Proteasomal degradation induced by DPP9-mediated processing competes with mitochondrial protein import

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Abstract

Plasticity of the proteome is critical to adapt to varying conditions. Control of mitochondrial protein import contributes to this plasticity. Here, we identified a pathway that regulates mitochondrial protein import by regulated N-terminal processing. We demonstrate that dipeptidyl peptidases 8/9 (DPP8/9) mediate the N-terminal processing of adenylate kinase 2 (AK2) en route to mitochondria. We show that AK2 is a substrate of the mitochondrial disulfide relay, thus lacking an N-terminal mitochondrial targeting sequence and undergoing comparatively slow import. DPP9-mediated processing of AK2 induces its rapid proteasomal degradation and prevents cytosolic accumulation of enzymatically active AK2. Besides AK2, we identify more than 100 mitochondrial proteins with putative DPP8/9 recognition sites and demonstrate that DPP8/9 influence the cellular levels of a number of these proteins. Collectively, we provide in this study a conceptual framework on how regulated cytosolic processing controls levels of mitochondrial proteins as well as their dual localization to mitochondria and other compartments.

Keywords adenylate kinase 2; dipeptidyl peptidase 9; MIA40; mitochondrial protein import; quality control

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Introduction

Mitochondria fulfill crucial functions in metabolism and signaling. They contain more than 1,500 proteins of which only a small number is synthesized in the organelle (Rhee et al, 2013; Hung et al, 2014; Calvo et al, 2016; Morgenstern et al, 2017). The remainder is translated on cytosolic ribosomes and relies on different mitochondrial import pathways to reach the mitochondrial subcompartments, namely the mitochondrial outer (OMM) and inner (IMM) membranes, the intermembrane space (IMS) and the mitochondrial matrix (Topf et al, 2016; MacPherson & Tokatlidis, 2017; Jackson et al, 2018; Habich et al, 2019b; Hansen & Herrmann, 2019; Planner et al, 2019).

Mitochondrial biogenesis and other cellular processes are tightly coordinated to adjust cells to different metabolic needs (Harbauer et al, 2014; Chandel, 2015; Finkel, 2015; Wai & Langer, 2016). Coordination thereby takes place on different levels including regulation of transcription and translation of mitochondrial proteins. Posttranslational regulation often directly targets mitochondrial precursor proteins in the cytosol and modulates their mitochondrial import. This includes the binding of metabolites or partner proteins to these precursors (Vongsamphanh et al, 2001; Dailey et al, 2005; Frechin et al, 2009) and the attachment of covalent modifications or the proteolytic processing of precursor proteins (Boopathi et al, 2008; DeRasmo et al, 2008; Avadhani et al, 2011; Bragoszewski et al, 2013). Moreover, competition between cytosolic folding and mitochondrial import has been reported as regulatory principle for a number of dually localized proteins (Strobel et al, 2002; Kloppel et al, 2011; Yogev & Pines, 2011; Suzuki et al, 2013). Folding in the cytosol prevents mitochondrial import, and thus, the majority of the
protein ends up localizing in the cytosol and not in mitochondria. Posttranslational modification of the translocase of the OMM (TOM) that constitutes the entry gate for the vast majority of mitochondria precursors can as well influence import of specific proteins (Schmidt et al., 2011; Gerbeth et al., 2013).

Precursor proteins are handed over to different import machineries after passage of the TOM pore. One such machinery is the mitochondrial disulfide relay that imports and oxidatively folds proteins of the IMS (Stojanovski et al., 2012; MacPherson & Tokatlidis, 2017; Habich et al., 2019b). The protein MIA40 (also coiled-coil-helix-coiled-helix domain containing protein 4, CHCHD4) is the main component of the mitochondrial disulfide relay and serves as oxidoreductase and import receptor (Petrungaro et al., 2015; Peleh et al., 2016; Habich et al., 2019a). Most substrates of the disulfide relay lack classical N-terminal mitochondrial targeting sequences (MTS) and instead contain conserved cysteine motifs (Fischer et al., 2013; Petrungaro et al., 2015; Modjahedi et al., 2016; Bragoszewski et al., 2017; Hansen & Herrmann, 2019). The majority of substrates contains four conserved cysteines in so-called twin CXXC or twin CXC motifs that become oxidized to two very stable disulfide bonds during substrate maturation by MIA40. Recently, few unconventional substrates of the disulfide relay with differing structures and cysteine patterns have been identified (Barchiesi et al., 2015; Petrungaro et al., 2015; Ramesh et al., 2016). One such putative substrate that occurred as interaction partner of MIA40 but has so far not been characterized was adenylate kinase 2 (AK2). AK2 catalyzes the reversible adenine nucleotide phosphoryl transfer in the reaction 2 ADP ↔ ATP + AMP in the IMS. Loss of AK2 leads to impaired mitochondrial function (Burkart et al., 2011; Six et al., 2015), hampers induction of the endoplasmic reticulum unfolded protein response (Burkart et al., 2011), and sensitizes cells to induction of apoptosis (Single et al., 1998; Kohler et al., 1999; Lee et al., 2007). Human patients with defects in AK2 suffer from an autosomal recessive form of severe combined immunodeficiency (SCID) named reticular dysgenesis (RD) (Lagresle-Peyrou et al., 2009; Pannicke et al., 2009). Although changes in AK2 activity can have severe consequences, the mechanism of AK2 IMS import and folding remains unclear.

In this study, we show that mitochondrial import of AK2 is mediated by the mitochondrial disulfide relay via an unusual intramolecular disulfide shuffling mechanism that makes use of the three conserved cysteines in AK2. We also demonstrate that N-terminal processing of AK2 by the dipeptidyl peptidases DPP8 and DPP9 controls its IMS import and cellular localization. Processed cytosolic AK2 is thereby targeted for rapid proteasomal degradation. Attenuation of processing results in increased amounts and activity of AK2 and induces dual localization to IMS and cytosol of previously exclusively IMS-localized AK2. Lastly, we show that processing by DPP8 and DPP9 is employed as general control mechanism of mitochondrial protein biogenesis and might affect more than 100 mitochondrial proteins to regulate mitochondrial proteome plasticity.

Results

AK2 is oxidized by MIA40 during its maturation

AK2 is a soluble IMS protein (Bruns & Regina, 1977; Kohler et al., 1999). It contains conserved cysteines (C40, C42, and C92) and lacks an N-terminal MTS (Fig 1A). Two AK2 isoforms (AK2-A and AK2-B) can be detected in HEK293 cells. These isoforms differ in the most C-terminal amino acids and thereby AK2-A contains one additional non-conserved cysteine (C232) (Fig 1A and Appendix Fig S1A). We identified AK2 as an interaction partner of MIA40 in proteomics screens (Petrungaro et al., 2015; Habich et al., 2019a). To confirm this interaction, we precipitated C-terminally Strep-tagged MIA40 variants from HEK293 cells and tested whether endogenous AK2 co-precipitated by immunoblot. We confirmed the AK2-MIA40 interaction but also observed that the interaction depended on the redox-active cysteines of MIA40 as no AK2 was co-precipitated with a MIA40 lacking these cysteines (Fig 1B). We also confirmed the interaction by performing the inverse experiment, precipitating C-terminally HA-tagged AK2 and testing for coprecipitation of endogenous MIA40 (Appendix Fig S1B), and by precipitating endogenous MIA40 and testing for coprecipitation of endogenous AK2 (Appendix Fig S1C). Lastly, we demonstrated by emetine chase and pulse-chase analyses that the AK2-MIA40 interaction was transient in nature (Appendix Fig S1D and E). This indicated that AK2 might undergo MIA40-catalyzed oxidative protein folding during its maturation. To test this, we assessed different parameters for such a maturation pathway.

First, we determined the redox state of the cysteines of mature endogenous AK2 using an inverse redox state assay (Erdogan et al., 2018) (Fig 1C and Appendix Fig S1F). In this assay, the thiol–disulfide redox state of proteins is “trapped” in intact cells by the addition of the membrane-permeable alkylating agent N-ethyl maleimide (NEM). This modifies all accessible (reduced) thiols. Thereafter, cells are lysed and previously oxidized cysteines are reduced using the reductant tris(2-carboxyethyl)phosphine (TCEP). Then, free thiols (the ones that had been previously oxidized) are modified using methyl-PEG-12-maleimide (mmPEG12). This compound introduces a molecular weight shift that can be detected on SDS–PAGE. Thus, proteins that contained oxidized cysteines in intact cells will migrate slower on SDS–PAGE compared to previously reduced proteins. When we applied this assay to HEK293 cells and analyzed the redox state of AK2, we found that both isoforms, AK2-A and AK2-B, migrated slower than the purely NEM-treated control indicating that they contained oxidized cysteines, most likely in one disulfide bond (Fig 1C, lanes 2 and 3, black arrow heads, and Appendix Fig S1F).

Next, we investigated whether the disulfide bond in AK2 formed during AK2 maturation (Fig 1D). To this end, we employed an in vivo oxidation assay (Fischer et al., 2013). In this assay, we combined a pulse-chase experiment with a redox state determination and immunoprecipitation (IP) of the protein of interest. Pulse labeling with 35S-methionine thereby allowed the labeling of a small fraction of newly synthesized proteins. In the subsequent chase period, the fate of this fraction can be followed. We did so by freezing the thiol–disulfide redox state by rapid acidification using trichloroacetic acid (TCA). Then, free thiols were modified with mmPEG12 before the protein of interest; in our case AK2, was immunoprecipitated. Subsequently, modified proteins were analyzed by SDS–PAGE and autoradiography. Reduced proteins thereby migrate slower than oxidized ones. When we employed this assay to AK2, we observed that both isoforms of endogenous AK2 migrate faster with increasing chase time. This indicates that they acquire an intramolecular disulfide bond with time (Fig 1D).
Finally, we assessed whether depletion of MIA40 by siRNA-mediated knockdown would interfere with AK2 oxidation in intact cells (Fig 1E and Appendix Fig S1G). We repeated the in vivo oxidation assay in cells depleted for MIA40. While cells treated with control siRNA exhibited normal oxidation kinetics of endogenous AK2, cells treated with siRNA directed against MIA40 failed to oxidize AK2 (Fig 1E). Taken together, we conclude that MIA40 oxidizes AK2 and thereby introduces one disulfide bond into the protein.
Disulfide bond formation between C42 and C92 is critical for mitochondrial AK2 accumulation

Many classical MIA40 substrates contain an even number of conserved cysteines in a specific defined spacing. AK2 seems to be an unconventional substrate of MIA40 as it is not only comparatively large (ca. 26 kDa) but has an odd number of conserved cysteines (C40, C42, and C92) with an unconventional spacing. Of these cysteines, only C40 and C92 are positioned in helices and thus constitute classical targets for initial MIA40 interaction (Milenkovic et al., 2009; Sideris et al., 2009; Banci et al., 2010; Koch & Schmid, 2014). We assessed the role of AK2 cysteines for oxidation and mitochondrial accumulation of the protein.

