Pathways Accessory to Proteosomal Proteolysis Are Less Efficient in Major Histo compatibility Complex Class I Antigen Production*

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Degradation of cytosolic proteins depends largely on the proteasome, and a fraction of the cleavage products are presented as major histocompatibility complex (MHC) class I-bound ligands at the cell surface of antigen presenting cells. Proteolytic pathways accessory to the proteasome contribute to protein turnover, and their up-regulation may complement the proteasome when proteasomal proteolysis is impaired. Here we show that reduced reliance on proteasomal proteolysis allowed a reduced efficiency of MHC class I ligand production, whereas protein turnover and cellular proliferation were maintained. Using the proteasomal inhibitor adamantane-acetyl-(6-aminohexanoyl)-3-(leucinyl)-3-vinyl-(methyl)-sulphone, we show that covalent inhibition of all three types of proteasomal β-subunits (β1, β2, and β3) was compatible with continued growth in cells that up-regulate accessory proteolytic pathways, which include cytosolic proteases as well as deubiquitinating enzymes. However, under these conditions, we observed poor assembly of H-2Dβ molecules and inhibited presentation of endogenous tumor antigens. Thus, the tight link between protein turnover and production of MHC class I ligands can be broken by enforcing the substitution of the proteasome with alternative proteolytic pathways.

Several cytosolic proteases, including the 26 S proteasome, bleomycin hydrolase, puromycin-sensitive amino peptidease and leucine-aminopeptidase, contribute to the generation of MHC class I ligands (1–3). However, the 26 S proteasome, a large multicatalytic protease complex, carries out the bulk of both cytosolic protein degradation and MHC class I ligand production (1, 2). This protease has a multisubunit 20 S core structure containing two sets of three distinct catalytic sites, X (β), Y (βi), and Z (βi), associated with one or two 19 S regulatory accessory complexes (1, 2). The proteasome generates a wide range of peptide cleavage products (3–24 amino acids in length) that are ultimately degraded into free amino acids (4–6). In mammalian cells, a minor subset of peptides is rescued from further degradation and is translocated from the cytosol into the endoplasmic reticulum for assembly with MHC class I molecules. The MHC class I pathway is thereby assured constitutive production of ligands through cytosolic proteolysis. In the case of an immunological challenge, mammalian cells express IFN-γ-inducible proteasomal β-subunits (LMP7/β5i, LMP2/β1i, and MECL-1/β2i) that replace the constitutively expressed subunits in newly synthesized proteasomes (1). Such replacement leads to increased proteosomal production of peptides with hydrophobic C termini, usually preferred for both TAP transport and MHC class I binding (7). However, the majority of potential MHC class I ligands, as deduced from their primary structure, are not efficiently processed, although the correct motifs for TAP transport and MHC class I binding are contained in the protein sequence (8, 9). Such failure may depend on proteolysis by cytosolic proteases inefficient at generating the requisite cleavage products. Thus, it is possible that MHC class I processing may be regulated by differential participation of non-proteasomal peptidases in cytosolic protein degradation. Impaired proteasomal activity can be functionally compensated, at least in part, by another large cytosolic peptidase, tripептидил-peptidase II (10–13). Despite covalent inhibition by NLVS (14) or lactacin (15), EL-4 lymphoma cells adapted to growth in the presence of this inhibitor (denoted EL-4ad) maintain cytosolic proteolysis and cell viability by a mechanism that includes compensatory up-regulation of tripептидил-peptidase II (10–12). However, it is unknown whether the adapted state has functional consequences at the level of MHC class I ligand generation and antigen presentation to CTLs.

We show that lymphoma cells with reduced reliance on proteasomal activity no longer efficiently produced MHC class I ligands, although cytosolic proteolysis continued, and proliferation was not altered compared with control cells. Assembly of H-2Dβ molecules was dramatically reduced, and endogenous tumor antigens were not presented efficiently under these conditions. This phenotype contributed to escape from tumor resistance.© 2003 by The American Society for Biochemistry and Molecular Biology, Inc.

