 24 hr) as indicated. Subtiligase reaction was conducted as described in Figure 1. Enrichment of biotin-labeled NatA and NatB substrates is observed in Bcl-xL cells compared to control cells. Biotin labeling is reduced to control levels by acetate or citrate treatment in Bcl-xL cells.

(D) Lysates generated from Bcl-xL+/+ or Bcl-xL−/− MEFs were subjected to subtiligase reaction as described in Figure 1. Bax is hypoacetylated in Bcl-xL+/+ MEFs compared to Bcl-xL−/− MEFs.

(E) Lysates generated from wild-type, Bax−/−, Bak−/−, or Bax/Bak−/− (DKO) MEFs were subjected to subtiligase assay as described in Figure 1. Blots are representative of at least three independent experiments. See also Figure S3.
Figure 5. Bcl-xL Expression Reduces Acetyl-CoA Production

(A) Identification of metabolites using an AB/Sciex 5500 QTRAP mass spectrometer separated with normal phase HILIC NH2 chromatography using positive/negative ion switching in Jurkat cells stably expressing GFP empty vector or Bcl-xL.

(B) Acetyl-CoA levels were measured in Jurkat cells stably expressing GFP empty vector or Bcl-xL by mass spectrometry. Peak areas of total ion current (TIC) were processed with MultiQuant 1.1 software. The average TIC for acetyl-CoA analyte is shown. Acetyl-CoA levels are lower in Bcl-xL cells relative to GFP cells.
metabolites increase acetyl-CoA levels in mammalian cells (data not shown). Under metabolite treatment, protein N-alpha-acetylation levels were restored in Bcl-xL-expressing cells to that of control levels (Figures 4A and 4B). Thus, a reduction in acetyl-CoA production in Bcl-xL cells may be responsible for the observed hypoacetylation.

**Metabolic Alterations CAUSED BY Bcl-xL Expression**

The expression of Bcl-xL is often elevated in tumors (Adams and Cory, 2007). To explore the effect of Bcl-xL on tumor metabolism, we conducted a systematic search using a combination of two-dimensional nuclear magnetic resonance (NMR) and mass spectrometry to identify metabolic changes associated with increased Bcl-xL expression. Principal component analysis (PCA) of one-dimensional proton spectra shows that the metabolome of Bcl-xL-expressing cells was significantly different from the metabolome of control cells (Figure S4).

We then used triple-quadruple mass spectrometry via selected reaction monitoring (SRM) to identify metabolite changes in Bcl-xL cells relative to GFP control cells as mass spectrometry is a more sensitive approach (Bajad et al., 2006; Lu et al., 2008). This is particularly relevant for intermediates of glucose metabolism as these metabolites are difficult to decipher by NMR due to their similar proton content. Thus, both NMR and mass spectrometry provide complementary approaches for a comprehensive understanding of the metabolic changes resulting from a specific perturbation. Indeed, we found that acetyl-CoA levels were decreased by 2-fold in Bcl-xL-expressing cells relative to GFP-expressing cells by mass spectrometry as well as an enzyme-based assay (Figures 5A and 5B). Conversely, acetyl-CoA levels were substantially increased in bcl-x−/− MEFs compared to bcl-x+/− MEFs (Figure 5C). These data provide strong evidence that Bcl-xL expression reduces the levels of acetyl-CoA, suggesting that reduced levels of acetyl-CoA in Bcl-xL-overexpressing cells leads to hypoacetylation.

Since bax/bak DKO cells are not defective in protein N-alpha-acetylation, we reasoned that Bcl-xL might be able to negatively regulate the levels of acetyl-CoA independent of Bax/Bak binding. Cheng et al. reported that certain Bcl-xL mutants, such as F131V/D133A and G148E, are unable to bind to Bax or Bak but nevertheless retain 70%–80% antiapoptotic activity of WT Bcl-xL (Cheng et al., 1996). We measured acetyl-CoA levels in cells expressing WT Bcl-xL, or these specific Bcl-xL mutants. A similar reduction in acetyl-coA levels was observed in cells expressing these Bcl-xL mutants and in cells expressing WT Bcl-xL (Figures 5D and 5E). Thus, Bcl-xL's metabolic function in regulating the levels of acetyl-CoA does not depend on its interaction with Bax/Bak.

