Cyclosporine A-Sensitive, Cyclophilin B-Dependent Endoplasmic Reticulum-Associated Degradation

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Abstract
Peptidyl-prolyl cis/trans isomerases (PPIs) catalyze cis/trans isomerization of peptide bonds preceding proline residues. The involvement of PPI family members in polypeptide refolding has been established in test tube experiments. Surprisingly, however, no data is available on the involvement of endoplasmic reticulum (ER)-resident members of the PPI family in protein folding, quality control or disposal in the living cell. Here we report that the immunosuppressive drug cyclosporine A (CsA) selectively inhibits the degradation of a subset of misfolded proteins generated in the ER. We identify cyclophilin B (CyPB) as the ER-resident target of CsA that catalytically enhances disposal from the ER of ERAD-L5 substrates containing cis proline residues. Our manuscript presents the first evidence for enzymatic involvement of a PPI in protein quality control in the ER of living cells.

Introduction
Formation and reduction of covalent bonds between cysteine side chains and cis/trans isomerization of peptide bonds preceding proline residues are rate-determining steps for the attainment of the native and functional 3D structure of polypeptides synthesized in the ER. The reactions might also be rate-limiting for the unfolding of aberrant polypeptides that require retro-translocation (dislocation) across the ER membrane for proteasomal degradation [1]. In vitro, these reactions are catalyzed by protein disulfide isomerase (PDIs) [2] and by PPIs [3,4], respectively. Extensive experimental evidence has shown the importance of PDI-assisted polypeptide folding and unfolding in living cells [5,6]. Despite 25 years of PPI catalysis experiments in vitro, a direct involvement of PPIs in catalysis of protein folding, in regulation of protein quality control or in clearance of misfolded polypeptides from the ER of living cells remains to be demonstrated [7,8,9].

Most polypeptides entering the ER lumen are covalently modified at asparagine side chains with glucose-1,2-mannosyl-α-N-acetylglucoexamine-2-oligosaccharides. Their maturation is assisted by a dedicated folding machinery comprising the oligosaccharide-binding chaperones calnexin and calreticulin and the oxidoreductase ERP57 [10]. Processing of oligosaccharides displayed on misfolded conformers by ER-resident α1,2-mannosidases, with removal of up to 4 terminal mannose residues, irreversibly extracts folding-defective polypeptides from the lectin-operated folding machinery [11]. In mammalian cells, two ER-associated degradation (ERAD) shuttles, OS-9 and XTP3-B [12,13,14], transport ERAD-L2 substrates (i.e. soluble, extensively de-mannosylated terminally misfolded glycopolypeptides) from the ER lumen to the site of dislocation across the ER membrane [15]. OS-9 and XTP3-B deliver ERAD-L5 substrates to a multi-protein complex comprising the membrane receptor SEL1L, the associated E3 ubiquitin ligase HRD1 and an elusive dislocation (retro-translocation) channel [16]. The stringent requirement for HRD1, SEL1L and OS-9/XTP3-B for disposal is bypassed when the same misfolded domains are tethered to the ER membrane (ERAD-L5 substrates) [15,17]. Thus, luminal misfolded polypeptides and membrane-tethered polypeptides with structural defects in the ER lumen have different requirements for efficient clearance from the ER.

Although the process of dislocation across the ER membrane is poorly defined, unfolding of aberrant polypeptide chains [18] and disassembly of disulfide-bonded protein aggregates [19] have been shown to facilitate protein clearance from the ER lumen. A role in ERAD has been demonstrated for several members of the PDI superfamily (e.g. PDI, ERP37, ERP72, ERP29, ERdj5), thus implying that reduction of inter- and intra-molecular disulfide bonds plays a crucial role in ERAD by eliminating tertiary and quaternary structures that could impair transport across a putative proteinaceous membrane dislocorn [reviewed in [5]]. On the same line, it is conceivable that the PPIs-catalyzed interconversion of cis into trans peptidyl-prolyl bonds could facilitate dislocation of ERAD substrates across the ER membrane by eliminating turns in the polypeptide secondary structure [9].