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Impaired assembly of MHC class I molecules in cells with reduced reliance on proteasomal activity. a, pulse-chase experiments were performed on control EL-4 cells, EL-4 cells treated with 50 μM NLVS for 16 h, or EL-4 ad cells. H-2D^b molecules were immunoprecipitated (IP) with anti-H-2D^b antibody B22.249.1 in the presence or absence of an H-2D^b-binding peptide, followed by SDS-PAGE analysis and autoradiography. b, RMA cells left untreated or treated with 50 μM NLVS for 3 h (3h), adapted (ad) to 50 μM NLVS (RMAad), or adapted to NLVS and washed (wash) and RMA-S cells were subjected to pulse-chase experiments. H-2D^b molecules were immunoprecipitated in the presence or absence of an H-2D^b-binding peptide, followed by SDS-PAGE analysis and autoradiography. Glycosylated heavy chains (GHC) were transported from the endoplasmic reticulum, heavy chains (HC), and β2-microglobulin (β2m) are indicated with arrows.

Materials and Methods

Cells and Transfections—EL-4 is a benzopyrene-induced thymoma cell line of the H-2^k haplotype, derived from C57Bl/6 mice. RMA is a Rauscher’s virus-induced T cell lymphoma cell line and is also derived from C57Bl/6. Adaptation to the proteasomal inhibitor NLVS was obtained by incubation of these cells in RPMI 1640 medium containing 5% fetal calf serum, 1% penicillin/streptomycin, 1% glutamine, and 10% C57Bl/6. Adaptation to the proteasomal inhibitor NLVS was obtained by incubation of these cells in RPMI 1640 medium containing 5% fetal calf serum, 1% penicillin/streptomycin, 1% glutamine, and 10% C57Bl/6. RMA is a Rauscher’s virus-induced T cell lymphoma cell line and is also derived from C57Bl/6.

Cytosolic Proteolysis without MHC Class I Ligand Production

Cytosolic proteolysis was measured in tumor graft experiments in syngeneic C57Bl/6 mice. Using a GFP reporter to measure proteolysis, we show in live cells that non-proteasomal serine peptidase activity participated in protein degradation, but inhibition of these enzymes failed to have a significant effect on the assembly of H-2K^b molecules. Continued proteolysis in proteasome-impaired cells afforded the cell the requisite housekeeping functions while preventing the full display of the usual set of MHC class I-restricted epitopes.

Proteasomal Inhibitors—NLVS (14) covalently modifies all catalytically active subunits of the proteasome, but with preference for the β-subunits with chymotryptic specificity. Several derivatives of NLVS were obtained by variations in the peptide scaffold: 4-hydroxy-5-iodo-3-nitrophenylacetyl-Ala-Ala-Phe-vinyl sulfone (AAF-VS) and 4-hydroxy-5-iodo-3-nitrophenylacetyl-Leu-Leu-Gly(cis)-vinyl sulfone (LLG-(cis)-VS). LLG-VS was obtained in the cis- and trans-isomers due to the absence of a side chain on the P1 glycine. Whereas the trans-form of LLG-VS modifies proteasomal β-subunits, the cis-form modifies yet uncharacterized targets in the cytosol distinct from the proteasome. Adamantane-acetyl-(6-aminohexanoyl)3-(leucinyl)3-vinyl-(methyl)-sulphone (Ada-Ahx3-Leu3-VS) is an N-terminally extended vinyl sulfone inhibitor that blocks all proteasomal β-subunits in a covalent manner (17).