As the majority of the cellular acetyl-group in acetyl-CoA is produced from glucose (DeBerardinis et al., 2007), we asked whether glucose metabolism might be altered in Bcl-xL-expressing cells. We fed Bcl-xL cells uniformly labeled 13C-glucose to differentiate glucose-derived metabolites from those derived from other carbon sources (Figure 6A and Table S1). We found that the levels of glucose-derived citrate were decreased by approximately 25% in Bcl-xL-expressing cells relative to control (Student’s t test, p value < 0.05; Figure 6B and Table S1). As citrate is the direct precursor of cytoplasmic pools of acetyl-CoA, the lower levels of glucose-derived citrate might explain the decrease in acetyl-CoA levels observed in Bcl-xL-expressing cells. Consistent with this hypothesis, levels of alpha-ketoglutarate, which is also derived from citrate, were lower in Bcl-xL-expressing cells relative to control (Figure 6B and Table S1).

Since metabolite addition rescues the defect on protein N-alpha-acetylation by Bcl-xL (Figures 4A and 4B), we asked whether these metabolites could alter cell survival that is supported by Bcl-xL expression. Bcl-xL expression effectively protects against a wide range doses of doxorubicin (Figures 7A–7D). Remarkably, increasing levels of citrate or acetate sensitized HeLa cells stably expressing Bcl-xL to doxorubicin-induced cell death compared to that of untreated cells (Figure 7B). This corresponds with a 2-fold increase in caspase activity (Figure 7D). Importantly, RNAi against acetyl-CoA synthetase or ATP citrate lyase completely suppressed the sensitization to doxorubicin elicited by addition of acetate or citrate, respectively (Figure 7E). This indicates that metabolite-induced apoptotic sensitization of cells expressing Bcl-xL specifically results from changes in acetyl-CoA production.

The above data suggest that Bcl-xL may mediate apoptosis resistance through two parallel pathways by inhibiting Bax/Bak oligomerization and by downregulating protein N-alpha-acetylation. We therefore directly tested whether the effects of inhibiting Bax and ARD1 are additive in protecting against apoptosis. We found that double knockdown of both ARD1 and Bax indeed provided enhanced protection against apoptosis compared to that of knockdown individually, which was especially significant at higher concentrations of doxorubicin (Figure 7F). This finding supports the notion that Bcl-xL has dual functions in regulating protein N-alpha-acetylation levels and Bax/Bak oligomerization.

**DISCUSSION**

The ability to rapidly assess protein modifications immunologically has been essential for exploring the significance and regulation of multiple protein posttranslational modifications such as phosphorylation, histone methylation, and acetylation. Since an antibody for protein N-alpha-acetylation does not exist, the ability to assess this modification was severely limited. In this regard, the subtiligase assay as described in the present study provides a powerful tool to allow us to rapidly assess the endogenous levels of protein N-alpha-acetylation. Using this assay,
we discovered that protein N-alpha-acetylation status is reduced in cells overexpressing Bcl-xL. Furthermore we show that protein N-alpha-acetylation is sensitive to acute changes in acetyl-CoA availability.

Our study directly links a specific metabolite, acetyl-CoA, to apoptotic sensitivity and supports a growing number of studies that describe a role for cell metabolism in controlling apoptosis (Nutt et al., 2005, 2009; Schafer et al., 2009; Vaughn and Deshmukh, 2008). Interestingly, the cellular levels of acetyl-CoA are sensitive to Bcl-xL status in a Bax/Bak-independent manner because expression of Bcl-xL mutants that are unable to bind to Bax or Bak (Cheng et al., 1996) can also affect acetyl-CoA levels to the same extent as that of wild-type Bcl-xL. Hardwick and colleagues demonstrated that these Bcl-xL mutants preserve 70%–80% antiapoptotic activity of WT Bcl-xL despite their inability to bind to Bax or Bak (Cheng et al., 1996). Thus, inhibition of acetyl-CoA production might provide an additional mechanism for Bcl-xL to protect against apoptosis in a Bax/Bak-independent manner. Taken together, these data suggest that Bcl-xL might protect against apoptosis through two parallel mechanisms: by directly binding and inhibiting Bax/Bak oligomerization and by regulating mitochondrial metabolism, which leads to reduced levels of acetyl-CoA and protein N-alpha-acetylation. We conclude that Bcl-xL integrates metabolism to apoptotic resistance by modulating acetyl-CoA levels.