First, we tested the thiol–disulfide redox state of the different AK2 cysteine variants (Fig 2A). To this end, we generated stable inducible cell lines, which express different C-terminally HA-tagged AK2 isoform A variants (AK2WT, AK2C40S, AK2C42S, AK2C92S, AK2C40S,C92S). Compared to the redox state determination experiment with endogenous AK2, we therefore only detect the isoform AK2-A. Wild-type AK2-HA and AK2C232S-HA were present in an overwhelmingly oxidized redox state. AK2C40S-HA was mainly oxidized but exhibited a considerable reduced fraction at steady state. Conversely, AK2C42S-HA and AK2C92S-HA were present in the completely reduced state indicating that in the wild-type protein a disulfide bond is present between C42 and C92 (Fig 2A). This is also in agreement with the AK2 crystal structure (Fig 2B, pdb: 2C9Y). The electron density and the deposited coordinates of this entry indicated a partial disulfide bond between C42 and C92 with an occupancy of about 0.5. The imperfect formation of the disulfide bond in the crystal structure is probably due to the fact that during the preparation of recombinant protein for crystallization no care was taken to preserve a native disulfide pattern and possibly a partial photoreduction during X-ray exposure may have taken place. Nevertheless, the structure together with our data provides strong evidence for the presence of a disulfide bond connecting C42 and C92.

Cysteine oxidation and mitochondrial accumulation are closely coupled events for MIA40 substrates without MTS (Habich et al., 2019a). Therefore, a lack of cysteines does not only result in the absence of oxidation but also in cytosolic mislocalization (Friederich et al., 2017; Mohanraj et al., 2019). We thus next investigated mitochondrial localization of AK2 cysteine variants by immunofluorescence microscopy (Fig 2C, and Appendix Fig S2A and B). Wild-type AK2-HA and AK2C232S-HA colocalized with a mitochondrial marker. Conversely, AK2C42S-HA and AK2C92S-HA did not colocalize with mitochondria and instead appeared to be cytosolic. This indicated that these two cysteines and the formation of the disulfide bond are critical for mitochondrial accumulation. Interestingly, AK2C40S-HA exhibited an intermediate behavior. It was colocalized with mitochondria to a significantly lower extent compared to wild-type AK2. Still the majority of the protein appeared to localize to mitochondria (Fig 2C).

Collectively, these data indicate that formation of a disulfide bond between C42 and C92 is critical for mitochondrial accumulation of AK2. C40 appears to exert an influence on mitochondrial accumulation and AK2 redox state as well.

C40 drives intramolecular disulfide shuffling dependent import of AK2

Intrigued by the behavior of AK2C40S-HA, we investigated this variant in more detail. In the AK2 structure, C40 is uniquely positioned in a motif that is suitable for interaction with MIA40 (Sideris et al., 2009; Koch & Schmid, 2014) (Appendix Fig S2C). This motif consists of an amphipathic helix with hydrophobic patches on the helix face that is marked by the presence of the MIA40-interacting cysteine. Although C92 is also positioned in a helix, this helix lacks the amphipathic character required for MIA40 interaction. Thus, the two cysteines (C42 and C92) that eventually form a disulfide bond in mature AK2 are not positioned in an environment that is predicted to facilitate efficient interaction with MIA40. Does this point to a role of C40 in initiating interaction between AK2 and MIA40? To answer this question, we immunoprecipitated endogenous MIA40 from cell lines expressing either AK2WT-HA or AK2C40S-HA, and analyzed the eluates by immunoblot against the HA tag. We thereby found that absence of C40 strongly interfered with the AK2-MIA40 interaction (Appendix Fig S2D). If C40 is so important for initiating the MIA40-AK2 interaction, its absence should also affect oxidative folding (i.e., maturation) of AK2. Indeed, when we analyzed the oxidation kinetics of AK2 variants, we found that AK2C40S has a delayed oxidation compared to wild-type AK2 (Appendix Fig S2E).

Our findings point to an initial interaction between C40 and MIA40. This mixed disulfide bond would have to isomerize to the final disulfide bond between C42 and C92. We hypothesized that this might take place via an intramolecular disulfide isomerization mechanism. Although such a process has never been described for mitochondrial disulfide relay substrates, it is well known for oxidative folding of complex proteins in the endoplasmic reticulum (ER) (Jansens et al., 2002; Land et al., 2003; Chakravarthi & Bulleid, 2004; Roberts et al., 2018). The temporal resolution of our in vivo pulse-chase experiments was not sufficient to address the question of intramolecular disulfide shuffling in AK2. We thus turned to an in vitro experiment (Appendix Fig S2F). With this experiment, we tested whether in principle disulfide formation between C40 and, e.g., C42 is possible. This disulfide bond would be a critical intermediate in such an isomerization process. We purified different cysteine variants of AK2 and analyzed their disulfide pattern as well as the stability of the disulfide bonds (Appendix Fig S2F). We indeed found that all tested variants except for the AK2C232S-HA contained a disulfide bond. This also included the AK2C232S,C232S variant that can only form a disulfide bond between C40 and C42. In line with this, also the C92S variant contained a disulfide bond. In contrast to the very stable disulfide bond found in the wild type, in AK2C42S and AK2C40S, the disulfide bonds in AK2C92S and AK2C92S,C232S were very labile toward treatment with the reductant DTT. Although these data were obtained with purified proteins and not with unfolded import intermediates as they would be found during vectorial import into the IMS, our data indicate that in principle, a disulfide bond between C40 and C42 can be formed. This disulfide bond is very labile, and thus, it would be thermodynamically favorable to rearrange it to the C42–C92 disulfide. The in vitro data also again support the presence of a very stable native disulfide bond between C42 and C92.
Taken together, our data support a model in which an initial interaction between C40 and MIA40 drives import of AK2 into the IMS. Subsequent formation of a C40–C42 disulfide and later intramolecular rearrangement to the native C42–C92 disulfide result in mature AK2. We propose that this C40-dependent reaction sequence is important to accelerate folding and import of AK2 compared to a situation without C40 (and thus without isomerization, Fig 2D).

Figure 2. Cysteine residues 42 and 94 are critical for mitochondrial accumulation of AK2.

A Influence of AK2 cysteines on the AK2 redox state. Cells expressing single cysteine to serine mutants of AK2-HA (isoform A) were lysed after NEM treatment and analyzed by inverse redox shift assay using mmPEG12, SDS–PAGE, and immunoblot against the HA tag. AK2*C42S and AK2*C92S are present in their completely reduced state indicating presence of a disulfide bond between both cysteines.

B Crystal structure of AK2 (pdb entry 2C9Y). In the crystal structure, a disulfide bond between C42 and C92 is visible with about 50% occupancy, while 50% of the Cys42/92 residues are in the reduced state. This disulfide bond is far away from the active site of AK2. The figure was created using PyMOL.

C Influence of AK2 cysteines on cellular localization. Cells expressing single cysteine to serine mutants of AK2-HA (isoform A) were fixed and analyzed by immunofluorescence. The colocalization of AK2-HA variants with MitoTracker was quantified using CellProfiler (Pearson’s correlation coefficient). C42 and C92 are critical for mitochondrial localization of AK2. The C40S mutant exhibits significantly decreased colocalization with mitochondria. A Kruskal–Wallis rank sum test/Dunn’s test was performed. Numbers of analyzed cells are indicated in the plot. *** represents P-value < 0.001 compared to wild type. The black horizontal line represents the median. Data information: Scale bars: 10 μm.

D Model for AK2 import and oxidative folding. See text for details.

Source data are available online for this figure.
AK2 import competes with proteasomal degradation

Diminished disulfide relay substrate levels are often a hallmark of MIA40-depletion (Hangen et al., 2015; Habich et al., 2019a). In line with this, AK2 was almost absent from cells when MIA40 was depleted by siRNA-mediated knockdown (Fig 3A, lanes 2, 3) or dominant-negative MIA40 variants were expressed ((Habich et al., 2019a), Appendix Fig S3A, lane 3). This was due to the short half-life of AK2 under these conditions as we demonstrated by pulse-chase experiments (Fig 3B).

MIA40 substrates that fail to become imported into the IMS are eventually degraded by the proteasome (Habich et al., 2019a; Mohanraj et al, 2019). To test whether this is also the case for AK2, we assessed the stability of an import-deficient cysteine variant of AK2 (AK2-C40S,C42S,C92S, localization see Appendix Fig S3B) in dependence of the proteasomal inhibitor MG132. Indeed, this AK2 variant was very unstable and could be stabilized by MG132 (Fig 3C). In line, steady-state levels of AK2-C40S,C42S,C92S-HA but also of AK2WT-HA were increased upon MG132 incubation or siRNA-mediated depletion of proteasomal subunits PSMB5/C3 (Fig 3D and E). Our findings thus indicate that cystosolic accumulation of (wild type and mutant) AK2 is actively prevented by competition between import and quick proteasomal degradation.