Peptide Substrates and Peptidase Assays—To assay the activity of the proteasome, we used the fluorogenic substrates succinyl-LLVY-AMC, benzoyloxy carbonyl-GGL-AMC, t-butoxy carbonyl-LRR-AMC, and benzoyloxy carbonyl-YVAD-AMC (Sigma). To assay tripeptidyl-peptidase II activity, we used AAF-AMC (Sigma). Cell extracts or proteasome-enriched fractions and substrate (100 μM) were mixed in 50 mM Tris (pH 7.5), 5 mM MgCl2, 1 mM dithiothreitol, and 2 mM ATP in a final volume of 100 μl. Peptide hydrolysis was monitored by fluorescence spectroscopy (PerSeptive Biosystems, Framingham, CT) with excitation at 380 nm and fluorescence reading at 460 nm.

Proteasome Purifications—Preparation of proteasome-enriched fractions was performed using 0.5–1 × 10^9 control or adapted EL-4 cells and C57Bl/6 livers. Cells were washed with phosphate-buffered saline and lysed by vortexing with glass beads in 50 mM Tris base (pH 7.5), 5 mM MgCl2, 1 mM dithiothreitol, and 2 mM ATP. Glass beads and cell debris were removed by sequential centrifugations at 3000 and 14,000 rpm, respectively. Microsomes were removed by centrifugation for 1 h at 100,000 x g, and large cytosolic proteins or protein complexes containing proteasomes and tripeptidyl-peptidase II were then sedimented at 100,000 × g for 5 h. The resulting pellet was dissolved in 50 mM Tris base (pH 7.5), 5 mM MgCl2, 1 mM dithiothreitol, 2 mM ATP, Glass beads and cell debris were removed by sequential centrifugations at 3000 and 14,000 rpm, respectively. Microsomes were removed by centrifugation for 1 h at 100,000 × g, and large cytosolic proteins or protein complexes containing proteasomes and tripeptidyl-peptidase II were then sedimented at 100,000 × g for 5 h. The resulting pellet was dissolved in 50 mM Tris base (pH 7.5), 5 mM MgCl2, 1 mM dithiothreitol, 2 mM ATP, and 30% glycerol.

Pulse-Chase Experiments—Cells were starved in methionine/cysteine-deficient medium for 45–60 min, pulsed with [35S]methionine for 15 min, and chased for the indicated times. Cells were collected by centrifugation and lysed in 0.5% Nonidet P-40 lysis buffer, and MHC class I molecules were immunoprecipitated with rabbit anti-p8 serum.
molecules were transported from the endoplasmic reticulum within 120 min after onset of the chase, as judged from their acquisition of Golgi-specific glycan modifications (Fig. 1a, left panel). In EL-4 cells treated with NLVS (50 μM, 3 h), only a minimal fraction of H-2D\(^b\) heavy chains were transported even after long chase times (Fig. 1a, middle panel), as reported previously (21). Stabilization of H-2D\(^b\) molecules in cell lysates of NLVS-treated EL-4 cells by addition of the influenza nucleoprotein-(366–374) peptide confirmed that most of these H-2D\(^b\) heavy chains were devoid of peptide ligand (+ lanes) (22). In EL-4ad cells, a fraction of H-2D\(^b\) resumed folding, although at much lower levels compared with control EL-4 cells. The majority of H-2D\(^b\) heavy chains in EL-4ad cells remained unassembled in the endoplasmic reticulum devoid of peptide, as indicated by the stabilizing effect of nucleoprotein-(366–374) added to lysates of these cells (Fig. 1a, right panel).

In RMAad cells, which were similarly adapted to NLVS, maturation of H-2D\(^b\) molecules was comparable to that observed in EL-4ad cell (Fig. 1b). Despite normal proliferation, tumor cells can therefore avoid production of most H-2D\(^b\) ligands by reduced reliance on proteasomal activity.

