Catalytically Active Proteasomes Function Predominantly in the Cytosol*

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The ubiquitin/proteasome pathway is a well characterized system for degrading intracellular proteins, although many aspects remain poorly understood. There is, for instance, a conspicuous lack of understanding of the site(s) where nuclear proteins are degraded because the subcellular distribution of peptidase activity has not been investigated systematically. Although nuclear proteins could be degraded by importing proteasomes into the nucleus, it is also evident that some nuclear proteins are degraded only after export to cytosolic proteasomes. Proteasomes and substrates are mobile, and consequently, the sites of degradation might not be static. We sought to identify the location of proteasomes to provide more conclusive evidence on the sites of protein degradation. We report that catalytically active proteasomes exist almost exclusively in the cytosol. The resulting lack of nuclear peptidase activity suggests that little, if any, degradation occurs in the nucleus. These and other studies suggest that the export of proteolytic substrates could define an important regulatory step in the degradation of nuclear proteins by cytosolic proteasomes.

The ubiquitin/proteasome pathway is a major mechanism for eliminating regulatory and damaged proteins. The key enzymology is well understood, and many targeting factors that attach ubiquitin to proteolytic substrates have been identified and characterized (1). Despite these advances, there remain areas of ambiguity. A detailed understanding of the assembly of mult ubiquitin chains, the transport of proteolytic substrates, and sites of intracellular protein turnover remain unclear.

Many proteins that are involved in cell cycle progression, DNA repair, and transcription are nuclear proteins that play a central role in cell growth and stress response. The stability of these proteins is controlled by the proteasome, and it is generally assumed that they are degraded inside the nucleus. This view is fostered by the detection of proteasome subunits in the nucleus (2, 3) and enrichment in the nuclear envelope (4). However, many of these studies examined the distribution of GFP-tagged proteins, which does not ensure that the tagged subunits are present in intact proteasomes. This is an important consideration because certain proteasome subunits and subcomplexes perform non-proteolytic roles in the nucleus (5, 6). We also note that certain GFP-tagged proteasome subunits are not efficiently assembled into intact complexes, and in some instances the fluorescence is reduced after assembly into the proteasome (7). Consequently, the signal observed could arise predominantly from the free form of proteasome subunits. Critically, there is no convincing evidence that peptidase activity is present in the nucleus.

We and others reported that the yeast DNA repair protein Rad4, DNA polymerase subunit Cdc17 (2), and HO endonuclease (8) are stabilized in nuclear export mutants, although proteasome assembly and catalytic activity are unaffected. The stabilizing effect of blocking export is not restricted to a few nuclear proteins because overall polyubiquitylated protein levels increased (2). Similarly, the degradation of mammalian nuclear proteins, including p53, β-catenin, TRIP-Br2, and hMSH5, requires nuclear export (9–12). Because import is unaffected in export mutants, proteasomes could have entered to degrade nuclear proteins. However, this was not observed, suggesting that nuclear proteins are exported and degraded by cytosolic proteasomes.

To test this hypothesis, we investigated the site of peptidase activity in both yeast and cultured human cells. We did not detect intact proteasomes in the nucleus, and virtually all peptidase activity was present in the cytosol. These findings diverge from the general opinion, which maintains that proteasomes degrade proteins inside the nucleus (4, 13–25). Our results predict an important regulatory role for export in the degradation of nuclear proteins.

Results

Purified Nuclei Lack Peptidase Activity—Actively growing yeast cells were lysed with zymolyase, and nuclei were separated by differential centrifugation. Prior to centrifugation, “total” cellular protein was prepared by lysing an aliquot of the unfractinated spheroplasts. We measured histone deacetylase (HDAC)³ activity to gauge the purity of the isolated nuclei (Fig. 1A). HDAC activity was detected only in purified nuclei (Nuc), whereas the hydrolysis of a fluorogenic proteasome substrate (LLVY-AMC) was found predominantly in the cytoplasm (Cyto). A low level of peptidase activity in the nuclear fraction is

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³ The abbreviations used are: HDAC, histone deacetylase; Nuc, nucleus; Cyto, cytoplasm; LMB, leptomycin B; NPC, nuclear pore complex; NLS, nuclear localization signal; Tricine, N-[2-hydroxy-1,1-bis(hydroxymethyl)ethyl]glycine; Suc, N-succinyl; AMC, 7-amino-4-methylcoumarin.
likely to originate from unlysed cells and endoplasmic reticulum that can co-sediment with nuclei. Fractionated yeast extracts were examined by immunoblotting to confirm that peptidase activity coincided with the location of proteasome subunits (Fig. 1B). High levels of proteasome subunits (Rpn11, Rpn10, and Rpt1) were detected in the cytosolic fraction, and significantly lower levels were present in purified nuclei. The nuclear localization of native Ubc4 (B), consistent with the localization of native Ubc4 (B). DIC, differential interference contrast.