Previous studies show that Bcl-xL directly binds to the voltage-dependent anion channel (VDAC), a component of the mitochondrial permeability transition pore, which controls mitochondrial metabolite exchange (Shimizu et al., 1999; Vander Heiden et al., 2001). It is possible that Bcl-xL expression may alter levels of acetyl-CoA by regulating mitochondrial membrane permeability. Citrate carrier (CiC), a nuclear-encoded protein located in the mitochondrial inner membrane and a member of the mitochondrial carrier family, is responsible for the efflux of acetyl-CoA from the mitochondrion to the cytosol in the form of citrate (Gnoni et al., 2009; Kaplan et al., 1993). We found that the levels of glucose-derived citrate were decreased by approximately 25% in Bcl-xL-expressing cells relative to the control. This reduction in citrate levels could explain the observed decrease in acetyl-CoA levels in Bcl-xL-expressing cells and contribute to the antiapoptotic function of Bcl-xL. Indeed, addition of citrate to Bcl-xL-expressing cells leads to increased protein N-alpha-acetylation and sensitization of these cells to apoptosis. Perturbations in acetyl-CoA production may extend to other oncogenic contexts beyond that of Bcl-xL. For example, the levels of glucose-derived acetyl-CoA were found to be approximately 20% higher in myc+/+ cells relative to myc-/- cells (Morrish et al., 2009). An increase in acetyl-CoA levels might contribute to enhanced apoptotic sensitivity of cells overexpressing c-Myc (Evan et al., 1992). We propose that the basal levels of acetyl-CoA may influence the apoptotic threshold in multiple oncogenic contexts.

The ability of Bcl-xL to control the levels of acetyl-CoA and protein-N-acetylation provides a clear example by which
metabolism is mechanistically linked with apoptotic sensitivity. Loss-of-function ard1 mutant yeast are specifically defective in alpha-factor response but not to a-factor (Whiteway and Szostak, 1985), indicating that protein N-alpha-acetylation status can dictate a specific cellular behavior or process. Since protein N-alpha-acetylation affects a large number of cellular proteins, we speculate that metabolic regulation of this process exerts its control on cellular processes through regulating a group(s) of proteins rather than individual proteins. ARD1-deficient mammalian cells are defective in the activation of caspase-2, caspase-3, and caspase-9 in response to DNA damage (Yi et al., 2007). Consistently, N-alpha-acetylation of multiple caspases, including caspase-2, caspase-3, and caspase-9, was reduced in Bcl-xL-overexpressing cells. It is possible that defects in N-alpha-acetylation of multiple caspases, which may negatively regulate their activation, contribute to apoptotic resistance of ARD1-deficient cells as well as Bcl-xL-overexpressing cells. Thus, the N-alpha-acetylation status of multiple proteins that are involved in a particular pathway may collectively determine a specific physiological outcome. In this regard, the cofactor for the Nat complexes, acetyl-CoA, serves as a signaling molecule that functions as an important liaison between metabolism and multiple cellular processes.

EXPERIMENTAL PROCEDURES

Cell Death Assays

For RNAi studies, low-passage HeLa cells were transiently transfected (5 x 10^5/well in 96-well plates) with a pool of four small interfering RNAs (siRNAs) (predesigned ON-TARGETplus siRNAs available from Dharmacon, 50 nM) with Oligofectamine transfection reagent (Invitrogen). After a 48 hr incubation, cells were treated with doxorubicin (concentrations and incubation times are noted in figure legends). siRNAs were tested in triplicate for each independent experiment. For detection of caspase cleavage by western blot, HeLa cells were transfected as described above (1.5 x 10^5/well in 6-well plates) followed by treatment with doxorubicin. Cells were lysed directly in SDS samples buffer and subjected to SDS-PAGE and western blot analysis via standard procedures.

For metabolite sensitization experiments, HeLa cells stably expressing GFP or Bcl-xL (3 x 10^5/well in 96-well plates) were pretreated with acetate (50 mM) or citrate (10 mM) for 24 hr as indicated. Cells were lysed in 0.2% Tween 20 and 0.2% Triton X-100 buffer, and the resulting lysates were used for subtiligase reaction that includes 1 mM purified subtiligase, 1 mM purified biotin-peptide, and 2 mM DTT as previously described (Mahrus et al., 2008). Reactions were allowed to proceed for 1 hr at room temperature. Biotinylated proteins were affinity purified with Neutravidin agarose at 4°C overnight (Thermo Scientific). The following day, agarose was pelleted and washed three times in lysis buffer. Purified proteins were eluted directly in 2x SDS sample buffer, and eluants were analyzed by SDS PAGE.