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Citation: Bernasconi R, Soldà T, Galli C, Pertel T, Luban J, et al. (2010) Cyclosporine A-Sensitive, Cyclophilin B-Dependent Endoplasmic Reticulum-Associated Degradation. PLoS ONE 5(9): e13008. doi:10.1371/journal.pone.0013008

Editor: Suzannah Rutherford, Fred Hutchinson Cancer Research Center, United States of America

Received June 24, 2010; Accepted September 1, 2010; Published September 28, 2010

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Funding: M.M. is supported by grants from the Foundation for Research on Neurodegenerative Diseases, the Fondazione San Salvatore, the Swiss National Center of Competence in Research on Neural Plasticity and Repair, the Swiss National Science Foundation and ONELIFE Advisors SA. J.L. is supported by grants from the Swiss National Science Foundation, the NIH (AI059159) and the EU project HIV-ACE. The funders had no role in study design, data collection and analysis, decision to publish, or preparation of the manuscript.

Competing Interests: The authors have declared that no competing interests exist.

These authors contributed equally to this work.
Here we report that the immunosuppressive drug CsA, a specific inhibitor of the cyclophilin family of PPIs, selectively delays the degradation of the ERAD-LS substrate BACE457Δ leaving unaffected disposal from the ER of the same polypeptide when tethered to the ER membrane (the ERAD-LM protein BACE457). This identifies CsA as the first inhibitor that selectively acts upon an ERAD-LS substrate and not upon the corresponding ERAD-LM polypeptide. We then extend this finding by showing that, among roughly 20 mammalian cyclophilin family members, CyPB is unique because it plays a crucial role in ERAD that requires its enzymatic activity. Importantly, CsA is not a general inhibitor of the ERAD-LS pathway and CyPB is not required for disposal of all ERAD-LS substrates. Rather, the presence of peptidyl-prolyl bonds in the cis configuration renders disposal of ERAD-LS substrates sensitive to CsA and dependent on CyPB intervention. Altogether, our manuscript presents the first evidence for the enzymatic involvement of a PPI in protein quality control in the ER of a living cell.

Results and Discussion

CsA selectively inhibits disposal of BACE457Δ

BACE457 and BACE457Δ are splice variants of the human beta-site amyloid precursor protein cleaving enzyme BACE501 [20], an aspartic protease involved in generation of the Aβ peptide that forms plaques in the brain of Alzheimer’s disease patients. A 44-residue deletion in the ectodomain prevents attainment of the native structure and results in degradation from the ER lumen when the proteins are ectopically expressed in cultured cells. Proteasome-dependent disposal of both proteins requires intervention of EDEM variants and extensive de-mannosylation of the 2 protein-bound N-glycans [19,21,22,23]. However, degradation of BACE457Δ, an ERAD-LS protein, strictly depends on HRD1, SELIL and OS-9/XTP3-B, while disposal of BACE457, an ERAD-LM protein, progresses efficiently even upon inactivation of the HRD1 pathway [15].

BACE457 and BACE457Δ contain 26 and 25 proline residues, respectively. It is impossible to establish if, and which one of the peptidyl bonds preceding these proline residues is converted from the trans to the cis configuration during the short retention of these folding-defective polypeptides in the ER lumen. It is of interest, however, that in the folding competent variant BACE501 the peptidyl bonds preceding Pro84, Pro146 and Pro390 are in the cis configuration (see below and Materials and Methods). To assess whether prolyl isomerases might facilitate disposal of BACE457 and BACE457Δ from the mammalian ER, we exposed cells transiently transfected for expression of either one of the two model substrates to CsA, a selective inhibitor of immunophilin members of the PPIs family [24]. CsA-treatment was compared with cell exposure to a series of well-characterized ERAD inhibitors (thapsigargin (Tg, which inhibits the SERCA pump with cell exposure to a series of well-characterized ERAD members of the PPIs family [24]. CsA-treatment was compared

Confirming published data [19,21,22,23], Tg, Kif and PS341 substantially delayed disposal of both BACE457 and BACE457Δ (Figs. 1A–1D). CsA did not inhibit degradation of BACE457 (Figs. 1A–1B, lane 3 vs lane 2 and Figs. 2B–2C, lanes 1–3), but substantially delayed the clearance from the ER lumen of BACE457Δ (Figs. 1C–1D, lane 3 vs lane 2 and Figs. 2D–2E, 3B–3C, lanes 1–3) as efficiently as the conventional ERAD inhibitors Tg, Kif and PS341 (Figs. 1C–1D, lanes 4–6). To summarize, we identify CsA as the first compound that selectively inhibits disposal of a soluble (ERAD-LS) but not of a membrane-tethered (ERAD-LM) variant of a misfolded polypeptide.