To demonstrate that peptidase activity was generated by intact proteasomes, we prepared total, cytosol, and nuclear fractions from HEK293T cells, as described above. We again measured peptidase activity using equal amount of protein (Fig. 3, conc.,) and proportional volumes of lysate (vol.). Both methods confirmed that the nuclear fraction lacked peptidase activity (Fig. 3A). Significantly, the peptidase activity in the cytosol was comparable with that in total extract (when we used proportional volumes), indicating that almost all peptidase activity is cytosolic. Data representing four independent experiments were standardized to the peptidase activity in total extract. To determine whether the peptidase activity was generated by intact proteasomes, and not free catalytic (20S) particles, we separated lysates in native polyacrylamide gels (Fig. 3B). In-gel hydrolysis of LLVY-AMC revealed two bands corresponding to single capped (19S + 20S), and double-capped (19S + 20S + 19S) proteasomes in both total and cytosolic extracts. A significantly lower level of free 20S particle was also detected.
Proteasome Catalysis Occurs Outside the Nucleus

FIGURE 2. Proteasome peptidase activity can be released into the medium. A, HEK293T cells were treated with varying concentrations of digitonin and pelleted. Following digitonin treatment, the peptidase activity that was released into the medium (M) was compared with that detected in the cell pellet (P). Prolonged exposure to digitonin (30 min) did not further increase peptidase activity in the medium. Total peptidase activity (T) was determined by lysing untreated cells and measuring LLVY-AMC hydrolysis. B, HEK293T cells were treated for 10 min with varying concentrations of digitonin (0, 25, and 50 μM), and immunoblotting was used to determine the proteasome subunit levels in the medium (M) and cell pellet (P). The detection of tubulin and histone H3 verified the location of a cytosolic and nuclear protein, respectively. Basal levels of Rpn2, Rpt6, and α-proteasome subunits were established using lysates prepared from untreated cells (lane 1). Histone H3 was recovered in the pellet in both treated and untreated cells (lanes 2, 4, and 6). Treatment with digitonin resulted in significant depletion of proteasome subunits from the pellet fraction (lanes 4 and 6) and their recovery in the medium (lanes 5 and 7). C, nuclear and cytosolic fractions were prepared from HEK293T and HCT116 cells. LLVY-AMC hydrolysis was measured in the fractionated lysates using either an equal amount of protein (Conc) or proportional volumes (Vol). Peptidase activity in both cell lines was predominantly cytosolic. The sum of cytosolic (C) activity was similar to total (T) activity, indicating that the proteasome activity is predominantly in the cytosol. D, protein extracts described in C were examined by immunoblotting. Rpn7 and multiple α-subunits (Alpha mix), as well as tubulin, were detected primarily in the cytosol (C) in both cell lines, whereas histone H3 was found in the nucleus (N). Analysis of equal protein (Conc) significantly over-represents the nuclear fraction, as evidenced by the high levels of histone H3 in the nuclear (N) fraction when an equal amount of protein (Conc) was examined. In contrast, proteasome subunits were not detected in the nuclear fraction when we examined proportional volumes of the fractionated lysates. Similar findings were observed in two independent cell lines (HEK293T and HCT116).

ingly, none of these complexes was present in the nuclear fraction (Fig. 3B, lane 3), even when 5-fold excess nuclear extract was examined (lane 4).

Purified nuclear and cytosolic fractions were separated in SDS-polyacrylamide gels and probed with antibodies to confirm the location of proteasome subunits. An immunoblot was stained with Ponceau S (Fig. 3C) and then probed with antibodies against tubulin and histone H3 to gauge the purity of the cytosolic and nuclear preparations, respectively (bottom panel). Ponceau S staining showed much lower amounts of protein in the nuclear fraction when proportional volumes were examined (Fig. 3C, lane 5). In contrast, loading an equal amount of protein (Fig. 3C, conc.) significantly over-sampled nuclear proteins (lane 2), and an asterisk points to a particularly over-represented nuclear protein (Fig. 3C, compare lanes 1 and 2). Despite this over-sampling of the nuclear fraction, we detected virtually no peptidase activity in purified nuclei (Fig. 3A).

The filters were probed with antibodies against 19S (Rpt1, Rpn2, Rpn10, and Rpn12) and 20S subunits (α1, α2, α3, α5, α6, α7, β1, and β2) (Fig. 3D). High levels of proteasome subunits were detected in the cytosol (Fig. 3D, Cyto, lanes 3 and 6), consistent with the results shown in Fig. 1B. These subunits were not present in purified nuclei (Fig. 3D, Nuc, lanes 2 and 5). The levels of proteasome subunits in the cytosol and total fractions were equivalent (Fig. 3D, compare lanes 1 and 3, 4 and 6), demonstrating that proteasomes located in the cytosol represent the total cellular quantity. Similar findings were obtained whether we examined equal protein concentration (Fig. 3D, conc) or equal volume (vol).

The nuclear substrates XPC and p53 were strongly enriched in the nucleus. The retention of these proteins and histone H3 in isolated nuclei and the exclusion of cytosolic tubulin indicated that the nuclei were undamaged. Based on its known association with the nuclear envelope, it is likely that prote-

FIGURE 2.
somes are embedded within the nuclear pore complex (NPC) but remain on the cytoplasmic side (4).

**Nuclear Proteins Are Retained in Purified Nuclei**—We considered several approaches to address the concern that proteasomes could have been released into the cytosol if the nuclei were damaged during purification. We expressed bacterial derivatives of β-galactosidase in HEK293T cells; NLS-βGal-GFP contained the SV40 nuclear localization signal, whereas βGal-GFP lacked this targeting motif. NLS-βGal-GFP showed strong nuclear localization (Fig. 4A), as reported previously (31), and was retained in purified nuclei (lower panels). We note that the tetrameric form of GFP-βgal is ~575 kDa, similar to the size of the 20S catalytic particle. βGal-GFP lacking the NLS was detected only in the cytosol and was not present in purified nuclei (Fig. 4B). Proteasome peptidase activity was measured in extracts prepared from cytosol and purified nuclei from cells expressing either NLS-βGal-GFP or βGal-GFP (Fig. 4C). Consistent with earlier results, LLVY-AMC hydrolysis was only seen in the cytosolic fraction, regardless of the location of the βGal derivatives.