Dipeptidyl peptidase 9 targets AK2 for proteasomal degradation

Protein stability in the cytosol is often determined by the N-terminal amino acids of proteins (Varshavsky, 2017). We thus analyzed the N-terminus of wild-type AK2-HA after its immunoprecipitation by proteomic approaches (Fig 4A). We thereby found that the most N-terminal peptid of the AK2 protein existed in four different versions: the native N-terminus (“MAPS-”), an N-terminus, which lacked the methionine and likely was processed by methionine aminopeptidase MAP (“APS-”), and two N-termini, which lacked the three most N-terminal amino acids (“S-”). Of the latter, one variant was N-terminally acetylated while the other was not. Notably, assessment of AK2 conservation in 163 species revealed that the three most N-terminal amino acids including the proline are highly conserved (Fig 4A, logo plot) indicating that these amino acid residues might fulfill an evolutionary conserved role.

The identification of different AK2 N-termini might constitute the regulatory mechanism to control cytosolic stability of AK2. Given the fact that the IMS does not harbor the respective degradation machinery, a protein that escapes the cytosolic N-end rule machinery by becoming imported would be stable in the IMS. Which protease facilitates N-terminal processing of AK2? Members of the dipeptidyl peptidase 4 (DPP4) protease family possess the unusual capacity to cleave a post-proline bond to release dipeptides from the N-terminus of proteins (Wilson et al., 2016; Sato & Ogita, 2017). Two members of this family are present in the cytosol, DPP8 and DPP9 (Wilson et al, 2016).

We tested whether DPP8 and/or DPP9 affected stability of cytosolic AK2 (AK2-C40S,C42S,C92S-HA, Fig 4B and C). Indeed, siRNA-mediated depletion of DPP8 and/or DPP9 revealed a stabilization of this AK2 variant at steady state (Fig 4B). The stabilization was thereby most prominent when DPP9 was depleted but also increased further upon double depletion of both DPPs. Since DPP8 alone did not affect AK2-C40S,C42S,C92S-HA stability, we reasoned that DPP9 might be the main processing enzyme with DPP8 serving as backup. We complemented the siRNA-based experiment with an experiment in which we employed 1G244, a specific inhibitor of DPP8 and DPP9 (Wu et al, 2009; Fig 4C). We thereby found a strong stabilization of AK2-C40S,C42S,C92S-HA upon DPP8/9 inhibition. Our experiments so far were performed with an HA-tagged AK2 cysteine variant. Next, we repeated the experiments with endogenous AK2 (Fig 4D and E). We thereby found that also endogenous AK2 levels increased upon DPP9 depletion or inhibition of DPP8/9 activity. Thus, DPP9 indeed affects cellular AK2 levels.

To test whether DPP8/9 directly process AK2, we introduced mutations into the N-terminus of AK2WT or AK2-C40S,C42S,C92S and analyzed the stability of the respective variants (Fig 4F and Appendix Fig S4A). The first mutation, replacement of alanine at position 2 by aspartate (A2D), hampers processing by MAP and as a consequence also the processing by DPP9. This AK2 variant would...
Figure 3.
carry a methionine at its N-terminus (Appendix Fig S4A). As second mutation, we replaced serine at position 4 by proline, which prevents cleavage by DPP9 and thus leaves alanine as the neo-N-terminus of this AK2-variant (S4P). When we analyzed the levels of these variants either in the context of wild-type AK2 or the C40S, C42S,C92S variant of AK2, we found a striking stabilization of AK2 upon modification of its N-terminus (Fig 4F). This effect was not further increased by application of 1G244 or siRNA-mediated depletion of DPP9 indicating that both, the A2D and the S4P mutation, prevented processing in the first place (Fig 4F and Appendix Fig S4B). Moreover, incubation with MG132 did not additionally further increased by application of 1G244 or siRNA-mediated depletion of C42S,C92S variant of AK2, we found a striking stabilization of AK2 N-terminus of this AK2-variant (S4P). When we analyzed the levels carried a methionine at its N-terminus (Appendix Fig S4A). As second mutant, DPP9 cleavage is allowed but instead of an N-terminal serine, AK2 will then carry valine or glycine as neo-N-terminus. Compared to serine, these two amino acids should provide AK2 with increased resistance to degradation. Indeed, both AK2 variants showed increased stability (Appendix Fig S4D). Taken together, DPP9 processes the N-terminus of AK2 in the cytosol. This targets AK2 for proteasomal degradation. Once AK2 reaches the IMS, it is protected from degradation.

DPP9 processing prevents cytosolic AK2 accumulation

Why is AK2 specifically targeted for rapid cytosolic degradation? One reason might be that upon cytosolic accumulation, AK2 could be capable of folding and acquiring enzymatic activity. We first investigated whether AK2 indeed accumulates in the cytosol when DPP9 processing does not take place. Using cell fractionation approaches, we found that a small amount of endogenous AK2 accumulated in the cytosolic fraction upon treatment with DPP9 inhibitor, while the IMS proteins CPOX and cytochrome c remained completely in the mitochondrial fraction (Fig 5A), thus excluding contamination. In agreement, by immunofluorescence analyses, we found that the DPP9-insensitive variant AK2S40T that otherwise can become normally imported into mitochondria was partially localized in the cytosol. AK2S40T fully localized to mitochondria (Fig 5B and Appendix Fig S5A). Lastly, AK2C40S, the variant that can become oxidized and imported into mitochondria albeit with slower kinetics compared to wild-type AK2, accumulated more strongly in the cytosol when proteasomal degradation was inhibited (Fig 5C and Appendix Fig S5B and C). In summary, inhibition of AK2 processing by DPP9 results in accumulation of a cytosolic AK2 species and the effects are more pronounced when using AK2 variants with delayed import kinetics.

Cytosolic AK2 is stable and enzymatically active

Mitochondrial proteins are imported in an unfolded state. “Mislocalization” of AK2 might thus result from premature folding in the cytosol. AK2 is one of the few disulfide relay substrates with inherent enzymatic activity. We thus next assessed whether cytosolic AK2 acquires enzymatic activity (Fig 6). We first established two activity assays for AK2 function, one with respect to its substrate AMP and one toward its substrate ADP (Fig 6A and B, and Appendix Fig S6A–C). Unfortunately, all AK2 variants still containing C232 tended to form non-physiological disulfide-bonded dimers involving this cysteine (Appendix Fig S6D). This was not the case for AK2S40T. To avoid the comparison of dimeric with monomeric AK2 variants, we decided to employ instead of the wild type the “wild-type-like” AK2 variant, AK2C40S,C232S that contains the C42-C92 structural disulfide bond but lacks the other cysteines. First we confirmed that AK2C40S,C232S and AK2S40T had similar activities

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Figure 4.
toward AMP and ADP as substrates (Appendix Fig S6E). We then compared AK2C40S,C232S (only containing the disulfide-forming C42 and C92) with a cysteine-less AK2 variant. We observed that purified AK2 exhibited typical substrate inhibition curves as also observed for other AKs (Liang et al., 1991; Sinev et al., 1996; Fig 6C and D). We observed maximal activity for 40 μM AMP and 1.5 mM ADP, respectively. With increasing AMP concentrations, AK2 activity dramatically decreased switching off the enzyme almost completely at about 0.5–1 mM AMP. For ADP, this decrease in activity featured a more gradual decline. Importantly, irrespective of the presence of the structural disulfide bond in AK2, enzyme kinetic profiles were the same (compare AK2C40S,C232S and AK2C40S,C42S,C92S,C232S, Fig 6C and D). Thus, the presence of the structural disulfide bond had no influence on AK2 enzymatic activity.

Next, we developed an assay that allowed the determination of AK2 activity from samples isolated from HEK293 cells (Fig 6E and Appendix Fig S6F). Cells stably expressing different AK2-HA variants were lysed using a mild detergent. Subsequently, AK2 was isolated using HA antibodies. With the AK2-HA variants still bound to the beads, the AK2 activity assay with the substrate ADP was performed. The enzymatic inactive K28Q variant of AK2 (Reinstein et al., 1990) thereby served as negative control to ensure that no endogenous AK activity was precipitated from cells. In the absence of DPP8/9 inhibitor, we found that mitochondrial AK2 variants showed similar activity irrespective whether they are DPP8/9 insensitive. The cytosolic AK2C40S,C42S,C92S showed very low activity; yet this activity was clearly higher than the one of the K28Q control. The DPP8/9-insensitive variants AK2A2D,C40S,C42S,C92S and AK2S4P,C40S,C42S,C92S exhibited activity almost as strong as wild-type AK2. Upon pretreatment of cells with DPP8/9 inhibitor, especially the activity of AK2C40S,C42S,C92S increased almost to wild-type levels. When we normalized these activities to precipitated protein levels, we found specific activities of cytosolic and IMS-AK2 to be relatively similar (Appendix Fig S6G). Taken together, AK2 can accumulate and fold in the cytosol and acquire activity. Preventing DPP9 processing and thereby proteasomal degradation increases the amount of cytosolic AK2 levels and activity.

Cytosolic AK2 is not oxidatively folded. This raises the question to which extent the native C42-C92 disulfide bond affects AK2 stability. To test this, we determined the melting temperature of AK2 variants (Appendix Fig S6H) and found it to be 58.4°C for AK2C40S,C232S and 45.5°C for a disulfide-free mutant. With this, the disulfide-free AK2 has the same stability against thermal unfolding as the cytosolic adenylate kinase, AK1 (Appendix Fig S6H). Thus, the disulfide-free (cytosolic) AK2 is stable and AK2 containing the C42-C92 disulfide bond has increased stability. In the light of recent findings on potentially increased temperatures in the IMS (Chretien et al., 2018), this implies that disulfide formation in AK2 not only stabilizes AK2 variants processed by DPP9 but can increase the stability of the native AK2 variant as well.

Collectively, our data support a model, in which DPP9 processes the N-terminus of AK2 in the cytosol to promote proteasomal degradation and to ensure that AK2 activity is absent from the cytosol under normal conditions. Importantly, in the IMS, N-terminally processed AK2 is protected from proteasomal degradation. Inhibition of DPP9 results in increased cellular AK2 levels and cytosolic AK2 activity. While we achieved inhibition by introducing mutations or by chemical inhibitors, DPP9 activity might also be lowered by transcriptional regulation or might be inhibited by posttranslational modifications, although very little is known about modes of DPP9 regulation (Fig 6F).