Subtiligase Biotinylation Experiments

Synthesis of Peptide Substrate

The biotinylated peptide substrate for subtiligase with a TEV protease cleavage site, biotin-ahx-ahx-GGTENLYFQSY-gluc-Y-NH2, was synthesized manually on a rink amide MBHA resin according to standard 9-Fluorenylmethoxycarbonyl (Fmoc) chemistry-based solid-phase peptide synthesis protocols (Mahrus et al., 2008). The crude peptide was purified with a C18 semipreparative reverse-phase column on a Waters HPLC system. The identity of the purified product was confirmed by ESI-MS (isotopic molecular weight [MW] calculated 1949.9 Da, m/z [M+H]^+ found 1951.0). The peptide substrate could be made more soluble by incorporating D-arginine residues into the sequence (Yoshihara et al., 2008). So a more soluble form of the substrate, biotin-ahx-ahx-dRdRdR-dR-ahx-ahx-GGTENLYFQSY-glgc-Y-NH2, was also synthesized, and its integrity was verified by HPLC and ESI-MS (MW calculated 2644.4 Da, m/z [M+H]^+ found 2645.9).

Expression and Purification of Subtiligase

The expression construct for subtiligase was prepared with the plasmid pBS42 (ATCC 37279) according to published procedures (Wells et al., 1983), except that a His6 tag was added to the C terminus (Tan et al., 2007). The gene of the subtiligase variant contains point mutations S221C, P225A, M124L, and S125A on wild-type subtilisin BPN’. The recombinant protein was expressed in B. subtilis RIK1285, which is deficient in the production of neutral and alka-line proteases. Purification of subtiligase was performed as previously described (Abrahmsén et al., 1991), except that a Co(II) affinity chromatography step was used instead of ion-exchange chromatography (Tan et al., 2007). The affinity-purified subtiligase was desalted using a PD-10 column (Amersham) with deionized H2O. Aliquots of the desalted enzyme solution were flash frozen, lyophilized, and stored at –80°C until utilized. The prepared subtiligase was analyzed by SDS-PAGE gel and MALDI-TOF MS, which confirmed its purity and identity. The ligase activity of the enzyme preparation was confirmed in model ligation reactions using known peptide substrates for subtiligase.

Subtiligase Ligation Reaction

Cells used for subtiligase assay were plated at 50%, 60%, and 70% confluency to support exponential growth, followed by pretreatment with acetate (50 mM) or citrate (10 mM) for 24 hr as indicated. Cells were lysed in 0.2% Tween 20 and 0.2% Triton X-100 buffer, and the resulting lysates were used for subtiligase reaction that includes 1 mM purified subtiligase, 1 mM purified biotin-peptide, and 2 mM DTT as previously described (Mahrus et al., 2008). Reactions were allowed to proceed for 1 hr at room temperature. Biotinylated proteins were affinity purified with Neutravidin agarose at 4°C overnight (Thermo Scientific). The following day, agarose was pelleted and washed three times in lysis buffer. Purified proteins were eluted directly in 2x SDS sample buffer, and eluants were analyzed by SDS PAGE.

12C Mass Spectrometry Sample Preparation

Three replicates of Jurkat cells stably expressing GFP or Bcl-xL (5 x 10^6 for each replicate) were washed twice in PBS and resuspended in RPMI medium with glutamine, 10% dialyzed NCS, and 10 mM uniformly labeled (3-13C)-glucose (Cambridge Isotope Laboratories), followed by incubation for 24 hr. Cells were washed twice, and metabolites were extracted in 3 ml 80% ice-cold methanol. Insoluble material in lysates was pelleted at 13,000 g for 10 min, and methanol from the resulting supernatant was evaporated. Samples were resuspended with 20 µL HPLC grade water for mass spectrometry.

