The N-terminal domain plays a crucial role in the structure of a full-length human mitochondrial Lon protease

Sami Kereîche\(^1\,\ast\), Lubomír Kováčik\(^1\,\ast\), Jan Bednár\(^1\,\ast\), Vladimír Pevala\(^1\), Nina Kunová\(^3\), Gabriela Ondrovičová\(^3\), Jacob Bauer\(^3\), Luboš Ambro\(^3\), Jana Bellová\(^3\), Eva Kutejová\(^3\,\ast\)\(^4\,\ast\) & Ivan Raška\(^1\)

Lon is an essential, multitasking AAA\(^+\) protease regulating many cellular processes in species across all kingdoms of life. Altered expression levels of the human mitochondrial Lon protease (hLon) are linked to serious diseases including myopathies, paraplegia, and cancer. Here, we present the first 3D structure of full-length hLon using cryo-electron microscopy. hLon has a unique three-dimensional structure, in which the proteolytic and ATP-binding domains (AP-domain) form a hexameric chamber, while the N-terminal domain is arranged as a trimer of dimers. These two domains are linked by a narrow trimeric channel composed likely of coiled-coil helices. In the presence of AMP-PNP, the AP-domain has a closed-ring conformation and its N-terminal entry gate appears closed, but in ADP binding, it switches to a lock-washer conformation and its N-terminal gate opens, which is accompanied by a rearrangement of the N-terminal domain. We have also found that both the enzymatic activities and the 3D structure of a hLon mutant lacking the first 156 amino acids are severely disturbed, showing that hLon’s N-terminal domains are crucial for the overall structure of the hLon, maintaining a conformation allowing its proper functioning.

Human Lon (hLon, P36776) is a mitochondrial AAA\(^+\) protein (ATPases Associated with diverse cellular Activities) belonging to the LonA protease subfamily\(^1\), which plays a crucial role in the maintenance of mitochondrial homeostasis. Its primary function is the degradation of misfolded, oxidatively modified and regulatory proteins\(^2\), but it also participates in the maintenance of mitochondrial DNA\(^3\) and possesses a chaperone activity important for the proper assembly of protein complexes\(^4\). Changes in hLon expression have been linked to severe diseases, including epilepsy, myopathy, paraplegia, and cancer\(^5\). In several cancerous tissues, overexpression of hLon promoted proliferation of cancer cells\(^6\) by remodeling their mitochondrial functions\(^7\) while its down-regulation led to apoptosis and cell death\(^8\). Silencing of hLon or pharmacologically inhibiting its activity has therefore been considered as a new target for the development of anticancer drugs\(^9\).

Like other ATPases, Lon’s activities are accompanied by conformational changes induced by ATP binding and hydrolysis\(^10\,\,11\). Early biochemical studies revealed that the binding of protein substrates by Lon stimulates its ATPase and peptidase activities and that this activation is likely to be allosteric\(^12\,\,13\). Menon and Goldberg\(^12\) first suggested a substrate-induced proteolytic mechanism, in which the default state of Lon is its inactive, ADP-bound form preventing accidental degradation of cellular proteins. Upon substrate binding, this form releases its ADP molecules and binds ATP, which is followed by its rapid hydrolysis and the cleavage of peptide bonds. In this mechanism, Lon can bind and hydrolyze ATP as long as the substrate binding sites are occupied. More recently, the idea that Lon’s ATPase and protease activities are under allosteric control has been supported by degron