To confirm that purified nuclei retained structural integrity after fractionation, we examined their permeability to FITC-dextran (Fig. 4D). FITC-conjugated to 10-kDa dextran entered the nucleus immediately (Fig. 4D, upper row). Much lower lev-
els of 70-kDa dextran entered the nucleus (Fig. 4D, middle row), consistent with the ~50-kDa diffusion limit for transit of a globular protein through the nuclear pore. In contrast, 2,000-kDa FITC-dextran (Fig. 4D, lower row), which is almost as large as the 26S proteasome, was entirely excluded from the nucleus after 5 min of incubation. We also incubated purified nuclei with FITC-dextran for a significantly longer duration (3 h) in the presence and absence of 1% Triton X-100 (data not shown). After 3 h of incubation in PBS, 2,000-kDa FITC-dextran was not detected in the nucleus. However, treatment with Triton X-100 caused nuclear entry at 3 h. Collectively, these studies showed that NLS-βGal-GFP is retained in purified nuclei and 2,000-kDa FITC-dextran does not enter purified nuclei. We believe these compelling results support our hypothesis that nuclei are undamaged following purification. Consequently, the absence of catalytically active proteasomes in purified nuclei after fractionation suggests that proteasomes function in the cytosol.

The detection of GFP-tagged proteasome subunits in the nucleus has fostered the view that protein degradation occurs in the nucleus. We expressed proteasome subunit β7-GFP (7) in HEK293T cells and detected fluorescence in both the nucleus and cytosol (Fig. 5A). Fluorescence was also detected in purified nuclei (Fig. 5A, bottom row). However, purified nuclei did not contain peptidase activity (Fig. 5B) consistent with other studies described here. These results indicate that the nuclear localization of a GFP-tagged proteasome subunit does not conclusively establish the site of catalytically active proteasomes. Lysates were separated by SDS-PAGE (Fig. 5C), and immunoblotting showed that β7-GFP was present in purified nuclei (Fig. 5C, lanes 2 and 5), in agreement with the imaging results (Fig. 5A). However, native β7 was only detected in total lysate and in the cytosol (lanes 3 and 6) and not in the nuclear fraction (Fig. 5C). Thus, the location of native β7 differed from its GFP-tagged derivative. (As noted previously, the high level of β7-GFP and histone H3 in purified nuclei (Fig. 5C, lane 2) is due to oversampling of nuclear proteins when equal protein levels are examined.)

**Localization of Functional Immunoproteasome Resembles 26S Proteasome**—A previous study (32) reported that chimeric proteasomes, comprising subunits from the 26S and immunoproteasome, can be detected in cultured cells. We therefore investigated immunoproteasome localization in HEK293T cells because the substrate selectivity of chimeric proteasomes is unclear. LMP2-GFP, encoding the β1i subunit of the immunoproteasome, was uniformly distributed in HEK293T cells (Fig. 6A) and was also recovered in purified nuclei (lower panels). Immunoblotting showed that both LMP2-GFP and native LMP2 were present in the nuclear fraction (Fig. 6B). In contrast, subunits of the 20S core particle, and LLVY-AMC hydrolysis, were detected primarily in the cytosolic fraction (Fig. 6B, lanes 3 and 6). We measured immunoproteasome peptidase activity and detected Ac-PAL-AMC hydrolysis only in the cytosolic fraction (Fig. 6C), similar to the hydrolysis of LLVY-AMC (Fig. 6D). Thus, peptidase activity corresponding to both 26S proteasome and the immunoproteasome was only detected in the cytosol, irrespective of the subcellular distribution of individual GFP-tagged subunits.

**Cytosolic Proteasomes Are Quantitatively Intact**—Cytosolic extracts were prepared from HEK293T cells and resolved in Sephacryl 300HR, and proteasome peptidase activity, subunit levels, and composition were examined. Peptidase activity was present in fractions representing high molecular weight species (Fig. 7A, fractions 13 and 14), consistent with the size of intact

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**FIGURE 4. Purified nuclei are structurally intact.** A, HEK293T cells were transfected with a vector expressing NLS-βGal-GFP, and fluorescence was detected predominantly in the nucleus (upper panels). Following fractionation, GFP fluorescence was retained in the purified nuclei (lower panels). B, βGal-GFP (lacking a nuclear localization signal) was similarly expressed in HEK293T cells, and cytosolic expression was observed (upper panels). Purified nuclei showed no evidence for GFP fluorescence (lower panels). C, we measured proteasome peptidase activity in cells described above. LLVY-AMC hydrolysis in the cytosol (C) and nucleus (N) was compared with activity present in total lysate (T). Error bars indicated S.D. nuclei were isolated from HEK293T cells and incubated with FITC-dextran in the sizes indicated. FITC 10-kDa dextran readily entered the nucleus, whereas FITC 2,000-kDa dextran was entirely excluded. Hoechst staining of nuclei and a merged differential interference contrast image are also shown.
Purified nuclei contained some activity is present in the cytosol. Peptidase activity (data not shown), indicating that all proteasomes contained 20S particles (Fig. 7). Peptidase activity was detected in fractions 16 and 17, which contained 26S proteasomes in fractions 13 and 14. Weak peptidase activity was detected in fractions 16 and 17, which contained 20S particles (Fig. 7C). We also separated 10-fold excess nuclear extract in the same column and detected no peptidase activity (data not shown), indicating that all proteasome activity is present in the cytosol.