DPP9 targets additional mitochondrial proteins

Our data support a critical role of N-terminal processing by DPP8/9 for the regulation of AK2 levels and cellular localization. To test whether other mitochondrial proteins are targeted by DPP8/9, we screened the MitoCarta2.0, which contains proteins with a strong support for mitochondrial localization (Calvo et al., 2016) for DPP8/9 recognition and processing motifs. These motifs include MXXPY and MXY (with X not D/E, and Y not P) (Wilson et al., 2013, 2016; Zhang et al., 2015; Fig 7A, Dataset EV1). We thereby identified 107 potential mitochondrial DPP8/9 substrates. Interestingly, IMS proteins and disulfide relay substrates appeared to be enriched among the potential DPP8/9 targets.

Among the potential DPP8/9 substrates were 13 disulfide relay substrates including AK2 (Fig 7B). Their N-termini are often conserved, indicating that DPP8/9 cleavage might be an evolutionary widely applied regulatory pathway regarding substrates and species. Moreover, analysis of the mouse N-terminome (Calvo et al., 2017) revealed two disulfide relay substrates, COX17 and NDUFA8 that as mature proteins carry N-termini, as they would occur after DPP9 cleavage. Disulfide relay substrates often are not detected in proteomic approaches, and also in this proteomics approach, no other disulfide relay substrates were identified.

To complement our bioinformatics analysis, we tested effects of DPP8/9 inhibition on the disulfide relay substrates, NDUFB10, NDUF8, NDUF5S, COA6, and CHCHD2 (Fig 7C). We cultured in the presence or absence of 1G244 HEK293 cells that stably and inducibly express HA-tagged variants of these proteins. We thereby found that indeed all proteins became stabilized upon 1G244 treatment. Conversely, the disulfide relay substrate TIMM9 that does not qualify as DPP8/9 substrate did also not become stabilized upon 1G244 treatment (Fig 7C).

We had previously identified a disease variant in NDUFB10, NDUF10C1075 that was impaired in mitochondrial import (Friederich et al., 2017). The majority of the mutant protein was degraded by the proteasome, and consequently proteasomal inhibition stabilized NDUF10C1075 (Habichi et al., 2019a). Proteasomal inhibition has recently been proposed as mode to rescue the mitochondrial accumulation of a disease variant of COA7 that was likewise impaired in its mitochondrial import (Mohannaji et al., 2019). We tested whether DPP9 inhibition would allow a similar rescue for NDUF810C1075 (Fig 7D and E). To this end, we cultured HEK293 cells expressing either NDUF810WT-HA or NDUF10C1075-HA in the presence or absence of 1G244. We thereby found that indeed NDUF10C1075-HA became stabilized upon 1G244 treatment (Fig 7D). Remarkably, the fraction of the protein that accumulated
Figure 5. Cytosolic AK2 accumulates when DPP9 processing is prevented.

A Cell fractionation upon inhibition of DPP9 activity. HeLa cells were incubated for 4 days with the DPP9 inhibitor 1G244 or as control with DMSO. Cells were subjected to fractionation into “cytosol” (supernatant) and “mitochondria” (pellet) by differential digitonin lysis and centrifugation. The IMS marker proteins CPOX and cytochrome c were exclusively localized to the pellet fraction at one specific digitonin concentration (lanes 2, 5, 8, 11) irrespective of 1G244 treatment. Likewise, the cytosolic marker protein LDH localized under all conditions to the same fractions. Conversely, endogenous AK2 appeared at this digitonin concentration in the “cytosol” fraction but only upon 1G244 treatment (compare lanes 2 and 5). Thus, inhibition of DPP9 results in appearance of AK2 in the cytosol. Quantification shows loading-control adjusted AK2-signal of 1G244-treated “cytosol” fraction (lane 5) over DMSO-treated “cytosol” fraction (lane 2). White arrow head indicates the endogenous AK2 appearing in the cytosolic fraction upon 1G244 treatment. Reported values are the mean of 3 independent experiments; error bars represent ± SD. Student’s t-test was performed.

B Localization of DPP9 insensitive AK2-HA variants. Experiment was performed as described in Fig 2C. Compared to the wild type, DPP9-insensitive and import-competent AK2-HA variants colocalized with cytosolic markers indicating partial cytosolic localization. The colocalization of AK2-HA variants with MitoTracker was quantified using Fiji (Pearson’s correlation coefficient). A one-way ANOVA with post hoc Tukey test was performed. Numbers of analyzed cells are indicated in the plot. ** represents P < 0.01. Data shown in Figs 2C and 5B, and Appendix Figs S2 and S5 are derived from the same experiments. The black horizontal line represents the median. Data information: Scale bars: 10 μm.

C Localization of AK2-C40S-HA upon proteasomal inhibition. Experiment was performed as described in Fig 2C. Upon MG132 treatment, colocalization of AK2-C40S-HA with the cytosolic marker increased. A one-way ANOVA with post hoc Tukey test was performed. Numbers of analyzed cells are indicated in the plot. *** represents P < 0.001. The black horizontal line represents the median. Data information: Scale bars: 10 μm.

Source data are available online for this figure.
AK2 variant

Cytosolic processing by DPP8/9 controls accumulation of active cytosolic AK2

Figure 6.
in mitochondria upon 1G244 treatment also increased, indicating that DPP8/9 inhibition indeed provided the additional time required for NDUFB10C107S to reach at least partially the IMS (Fig 7E). Notably, also NDUFB10WT became stabilized and its mitochondrial levels increased strengthening the notion that NDUFB10 is a target of DPP9. Collectively, our data indicate that DPP8/9 processing might be a widely applied mechanism to influence mitochondrial protein import.

Discussion

Cytosolic processing by DPP8/9 controls cellular AK2 levels and dual AK2 localization

In this study, we identified a novel import regulation and quality control pathway for mitochondrial proteins. This regulation relies on cytosolic processing by DPP8/9. We propose that this DPP8/9 processing serves as a timer that prevents aberrant cytosolic accumulation of mitochondrial precursors, regulates cellular protein levels, and enables rapid control of dual localization.

Our findings suggest that the DPP8/9 quality control pathway employs the N-end-rule degradation pathways to modulate the dwelling time of precursors in the cytosol. Processing of human AK2, for example, exposes a serine residue. An N-terminal serine residue targets cytosolic proteins to the Ac/N-end rule pathway (Varshavsky, 2017; Nguyen et al., 2018). In this pathway, the N-terminal amino group becomes acetylated by N-acetyl transferases, which generates recognizable degrons. In line, we identified an N-acetylated N-terminal peptide in our proteomic approach (Fig 4A) and found AK2 variants with mutated serine 4 to be more stable (Appendix Fig S4A and D). Notably, while the “MAP” motif in AK2 is highly conserved, the amino acid residue at position 4 is not. Besides serine, e.g., asparagine and arginine residues are present at this position. These (neo-) N-terminal residues target AK2 to different branches of the N-end rule pathway. For example, an N-terminal asparagine would thereby become deaminated to an aspartate residue. For bovine AK2 (containing an asparagine at position 4), this was indeed observed. In an analysis of its N-terminus, an aspartate was observed at position 4 for some molecules in line with a preceding deamination step (Schlauderer & Schulz, 1996).

AK2 was in a previous high-throughput study identified as potential target for both DPP8 and DPP9 (Wilson et al., 2013). We demonstrate that DPP8 and DPP9 processing of AK2 indeed occurs in cells and that this processing regulates AK2 levels and localization. DPP8-dependent processing became however apparent only in the absence of DPP9 indicating that at least in HEK293 cells DPP9 is the dominant cytosolic protease of the two (Fig 4B). This is in line with the high sequence similarity, structure, and determined cleavage motif for DPP8 and DPP9 (Ross et al., 2018). It also indicates the potential for specific regulation by differential expression or post-translational modifications of both proteases.

What are the consequences of differential DPP8/9 activities for AK2? We demonstrated that absence of AK2 processing results in increased cellular levels of AK2 and in the accumulation of folded, active, cytosolic AK2 (Figs 5 and 6E). AK2 activity is linked to metabolic signaling (Dzeja et al., 2004, 2011; Dzeja & Terzic, 2009), maintenance of OXPHOS efficiency (Burkart et al., 2011; Six et al., 2015), and to induction of the ER unfolded protein response (Burkart et al., 2011) as well as apoptosis (Single et al., 1998; Kohler et al., 1999; Lee et al., 2007). Control of DPP8/9 activity might thus serve in modulating AK2 function, e.g., during differentiation or mitochondrial biogenesis. Indeed, using a murine primary B-cell differentiation model, we found that DPP8/9 impacts AK2 levels (Fig 4G). AK2 has together with the DUSP26 phosphatase been reported to play a role in regulating cell proliferation and activation of apoptosis (Kim et al., 2014). This would require cytosolic stabilization of AK2. Attenuation of DPP8/9 processing would provide an explanation on how AK2 could remain stable in the cytosol to fulfill these functions.

Cytosolic AK2 has the same enzymatic properties as its IMS counterpart, including the stringent regulation by AMP inhibition in a range that might become relevant under starvation conditions (Fig 6C and D; Soboll et al., 1978, 1980). AK2 activity is however considerably lower than the one of its cytosolic counterpart AK1 (Appendix Fig S7A and B) indicating that cytosolic AK2 activity becomes relevant only in cells with low or absent AK1 activity. Cells of the hematopoietic system lack AK1 activity, explaining the
A MitoCarta2.0 - DPP8/9 motif

Subcellular localization

matrix
IMM
IMS
OMM
unknown

Fraction with DPP8/9 motif

% of dataset

0 10 20 30 40 50

protein number

0 400 800 1200

B MIA40 substrates

human protein

AK2
NDUFB10
CMC2
COA5
CHCHD2
NDUFS5
NDUFA8
CHCHD7
COX17
CMC4
CHCHD10
COA6
MIA40 isoform 2
DPP8

C

Figure 7.