Tumor cells often acquire deficiencies in MHC class I antigen presentation to escape from host immune detection. To test whether reduced reliance on proteasomal activity has functional consequences, we tested presentation of endogenous tumor antigens to CTLs by EL-4ad and RMAad cells. Both EL-4 and RMA cells express H-2K\(^b\)-restricted (gagL75–83) as well as H-2D\(^b\)-restricted (env195–203) murine leukemia virus-derived peptides and, in addition, an endogenous H-2D\(^b\)-restricted tumor epitope (23). We generated antitumor CTLs by priming C57Bl/6 mice and subsequent in vitro restimulation of splenocytes with B7.1-transfected EL-4 cells. We found that tumor antigen-specific CTLs performed efficient killing of control EL-4 cells, whereas EL-4ad cells were not efficiently recognized, although the latter were killed significantly better than
We also found that RMAad cells likewise had a reduced ability to present endogenous tumor antigens compared with control RMA cells. Although RMAad target cells were killed at higher levels than TAP-deficient RMA-S cells, 5–10 times more CTLs were required to obtain the same degree of killing as seen on RMA target cells (Fig. 2b). Even more pronounced differences were obtained using CTL clone ln17, specific for tumor antigen-specific peptide NKGENAQAI restricted by H-2Db (20). In line with previous data, we found that ln17 detected the presence of the tumor-specific epitope on RMA cells, but failed to recognize RMA-S cells (Fig. 2c). Furthermore, no recognition of RMAad cells was observed. MHC class I-restricted presentation of a tumor antigen-specific peptide can thereby be inhibited when proteasomal proteolysis is inhibited in a suitable manner. Because we used IFN-γ secretion as readout for antigen detection by ln17, these data also exclude that the differences in CTL killing were due merely to differences in target cell apoptosis when comparing control and NLVS-adapted target cells. Whereas the first and third compounds efficiently labeled the β3- and β5-subunits of the proteasome, [125I]LLG-VS labeled other yet uncharacterized cellular proteases.

Up-regulation of Deubiquitinating Enzymes and Non-proteasomal Peptidases in EL-4ad Cells—The activity of the ubiquitin-specific protease USP14 is associated with the 19 S cap of the proteasome and is up-regulated when the proteasome is inhibited (24). More generally, inhibition of proteasomal proteolysis should lead to accumulation of ubiquitin-conjugated substrates. Adaptation to proteasomal inhibitors might well include increased activity of deubiquitinating enzymes to deal with such accumulation. We therefore examined whether this was the case also in EL-4ad cells using 125I-labeled ubiquitin-vinyl sulfone (24). Cellular fractions of control EL-4 and EL-4ad cells (cytosolic as well as proteasome-enriched fractions) were incubated with ubiquitin-vinyl sulfone, and covalently modified polypeptides were separated by SDS-PAGE. We found increased labeling of IsoT1, USP14, and UCH-L1 in EL-4ad cells compared with control EL-4 cells (Fig. 3a), in line with what was observed in acutely treated EL-4 cells (24). More
active ubiquitin removal could prepare these substrates for degradation by other proteases.

Two additional active-site probes with different peptide scaffolds were used (25), [125I]AAF-VS and [125I]LLG-VS, to examine whether residual proteasomal activity is mediated by the $\beta_2\beta_5i$-subunits (X/LMP7). None of these probes labeled $\beta_2\beta_5i$-subunits (X/LMP7) in lysates of EL-4ad cells, whereas strong labeling was detected in lysates of control EL-4 cells (Fig. 3b). This confirms that virtually no catalytic activity remains for the $\beta_2\beta_5i$-subunits (X/LMP7) in EL-4ad cells, which is important in view of the fact that small amounts of peptide ligand suffice to load MHC class I molecules with peptide (26). Interestingly, using [125I]LLG(cis)-VS, we detected a series of modified polypeptides distinct from proteasomal $\beta$-subunits. Because vinyl sulfones are mechanism-based probes (14, 27), we conclude that these polypeptides correspond to additional, yet to be identified, proteases. This activity is not inhibited by NLVS, further supporting the alteration in proteolytic specificity in EL-4ad cells (Fig. 3b, middle panel). Labeling of the $\beta_2\beta_5i$-subunits (X/LMP7) with [125I]NLVS was likewise inhibited in RMAad cells when tested with these peptide vinyl sulfones (Fig. 3c).