\(^1\)Institute of Cellular Biology and Pathology, First Faculty of Medicine, Charles University in Prague, Albertov 4, 128 01 Prague 2, Czech Republic. \(^2\)Université de Grenoble Alpes,CNRS UMR 5309, 38042 Grenoble Cedex 9, France. \(^3\)Department of Biochemistry and Structural Biology, Institute of Molecular Biology, Slovak Academy of Sciences, Bratislava, Slovakia. \(^4\)Institute of Microbiology, Academy of Sciences of the Czech Republic, Prague, Czech Republic. \(^5\)Biomedicine Center of the Academy of Sciences and Charles University in Vestec, Czech Republic. \(^\ast\)These authors contributed equally to this work. Correspondence and requests for materials should be addressed to L.K. (email: Lubomir.Kovacik@lf1.cuni.cz) or E.K. (email: Eva.Kutejova@savba.sk) or I.R. (email: iraska@lf1.cuni.cz)
binding studies\textsuperscript{14,15}, and Su et al.\textsuperscript{16} have very recently found that binding of Mg\textsuperscript{2+} ions to Lon protease from \textit{M. taiwanensis} (MtaLon) induces conformational changes in Lon's AP-domain that accompany ATP-independent partial proteolysis of unfolded proteins and cleavage of specific peptides. Therefore, the Lon protease is a dynamic protein whose functional domains (proteolytic, ATPase-, and N-terminal domains) are present on a single polypeptide chain\textsuperscript{1}. This arrangement distinguishes it from most other AAA\textsuperscript{+} proteases; the exceptions are the bacterial FtsH and m-AAA proteases\textsuperscript{17–19}. To date, a full-length three-dimensional Lon structure has not been determined, and therefore its complex mechanisms of action are only partially understood. Although several LonA X-ray crystal structures have been reported to date, some of them representing a substantial portion of the molecule\textsuperscript{16,20–25}, none of them have shown the complete oligomeric protein complex. The most complete structural study reported the determination of two sub-structures of \textit{B. subtilis} Lon (BsLon), which covered part of its N-terminal domain (PDB ID: 3M65) and both its ATPase and proteolytic domains (PDB ID: 3M6A)\textsuperscript{20} (Fig. 1a). The region linking these two domains was not determined, however, the amino acid sequence analysis suggested that it was likely to be formed by coiled-coils. This linker region seems to be crucial: an \textit{E. coli} Lon mutated in or lacking this region exhibits severely decreased ATPase activity and disruption to both its substrate translocation and degradation abilities\textsuperscript{26,27}. Recently, four X-ray crystal structures of \textit{M. taiwanensis} LonA have been resolved, including the structure of the AP-domain bound to three ADP molecules (PDB ID: 4YPL). Interestingly, the BsLon AP-domain crystallized as an open, helically arranged hexamer when bound to six molecules of ADP, but the MtaLon's AP-domain adopted a planar conformation. Unfortunately, electron microscopy (EM) studies of Lon and its bacterial and yeast counterparts FtsH and m-AAA proteases have likewise failed to provide a clear picture of the overall organization of the protease\textsuperscript{16–18,28–32}.

In this work, we present the first full-length structures of a LonA protease and show the importance of the N-terminal domain for its integrity. Using cryo-electron microscopy, we studied the structure of a proteolytically inactive \textit{hLon} S855A mutant, which retains near wild-type levels of ATPase activity. We determined the structure of this mutant after its incubation with the non-hydrolyzable ATP-analogue AMP-PNP and with ADP at resolutions of 15 Å and 21 Å, respectively. In order to study the role of \textit{hLon}'s N-terminal domain, we also determined the enzymatic properties of a truncated \textit{hLon} protein lacking its first 270 amino acids (\textit{hLon}\text{Δ}270) and acquired insight into its structure by a cryo-EM analysis. The deletion includes the 114 amino acids of the mitochondrial targeting sequence and 156 amino acids from the N-terminal domain (Fig. 1a). Since both the enzymatic activities of \textit{hLon}\text{Δ}270 were severely disturbed and its structure showed high variability, the missing 156 N-terminal residues are essential for the stability and proper functioning of the \textit{hLon} hexamer.
Results and Discussion

Structure of the proteolytically inactive \( h\text{Lon} \) incubated with AMP-PNP. The three-dimensional cryo-EM analysis of the S855A Lon at 15 Å revealed that \( h\text{Lon} \), in the presence of AMP-PNP, forms an asymmetric hexamer ~230 Å long by 143 Å across. The structure can be divided into three regions, the “head”, the “neck”, and the “legs” (Fig. 2, Supplementary Figs S1 and S2a). Individual cross-sections perpendicular to the \( z \)-axis show signs of six-fold symmetry in the head and three-fold symmetry in the neck and legs (Fig. 2a,b), while interconnection of their centers of mass gives rise to a curve and the reconstructed structure appears slightly bent (Fig. 2c).