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immune phenotype of AK2 patients (Lagresle-Peyrou et al., 2009; Panniccier et al., 2009), and it is tempting to speculate that here DPP9 processing might affect localization and cellular levels of AK2. This would also be in line with gene expression data (Bagger et al., 2019). Granulocyte/monocyte progenitors can differentiate into either monocytes or myelocytes. DPP9 transcripts were maintained at low levels during differentiation into early promyelocytes, while they were kept at higher levels during differentiation into monocytes (Appendix Fig S7C and Bagger et al., 2019). AK2 expression was inversely regulated which might point to increased AK2 levels and a cytosolic AK2 fraction in the case of differentiation toward myelocytes. Likewise, in bone marrow plasma cells, a striking inverse correlation of DPP8/9 expression and expression of their substrates can be observed (Appendix Fig S7D).

**DPP8/9 as general regulator of mitochondrial biogenesis and cellular protein localizations**

The role of DPP8/9 processing for mitochondrial biogenesis extends beyond its role in AK2 processing. Our *in silico* predictions show that more than 100 MitoCarta2.0 proteins contain the DPP8/9 processing site and might thus be potential targets of these peptides (Fig 7). These predictions are supported by protein N-terminome data (Calvo et al., 2017). The proteins COX17 and NDUF8A were thereby identified with N-termini that correspond to the N-terminome predicted N-termini expected to occur after DPP9 cleavage (“GL-” and “GI-,” respectively; (Calvo et al., 2017)).

The *in silico* prediction also indicated that substrates of the disulfide relay are overrepresented among the potential mitochondrial DPP8/9 targets. Disulfide relay substrates usually do not contain an N-terminal MTS, and they are imported slowly most likely in a post-translational manner (Fischer et al., 2013). This slow import might therefore result in extended dwelling times of fully translated precursors in the cytosol and the risk of premature (mis-)folding (and acquisition of activity) in the wrong compartment. However, these properties would also allow efficient modulation of imported protein amounts by cytosolic N-terminal processing. We indeed observe that for some potential DPP8/9 targets levels change upon DPP8/9 inhibition in HEK293 cells (Fig 7). AK2 levels appear to be most affected by DPP8/9 inhibition, and this might at least in part also be explained by its slow and complex import and folding pathway that we propose involves intramolecular disulfide isomerization. Isomerization was so far not reported for disulfide relay substrates but frequently occurs during oxidative protein folding in the ER (Jansens et al., 2002; Land et al., 2003; Chakravarthi & Bulleid, 2004; Roberts et al., 2018).

Importantly, DPP8/9-mediated processing might not only influence the stability of its targets. For example, some disulfide relay substrates would expose a glycine after processing that might facilitate myristoylation. Myristoylation has been recently reported to affect mitochondrial import kinetics and multi-subunit complex assembly (Ueda et al., 2019).

DPP8/9 processing could also become relevant during pathophysiological conditions. Recent reports indicated that proteasome inhibition might be a promising therapeutic approach for patients that carry protein variants that delay protein import by the disulfide relay (Habich et al., 2019a; Mohanraj et al., 2019). One of these proteins, NDUFB10 (patient mutation: C107S), which is a part of complex I, exhibited a striking tissue heterogeneity: While the mutant protein was absent from liver and heart, in fibroblasts it was present and functional in mitochondria (Friederich et al., 2017). In the present study, we demonstrated that indeed DPP8/9 inhibition led to increased amounts of mitochondrial NDUFB10(C107S) (Fig 7). DPP8/9 activity was previously shown to be highest in liver, with skin cells harboring only little DPP8/9 activity (Yu et al., 2009). Thus, one might speculate that due to efficient DPP8/9 processing, NDUFB10 in liver cells has little chance to become imported when its import mechanism is impaired, as is the case with the C107S variant.

For AK2, we provide evidence that DPP9 provides a novel mode to control dual localization. This might expand to other proteins like, e.g., COX17 for which a dual localization to cytosol and mitochondria has previously been reported (Beers et al., 1997; Heaton et al., 2001). Compared to another major mode of cytosol-IMS dual localization, competition between folding and import (resulting in high cytosolic and low IMS protein levels), DPP8/9-dependent regulation ensures high IMS protein levels and only a small protein fraction to accumulate in the cytosol. It also allows depleting the cytosol completely of unwanted localized IMS proteins.
To serve in regulation of mitochondrial biogenesis, cellular DPP8/9 activity needs to be tunable. High-throughput studies indicated the potential to regulate DPP9 activity posttranslationally by cysteine residue modification, lysine acetylation, and malonylation or binding of sumoylated proteins (Gorrell et al., 2009). One of the modified lysine residues, K43 in DPP9, is situated very close to the substrate-binding site and its modification might well affect substrate binding and link DPP9 activity to the energy state of the cell. Likewise, the conserved cysteine 844 in DPP9 (which is not found in DPP8) is close to the substrate-binding site. Modification of cysteine residues with NEM or hydrogen peroxide inhibited DPP9 activity (Gorrell et al., 2006). Thus, it is tempting to assume that specifically DPP9 activity is susceptible to oxidative stress or redox signaling. Lastly, the activity of DPP9 is increased by binding to sumoylated proteins (Pilla et al., 2012), integrating the DPP9 processing pathway into other cellular signaling networks. DPP8/9 variations in DPP8/9 activity.

Inflammation, in preadipocyte differentiation, and during cancer progression. We demonstrate here an important molecular role for DPP8 and DPP9 in controlling mitochondrial biogenesis. Given the crucial role of mitochondria in many of the above processes, our findings might well be closely integrated into many cellular processes.

DPP8 and DPP9 have been assigned important roles in different physiological settings including roles in the immune system, in inflammation, in preadipocyte differentiation, and during cancer progression. We demonstrate here an important molecular role for DPP8 and DPP9 in controlling mitochondrial biogenesis. Given the crucial role of mitochondria in many of the above processes, our results in the future allow detailing the role of DPP8/9-modulated mitochondrial import in understanding physiological outcomes of variations in DPP8/9 activity.

Materials and Methods

Plasmids and cell lines

For plasmids and cell lines used in this study, see Appendix Table S1. For the generation of stable, inducible cell lines the HEK293 cell line-based Flp-In T-Rex-293 cell line was used with the Flp-In T-Rex system (Invitrogen). For siRNA-mediated knockdown experiments and analysis of endogenous protein levels upon 1G244-treatment, HeLa cells were used. Cells were cultured in DMEM supplemented with 8% fetal bovine serum and penicillin/streptomycin at 37°C under 5% CO2.

Digitonin-fractionation

To assess the localization of AK2 upon inhibition of DPP8/9, 1G244-treated HeLa cells were trypsinized and washed with ice-cold PBS. The PBS was removed, and cells were resuspended in 200 μl fractionation buffer (20 mM HEPES pH 7.4, 250 mM sucrose, 50 mM KCl, 2.5 mM MgCl2, 1 mM DTT). Then, the cell suspension was supplemented with 800 μl fractionation buffer containing either 0, 0.003 or 0.03% digitonin and 1× protease inhibitor cocktail (Sigma-Aldrich). To degrade DNA, 25 U Benzonase (Sigma-Aldrich) was added. The samples were incubated for 30 min on ice while being inverted every 5 min. Samples were centrifuged for 10 min at 9,000 g at 4°C. Proteins of the supernatant were precipitated using ice-cold TCA. The pellet was resuspended in 800 μl fractionation buffer containing 25 μg/ml trypsin, and samples were incubated for 30 min on ice while being inverted every 5 min. Then, proteins of the pellet were precipitated using ice-cold TCA. TCA pellets were dissolved in 80 μl resolubilization buffer (0.2 M Tris–HCl pH 7.5, 6 M urea, 10 mM EDTA, 2% SDS). Samples were sonicated until pellets were entirely dissolved, supplemented with 4× Laemmlili buffer containing 200 mM DTT, and analyzed via SDS–PAGE and immunoblotting.

Homogenizer fractionation

For separation of cytosol, mitochondria and nuclei, a fractionation protocol based on mechanical shear forces was performed. The whole procedure was carried out at 4°C and with pre-cooled buffers. HEK293-cells were washed with PBS, incubated in hypoosmotic PBS and afterward resuspended in 200 μl STM buffer (250 mM sucrose, 50 mM Tris–HCl pH 7.4, 5 mM MgCl2, protease inhibitor cocktail (Sigma-Aldrich, S8820)). The cells were disrupted in reaction tubes using a Potter–Elvehjem style homogenizer for 1 min. 20% of the lysate were taken as control. Nuclei were separated from the mixture by centrifugation for 3 min at 300 g. The nuclear fraction was subsequently washed twice in STM buffer and resuspended in NET-buffer (20 mM HEPES pH 7.9, 5 mM MgCl2, 0.5 M NaCl, 0.2 mM EDTA, 20% glycerol, 1% Triton X-100, protease inhibitor cocktail (Sigma-Aldrich, S8820)). The cells were disrupted in reaction tubes using a Potter–Elvehjem style homogenizer for 1 min. 20% of the lysate were taken as control. Nuclei were separated from the mixture by centrifugation for 3 min at 300 g. The nuclear fraction was subsequently washed twice in STM buffer and resuspended in NET-buffer (20 mM HEPES pH 7.9, 5 mM MgCl2, 0.5 M NaCl, 0.2 mM EDTA, 20% glycerol, 1% Triton X-100, protease inhibitor cocktail (Sigma-Aldrich, S8820)). The cells were disrupted in reaction tubes using a Potter–Elvehjem style homogenizer for 1 min. 20% of the lysate were taken as control. Nuclei were separated from the mixture by centrifugation for 3 min at 300 g. The nuclear fraction was subsequently washed twice in STM buffer and resuspended in NET-buffer (20 mM HEPES pH 7.9, 5 mM MgCl2, 0.5 M NaCl, 0.2 mM EDTA, 20% glycerol, 1% Triton X-100, protease inhibitor cocktail (Sigma-Aldrich, S8820)). The cells were disrupted in reaction tubes using a Potter–Elvehjem style homogenizer for 1 min. 20% of the lysate were taken as control. Nuclei were separated from the mixture by centrifugation for 3 min at 300 g. The nuclear fraction was subsequently washed twice in STM buffer and resuspended in NET-buffer (20 mM HEPES pH 7.9, 5 mM MgCl2, 0.5 M NaCl, 0.2 mM EDTA, 20% glycerol, 1% Triton X-100, protease inhibitor cocktail (Sigma-Aldrich, S8820)).