Proteolysis Accessory to the Proteasome Supports Protein Degradation, but Is Relatively Ineffective in MHC Class I Ligand Production—We next made stable EL-4 transfectants expressing Ub-R-GFP to monitor proteasomal degradation of a protein substrate in live cells (16). GFP was converted into an N-end rule substrate and was degraded in EL-4 cells due to its

Fig. 4. Measuring proteasomal degradation by fluorescent ubiquitinated substrates in live cells. Shown are the results from FACS analysis of EL-4 cells stably transfected with Ub-M-GFP (a) or Ub-R-GFP (b and c). EL-4.Ub-R-GFP cells were left untreated (b) or were treated with 10 $\mu$M NLVS (c). The level of proteasomal activity in EL-4.Ub-R-GFP cells treated with 10 $\mu$M NLVS was measured by cytosolic high molecular mass protein (100,000 $\times$ g, 5 h) cleavage of the fluorogenic peptide substrates succinyl (succ)-LLVY-AMC, t-butyloxycarbonyl (boc)-LRR-AMC, and benzyloxycarbonyl (z)-YVAD-AMC (d). Also shown are the results from FACS analysis of EL-4.Ub-R-GFP cells treated with up to 50 $\mu$M NLVS (e–h).
N-terminal arginine, whereas Ub-M-GFP was comparatively stable (Fig. 4, a–c) because of the presence of a methionine residue. NLVS treatment of EL-4.Ub-R-GFP cells led to accumulation of fluorescence, as detected by FACS. In line with data from yeast mutants (28, 29), efficient inhibition of primarily the chymotryptic proteasomal activity was sufficient for accumulation of fluorescence (Fig. 4, c and d). These data further confirm that NLVS is indeed an efficient inhibitor of proteasomal protein degradation in live cells.

We next tested whether inhibition of accessory pathways has any effect on cytosolic proteolysis. To do this, we accumulated high levels of the R-GFP substrate in live EL-4.Ub-R-GFP cells and then blocked protein synthesis to study changes in the steady state of the substrate (Fig. 5, a–d). This revealed residual substrate degradation in the continued presence of 10 μM NLVS because a substantial fraction of the substrate was removed after 8 h. However, this was inhibited by treatment with AAF-CMK, an efficient inhibitor of tripeptidyl-peptidase II and other serine oligopeptidases (Fig. 5, c and d). Although inhibition of oligopeptidases by AAF-CMK had minor effects on untreated EL-4.Ub-R-GFP cells, we observed a significant effect.
Cytosolic Proteolysis without MHC Class I Ligand Production