26S proteasomes. Lower activity was present in subsequent fractions corresponding to free 20S particles (Fig. 7A, fractions 16 and 17). All proteasome activity that was applied to the column was recovered in these fractions, and none was detected in the trailing column fractions. Because proteasome subunits are present predominantly in intact complexes in vivo (18), the weak 20S, specific activity observed in fractions 16 and 17, could reflect post-lysis dissociation of the 26S proteasome. The fractionated proteins were examined by immunoblotting, and 20S subunits (α4, β7) were found in all fractions that contained peptidase activity (Fig. 7B). Subunits in the 19S regulatory particle (Rpn2, Rpt1, and Rpn7) were also detected primarily in fractions 13 and 14. To confirm that these subunits were present in intact 26S proteasomes, we separated proteins in a native polyacrylamide gel. In-gel hydrolysis of LLVY-AMC confirmed the presence of 26S proteasomes in fractions 13 and 14. Weak peptidase activity was detected in fractions 16 and 17, which contained 20S particles (Fig. 7C). We also separated 10-fold excess nuclear extract in the same column and detected no peptidase activity (data not shown), indicating that all proteasome activity is present in the cytosol.

The degradation of a number of nuclear proteins in yeast and human cells requires export. The proposed cytosolic localization of catalytically active proteasomes imposes an unexpected requirement for transporting substrates out of the nucleus. We examined the effect of nuclear export on p53 stability using both pharmacological and genetic approaches. Native p53 was not detected in untreated cells but was readily detected after 18 h of treatment with leptomycin B (LMB) (data not shown). Based on these findings, we compared the stability and localization of an export-deficient p53 mutant (GFP-p53nes−). The level and stability of native p53 were similar in HCT116 cells expressing either GFP-p53 (Fig. 8A, lanes 1–5), or GFP-p53nes− (lanes 7–11). Native p53 was undetected in the absence of LMB (Fig. 8A, lanes 6 and 12; UNT) but was readily detected after exposure to LMB (lanes 1 and 7; + LMB, 18 h). Both an export defect and treatment with LMB had a strong stabilizing effect on GFP-p53nes− levels (Fig. 8A, lanes 7 and 11). HCT116 cells were suspended in fresh medium either containing (+) or lacking (−) LMB, and protein levels were followed. In the presence of LMB, noticeably higher levels of native p53 remained after 24 h (Fig. 8A, lane 5). In contrast, the level of native p53 declined in the absence of LMB (Fig. 8A, lane 4). GFP-p53nes− levels did not decrease after the removal of LMB (Fig. 8A, lanes 8 and 10), as this mutant is already export-deficient. Protein levels were quantified by densitometry and adjusted to the abundance of the α7 proteasome subunit. Numerical values in parentheses indicate the fold change, compared with time 0 (Fig. 8A, lanes 1 and 7).

We investigated where stabilized GFP-p53nes− accumulated in the cell. Fluorescence microscopy showed that both GFP-p53 and GFP-p53nes− were localized predominantly in the nucleus (Fig. 8B and C, 18 h + LMB). To examine the effect of inhibiting export, we compared the levels of GFP-p53 and GFP-p53nes−, and after removal of LMB from the growth medium GFP-p53 was entirely lost within 24 h (Fig. 8B, lane 7 – LMB). In contrast, if LMB was present in the medium, the nuclear levels of GFP-p53 persisted (Fig. 8B, lane J, 24 h + LMB). In contrast, the nuclear levels of GFP-p53nes− remained high even after LMB was removed (Fig. 8C, lane K, 24 h – LMB), consistent with the export deficiency of this mutant protein.

Discussion

The presence of proteasome subunits and subcomplexes in the nucleus (2–4, 16, 17, 22, 24, 25) is consistent with the view that proteins can be degraded in the nucleus (14, 18–21, 23). However, the site of proteasome-specific peptidase activity has not been examined comprehensively. Although GFP-tagged proteasome subunits are detected in the nucleus (2, 3) and are enriched in the nuclear envelope (4, 33), we detected peptidase activity only in the cytosol. One interpretation of this result is that proteasomes are present on the nuclear surface and are dissociated during fractionation. This arrangement supports our hypothesis that the degradation of nuclear substrates requires an export mechanism (2).