CsA is not a general inhibitor of the ERAD-LS pathway

To determine whether CsA is a general inhibitor of the ERAD-LS pathway, we next checked whether cell exposure to CsA delayed disposal of CD3Δα, Like BACE457Δ, this tris-glycosylated, soluble and folding-defective ERAD-LS protein stringently depends on HRD1, SELIL and OS-9/XTP3-B for efficient disposal [15]. As expected for an ERAD substrate, disposal of CD3Δα was substantially delayed upon inactivation of protein de-mannosylation and upon inactivation of 26S proteasomes (Figs. 1E–1F, lanes 4 and 5, respectively). However, CsA was ineffective in preventing clearance of CD3Δα from the ER (lane 3). Thus, even though CsA substantially delayed disposal of the ERAD-Ls protein BACE457Δ (and of other canonical ERAD-Ls substrates such as BACE476Δ [Figs. 1G–1H] and NHK, a folding-defective version of the secretory protein ζ1-antitrypsin (Figs. 11–1L)), the incapacity of CsA to prevent CD3Δα disposal showed that CsA is not a general inhibitor of the ERAD-Ls pathway.

Why is the disposal of CD3Δα insensitive to CsA and the disposal of other ERAD-Ls substrates efficiently delayed by this PPI inhibitor? It is possible that none of the peptidyl-prolyl bonds of the misfolded CD3Δα retained in the ER lumen is in the cis configuration, while one or more peptidyl-prolyl bonds of the misfolded BACE variants are in cis and must be isomerized to the trans conformation to promote efficient clearance from the ER. Of some relevance in this context could be that the corresponding native proteins do not have (the CD3δ in the functional T cell receptor) or do have peptidyl-prolyl bonds in the cis configuration (the native BACE501, Materials and Methods). We therefore hypothesized that the presence of peptidyl-prolyl bonds in the cis configuration determines CsA-sensitivity for the disposal of ERAD-Ls polypeptides from the mammalian ER (see next sections).

CyPB is the luminal CsA target involved in ERAD

CsA is a cyclic undecapeptide produced by the fungus Toxoplasma gondii [28]. It is used in the clinic as an immunosuppressant to reduce the risk of graft reaction upon allogenic transplant and to improve short-term allograft survival [28]. The PPI family member CyPB is the ER-resident target of CsA [29]. A role for CyPB (or of any other PPI family member) in catalysis of peptidyl-prolyl cis/trans isomerization in protein biogenesis and/or quality control in the ER of living cells is not supported by experimental data. To determine whether CyPB intervenes in protein disposal from the ER lumen, we compared degradation of the ERAD-LM, CsA-insensitive substrate BACE457 [Figs. 2B–2C] and of the ERAD-Ls, CsA-sensitive substrate BACE457Δ [Figs. 2D–2E] in cells with normal level of CyPB (lanes 1–3), with reduced level of CyPB (lanes 4–6) or with reduced level of CyPA, a cytosolic target of CsA (lanes 7–9). Down-regulation of the target proteins upon specific RNA interference is shown in Fig. 2A. The data shown in Figs. 2B–2C confirmed that cell exposure to CsA does not significantly affect the disposal of CyPB. However, the CyPB levels were not restored to normal by either the transfection of CyPB (Fig. 2D), suggesting possible inhibitory role of CyPB on the disposal of BACE457Δ. 

Results and Discussion

CyPB and ERAD

CyPB is the luminal CsA target involved in ERAD

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delay disposal of the membrane-tethered BACE457 from the ER lumen (compare lane 2 with 3). Down-regulation of CyPB (Figs. 2B–2C, lanes 4–6) or of CyPA (lanes 7–9) had no significant consequences on BACE457 disposal. Thus, CyPB is dispensable for disposal of this ERAD-LM substrate.

As shown in Figs. 1C–1D, CsA substantially inhibited disposal of BACE457Δ (Figs. 2D–2E, lane 3 vs lane 2). Consistent with the identification of CyPB as the intracellular target of CsA modulating disposal of this ERAD-LM substrate, the down-regulation of CyPB substantially delayed BACE457Δ disposal (Figs. 2D–2E, lane 5 vs lane 2). Exposure of cells with low intraluminal content of CyPB to CsA had a minor, additional inhibitory effect on BACE457Δ disposal (lane 6 vs lane 5) possibly due to the inhibition of the residual CyPB remaining in the cells subjected to specific RNAi (Fig. 2A, lane 2). In contrast, the down-regulation of CyPA did not delay BACE457Δ disposal (lane 8 vs lane 2) and CsA maintained the inhibitory effect on BACE457Δ disposal in cells with low levels of CyPA (lane 9). The disposal of CD3δΔ that was insensitive to CsA (Figs. 1E–1F) was also not inhibited upon variations in the intracellular levels of CyPB and of CyPA (Figs. 2F–2G).