The barrel-like “head” has pseudo six-fold symmetry, with individual subunits arranged in a planar conformation. It could be fitted without steric clashes with six subunits from the crystal structure of \( B. \text{subtilis} \) Lon (PDB ID 3M6A, chain A) containing the proteolytic and the ATP-binding domains (AP-domain). The “head” therefore contains the catalytic chamber of the protein complex, which has a 20 Å diameter opening at its C-terminal side, but no opening at its N-terminal side (Fig. 2b, cross-section 3). The fitted 3M6A subunits revealed that this closure corresponds to \( h\text{Lon} \)’s large ATPase domain containing the aromatic-hydrophobic motif (Ar-Φ) in the axial pore loops (RTYVG), which are characteristic for ATP unfoldases and form an entry gate to the catalytic chamber in structures of compartmental proteases and \( T. \text{onnurineus} \) LonB, and in the recently acquired structure of MtaLon’s AP-domain. Interestingly, the MtaLon 4YPL crystal structure adopted a planar conformation as well, even if it contains three ADP-bound and three nucleotide-free monomers, whereas our \( h\text{Lon} \) S855A was incubated with an excess of ATP-analogue AMP-PNP.

The “head” narrows into a tight trimeric neck, 93 Å across and ~10 Å long, where the six Lon monomers join into three pairs (Fig. 2a,b, cross-section 4), forming a narrow, ~15 Å channel (Fig. 2b, cross-section 3, Fig. 3a). The COILS program predicted that the residues expected to occupy this region, 396–423, are likely to possess a coiled-coil arrangement (Fig. 1b). Three pairs of legs emerge from this neck in a trimer-of-dimers arrangement,
proteasome33,34,40,41, and suggested also in the recent study ofstrate translocation into the catalytic chamber, as observed in the ClpXP proteolytic machine and in the 26S americ cryo-EM structure of the 26S proteasome ATPase domain39, and re-projections of the reconstructed resolution map of the reconstructed structure shows that the N-terminal domains are its most flexible part individual N-terminal domains to their corresponding AP-domains (Supplementary Fig. S1), indeed, the local predicted coiled-coil pattern of the domain linker region (Fig. 1a), is predicted to be predominantly to around residue 270 (valine), contains many in Figs 2b and 3), which suggests that ATP hydrolysis by the gate to the catalytic chamber formed by the Ar-cent to the ring opening are axially displaced, and the opening angle between them increased from ~58° to ~81° Supplementary Figs S2b and S3). In contrast to the AMP-PNP incubated structure, the curvature of the whole formation, in which six S855A at 21 Å in excess of ADP showed that the barrel-shaped “head” adopted an open-ring hexameric con -Figure 3. Central YZ-cross-section through the reconstructed catalytic chamber of hLon S855A in binding with AMP-PNP and ADP. Orientation of the particle is identical to Fig. 2. The N-terminal tail of the catalytic chamber seems to be closed in the AMP-PNP incubated structure but it is open in the ADP-bound hLon S855A structure (arrows). Two of the six fitted B. subtilis 3M6A monomers intersected by the plane of the cross-section, represented by blue ribbons, indicate that the density of the closure corresponds to the large ATPase domains (positions of the bridges connecting the termination points of the axial pore loops that aren’t included in the B. subtilis 3M6A crystal structure are indicated by red arrowheads). Scale bar: 5nm. Structure and enzymatic activities of the hLonΔ270 mutant. In order to study the role of Lon’s N-terminal domain in its activities, we constructed a hLon mutant lacking the first 270 residues, i.e. missing the 114 amino acids of the mitochondrial targeting pre-sequence and the first 156 amino acids of the mature protein. Secondary structure prediction algorithms of hLon’s N-terminal domain indicated that it could be divided into two subdomains separated by an unstructured region49. The first subdomain, from the beginning of the protein to around residue 270 (valine), contains many β-sheets, while the region after this residue, including also the predicted coiled-coil pattern of the domain linker region (Fig. 1a), is predicted to be predominantly α-helical.

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We found that this shortened hLon still forms a multimeric complex (Fig. 4a) and preserves a small degree of the ATPase and peptidase activities of the full-length protein, but that it has almost no proteolytic activity (Fig. 4).

During the cryo-EM image analysis of this protein, we encountered a large amount of structural heterogeneity.
and could not detect any hint of the N-terminal domain in the acquired 2D class averages (Fig. 5a, Supplementary Figs S4 and S5). In order to interpret the class averages, we compared them to matching projections of crystal structures consisting of five and six 3M6A subunits fitted into the ADP-bound full-length Lon protein after the conversion of the x-ray structures to electron density maps (Fig. 5). We observed features corresponding to the hexameric complex as well as markedly decreased intensities at the expected positions of the sixth subunit, which implies its large flexibility. Therefore, the structure of this shortened hLon mutant is highly variable, which indicates that the first 156 N-terminal amino acids of hLon are crucial for the stability and proper assembly of the hLon hexamer.