The degradation of misfolded nuclear and cytosolic proteins has been investigated using CPY* and other engineered proteins that display a strong tendency to misfold (19–21). The degradation of some of these proteins, such as ΔsPrA, Δ2GFP...
Although proteins can be conjugated to ubiquitin in the nucleus, there is insufficient evidence to conclude that they are degraded there, especially because export is required for the turnover of many nuclear proteins (2, 9–12, 35). Therefore, although misfolded cytosolic proteins may enter the nucleus, it less clear that they are degraded there. Intriguingly, the degradation of ΔssPrA, Δ2GFP, and CPY-GFP required the activity of nuclear ubiquitin E3 ligases (San1) and cytosolic E3 ligase (Ubr1) (14, 20, 34). Δ2GFP was strongly stabilized in san1 and accumulated in the nucleus. In contrast, Δ2GFP was only partially stabilized in ubr1 and was detected in both the nucleus and cytosol. Although San1 and Ubr1 may function independently, they could represent a two-step targeting paradigm in which San1 initiates ubiquitylation of a substrate in the nucleus that is followed by its cytosolic ubiquitylation by Ubr1. In agreement with this model, the attachment of an export signal rendered CPY-GFP vulnerable to ubiquitylation by cytosolic Ubr1 (34). Increased export could reduce nuclear retention and favor CPY-GFP ubiquitylation by cytosolic Ubr1. San1 might represent a rate-limiting or commitment step for the elimination of misfolded nuclear proteins. The distinct but coupled ubiquitylation of nuclear substrates by San1 and Ubr1 offers a coordinated mechanism for degrading misfolded nuclear proteins and mislocalized cytosolic proteins (34). It remains to be determined whether these E3 ligases function in concert to degrade physiological nuclear substrates.

The NPC plays a key role in DNA-related activities, including the degradation of transcription factors (36), DNA repair proteins (37), and in transcription silencing (38). Components in the NPC interact directly with chromatin (39) and gene promoters (16). Chromatin immunoprecipitation studies showed that proteasome subunits can be co-purified with the GAL10 gene and promoter (16), although the presence of proteasome peptidase activity was not verified. Whereas fluorescence imaging of nuclear pore subunits, such as Nup49-mCherry, reveals a ring circumscribing the nucleus, GFP-tagged proteasomes display a more dispersed signal across the nucleus, with intensification at the periphery. We note that certain individual proteasome subunits can perform non-proteolytic roles inside the nucleus, apart from intact proteasomes. In addition, proteasomes may partly enter into the volume of the nucleus without entering the nucleoplasm. These two observations could explain the diffuse nuclear localization of proteasome subunits using GFP-tagged subunits.

Niepel et al. (41) reported that a proteinaceous basket that is associated with the nuclear pore can extend ~80 nm into the interior of the nucleus. Myosin-like proteins Mlp1 and Mlp2 are located at the base of this basket and provide a gateway between the nucleus and the cytosol (40). Mlp1 interacts with...
Esc1, a peripheral nuclear protein that binds proteasomes (41). This is significant because Mlp1 also binds promoters of many inducible genes, including \textit{GAL10} (40). The juxtaposition of active transcription units and proteasomes through their interactions with Mlp1 and Esc1 provides a path for degrading transcription factors (2, 36) and DNA repair proteins (2). In addition, both Mlp1 and Mlp2 bind Cdc31 (centrin) (41), which we reported can bind the proteasome and multiubiquitylated proteins (26). Cdc31 (centrin) is also linked to the nuclear export machinery (42). Collectively, these findings suggest that multiple NPC components could promote the degradation of nuclear proteins (see model, Fig. 9). Proteasomes are mobile and can be recruited from cytosolic storage granules to the nucleus within minutes (18, 43). Because proteasomes are assembled in the cytosol (18), their rapid mobilization to the nucleus might indicate localization within the NPC and not entry into the nucleoplasm. Based on this hypothesis, nuclear export could guide proteolytic substrates to proteasomes in the NPC-associated basket.

Although we propose that proteasomes do not operate in the nucleus, we consider below alternative interpretations. We recognized that if nuclei were damaged during fractionation proteasomes could have leaked into the cytosol. To address this concern, we isolated nuclei using both isotonic and hypertonic lysis methods, and both approaches showed proteasome activity exclusively in the cytosol. We also used digitonin to selectively permeabilize the plasma membrane (31), and we determined that peptidase activity was released into the culture medium (Fig. 2, \textit{A} and \textit{B}). A low level of the free 20S particle was detected in fractions 16 and 17. Both total (\textit{T}) and nuclear (\textit{Nuc}) extracts showed peptidase activity in the high molecular weight fractions. In contrast, 10-fold excess nuclear extract showed no peptidase activity.

**FIGURE 7.** Cytosolic proteasomes are intact. \textit{A}, cytoplasmic protein extracts prepared from fractionated HEK293T cells were resolved in Sephacryl 300HR. Two peaks of proteasome activity were detected. Fractions 13 and 14 were coincident with the exclusion volume. The data represent one of three independent fractionation experiments. \textit{B}, immunoblotting showed the presence of both 19S (Rpn2, Rpt1, and Rpn7) and 20S subunits (\textit{α}4 and \textit{β}7) in fractions 13 and 14, which displayed peak LLVY-AMC peptidase activity. Very low levels of the 20S particle were detected in subsequent fractions. \textit{C}, Sephacryl 300HR fractions 12–18 were resolved in a native polyacrylamide gel. An "in-gel" assay, performed by overlaying the gel with fluorogenic substrate, revealed intact proteasomes in fractions 13 and 14, consistent with \textit{A} and \textit{B}. A low level of the free 20S particle was detected in fractions 16 and 17. Both total (\textit{T}) and nuclear (\textit{Nuc}) extracts showed peptidase activity in the high molecular weight fractions. In contrast, 10-fold excess nuclear extract showed no peptidase activity.
We examined the localization of nuclearly targeted bacterial β-galactosidase (31) to address the concern that proteins that were retained in the nucleus may have been pelleted in association with chromatin. βGal-GFP was attached to NLS, and NLS-βGal-GFP was quantitatively localized in the nucleus (Fig. 4A), whereas βGal-GFP was cytosolic (Fig. 4B). The retention of NLS-βGal-GFP, which is ~575 kDa, in purified nuclei lends strong support to our view that if proteasomes (2,600 kDa) were present in the nucleus they would not have leaked out during fractionation. In addition, if proteasomes did function in the nucleus, peptidase activity should have been detected in purified nuclei. However, the absence of nuclear peptidase activity disputes this view. We propose that nuclear substrates are exported and degraded in the cytosol.