Cell treatments

To inhibit the proteasome, cells were cultured with medium supplemented with MG132. For Western blot analysis, cells were treated with 1 μM MG132 for 16 h, while in pulse-chase experiments treatment with 5 μM MG132 started during the starvation period. To inhibit DPP8/9, cells were supplemented with medium containing 1G244. To assess the effect on HA-tagged proteins, cells were treated with 10 μM 1G244 for 16 h. The effect on endogenous proteins was assessed after incubation of cells with 1 μM 1G244 for 3 or 5 days and renewal of medium each day. Exogenous protein expression using the Flp-In T-Rex-293 cell line was induced by incubation of cells for 16 h with medium containing doxycycline (0.01 μg/ml or 1 μg/ml for AK2-variants, 1 μg/ml for all other proteins). For siRNA-mediated knockdown, cells were transfected with siRNA (Qiagen) (Appendix Table S1) using Interferin (Peqlab)
according to the manufacturer’s protocol and incubated for 72 h. Emetine chase experiments were performed by addition of 100 μg/ml emetine for the indicated times.

**Steady-state protein levels**

To assess steady-state protein levels, cells were washed with ice-cold PBS and lysed either by addition of Laemmli buffer (2% SDS, 60 mM Tris–HCl pH 6.8, 10% glycerol, 0.0025% bromophenol blue) or either LCW buffer (0.5% Triton X-100, 0.5% Na-deoxycholate, 150 mM NaCl, 20 mM Tris pH 7.5, 10 mM EDTA, 30 mM Na-pyrophosphate) or RIPA buffer (50 mM Tris pH 8, 150 mM NaCl, 1% Triton X-100, 0.5% sodium deoxycholate, 0.1% SDS). RIPA and LCW buffers were supplemented with 1× protease inhibitor cocktail (Sigma-Aldrich). Laemmli lysis samples were boiled for 10 min at 96°C and analyzed via SDS–PAGE and immunoblotting. RIPA/LCW lysis samples were incubated for 30 min on ice and then centrifuged for 30 min at 20,000 g at 4°C. The protein concentration of the supernatant was determined via Roti-Quant Universal Kit (Carl Roth). Samples of 50 μg protein were prepared, boiled for 5 min at 96°C, and analyzed via SDS–PAGE and immunoblotting.

**Viability assay**

Cell viability was assessed with the PrestoBlue Cell Viability Reagent (Thermo Fisher). To this end, cells were grown in 96-well plates and subjected to different treatments. The PrestoBlue reagent was applied to the cells in a 1:10 dilution and incubated for 1 h at 37°C. Afterward, the fluorescence was measured in a CLARIOstar plate reader with excitation at 560 nm and emission at 595 nm. After the measurement, medium was exchanged to fresh medium without PrestoBlue for continued culture. For analysis, all values were blank-corrected and normalized to their respective untreated control.

**Immunoprecipitation**

Immunoprecipitations were carried out either combined with native lysis or with denaturing lysis. For both, cells were washed with PBS containing 20 mM NEM (N-Ethylmaleimide) and afterward incubated at 4°C in PBS-NEM for 10 min. Afterward, the cells were scratched off and sedimented (500 g, 5 min). For native lysis, the cells were resuspended in native lysis buffer (100 mM NaPi pH 8.1; 1% Triton X-100) and lysed on ice for 1 h. Cell debris was sedimented (20,000 g, 4°C, 1 h), and the supernatant was transferred to beads. Denaturing lysis started with resuspension of the cells in denaturing lysis buffer (30 mM Tris pH 8.1, 150 mM NaCl, 1 mM EDTA). For cell lysis, 1.6% SDS was added and the sample boiled at 95°C for 5 min. To lower sample viscosity, 1 μl Benzonase (Sigma-Aldrich, E1014) was added. After lysis, the samples were filled up to 1 ml with denaturing lysis buffer containing 2.5% Triton X-100. The samples were incubated on ice for 1 h followed by centrifugation for 1 h at 20,000 g and 4°C. Supernatants were then transferred to beads. Beads were tumbled with the samples for 16 h at 4°C for both IP variants. After binding, the beads were washed four times with the respective lysis buffer with Triton X-100 and one last time with the respective lysis buffer without Triton X-100. For elution, washing buffer was removed completely and 1× Laemmli buffer was added, followed by 3 min boiling at 95°C.

**Protein purification**

All AK2 variants as well as AK1 were heterologously expressed with a C-terminal 6x-His tag in Escherichia coli Rosetta 2 cells. The proteins were purified by Immobilized Metal Affinity Chromatography using Ni-NTA columns (Qiagen), followed by size exclusion chromatography with a 16/600 Superdex 75 pg column (GE). Protein concentration was determined by measuring absorbance at 280 nm and using a calculated extinction coefficient (ProtParam).

**Assay to address inverse and direct redox states of protein thiols**

The steady-state redox state assay was performed similarly to what was described before (Erdogan et al., 2018). Reduced thiols were blocked by treatment with 20 mM NEM (N-Ethylmaleimide) dissolved in PBS at 4°C for 15 min for minimal shift (min) and steady-state (ss) samples. Maximum shift (max) samples were treated with PBS at 4°C for 15 min. The cells were washed twice with PBS and finally scratched off in PBS. The cells were sedimented by centrifugation at 500 g for 5 min and lysed in native IP lysis buffer. Immunoprecipitation was performed as described above. After elution of proteins by addition of 1× Laemmli buffer, the proteins were treated with 40 mM TCEP (tris(2-carboxyethyl)phosphine) for 15 min at 65°C before mmPEG12 modification (15 μM final concentration). For min shift controls, mmPEG12 was omitted. After modification, samples were loaded and analyzed on Tris–Tricine–PAGE, Western blotting, and immunodetection.

For determination of progressive oxidation during protein matura- tion, the in vivo oxidation assay was performed. After pulse labeling, the proteins were modified by addition of mmPEG12 (ss samples). The max shift samples were reduced by incubation with 40 mM TCEP for 15 min at 65°C and modification afterward. Min shift samples were left untreated. After modification, the samples were subjected to native immunoprecipitation as described above followed by elution from the beads and analysis on Tris–Tricine–PAGE, Western blotting, and autoradiography.

**Pulse-Chase biogenesis assay**

Pulse-chase assays to assess protein oxidation in intact cells (oxida- tion assay) were performed as described previously (Fischer et al., 2013). Newly synthesized proteins were pulse-labeled with EasyTag EXPRESS 35S Protein Labeling Mix (Perkin Elmer) at a concentration of 200 μCi/ml. The pulse labeling was stopped by addition of chase medium containing 20 mM methionine and incubation for indicated times. The chase was stopped by adding ice-cold 8% TCA. After TCA precipitation, the pellets were dissolved in resolving buffer (0.2 M Tris pH 7.5, 10 mM EDTA, 6 M Urea, 2% SDS, 0.0025% bromophenol blue) and either used for thiol modification or directly for immunoprecipitation.

**SILAC and mass spectrometry**

Stable Isotope Labeling in Cell Culture (SILAC) was performed as described in Petruangaro et al. (2015). Cells were subcultured and
passaged in SILAC-DMEM (Thermo Fisher), supplemented with 10% dialyzed FBS (Gibco, Invitrogen), 1% L-glutamine (PAN Biotech), containing either L-arginine or L-arginine-13C6-15N2 (42 mg/l), and L-lysine or L-lysine-13C6 (73 mg/l), and 27.3 mg/l proline. Cells were lysed by adding native lysis buffer (100 mM NaPi pH 8.0, 100 mM NaCl, 1% Triton X-100). Lysates were incubated for 1 h at 4°C and then centrifuged for 1 h at 25,000 g. The supernatant was subjected to immunoprecipitation with anti-HA beads (Sigma-Aldrich) at 4°C overnight. Samples were washed four times using native lysis buffer and once with lysis buffer without Triton X-100. The collected beads were heated in a Laemmli buffer containing 1 mM DTT for 10 min at 95°C. After alkylation using 5.5 mM iodoacetamide for 10 min at room temperature, the samples were centrifuged and the supernatants were loaded on 4–12% gradient gels (NuPAGE, Thermo Fisher) for protein separation. After staining, each gel lane was cut into 5–10 slices, the proteins were in-gel digested with trypsin (Promega) and the resulting peptide mixtures were processed on STAGE tips and analyzed by LC-MS/MS. The LC-MS measurements were performed on a QExactive Plus or HF-X mass spectrometer coupled to an EasyLC 1000/1200 nanoflow-HPLC. Peptides were separated on fused silica HPLC-column tip (i.D. 75 μm, New Objective, self-packed with Reprosil-Pur 120 C18-AQ, 1.9 μm (Dr. Maish) to a length of 20 cm) using a gradient of A (0.1% formic acid in water) and B (0.1% formic acid in 80% acetonitrile in water); loading of sample with 0% B with a flow rate of 600 nl/min; separation ramp from 5 to 30% B within 85 min with a flow rate of 250 nl/min). For nanoESI, the spray voltage was set to 2.3 kV and ion-transfer tube temperature to 250°C, no sheath and auxiliary gas was used. The mass spectrometer was operated in the data-dependent mode; after each MS scan (mass range m/z = 370-1,750; resolution: 70,000 for the Plus and HF-X) a maximum of ten and 12 MS/MS scans were performed using a normalized collision energy of 25%, a target value of 1,000, and 5,000 and a resolution of 17,500 and 30,000, for the Plus and HF-X mass spectrometers, respectively. The MS raw files were analyzed using MaxQuant Software version 1.4.1.2 (Cox & Mann, 2008) for peak detection, quantification, and peptide identification using a full-length UniProt human database (March, 2016) and common contaminants such as keratins and enzymes used for in-gel digestion as reference. Carbamidomethylcysteine was set as fixed modification and protein amino-terminal acetylation and oxidation of methionine were set as variable modifications. The MS/MS tolerance was set to 20 ppm, and three missed cleavages were allowed using trypsin/P as enzyme specificity. Peptide and protein FDR based on a forward-reverse database were set to 0.01, minimum peptide length was set to 7, and minimum number of peptides for identification of proteins was set to one, which must be unique. The “match-between-run” option was used with a time window of 1 min.