These data are consistent with a selective involvement of the luminal immunophilin CyPB in clearance of ERAD-LS substrates characterized by the presence of cis proline residues.

The enzymatic activity of CyPB is required to regulate ERAD

The data shown so far are consistent with a model in which CyPB facilitates BACE457Δ disposal by assisting the enzymatic...
conversion of peptidyl-prolyl bonds of the misfolded substrate from the *cis* into the *trans* configuration. This could eliminate turns in the polypeptide chain thus facilitating protein dislocation across the ER membrane, which is required for ERAD and occurs through an elusive proteinaceous channel [16]. Alternatively, peptidyl-prolyl isomerization could facilitate another rate-determining step in the disposal pathway of ERAD-LS substrates, for example their dissociation from a luminal retention factor. To assess whether the catalytic activity of CyPB is required for BACE457 disposal, an active and a catalytically inactive CyPB carrying a R62A mutation...
that substantially reduces the prolyl isomerization activity in vitro [30] were back transfected in cells with a reduced content of endogenous CyPB (Fig. 3A). Both recombinant CyPB and CyPB_R62A carried three silent mutations in their coding sequence to render their transcripts resistant to the small interfering RNA used to down-regulate endogenous CyPB. Ectopic expression of CyPB in cells with reduced level of the endogenous protein re-established efficient disposal of BACE457Δ (Figs. 3B–3C, lane 8 vs lane 5). In these cells, like in cells with normal content of endogenous CyPB (lanes 1–3), exposure to CsA substantially delayed BACE457Δ disposal (lane 9 vs lane 8). In contrast, ectopic expression of the enzymatically inactive CyPB R62A was not sufficient to recover BACE457Δ disposal in cells depleted of the endogenous enzyme (lane 11 vs lanes 2 and 8). This indicates that the enzymatic activity is required for CyPB-assisted acceleration of BACE457Δ disposal and implies that enzymatic conversion of one or more of the cis peptidyl-prolyl bonds of BACE457Δ facilitates disposal of the terminally misfolded polypeptide. These results are also consistent with the finding that CyPB is dispensable for efficient disposal of CD3δΔ, an ERAD-L substrate lacking proline residues in the cis configuration (Figs. 1E–1F and Figs. 2F–2G).

Cis proline replacement abrogates CsA-sensitivity and CyPB-dependency of ERAD
BACE457Δ has 24 proline residues. It is impossible to establish which peptidyl-prolyl bond needs to be interconverted from the cis into the trans configuration during the short retention in the ER lumen that precedes dislocation into the cytosol of this folding-defective polypeptide. However, we determined whether replacement of Pro84, 146 and 390 (which are in cis in the stable BACE501 splice variant) with alanine residues relieved the CyPB-dependency for efficient disposal.

Consistent with a CsA-insensitive ERAD pathway for ERAD-Lₘ proteins lacking cis proline residues (Figs. 1E–1F), disposal of BACE457Δ_P84,146,390A was not inhibited by cell incubation with CsA (Figs. 4A–4B). Similarly, while reduction in the intralumenal level of endogenous CyPB substantially delayed disposal of the wt BACE457Δ (Figs. 2–3), degradation of BACE457Δ_P84,146,390A remained unperturbed upon depletion of the ER-resident immunophilin (Figs. 4C–4D, lanes 4–6 vs 1–3). Taken together, these data show that the enzymatic activity of CyPB is only required for disposal of non-membrane tethered BACE457Δ containing cis peptidyl-prolyl bonds.