FIGURE 8. Export defect stabilizes p53 and causes accumulation in the nucleus. A, we expressed GFP-p53 and GFP-p53(nes−), a mutant that is poorly exported, in HCT116 cells. GFP-p53 and native p53 are poorly detected in untreated cells (lane 7). Treatment of cells with leptomycin B (+LMB 18 h) resulted in elevated levels of native p53 and the GFP-tagged derivatives. The cells were placed in fresh medium either containing (+) or lacking (−) LMB, and extracts were prepared after 14 and 24 h. Native p53 levels decreased after 24 h in the absence of LMB but remained elevated in the presence of LMB (compare lanes 4 and 5). Similarly, we detected a modest decrease in the levels of GFP-p53 at 24 h (−LMB). In contrast, the levels of GFP-p53(nes−) were strongly increased following treatment with LMB, and removal of LMB did not result in any appreciable decrease in its levels, even after 24 h (lane 10). B and C, HCT116 cells were pretreated with LMB for 18 h, and the localization of GFP-p53 and GFP-p53(nes−) was examined. Both GFP-p53 and GFP-p53(nes−) were strongly enriched in the nucleus (upper panels). Cells were incubated in fresh medium either containing (+) or lacking (−) LMB, and the protein levels were examined after 24 h. GFP-p53 was entirely eliminated, whereas high levels of nuclear GFP-p53(nes−) remained. Unt, untreated.

We examined the localization of nuclearly targeted bacterial β-galactosidase (31) to address the concern that proteins that were retained in the nucleus may have been pelleted in association with chromatin. βGal-GFP was attached to NLS, and NLS-βGal-GFP was quantitatively localized in the nucleus (Fig. 4A), whereas βGal-GFP was cytosolic (Fig. 4B). The retention of NLS-βGal-GFP, which is ~575 kDa, in purified nuclei lends strong support to our view that if proteasomes (2,600 kDa) were present in the nucleus they would not have leaked out during fractionation. In addition, if proteasomes did function in the nucleus, peptidase activity should have been detected in purified nuclei. However, the absence of nuclear peptidase activity disputes this view. We propose that nuclear substrates are exported and degraded in the cytosol.
In a complementary approach, we incubated purified nuclei with FITC-dextran and found that 10-kDa dextran readily entered the nucleus, whereas the 2,000-kDa FITC-dextran was entirely excluded (Fig. 4D). However, high molecular mass dextran entered the nucleus after we permeabilized the nuclear envelope with detergent (data not shown). Using the yeast experimental model, we found HDAC activity only in purified nuclei, whereas peptidase activity was present in the cytosol (Fig. 1A), consistent with our hypothesis.

The lack of nuclear peptidase activity could be the result of selective proteasome inactivation in the nucleus. Regulatory factors are known to modulate proteasome function, and therefore, it was possible that nuclear proteasomes are only conditionally activated. To investigate this possibility, we separated 10-fold excess nuclear extract by gel exclusion chromatography to detect intact proteasomes. We found that none of the fractions corresponding to high molecular weight complexes contained proteasome subunits. We note that cytosolic peptidase activity is essentially identical to total activity. We conclude that cytosolic proteasomes contribute the major fraction of peptidase activity.

An important distinction between our findings and recent reports (19–21) is that we examined proteasome localization and not substrate turnover. Although native and misfolded proteins could be degraded by mechanistically distinct paths, it is germane that catalytically active proteasomes were not detected in the nucleus, and thus export must contribute significantly to the turnover of many types of nuclear proteins. Because multiple pathways facilitate nuclear export, the degradation of specific classes of nuclear substrates might involve distinct mechanisms. Consequently, inhibiting a single export pathway might not unequivocally demonstrate whether a nuclear substrate is degraded in the cytosol. For instance, we detected GFP-tagged proteasome subunits in both the nucleus and cytosol, but found peptidase activity only in the cytosol in fractionated HeLa cells (7). In agreement, we detected GFP-β7 in the nucleus and cytosol, but found peptidase activity only in the cytosol, in both HEK293T and HCT116 cells (Fig. 2, C and D). These findings suggest that β7 in the nucleus is not present in intact proteasomes. It is also significant that multiple proteasome subunits were detected only in the cytosol in primary mouse cardiac myocytes (49), demonstrating that our findings in yeast and cultured cells are conserved in other animal models.