In vitro AK2 activity assays

For the forward activity assay, AK activity was coupled to pyruvate kinase (PK) and lactate dehydrogenase (LDH, PK/LDH mix from Sigma-Aldrich). In this coupled reaction, ADP produced by AKs is used by PK to convert phosphoenolpyruvate to pyruvate, which is used by LDH to oxidize NADH, thus decreasing the absorbance at 340 nm. The following reaction conditions were used: 50 mM Tris/HCl pH 7.5, 50 mM KCl, 4 mM MgCl2, 0.006% BSA, 0.25 mM NADH, 1 mM PEP, 1 mM ATP. The concentration of AK2 was set to 0.8 nM, AK1 to 0.4 nM.

The reverse activity assay was carried out similarly to the forward activity assay, by coupling the AK reaction to hexokinase (HK) and glucose-6-phosphate dehydrogenase (G6PDH, HK/G6PDH mix from Roche). In this assay, AKs provide the ATP for glucose phosphorylation by HK, followed by NADP+ reduction to NADPH and an increase in absorbance at 340 nm. The reaction conditions were as follows: 58 mM glycyglycine pH 7.4, 10 mM MgCl2, 0.006% BSA, 0.25 mM NADP+, 20 mM glucose. The concentration of AK2 was set to 4 nM, AK1 to 0.8 nM. All measurements were performed in triplicates in 96-well plates and read in a CLARIOstar microplate reader set to 25°C. A measurement without AK was performed simultaneously to all measurements to allow subtraction of the background reaction.

Cellular AK2 activity assay

AK2 activity in HEK293 cells was assessed by performing the aforementioned coupled enzyme assay with immunoprecipitated AK2-HA. HEK293 cells overexpressing AK2-HA variants were lysed in native lysis buffer (100 mM NaPi pH 8.1, 0.2% N-Dodecyl-D-maltoside) for 20 min on ice. AK2-HA was purified from the supernatants by incubation for 2 h with anti-HA agarose beads (Sigma). After washing, the beads were resuspended in ADP assay buffer containing 0.38 mM NADP+ and 1 mM ADP. The reaction mix was then incubated and shaken at 25°C. For each timepoint, some supernatant was taken after a short centrifugation and transferred to 96-well plates for absorbance measurement. Additionally, a reaction mix without added beads was used as negative control.

Immunofluorescence and colocalization analysis

HEK293 Flp-In T-REx cells expressing HA-tagged AK2 mutants were cultured on poly-L-lysine-coated cover slips for 48 h. Protein expression was induced by addition of 1 μg/ml doxycycline after 24 h of culture. Before fixation, the cells were incubated with MitoTracker Red (Thermo Fisher) for 1 h at 37°C. Fixation was performed with 4% paraformaldehyde for 15 min. Then, cell membranes were permeabilized with blocking buffer (20 mM HEPES pH 7.4, 3% BSA, 0.3% Triton X-100) for 1 h. Cells were washed and incubated with primary (anti-HA, 3F10, Roche) and secondary antibodies (anti-rat, AlexaFluor 488) for 16 h at 4°C and 1 h at room temperature, respectively. Then, the cells were washed and the cover slips transferred to microscope slides and mounted in ProLong Gold Antifade mounting DAPI. Afterward, the cells were analyzed by confocal fluorescence microscopy. The images were analyzed for correlation of the HA signals and the MitoTracker signals with CellProfiler (Carpenter et al, 2006). In the analysis, cellular outlines are defined by thresholding and watershedding followed by measurement of the Pearson’s correlation coefficient in the defined outlines. The results of the pipeline were checked manually for suitability afterward (e.g., clumped cells which are indistinguishable were excluded). Representative pictures were adjusted for the respective intensity ranges with Fiji (Schindelin et al, 2012). Statistical analysis was performed in RStudio using the Kruskal–Wallis
rank sum test followed by Dunn’s test adjusted with the Benjamini-Hochberg method.

**Thermofluor assay**

The thermal stability of proteins was determined by using the thermal shift fluorescence assay (Niesen et al., 2007). The assay was performed in a CFX 96 Real-Time PCR Detection Systems (Bio-Rad Laboratories). Sypro Orange (SS692, Sigma-Aldrich) was used in a 1:5,000 dilution. Melting curves were obtained while ramping the temperature from 20 to 95°C at a rate of 1°C/min in steps of 0.5°C every 30 s. Fluorescence was monitored using “FRET” scan mode and was measured after each step. The melting temperature (Tm) of a protein is defined as the inflection point of the unfolding transition; or in other words, as the minimum of the negative first derivative of the melting curve. Tm was determined by using the internal algorithms of the data analysis software (Bio-Rad CFX Manager™, Version: 3.1.1517.0823).

**Conservation and logo plots**

Amino acid conservation analysis was performed by using protein sequences provided by UniProt and aligning them using Jalview (Waterhouse et al., 2009). Alignment of sequences was achieved with help of the algorithm of MUSCLE (Edgar, 2004). For the AK2 N-terminus comparison, sequences from 163 species were aligned, while for the other proteins 11 species each were compared. The conservation is depicted as logo plot, where the relative sizes of the letters correspond to the relative abundance of the respective residue at the given position. Additionally, the consensus sequence is shown below the logo plots.

**Quantification and statistical analysis**

Intensity of autoradiography and immunoblot signals were quantified using ImageQuantTL (GE) and Image Laboratory (Bio-Rad Laboratories), respectively. For SILAC MS data interpretation, Perseus software was used (Tyanova et al, 2016). Correlation analysis was performed with CellProfiler (Carpenter et al, 2006) and statistical analysis of correlation analyses with RStudio. Analysis of amino acid conservation was performed with Jalview. Error bars in figures represent standard deviation. The number of experiments is reported in the figure legend.

**Data availability**

The authors declare that there are no primary datasets and computer codes associated with this study.

**Expanded View** for this article is available online.

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**Author contributions**

JR, MH, and YF designed the study and planned experiments. MH, YF, SG, JK, LS, KJL, Evdl, MA, CP, SLS, CP, and SU performed experiments and analyzed data. KH performed in silico analysis. JP performed proteomic analysis. JR, MH, and YF designed figures. JR, MH, and YF wrote the manuscript with critical input from UB, BB, KH, and DM.

**Conflict of interest**

The authors declare that they have no conflict of interest.

**References**

Avadhani NG, Sangar MC, Bansal S, Bajpai P (2011) Bimodal targeting of cytochrome P450s to endoplasmic reticulum and mitochondria: the concept of chimeric signals. *FEBS J* 278: 4218 – 4229

Bagger FO, Kinalis S, Rapin N (2019) BloodSpot: a database of healthy and malignant haematopoiesis updated with purified and single cell mRNA sequencing profiles. *Nucleic Acids Res* 47: D881 – D885

Banci L, Bertini I, Cefaro C, Casacchi L, Ciofi-Raffoni S, Felli IC, Gallo A, Connelli L, Luchinat E, Sideris D et al (2010) Molecular chaperone function of Mia40 triggers consecutive induced folding steps of the substrate in mitochondrial protein import. *Proc Natl Acad Sci USA* 107: 20190 – 20195

Barchiesi A, Wasilewski M, Chacinska A, Tell G, Vascotto C (2015) Mitochondrial translocation of APE1 relies on the MIA pathway. *Nucleic Acids Res* 43: 5451 – 5464

Beers J, Glerrum DM, Tzagoloff A (1997) Purification, characterization, and localization of yeast Cox17p, a mitochondrial copper shuttle. *J Biol Chem* 272: 33191 – 33196

Boopathi E, Srinivasan S, Fang JK, Avadhani NG (2008) Bimodal protein targeting through activation of cryptic mitochondrial targeting signals by an inducible cytosolic endoprotease. *Mol Cell* 32: 32 – 42

Bragoszewski P, Gornicka A, Sztolsztener ME, Chacinska A (2013) The ubiquitin-proteasome system regulates mitochondrial intermembrane space proteins. *Mol Cell Biol* 33: 2136 – 2148

Bragoszewski P, Turek M, Chacinska A (2017) Control of mitochondrial biogenesis and function by the ubiquitin-proteasome system. *Open Biol* 7: 170007

Bruns GA, Regina VM (1977) Adenylate kinase 2, a mitochondrial enzyme. *Biochem Genet* 15: 477 – 486

Burkart A, Shi X, Chouinard M, Corvera S (2011) Adenylate kinase 2 links mitochondrial energy metabolism to the induction of the unfolded protein response. *J Biol Chem* 286: 4081 – 4089

Calvo SE, Clauser KR, Mootha VK (2016) MitoCarta2.0: an updated inventory of mammalian mitochondrial proteins. *Nucleic Acids Res* 44: D1251 – D1257

Calvo SE, Julien O, Clauser KR, Shen H, Kamer KJ, Wells JA, Mootha VK (2017) Comparative analysis of mitochondrial N-termini from mouse, human, and yeast. *Mol Cell Proteomics* 16: 512 – 523

Carpenter AE, Jones TR, Lamprecht MR, Clarke C, Kang IH, Friman O, Guertin DA, Chang JH, Lindquist RA, Moffat J et al (2006) Cell Profiler: image
analysis software for identifying and quantifying cell phenotypes. Genome Biol 7: R100