12C and 13C Mass Spectrometry Data Acquisition and Analysis

Seven microliters of 20 µl were injected using a 5500 QTRAP mass spectrometer (Applied Biosystems/MDS Sciex) equipped with a Prominance UFLC HPLC system (Shimadzu) via SRM of a total of 249 endogenous metabolites for 12C analyses of GFP and Bcl-xL samples. For analyses of 13C-labeled GFP and Bcl-xL samples, 153 endogenous metabolites were targeted via SRM. Reliable quantitative data are only acquired from approximately 60% of the targeted metabolites (~160 metabolites from the 12C method and ~55% from the 13C method (~80 metabolites). Some metabolites were targeted in both positive and negative ion mode, including acetyl-CoA. ESI voltage was 5000 V in positive ion mode and −4500 V in negative ion mode. The dwell time was 5 ms per SRM transition, and the collision energy was optimized for each SRM transition. Total cycle time was 2.09 s for the 12C method and 1.24 s for the 13C method. Scheduled SRMs were not utilized. Samples were delivered to the MS via normal phase chromatography using a 2.0 mm i.d. × 10 cm HILIC Luna NH2 column (Phenomenex) at 250 µl/min at basic pH using positive and negative ion switching within the same 30 min LC/MS/MS analytical run. Gradients were run starting from 85% buffer B (HPLC grade acetonitrile) to 42% B from 0–5 min; 42% B to 0% B from 5–16 min; 0% B was held from 16–24 min; 0% B to 85% B from 24–25 min; 85% B was held for 7 min to re-equilibrate the column. Buffer A was composed of 20 mM ammonium hydroxide/20 mM ammonium acetate in 95/5 water/acetonitrile. All metabolomic measurements were performed in triplicate. Peak areas from the total ion current (TIC) for each metabolite SRM transition were integrated using MultiQuant v1.1 software (Applied Biosystems).
Figure 7. Metabolic Function of Bcl-xL Contributes to Cell Survival

(A–D) HeLa cells stably expressing empty GFP vector or Bcl-xL were treated with doxorubicin as indicated. Cell viability was determined by measurement of cellular ATP levels with a luminescence-based assay (Promega). Caspase activity was determined by measurement of the cleavage of a luciferin substrate containing the caspase cleavage site, DEVD (Promega).

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Graphs are representative of three independent experiments. Data are represented as mean ± SD (n = 3). (Student’s t test: *p < 0.05; **p < 0.01; ***p < 0.001.)

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**REFERENCES**


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Supplemental Information

EXTENDED EXPERIMENTAL PROCEDURES

Antibodies
The following antibodies were used for immunobloting: Caspase-9 (R&D systems), Caspase-2 (Alexis), Caspase-3 (Cell Signaling), cleaved Caspase-3 (Cell Signaling), ARD1 (Santa Cruz), NATH (provided by T. Arnesen and J. Lillehaug), γ-tubulin (Sigma), Bax (Santa Cruz), Bak (Santa Cruz), Msh2 (R&D Systems), Chk1 (Cell Signaling), pan-pyruvate kinase M (Abcam), Bcl-xL (provided by Craig Thompson), β-actin (Sigma), FLAG (Sigma), and 14-3-3 (Santa Cruz).

Cell Line Construction
Jurkat cells stably expressing GFP or Bcl-xL were generated by retroviral-mediated expression of pMIG bicistronic vector containing GFP only or GFP and Bcl-xL (Addgene) using standard procedures. High GFP expressing cells were purified by fluorescence activated cell sorting. HeLa cells stably expressing GFP or Bcl-xL were generated using the same methods without sorting. Cells that showed greater than 90% infection efficiency were used for experiments described in the study. HeLa cells were maintained in DMEM with glutamine (Invitrogen) and 10% NCS. Jurkat cells were maintained in RPMI with glutamine and 10% NCS.

HEK293T cells stably expressing shRNA against human ARD1 or NATH were generated by retroviral-mediated expression of pSRP-puro vector using standard procedures. The sequences used for the oligonucleotides inserted into the pSRP vector correspond to the individual siRNA sequences provided by Dharmacon (ARD1, J-009606-05; NATH, J-012847-05). Following puromycin selection, cells were passaged for no more than 1 week. Following puromycin selection, 293T cells were transiently transfected with full-length mouse caspase-2 containing active cysteine mutation (C320G) to reduce cell death using calcium phosphate.

HEK293T cells were transiently transfected with pMIG GFP empty vector or pMIG GFP and Bcl-xL vector using calcium phosphate.

All cells described above were maintained in the presence of penicillin and streptomycin (Invitrogen) and passaged for no more than 4 weeks with exceptions noted.