The ATPase and peptidase activities of hLonΔ270 are much lower than those of the wild type, but they could still be stimulated nearly 2× by β-casein binding (Fig. 4). This suggests that β-casein is still able to bind to hLonΔ270 and stabilize the formed complex, which then becomes more efficient. However, hLonΔ270 is almost completely unable to cleave β-casein (Fig. 4), even though Mg2+ ions and ATP are present and it’s ATPase and proteolytic compartments preserve a significant portion of their functionality. Since the loss of functionality of hLonΔ270 seems to be linked to its high structural variability, the cleavage of β-casein likely requires a cooperation of all six hLon subunits, which supports the recently proposed models of a coordinated substrate translocation proposed in M. taiwanensis Lon25 and the ClpX unfoldase45.

**Conclusions**

In this work, we presented the first two structures of a Lon protease in full length, as acquired by cryo-electron microscopy, which show that ATP hydrolysis by human mitochondrial Lon protease induces conformational changes to the whole hexameric complex and stress the important role of its N-terminal domains. In particular, the N-terminal gate to its catalytic chamber appears closed by the axial pore loops when hLon is incubated with AMP-PNP, but opens up at ADP binding. In addition, the hLonΔ270 mutant lacking the first 156 amino acids could not cleave β-casein and its 2D class averages indicated large flexibility of its sixth subunit. Therefore, proper assembly and functioning of the hLon complex are guaranteed only if the first 156 amino acids of hLon’s N-termini domains are present.

**Methods**

**Expression and purification of recombinant proteins.** Expression, purification, and in vitro mutagenesis of hLon were performed as described in Ambro et al.46. The hLonΔ270 mutant (Δ1–270) was prepared by in-fusion cloning into a pOPINJ vector47 as an N-terminal 6× His-GST tagged protein; the following primers were used:

**Lon_271_FW:** AGTTTCTGTTCAGGTACCATGTTGGTAGAGAACGTTGTC

**Lon_960_RV:** CTGGTCTAGAAAGCTTTCACCGTTCCACGACCGCAG

The construct was verified by DNA sequencing (Macrogen). Protein expression and purification was performed following the protocol for wt hLon46. The 6× His-GST tag was removed by overnight incubation with PreScission protease at 6°C during protein purification according to the GE Healthcare protocol.
Gel Filtration Analysis. Analytical gel filtration was performed with a Superose 6 10/300 GL (GE Healthcare) column using buffer A (20 mM HEPES, pH 8.0, 150 mM NaCl, 20 mM MgCl₂, 10% (v/v) glycerol). The flow rate was 0.4 ml/min. The peak fractions were analyzed by SDS-PAGE, concentrated on Microsept Advance 100 K columns (Pall, USA) and stored at −20 °C. The concentration of protein was determined using the BCA kit (Thermo Scientific, USA).

Crosslinking of hLon. Crosslinking of hLon proteins was performed as described in Ambro et al.46. Briefly, 5 μg of protein was crosslinked with 0.1% (v/v) glutaraldehyde for 30 min on ice in 50 mM HEPES, pH 8.0, 10 mM MgCl₂, 2 mM ATP and then visualized on a 5.5% SDS-PAGE gel stained with Coomassie Brilliant Blue.

ATPase, peptidase and protease activities of hLon. ATPase, peptidase and protease assays were performed as described in Ambro et al.46. In short, to measure the ATPase activity, 5 μg of wt hLon, its S855A and Δ270 mutants was incubated at 37 °C in 50 mM Tris-HCl pH 8.0, 40 mM MgCl₂, 0.5 mM ATP and measured once a minute from 0 to 11 min. Substrate stimulation of the ATPase activity was determined by assaying the ATPase reaction in the presence of 25 μg 3-casein. To estimate the peptidase activity, 5 μg of wt hLon and its Δ270 mutant was incubated at 37 °C in 50 mM Tris-HCl pH 8.0, 40 mM MgCl₂, 0.5 mM ATP with 1 mM of the fluorogenic peptide glutaryl-Ala-Ala-Phe-MNA and in the absence or presence of 25 μg 3-casein. Measurements were taken every 40 s for 20 min. To measure the protease activity, 15 μg of FITC-casein was cleaved by 5 μg of wild type hLon and its S855A and Δ270 mutants in 50 mM Tris-HCl pH 8.0, 40 mM MgCl₂, 0.5 mM ATP at 37 °C. Measurements were taken every 30 s for 20 min. To measure the proteolytic activity of hLon and the Δ270 mutant, 1 μg of 3-casein was cleaved by 4 μg hLon in 50 mM Tris-HCl pH 8.0, 10 mM MgCl₂, 2 mM ATP for 0, 15, and 30 minutes at 37 °C. The reaction mixtures were separated on a 12% SDS-PAGE gel.