GFP tags are valuable for identifying the subcellular localization of proteins. However, it is less clear whether the location of a GFP-tagged proteasome subunit provides unambiguous evidence for the site of peptidase activity, because imaging studies cannot distinguish between intact proteasomes and free subunits. This concern is relevant because proteasome subunits, such as Rpn4, can perform non-proteolytic roles in transcription (6); other components (Rpn10) are expressed at super-stoichiometric levels (44, 45) and are not present exclusively in the proteasome (46). Consequently, proteasome subunits that are detected in the nucleus may be engaged in non-proteolytic functions. It has also been reported that certain GFP-tagged subunits (α7, α5, and β7) are unable to assemble efficiently into proteasomes (47), and in some instances, the fluorescence generated is reduced following assembly (7). These observations raise concern that GFP fluorescence arises disproportionately from free subunits. Pack et al. (18) expressed fusions of regulatory particle (RP) and catalytic particle (CP) proteasome subunits and found that the chimeras could replace the wild type proteins. The co-localization of these chimeras with the nucleus suggested that intact proteasomes operated in the nucleus. However, different proteasome subunits were tagged to GFP to define proteasome localization, and it is unclear whether they were entirely localized with intact proteasomes. These and other studies did not establish whether proteasomes entered the nucleus or remained at the nuclear surface. We report here that peptidase activity is detected in the cytosol, suggesting that proteasomes may be bound to the nuclear surface.

The presence of NLS motifs in certain proteasome subunits suggested that they facilitate entry into the nucleus (25). However, we reported that the targeting of proteasomes to the nucleus is entirely blocked in srp1-49, although Srp1 plays no role in nuclear import. The targeting of proteasomes to the nucleus also requires Srp1, a yeast importin-α protein. However, in contrast to other reports we determined that srp1-49 can successfully import NLS-bearing proteins but is specifically defective in targeting proteasomes to the nucleus (48). We speculate that Srp1 guides proteasomes to the nuclear pore complex but does not facilitate entry into the nucleus.

Our studies agree with the results of Kulichkova et al. (7), who detected human GFP-β7 in both the nucleus and cytosol. They also reported that intact proteasomes were present exclusively in the cytosol in fractionated HeLa cells (7). In agreement, we detected GFP-β7 in the nucleus and cytosol, but found peptidase activity only in the cytosol, in both HEK293T and HCT116 cells (Fig. 2, C and D). These findings suggest that β7 in the nucleus is not present in intact proteasomes. It is also significant that multiple proteasome subunits were detected only in the cytosol in primary mouse cardiac myocytes (49), demonstrating that our findings in yeast and cultured cells are conserved in other animal models.

Proteasome trafficking is dynamically regulated in response to growth conditions (43). Similarly, the degradation of certain substrates requires movement into and out of the nucleus. As many nuclear functions occur in the vicinity of the nuclear pore, a model envisioning proteolysis in the NPC would not be unorthodox. Stationing proteasomes in the nuclear pore com-
plex would allow degradation of nuclear proteins without requiring entry into the nucleus. To reconcile the divergent views on nuclear protein turnover, we propose that export pathways traffic substrates to the NPC where proteasomes are located (Fig. 9). Further study will be required to establish the veracity of this paradigm.

Experimental Procedures

Yeast Strains and Methods—Exponential phase wild type cells expressing FLAG-H2B were pelleted, suspended at 1 ml/g in pretreatment buffer (PB: 50 mM Tris-HCl, pH 7.5, 10 mM MgCl₂, 1 m sorbitol, and 60 mM 2-β mercaptoethanol), and incubated at room temperature for 15 min with gentle mixing. Cells were collected by centrifugation for 5 min at 1,500 × g, suspended in 3 ml/g digestion buffer (DB: PB + 5 mM 2-β mercaptoethanol, 2 mg/ml zymolyase 20T), and incubated at 37 °C for 60 min. Spheroplasts were collected by centrifugation at 4,000 × g for 10 min and then washed three times with cold DB. Spheroplasts were suspended in 0.5 ml/g DB and lysed by dropwise addition to 20 volumes of cold lysis buffer (LB: 18% Ficoll, 10 mM Tris-HCl, pH 7.5, 20 mM KCl, 5 mM MgCl₂, 5 mM 2-β mercaptoethanol, 1 mM EDTA and protease inhibitors). Unlysed cells and debris were removed by centrifugation for 30 min at 3,000 × g. An aliquot of the suspension (3 ml) was removed and lysed by sonication to provide a sample representing total (Nuc + Cyto) cellular protein. The remaining supernatant was centrifuged for 20 min at 20,000 × g to pellet the nuclei. The nuclear pellet was suspended in LB at ~1/7th of the cytoplasmic volume, as the yeast nucleus represents ~15% of total cell volume (50). Nuclear proteins were recovered following disruption by sonication (Nuc). An equal volume (20 μl) of total, Nuc, and Cyto was used to measure HDAC (Sigma catalog no. CS1010) and proteasome activities (see below and Ref. 51).

One hundred microliters of each fraction was resolved in a 12% SDS-Tricine/polyacrylamide gel and examined by immunoblotting.

Human Cell Lines and Plasmids—HEK293T cells were provided by B. Firestein (Rutgers University). HCT116 cells were provided by W. Hu (Cancer Institute of New Jersey). A plasmid encoding human pQCXIP provided by W. Hu (Cancer Institute of New Jersey). A plasmid provided by B. Firestein (Rutgers University). HCT116 cells were cultured at 37 °C in Dulbecco’s modified Eagle’s medium or PBS (containing protease inhibitors) at 4 °C. Western Blotting Analysis and Immunoprecipitation—Protein samples were resolved in a 12% SDS-Tricine-polyacrylamide gel, transferred to 0.45-μm nitrocellulose, and examined by immunoblotting. Chemiluminescence was captured using Kodak GelLogic Imaging software. Antibodies against histone H3 and p53 were purchased from BioLegend and Origene, respectively. Tubulin antibody was purchased from Life Technologies, Inc., catalog no. R37605) were added to 5% of the purified nuclei (5 μl) and incubated on ice for 5 min. Microscopy images were captured using appropriate optical filters (excitation 430 nm; emission 535 nm). Derivatives of FITC-dextran (representing 10, 70, and 2,000 kDa of dextran) were purchased from Sigma.