Unlabeled NMR Sample Preparation
10 replicates of Jurkat cells stably expressing GFP or Bcl-xL (100x10⁶ for each replicate), and metabolites were extracted in 4mL 100% ice-cold methanol followed by addition of 4mL chloroform and 4mL water. Lysates were spun at 10,000 g for 30 min at 4°C to separate the aqueous and inorganic phases. Methanol was evaporated from aqueous fractions by nitrogen gas and lyophilized. Samples were prepared for NMR spectroscopy by dissolving the lyophilized cell extract pellets in 400 ml of sample buffer, containing 50mM HEPES (pH = 7.0) and 2mM DSS (as a chemical shift reference). The samples were immediately transferred into 5mm Shigemi NMR tubes (Shigemi, Inc. Suite 21, 4790 Route 8, Allison Park, PA 15101) for data acquisition.

NMR Spectroscopy
All NMR spectra were acquired on a Varian INOVA 600MHz spectrometer (Varian, Inc., Palo Alto, CA.) using a 5mm triple resonance (H, C, N) Salt Tolerant Cold Probe. The sample temperature was 25 degrees centigrade for all samples. One-dimensional proton spectra were acquired using a one pulse sequence with continuous-wave (CW) pre-saturation of the residual water signal during the relaxation delay (2 s), with an RF field of 50Hz. 16K complex data points were acquired, with 256 transients and 4 dummy transients. The spectral width was 8KHz.

Mass Spectrometry (MS) Analysis and Quantification of Caspase-2
FLAG-tagged caspase-2 or mutant (A2S and A3P) was transiently expressed in HEK293T cells. After lysis in 0.2% Tween 20 and 0.2% Triton X-100 buffer, FLAG-tagged caspase-2 was affinity purified with FLAG agarose beads (Sigma) and eluted with FLAG peptide. Eluants were resolved on SDS-PAGE and visualized by GelCode Blue stain. The band containing caspase-2 or mutant protein was excised and subjected to in-gel Endoproteinase Lys-C digestion. Peptides were extracted from the gel, differentially labeled with reductive dimethylation as previously described (Hsu et al., 2003; Khidekel et al., 2007), and heavy- and light-dimethylated peptides were combined 1:1 (wild-type: mutant) for MS analysis.

The peptide mixture was analyzed by liquid chromatography coupled to tandem mass spectrometry (LC-MS/MS). Peptides were separated across a 37 min gradient ranging from 4% to 27% (v/v) acetonitrile in 0.1% (v/v) formic acid in a microcapillary (125 μm x 18 cm) column packed with C18 reverse-phase material (Magic C18AQ, 5 μm particles, 200 Å pore size, Michrom Bioresources) and on-line analyzed on an LTQ Orbitrap XL hybrid FTMS (Thermo Electron). For each cycle, one full MS scan acquired on the Orbitrap at high mass resolution was followed by ten MS/MS spectra on the linear ion trap XL from the ten most abundant ions. Single ion chromatograms from the Orbitrap MS scans were generated for N-terminal peptides. The area under the SIC curve was used to determine relative abundance between wild-type and mutant. Internal peptides common to both wild-type and mutant were used to normalize mixing ratio to 1:1. This isotope-labeling strategy allowed for a comparative analysis of a wild-type and a mutant sample together in a single LC-MS/MS experiment.
SUPPLEMENTAL REFERENCES


Figure S1. Total Protein Levels and General Protein Synthesis Is Unaffected in Cells Deficient in NatA Components, Related to Figure 1

(A) HeLa cells were transfected with a pool of siRNAs targeting ARD1 or non-targeting control siRNA as indicated and treated with cyclohexamide (100 ng/mL) for 5 hr. Caspase-2, -9, -3, and Chk1 but not Chk2 are substrates of NatA complex (Figure 2). This experiment is representative of 2 independent experiments. Asterisk indicates non-specific band.

(B) HeLa cells were transfected as described in (A). Total protein lysates were analyzed by SDS PAGE, and the gel was stained with GelCode Blue (Pierce). Representative digital images were generated by an infrared imager (Li-Cor).

(C) HeLa cells were transfected with siRNAs targeting ARD1 or NATH followed by 48h incubation to allow for protein turnover. Newly synthesized proteins were labeled with [35S]-Cys and [35S]-Met for 1h followed by a 24h chase with cold medium. Cells were lysed directly in SDS sample buffer and lysates were analyzed by SDS-PAGE and autoradiography.