YVAD-AMC) (upper three panels) or labeled with Ada-[125I-Z
upper panels] untreated (left panels) or EL-4ad (right panels) cells were left
proteolysis. EL-4 (panel b). GGL-AMC, reporter substrates succinyl (Suc)-LLVY-AMC, benzyloxy-carbonyl-
somes. The samples were either tested for cleavage of the peptide
mitted to differential centrifugation for partial purification of protea-
or EL-4ad cells were incubated with either 50
/H9262 or 10
left panel
/H9262 or 50
middle right panel
/M
 (30). NLVS fails to block
all proteasomal active sites were covalently modified during
assays using fluorogenic peptide substrates, labeling of
inhibitor to the X/LMP7 site (32). Consistent with the enzyme
caspase-like specificities during inhibitor treatment, possibly
early time points, we observed an induction of the trypsin- and
caspase-like activity was 70% inhibited (Fig. 6 a). Initially, at
treatment of live cells with the Ada-Ahx3-Leu3-VS inhibitor
all proteasomal active sites were covalently modified during
and epoxomicin (31). To examine if residual proteasomal activity
influence the viability of EL-4ad cells, we used Ada-Ahx3-
Leu3-VS, a cell-permeable tripeptide vinyl sulfone that co-
valently modifies all proteasomal β-subunits with comparable
efficiency (17). We found that proteasome-enriched fractions
from control EL-4 or EL-4ad cells treated with either NLVS or
Ada-Ahx3-Leu3-VS had almost completely blocked chymotryp-
tic and trypsin-like proteasomal activities, whereas the
caspase-like activity was 70% inhibited (Fig. 6a). Initially, at
early time points, we observed an induction of the trypsin- and
caspase-like specificities during inhibitor treatment, possibly
due to allosteric effects on the proteasome upon binding of the
inhibitor to the X/LMP7 site (32). Consistent with the enzyme
assays using fluorogenic peptide substrates, labeling of β-sub-
units with Ada-[125I-Tyr]Ahx3-Leu3-VS in cell lysates followed
by separation of the β-subunits by SDS-PAGE confirmed that
all proteasomal active sites were covalently modified during
treatment of live cells with the Ada-Ahx3-Leu3-VS inhibitor
(Fig. 6a, lower panel). Furthermore, EL-4ad cells proliferated
regardless of the presence of Ada-Ahx3-Leu3-VS, whereas con-
trol EL-4 cells died within 48 h (Fig. 6b). To confirm that
proteasomes of proliferating EL-4ad cells were indeed modi-
fied, we prepared proteasome-enriched fractions from cells
treated for several days with Ada-Ahx3-Leu3-VS. This analysis
revealed results similar to those observed in acutely treated
cells. Essentially no residual tryptic and chymotryptic activi-
ties and inhibited caspase-like activity were detected (data not

EL-4ad cell proliferation despite inhibition of all catalytic sites of the proteasome by Ada-Ahx3-Leu3-VS. a, control EL-4 or EL-4ad cells were incubated with either 50 μM NLVS or 50 μM Ada-Ahx3-Leu3-VS for the indicated times, and cell lysates were sub-
mitted to differential centrifugation for partial purification of protea-
somes. The samples were either tested for cleavage of the peptide
reporter substrates succinyl (Suc)-LLVY-AMC, benzyloxy-carbonyl-
GGL-AMC, t-butyloxy-carbonyl (Boc)-LRR-AMC, and benzyloxy-carbonyl
(Z)-YVAD-AMC (upper three panels) or labeled with Ada-[125I-
Tyr]Ahx3-Leu3-VS, followed by SDS-PAGE and autoradiography (lower panel). b, EL-4ad cell viability was mostly independent of proteasomal
proteolysis. EL-4 (left panels) or EL-4ad (right panels) cells were left
untreated (upper panels) or were incubated with 50 μM NLVS (middle
left panel) or 10 μM (middle right panel) or 50 μM (lower panels) Ada-
Ahx3-Leu3-VS for the indicated times. Live (○) and dead (●) cells were
counted by trypan blue exclusion.

FIG. 6. EL-4ad cell proliferation despite inhibition of all catalytic sites of the proteasome by Ada-Ahx3-Leu3-VS. a, control EL-4 or EL-4ad cells were incubated with either 50 μM NLVS or 50 μM Ada-Ahx3-Leu3-VS for the indicated times, and cell lysates were sub-
mitted to differential centrifugation for partial purification of protea-
somes. The samples were either tested for cleavage of the peptide
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Ahx3-Leu3-VS for the indicated times. Live (○) and dead (●) cells were
counted by trypan blue exclusion.

on R-GFP accumulation upon treating EL-4Ub-R-GFP cells
with both NLVS and AAF-CMK in combination (Fig. 5e). These
data show that non-proteasomal oligopeptidase activity indeed
contributes to cytosolic proteolysis, especially during situations
of limiting or insufficient proteasomal activity.