Gel Exclusion Chromatography—Purified cytosolic proteins (~2 mg) were clarified by brief centrifugation and resolved in Sephacryl S-300 HR at 4 °C. The column was equilibrated in 50 mM potassium phosphate, 5 mM EDTA, 100 mM NaCl, 10 mM MgCl₂, 2 mM ATP, 1 mM DTT, and 5% glycerol. The sample represents ~1% of column volume, and 1-ml fractions were collected at a rate of ~0.5 column volume/h. Proteasome peptidase activity was measured in the fractions, and those containing peak activity were further characterized in native in-gel assays.

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a final concentration of 0.25, and 50 μg/ml. Samples were incubated at room temperature for 10, 20, and 30 min and centrifuged at 500 × g for 5 min at 4 °C, following which the supernatant was removed to assay proteasome peptidase activity.

Nuclei Purification—HEK293T and HCT116 cells were grown to ~70–80% confluence. Media were removed, and cells were washed twice with PBS. EZ-lysis buffer (500 μl; Sigma catalog no. NUC-101), containing 5 mM ATP, 1 mM DTT, and protease inhibitors, was added to each flask. Lysed cells were dislodged and transferred to a single microcentrifuge tube (yielding ~1.7 ml of cell suspension). Lysis was completed on ice for 5 min, following which 500 μl was removed and sonicated, serving as a total protein sample. The remainder was centrifuged at 500 × g for 5 min at 4 °C. The supernatant, containing cytoplasmic proteins, was withdrawn, and the pellet was washed in lysis buffer to remove residual cytoplasm. The pellet containing purified nuclei was suspended in 120 μl of lysis buffer and sonicated. Total, cytosolic, and nuclear fractions were examined biochemically and by gel filtration and electrophoresis.

Nuclear Permeabilization Studies—HEK293T cells were grown to ~70% confluence in a 150-cm² flask. Cells were collected and fractionated to separate nuclei from cytosol, as described above. Purified nuclei were suspended in 100 μl of PBS containing a protease inhibitor mixture. FITC-dextran (5 μl at 1 mg/ml) and Hoechst-33342 (1 μl; NucBlue Live Stain from Life Technologies, Inc., catalog no. R37605) were added to 5% of the purified nuclei (5 μl) and incubated on ice for 5 min. Microscopy images were captured using appropriate optical filters (excitation 430 nm; emission 535 nm). Derivatives of FITC-dextran (representing 10, 70, and 2,000 kDa of dextran) were purchased from Sigma.
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Microscopy—Transfected cells were fixed and permeabilized with 4% paraformaldehyde in culture medium and 0.15% Triton X–100 in PBS, respectively. Hoechst was added directly to each slide prior to microscopy. Images were captured using Zeiss Imager.M1 microscope, an AxioCam MRm camera, Plan-Apochromatic 100×/1.4 objective lens, and Zeiss AxioVision software.

Biochemical Assays—For the proteasome activity measurement, 5 μl of epoxomicin (prepared in 50% DMSO at 50 ng/μl) was preloaded in duplicate in a 96-well plate. Similarly, 5 μl of vehicle (50% DMSO) was loaded in duplicate. Protein fractions (25 μl) were added to the aforementioned four wells. The proteasome assay buffer (200 μl: 25 mM Tris, pH 7.5, 1 mM EDTA) containing 40 μM Suc-LLVY-AMC was added to each well. A Tecan fluorometer was used to detect fluorescence generated by the hydrolysis of Suc-LLVY-AMC at 37 °C, using excitation wavelength 360 nm and emission wavelength 465 nm. Results were captured at 5-min intervals for at least 30 min. The background value was determined by measuring epoxomicin-insensitive activity, which was typically ~5% of total fluorescence. GFP fluorescence was measured using a Tecan fluorometer using excitation wavelength 430 nm and emission wavelength 535 nm.

Immunoproteasome Activity Measurement—Total, cytosolic, and nuclear protein extracts were prepared and examined as described above. Immunoproteasome assay buffer contained 14.38 μM Ac-PAL-AMC, and fluorescence was detected using the Tecan fluorometer. Data were captured at 5-min intervals (excitation wavelength 360 nm/emission wavelength 465 nm).

p53 Chase with Leptomycin B—HCT116 cells were grown to ~70% confluence. The growth medium was removed and replaced with fresh media containing 10 nM LMB. Following incubation for 18 h the medium was replaced and half the dishes were supplemented with LMB. Cells were harvested, pelleted, and frozen at the times indicated. GFP derivatives of wild type p53 and an export mutant (p53nes−) were transfected into HCT116 cells 48 h prior to treatment with LMB.

Author Contributions—K. M. and L. C. designed and coordinated the study. The manuscript was prepared by K. M. Figures were prepared by F. D. Data shown in Fig. 1 were generated by L. C. All other data were generated by F. D. Protocols for in-gel analysis were refined by L. C. K. M. guided studies involving gel filtration. All authors proofread and approved the final version of this manuscript. We thank members of the laboratory for critical review of this manuscript.

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