(D) The ability of NatA deficient cells to progress through the cell cycle normally is shown. Following RNAi treatment, cells were resuspended in PBS with propidium iodide for FACS analysis. Cells treated with RNAi against ARD1 or NATH do not show any appreciable change in the cell cycle compared to control. Altogether these results suggest that the defect in Chk1 activation is likely due to the decrease in acetylation of Chk1 in NatA deficient cells as assessed by subtiligase in Figure 1. Blots are representative of at least 3 independent experiments.
Figure S2. Caspase-2 Is N-Terminally Acetylated, Related to Figures 2 and 3

N-terminal peptides of WT, A2S, and A3P caspase-2 were identified by mass spectrometry. A2S substitution of caspase-2 remains N-alpha-acetylated, whereas N-alpha-acetylation of A3P caspase-2 is not detected by mass spectrometry. Schematic of differential isotope labeling to detect acetylated or unacylated N-terminal peptides of caspase-2 by mass spectrometry is shown in Figure 1C.

(A) Analysis of differentially isotope labeled WT and A2S caspase-2 shows that the N-terminal Met is removed and the newly exposed residue is acetylated. N-terminal peptides from both WT and A2S mutant caspase-2 were found to co-elute and therefore shown in one MS spectrum. N-terminal peptide corresponding to acetylated wild-type caspase-2 (heavy dimethyl modification) is A(Ac)APSGRSQSSLHRK(C2D6) (3+) at m/z = 519.96. N-terminal peptide corresponding to acetylated A2S caspase-2 (light dimethyl modification) is S(Ac)APSGRSQSSLHRK(C2H6) (3+) at m/z = 523.28.

(B and C) Analysis of differentially isotope labeled wild-type (B) and A3P caspase-2 (C) shows that only wild-type caspase-2 is acetylated at the N-terminus. N-terminal peptide corresponding to acetylated wild-type caspase-2 (heavy dimethyl modification) is A(Ac)APSGRSQSSLHRK(C2D6) (3+) at m/z = 519.96. The non-acetylated N-terminal mutant peptide (A3P) that is consequently dimethylated was identified at an earlier retention time on the reverse phase column. This is expected from the lower hydrophobicity of the molecule with a dimethyl group instead of an acetyl modification at the N-terminus for the A3P peptide. Thus the A3P and WT caspase-2 peptides are shown in two different MS spectra. N-terminal peptide corresponding to unacetylated A3P caspase-2 (light dimethyl modification) is A(C2H6)PPSGRSQSSLHRK(C2H6) (3+) at m/z = 521.96, and 391.71 (4+) (not shown).

(D) FLAG tagged caspase-2 (C320G) was immunopurified in control or NATH knockdown cells. Peptides were generated by Lys-C followed by chemical modification using NaCNBH3 or NaCNBD3 and formaldehyde treatment (formaldehyde-H2 or formaldehyde-D2 respectively). Ratios for levels of N-terminal acetylation were determined by normalizing to 1:1 ratio of shared internal peptides. The following internal peptides were used for normalization: EELMK, VNALIK, NHTQSPGCEESDAGK. Data are represented as mean +/- s.d. (n = 3).
Figure S3. Histone Acetylation Levels Are Unaffected in Bcl-xL Expressing Cells, Related to Figure 4
Jurkat cells stably expressing Bcl-xL or GFP were lysed directly in SDS sample buffer. Lysates were analyzed by SDS-PAGE. Levels of acetylated histone-H3 and histone-H4 were analyzed by immunoblot.
Figure S4. Metabolomic Profiling of Bcl-xL Expressing Cells by NMR, Related to Figure 5
Metabolite levels were compared between cells expressing Bcl-xL or green fluorescence protein (GFP). Principal components analysis (PCA) of 1D Proton NMR spectra of 10 Bcl-xL expressing, and 9 GFP expressing Jurkat cell extract samples using AMIX software (version 3.7.10, Bruker-BioSpin Corp.). On the left is a scores plot of PC1 versus PC2 representing the corresponding loadings plot shown on the right. All spectra were corrected for dc offset, and calibrated to DSS at 0 ppm prior to binning. Spectra were binned using equally spaced buckets of 0.04 ppm width, covering the range from 0.5 to 10 ppm. The water region was excluded. Data were pareto scaled.