To further study whether oligopeptidases inhibitable by
AAF-CMK are important in generating MHC class I ligands,
we performed a pulse-chase experiment with [35S]methionine
metabolic labeling and precipitation of H-2Kb molecules.
A substantial fraction of H-2Kb molecules continue to assemble
in EL-4ad cells (10). We examined whether this may be due to
ligands produced by oligopeptidases inhibitable by AAF-CMK.
We found that this treatment had minor effects on the assembly
and transport of H-2Kb molecules in EL-4ad cells and also in
control EL-4 cells with active proteasomes (Fig. 5f). We con-
clude that pathways accessory to proteasomal proteolysis
that are inhibited by AAF-CMK support protein degradation,
but reveal poor yields of MHC class I ligands.

Evidence for Continued Cell Survival and Growth without
Significant Proteasomal Activity—EL-4ad cells continue to
depend on proteasomal β-subunit activity, at least to some extent
(30). NLVS fails to block β2- and β5-subunits (Z/MECL-1) in
vivo, a pattern of inhibition that is shared between NLVS and
other covalent proteasomal inhibitors such as lactacystin (15)
and epoxomicin (31). To examine if residual proteasomal activity
influences the viability of EL-4ad cells, we used Ada-Ahx3-
Leu3-VS, a cell-permeable tripeptide vinyl sulfone that co-
valently modifies all proteasomal β-subunits with comparable
efficiency (17). We found that proteasome-enriched fractions
from control EL-4 or EL-4ad cells treated with either NLVS or
Ada-Ahx3-Leu3-VS had almost completely blocked chymotryp-
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caspase-like activity was 70% inhibited (Fig. 6a). Initially, at
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(Fig. 6a, lower panel). Furthermore, EL-4ad cells proliferated
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fied, we prepared proteasome-enriched fractions from cells
treated for several days with Ada-Ahx3-Leu3-VS. This analysis
revealed results similar to those observed in acutely treated
cells. Essentially no residual tryptic and chymotryptic activi-
ties and inhibited caspase-like activity were detected (data not

FIG. 7. Increased in vivo tumorigenicity of cells with reduced reliance on proteasomal proteolysis. Control EL-4 (open bars) and
EL-4ad (closed bars) cells were grafted at 10⁴ to 10⁶ cells into the right
flanks of syngeneic C57Bl/6 mice (a) or perforin/RAG-1⁻/⁻ mice (PKOB/
RAG⁻⁻) (b). Frequency of tumor growth is displayed.
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...shown). Thus, tumor cells may adapt and proliferate normally even when proteasomal β-subunit activity is almost absent.

Escape from in Vivo Immune Detection by Tumor Cells with Reduced Reliance on Proteasomal Activity—In vivo tumors frequently acquire mutations in genes encoding proteins of the MHC class I-processing pathway, and these are believed to be the result of immune selection (33). Therefore, we explored whether an adapted phenotype similar to that of EL-4ad cells was selected during growth in vivo. We inoculated control EL-4 and EL-4ad cells into syngeneic C57Bl/6 mice and observed tumors in most mice inoculated with EL-4ad cells at both 10^6 and 10^4 cells/animal, whereas control EL-4 cells failed to grow and produce tumors (Fig. 7a). The tumor-forming ability of EL-4ad cells was dependent, at least in part, on escape from immune recognition because both EL-4ad and control EL-4 cells formed tumors in mice with a deficiency of perforin and RAG-1 (PKOB/RAG^-/-) (Fig. 7a). Furthermore, after low dose irradiation (400 rads) of the C57Bl/6 mice, commonly used to reduce in vivo transplantation barriers (34), both control EL-4 and EL-4ad cells were able to grow after inoculation of 10^6 cells/animal, whereas only EL-4ad cells grew at 10^4 cells/animal (Fig. 7b). In conclusion, we have shown that the ability of NLVS-adapted cells to proliferate independently of proteasomal β-subunit activities leads to reduced immune recognition in vivo.