Specimen preparation for cryo-electron microscopy. All three specimens (hLon S855A incubated with 1 mM ADP or AMP-PNP, and hLonΔ270), concentrated to 1 mg/ml, were diluted 5 × in buffer B (20 mM HEPES, pH 8.0, 150 mM NaCl, 5 mM MgCl₂) and 3 μl of the dilution were applied to freshly glow-discharged Quantifoil 2/2 grids (EMS, Hatfield, PA, USA). The grids were vitrified by plunge-freezing into liquid ethane in a Vitrobot machine (FEI, Hillsboro, OR, USA) after 2 s blotting (blotting force 2) at 4 °C and 100% humidity.

Data acquisition. Datasets of both hLon S855A samples were acquired with a FEI Polara TEM (FEI, Hillsboro, OR, USA) at an operating voltage of 100 kV with a FEI Falcon I 4 k × 4 k direct detection camera at IGBMC Strasbourg, France. Since the overall organization of the Lon protease has been intriguing so far, we made use of the markedly better performance of the Falcon I DED at voltages lower than 200 kV in the low-resolution range of 0−0.25 × Nyquist frequency46. Images were automatically recorded by the EPU software at a nominal magnification of 59000 ×, yielding a final image pixel size of 1.8 Å. At this magnification, the 0.25 × Nyquist frequency corresponds to the resolution of ~14.4 Å. Image underfocus was set in EPU to vary between −1.2 μm and −2.6 μm, the total electron dose used to acquire a single image was ~10 electrons/Å². In total, 1074 images of ADP-incubated hLon S855A and 575 images of AMP-PNP-incubated hLon S855A were recorded.

The hLonΔ270 dataset was recorded with a C₃-corrected FEI Titan Krios TEM at an accelerating voltage of 300 kV at IGBMC Strasbourg, France, on a Falcon II 4 k × 4 k direct-detection camera operated in movie mode. The nominal acquisition magnification was 47000 ×, resulting in a final image pixel size of 1.42 Å. Seven of the recorded frames (frames 2–8) covering an accumulated electron dose of ~20 electrons/Å² were saved for further processing. The EPU-driven image underfocus varied between −1.3 μm and −2.4 μm; in total 2980 stacks of 7 frames were recorded.

Image processing. After CTF and ice quality inspection, 498 images of the ADP-incubated hLon S855A, 458 images of the AMP-PNP incubated hLon S855A, and 2739 images of the hLonΔΔ270 specimens were selected for further processing. The hLonΔ270 frames were aligned and summed with the GPU-based dosegpu_driftcorr49 program prior to further analysis. Image CTFs were estimated by CTFFIND30, the resolution of our micrographs by the GCTF’s EPA procedure51, which revealed that reliable signal can be extracted from these micrographs up to a resolution, on average, of 6.74 Å for the S855A-AMP-PNP sample, 8.1 Å for the S855A-ADP sample, and 4.85 Å for the frame-aligned hLonΔΔ270 specimen. The value of the C₃ coefficient of spherical aberration was set to 0.1 in the processing of the hLonΔ270 dataset. Particles were picked semi-automatically using the e2boxer program from EMAN2. Altogether, 38100 hLon S855A ADP-incubated particles, 97500 hLon S855A AMP-PNP-incubated particles, and 39800 hLonΔ270 particles were extracted. The box size was set to 224 pixels for the hLon S855A samples, and to 192 pixels for the hLonΔΔ270 specimen.

The collected particle sets were fully processed in Relion 1.353. After three runs of 2D classification with 300 classes, 73000 hLon S855A AMP-PNP-incubated particles, 32500 hLon S855A ADP-incubated particles, and 12450 hLonΔ270 particles were selected for subsequent 3D classification. Inspection of the resulting 2D class averages of full-length specimens revealed top, side, and tilted views (Supplementary Figs S1–S3). Top views were identified according to oligomeric features51, while side and tilted views according to hints of the N-terminal domains.