All in all, CsA was identified as the first selective inhibitor of disposal of a soluble (ERAD-Lₘ) but not of a membrane-tethered (ERAD-Lₘ) version of a misfolded polypeptide with luminal structural defects. This confirms that tethering at the ER membrane changes the requirements for efficient polypeptide clearance from the mammalian ER lumen [15]. The CsA-sensitive step of ERAD occurs after substrate de-mannosylation and before intervention of cytosolic proteasomes (both progressing unperturbed in cells exposed to CsA [legend of Fig. 1C and [31]). We identify CyPB as the ER-resident target of CsA involved in disposal from the mammalian ER of some (e.g., BACE457Δ, BACE476Δ, NHK) but not all (e.g., CD3δΔ) ERAD-Lₘ substrates. We provide evidence that the intervention of CyPB in ERAD requires a functional active site. As such, our data are the first demonstration of enzymatic intervention of a member of the PPI superfamily in protein quality assurance.
control in the ER of living cells. We hypothesize that the presence of peptidyl-prolyl bonds in the cis configuration is a characteristic of those ERAD-LS substrates that show CsA-sensitive, CyPB-dependent disposal. For these misfolded polypeptides, consequences of CsA exposure or of reduction in the intralumenal level of CyPB are comparable to consequences of inactivation of components of the dislocon complex built around the membrane-embedded E3 ubiquitin ligase HRD1 that are stringently required for disposal of ERAD-LS proteins [15]. Our hypothesis that CyPB participates in the HRD1/ERAD-LS pathway is consistent with a recent report showing that CyPB forms a functional complex with GRP94, another component of the HRD1 pathway [14], to protect cells against ER stress [32]. Finally, our data imply that unfolding of non-native polypeptides upon cis to trans isomerization of peptidyl-prolyl bonds might facilitate dislocation across the ER membrane [9] similarly to what has been proposed for polypeptide unfolding upon PDI-catalyzed reduction of intra- and inter-molecular disulfide bonds [5]. Alternatively, it could promote dissociation of misfolded polypeptides from ER retention factors thus facilitating dislocation across the ER membrane.

Materials and Methods

Expression plasmids, antibodies and inhibitors

Plasmids and antibodies for NHK, CD3δΔ and BACE variants are described in [15,23]. The plasmid for CyPB expression is described in [33]. Primers for silent mutations that protect ectopic CyPB from siRNA (CyPB, 5′-AAAGA CTGTTCACACGCGCAGC-TAGACAATTTTGTGGCCTTAGCT-3′). Primers for generation of inactive CyPBΔG62A (5′-GGCTACAAAAACAGCAAATTC- CATGCTGTAAT CAAGGACTTCATG-3′). Primers for generation of BACE457DΔP84,146,390A, which lacks cis prolines (5′- CCCTGGGCGG GCCCCGCAAGCAG-3′, 5′-GCCACGCACCGC-CTGGCTGACGCAC TCCC-3′, 5′-CAGCGGTGGAAGGCGCTTTTGTCACCTTG-3′). Mutants were generated using the site-directed mutagenesis kit (Stratagene). DNA preparations were obtained using commercially available purification kits (Sigma). The nucleotide sequences of all plasmids were verified on both strands. Antibodies against CyPB, CyPA and Tubulin were from ABR, Biomol and ABM. The proteasome inhibitor PS341 was a kind gift of Millenium Pharmaceuticals Inc and was used at a concentration of 9 μM. Kifunensine (Toronto Research Chemicals Inc), thapsigargin (Sigma) and CsA (Bedford Labs) were used at a concentration of 100 μM, 300 nM and 20 μM, respectively. All inhibitors were only included in the chase media.

Cell Lines, transient transfections, RNA interferences, metabolic labelling, immunoprecipitations, immunoblots and analysis of data

HeLa cells (from ATCC) were grown in MEM Alpha supplemented with 10% FBS. Cells at 80–90% confluence in a
6 cm tissue culture plate were transected with the expression plasmid of interest (4 µg for single transfections, 6 µg total DNA for double transfections) using Lipofectamine2000 (Invitrogen) according to the manufacturer instructions. Experiments were normally performed 17 hours after transfection. For siRNA-based interference, HeLa cells at 50% confluence in a 3.5 tissue culture dish were transfected with siRNA duplex (Ambion Inc, 50 pmol/dish) using Lipofectamine2000 according to the manufacturer instructions. Four hours after transfection, the medium was replaced with MEM Alpha supplemented with 1% of non-essential amino acids (GIBCO). Thirty hours after siRNA transfection, cells were transfected with the expression plasmids of interest. Experiments were performed 48 hours post-siRNA transfection. siRNA targeting sequences: CyPB: CAAAAACA-Ci

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