DISCUSSION

This study shows that tumor cells may avoid efficient production of MHC class I ligands and hence immune recognition by modulation of proteasomal activity. Pathways accessing to proteasomal proteolysis can reduce the extent to which cells depend on proteasomal activity. In our case, cells adapted to growth in the presence of proteasome inhibitors were unable to maintain normal levels of MHC class I ligand production. In addition, using the vinyl sulfonyl inhibitor Ada-Ahx6-Leu9-VS, we observed that inhibition of all catalytic β-subunit activities of the proteasome (more efficiently than achieved with NLVS) was compatible with continued cell growth of EL-4ad cells. These results indicate that it is possible for mammalian cells to partly escape from production of MHC class I ligands by averting to pathways of protein degradation involving proteasomes other than the proteasome.

Although MHC class I processing is a rather inefficient process overall, in which most (>99%) of the cleaved peptides are never displayed at the cell surface, it is an adequate method for screening of the bulk of cellular protein content for the presence of foreign antigens (26). The steady-state level of MHC class I at the surface of cells depends on both its transport and removal from the cell surface (35), and transport of H-2Db is substantially inhibited in EL-4ad cells. Despite this, the cell-surface H-2Db (and also H-2Kb) levels detected by FACS are almost normal, suggesting that the rate of decay at the cell surface may be reduced when transport is slow (data not shown). Earlier data on MHC class II transport in cathepsin S^-/- mice have revealed a similar feature; and also in this case, an absence of T cell detection is observed for certain antigens (36, 37). In the course of an immune response, the proteolytic specificity in antigen processing has profound influence on the generation of MHC class I ligands, as illustrated by the IFN-γ-dependent substitution of proteasomal β-subunits (7). The fact that pathways accessible to proteasomal proteolysis, such as tripeptidyl-peptidase II, can contribute to maintaining proteolysis when the proteasome is inhibited allows for mammalian cells to alter the spectrum of cleavage fragments in the cytosol more dramatically (10–13). This notion is supported by the up-regulation of several deubiquitinating enzymes in EL-4ad cells, observed otherwise in cells suffering from acute proteasomal inhibition (24). USP14 is associated with the 19 S regulatory complex, and its precise role in proteasomal proteolysis remains to be established. Other deubiquitinating enzymes are also up-regulated in EL-4ad cells, such as IsoT1 and UCH-L1, which participate in the disassembly of free polyubiquitin chains (38, 39). When the proteasome is blocked, the removal of ubiquitin from ubiquitin-conjugated substrates may be a crucial step to engage alternative proteolytic pathways.

Reduced expression of several components of the MHC class I antigen-processing pathway is often observed in human tumors. This includes down-regulation of the IFN-γ-inducible proteasomal β-subunits (β1, β2, and β5) (40), important for production of MHC class I ligands, as well as down-regulation of other gene products involved in antigen processing (41–43). Tumors may fail to produce certain immunodominant ligands due to altered proteasomal specificity (40, 44). Our data show that reduced reliance on proteasomal proteolysis biases cytosolic proteolysis to produce peptides that are less fit for MHC class I assembly, thereby down-regulating the pool of potential MHC class I-restricted epitopes. Such down-regulation is stabilized in rapidly proliferating cells and can be induced at fairly high frequency (10). Furthermore, EL-4ad cells appear to use this phenotype to avoid immunological rejection during the formation of tumors in vivo. An EL-4ad-like phenotype may be preferentially selected in tumors that are poorly antigenic, a trait observed in many types of tumors. Interestingly, in Burkitt’s lymphomas, the oncogene c-myc is known to induce down-regulation of a number of components of the MHC class I-processing pathway, including down-regulation of proteasomal chymotryptic activity and up-regulation of tripeptidyl-peptidase II, thus linking the deficiency in antigen processing directly to oncogene expression (45, 46). This study reveals a new strategy for regulation of MHC class I processing: reduced reliance on proteasomal activity to down-regulate generation of MHC class I ligands.

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REFERENCES

Pathways Accessory to Proteasomal Proteolysis Are Less Efficient in Major Histocompatibility Complex Class I Antigen Production

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