In order to create an initial model for the 3D classification of both full-length hLon S855A datasets, the identified side-view class average of the hLon S855A AMP-PNP-incubated specimen was low-pass filtered to 120 Å and extended to a rotationally symmetric 3D model by rotation of its 2D Fourier transform about its long axis, followed by inverse 3D Fourier transformation. The rotation of the 2D Fourier spectrum was performed by bilinear interpolation between neighboring points of Fourier profiles at each Z-level of the Fourier-transformed side view. All these computations were performed in Matlab (Mathworks, Inc., Natick, MA, USA).

In Relion, these initial 3D models were again low-pass filtered to 120 Å and subjected to grey-scale invariant cross-correlation in the first iteration of the 3D classifications. All 3D classification runs were performed with...
the initial starting model, and the number of classes in each 3D classification run was selected so that at least one “empty”, i.e. very sparsely populated class, resulted.

The first 3D classification of the hLon S855A AMP-PNP-incubated dataset was performed with 5 classes; 30000 particles from the most populated class were then passed through another classification run with 3 classes. From this 2nd run, 23170 particles were selected for final 3D refinement. 3D classification of the hLon S855A ADP-incubated dataset was performed in 3 runs with 6, 3, and 6 classes, respectively. After the 3rd run, 10809 particles were available for 3D refinement. In each 3D classification run on the full-length Lon datasets, only those particles contributing to classes with well-formed N-terminal domains were selected for further 3D processing.

The 3D maps of the major classes acquired from the final 3D classification run were low-pass filtered to 40 Å and used as initial models for 3D refinement, giving an 18.3 Å resolution hLon S855A–AMP-PNP structure, and 22.4 Å hLon S855A–ADP structure. The reported resolutions were estimated using Relion’s gold-standard Fourier Shell Correlation at the level of 0.143. Both full-length hLon S855A structures were then masked, resulting in a resolution of 15 Å for the AMP-PNP-incubated structure and 21 Å for the ADP-incubated structure. Local resolution maps were computed using ResMap. The handedness of all reconstructions was adjusted to match the orientation of the B. subtilis Lon hexamer (PDB ID: 3M6A).

In order to check the influence of the starting model on the resulting quaternary structure of the full-length protein, we processed the 23170 hLon S855A AMP-PNP-incubated particles against a synthetic soft-edged cylinder with a length of 220 Å and a diameter of 140 Å. The resulting structure was identical to the one showed in Fig. 2a, albeit its resolution was slightly lower (15.5 Å).

Fitting of X-ray structures. Rigid-body fitting of the AP-domains of the reconstructed structures was performed by the computation of cross-correlation coefficients between the reconstructed cryo-EM maps and chain A of the B. subtilis Lon AP-domain (PDB ID: 3M6A) in the colores package of the Situs program suite, followed by interactive docking in Sculptor. Six AP-domain monomers could be fitted into both full-length cryo-EM S855A Lon maps without clashes (Fig. 2). The N-terminal domain of both full-length S855A Lon structures was fitted in UCSF Chimera with six copies of a fragment of the E. coli Lon N-terminal domain crystal structure (PDB ID: 3LJC) containing residues 1–219.

Measurements in the fitted PDB structures were performed in UCSF Chimera. In order to compare the two nucleotide-state structures of hLon S855A, both reconstructed cryo-EM maps were loaded into Chimera, aligned, and fitted with the six corresponding B. subtilis AP-domain monomers from the Sculptor docking. The positions of the ADP molecules in the hLon S855A AMP-PNP Lon reconstruction could be approximated by a virtual plane, which served as a reference also for the measurement of both vertical and angular distances between the ADP molecules in the hLon S855A ADP Lon reconstruction. Distances in the cryo-EM maps were measured using ImageJ. Conversion of the x-ray structures to electron density maps was performed by the e2pdb2mrc.py programme of EMAN2.

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Author Contributions
S.K. and L.K. acquired, processed, and interpreted cryo EM-datasets, and wrote the paper. J. Bednár acquired and interpreted data, V.P., N.K., G.O., L.A., J. Bauer, J. Bellová and E.K. purified hLon proteins and analysed their activities. E.K., J. Bednár, J. Bauer, V.P. and I.R. contributed to manuscript text, E.K. and I.R. supervised the research.

Additional Information
Accession codes: The reconstructed density maps of the cryo-EM hLon S855A incubated with AMP-PNP and ADP have been deposited in the Electron Microscopy Data Bank under accession codes EMD-3275 and EMD-3274.

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