Chakravarthi S, Bulleid NJ (2004) Glutathione is required to regulate the formation of native disulfide bonds within proteins entering the secretory pathway. J Biol Chem 279: 39872 – 39879

Chandel NS (2015) Evolution of mitochondria as signaling organelles. Cell Metab 22: 204 – 206

Chretilen D, Benit P, Ha HH, Keipert S, El Khoury R, Chang YT, Jastrow M, Jacobs HT, Rustin P, Rak M (2018) Mitochondria are physiologically maintained at close to 50 degrees C. PLoS Biol 16: e2003992

Cox J, Mann M (2008) MaxQuant enables high peptide identification rates, individualized p.p.b.-range mass accuracies and proteome-wide protein quantification. Nat Biotechnol 26: 1367 – 1372

Dailey TA, Woodruff JH, Dailey HA (2005) Examination of mitochondrial protein targeting of haem synthetic enzymes: in vivo identification of three functional haem-responsive motifs in 5-aminolevulinate synthase. Biochem J 386: 381 – 386

Delfasmo D, Panelli D, Sardenelli AM, Papa S (2008) CAMP-dependent protein kinase regulates the mitochondrial import of the nuclear encoded NDUFS4 subunit of complex I. Cell Signal 20: 989 – 997

Dzeja PP, Terzic A (2012) The metabolic regulation of aging. Adv Exp Med Biol 752: 109 – 132

Dzeja P, Terzic A (2013) Intracellular copper metallochaperone Cox17 exists as an oligomeric, polycopper complex. Biochemistry 42: 743 – 751

Habich M, Salscheider SL, Schmahl LM, Hoehne MN, Fischer M, Schom F, Petrungrao C, Ali M, Erdogan Aj, Abou-Eid S et al (2019a) Vectorial import via a metastable disulfide-linked complex allows for a quality control step and import by the mitochondrial disulfide relay. Cell Rep 26: 759 – 774

Habich M, Salscheider SL, Riemer J (2019b) Cysteine residues in mitochondrial intermembrane space proteins: more than just import. Br J Pharmacol 176: 514 – 531

Hupf V, Zou P, Rhee HW, Udedi ND, Cracan V, Svinkina T, Carr SA, Mootha VK, Ting AY (2014) Proteomic mapping of the human mitochondrial intermembrane space in live cells via ratiometric APEX tagging. Mol Cell 55: 332 – 341

Jackson TD, Palmer CS, Stojanovski D (2018) Mitochondrial diseases caused by dysfunctional mitochondrial protein import. Biochem Soc Trans 46: 1225 – 1238

Jansens A, van Duijn E, Braakman I (2002) Coordinated nonvectorial folding in a newly synthesized multidomain protein. Science 298: 2401 – 2403

Kim H, Lee HJ, Oh Y, Choi SG, Hong SH, Kim HJ, Lee SY, Choi JW, Su Hwang D, Kim KS et al (2014) The DUSP26 phosphatase activator adenylosuccinate kinase 2 regulates FADD phosphorylation and cell growth. Nat Commun 5: 3351

Kloppel C, Suzuki Y, Kojer P, Petrungrao C, Longen S, Fiedler S, Keller S, Riemer J (2011) Mia40-dependent oxidation of cysteines in domain I of Ccs1 controls its distribution between mitochondria and the cytosol. Mol Biol Cell 22: 3749 – 3757

Koch JR, Schmid FX (2014) Mia40 targets cysteines in a hydrophobic environment to direct oxidative protein folding in the mitochondria. Nat Commun 5: 3041

Kohler C, Gamh A, Noma T, Nakazawa A, Orrenius S, Zhivotovsky B (1999) Release of adenylate kinase 2 from the mitochondrial intermembrane space during apoptosis. FEBS Lett 447: 10 – 12

Lagesle-Peyrou C, Sicsu CM, Picard C, Rieux-Laucat F, Michel V, Ditadi A, Demerens-de Chapellelaine C, Morillon E, Valensi F, Simon-Stos KL et al (2009) Human adenylate kinase 2 deficiency causes a profound hematopoietic defect associated with sensorineural deafness. Nat Genet 41: 106 – 111

Land A, Zonneveerd D, Braakman I (2003) Folding of HIV-1 envelope glycoprotein involves extensive isomerization of disulfide bonds and conformational-dependent leader peptide cleavage. PASEB J 17: 1058 – 1067

Lee HJ, Pyo JO, Oh Y, Kim HJ, Hong SH, Jeon YJ, Kim H, Cho DH, Woo HN, Song S et al (2007) AK2 activates a novel apoptotic pathway through formation of a complex with FADD and caspase-10. Nat Cell Biol 9: 1303 – 1310

Liang P, Phillips GN Jr, Glaser M (2019) Assignment of the nucleotide binding sites and the mechanism of substrate inhibition of Escherichia coli adenylate kinase. Proteins 9: 28 – 36

MacPherson L, Tokatlidis K (2019) Protein trafficking in the mitochondrial intermembrane space: mechanisms and links to human disease. Biochem J 474: 2533 – 2545

Mendillo ML, Santagata S, Koeva M, Bell GW, Hu R, Tamimi RM, Fraenkel E, Ince TA, Whitesell L, Lindquist S (2012) HSF1 drives a transcriptional...
program distinct from heat shock to support highly malignant human cancers. Cell 150: 549 – 562
Milenkovic D, Ramming T, Muller JM, Wenz LS, Gebert N, Schulze-Specking A, Stojanovski D, Rospert S, Chacinska A (2009) Identification of the signal directing Tim9 and Tim10 into the intermembrane space of mitochondria. Mol Biol Cell 20: 2530 – 2539
Mohranjan K, Wasielski M, Benincà C, Cysewski D, Poznanski J, Sakowska P, Bugajska Z, Deckers M, Dennerlein S, Fernandez-Vizarra E et al (2019) Inhibition of proteasome rescues a pathogenic variant of respiratory chain assembly factor COA7. EMBO Mol Med 11: e9561
Nguyen KT, Mun SH, Lee CS, Hwang CS (2018) Control of protein degradation by N-terminal acetylation and the N-end rule pathway. Exp Mol Med 50: 91
Niesen FH, Berglund H, Vedadi M (2007) The use of differential scanning fluorimetry to detect ligand interactions that promote protein stability. Nat Protoc 2: 2212 – 2221
Pannicke U, Tokatidis K, Dessen P, Kroemer G (2016) Mitochondrial proteins containing Coiled-Coil-Helix-Coiled-Coil-Helix (CHCH) domains in health and disease. Trends Biochem Sci 41: 245 – 260
Peleh V, Cordat E, Herrmann JM (2013) Mia40 is a trans-site receptor that drives protein import into the mitochondrial intermembrane space by hydrophobic substrate binding. Elife 5: e16177
Petrungaro C, Zimmermann KM, Kuttner V, Fischer M, Dengjel J, Bogeski I, Riemer J (2015) The Ca(2+)-dependent release of the Mia40-induced MICU1-MICU2 dimer from MCU regulates mitochondrial Ca(2+) uptake. Cell Metab 22: 721 – 733
Pfanner N, Warscheid B, Wiedemann N (2013) The Ca(2+)-specific interacting motif in dipeptidyl peptidase 8 affects redox homeostasis in the endoplasmic reticulum. EMBO J 32: 3773
Ramesh A, Peleh V, Martinez-Caballero S, Wollweber F, Sommer F, van der Veen I, Mia et al (2012) Mia40 is a trans-site receptor that drives protein import into the mitochondrial intermembrane space by hydrophobic substrate binding. Elife 5: e16177
Reinstein J, Schlichting I, Wittinghofer A (1990) Structurally and catalytically important residues in the phosphate binding loop of adenylate kinase of Escherichia coli. Biochemistry 29: 7451 – 7459
Rhee HW, Zou P, Udeshi ND, Martell JD, Motha VK, Carr SA, Ting AY (2013) Proteomic mapping of mitochondria in living cells via spatially restricted enzymatic tagging. Science 339: 1328 – 1331
Roberts BS, Babilonia-Rosa MA, Broadwell LJ, Wu MJ, Neher SB (2018) Lipase maturation factor 1 affects redox homeostasis in the endoplasmic reticulum. EMBO J 37: e97379
Ross B, Krapp S, Augustin M, Kierfussaer R, Arciniega M, Geiss-Friedlander R, Huber R (2018) Structures and mechanism of dipeptidyl peptides 8 and 9, important players in cellular homeostasis and cancer. Proc Natl Acad Sci USA 115: E1437 – E1445
Santagata S, Mendillo ML, Tang YC, Subramanian A, Perley CC, Roche SP, Wong B, Narayan R, Kwon H, Koeva M et al (2013) Tight coordination of protein translation and HSF1 activation supports the anabolic malignant state. Science 341: 1238303
Sato A, Ogita H (2017) Pathophysiological implications of dipeptidyl peptidases. Curr Protein Pept Sci 18: 843 – 849
Schindelin J, Arganda-Carreras I, Frise E, Kaynig V, Longair M, Pietzsch T, Preibisch S, Rueden C, Saalfeld S, Schmid B et al (2012) Fiji: an open-source platform for biological-image analysis. Nat Methods 9: 676 – 682
Schlaurer GJ, Schulz GE (1996) The structure of bovine mitochondrial adenylate kinase: comparison with isoenzymes in other compartments. Protein Sci 5: 434 – 441
Schmidt O, Harbauer AB, Rao S, Eyrich B, Zahedi RP, Stojanovski D, Schonfisch B, Giaid B, Sickmann A, Pfanner N et al (2011) Regulation of mitochondrial protein import by cytosolic kinases. Cell 144: 227 – 239
Sides DP, Petrakis N, Katakii N, Mikropoulou D, Gallo A, Cioffi-Baffoni S, Banci L, Bertini I, Tokatidis K (2009) A novel intermembrane space-targeting signal docks cysteines onto Mia40 during mitochondrial oxidative folding. J Cell Biol 187: 1007 – 1022
Sineva MA, Sineva EV, Itah V, Haas E (1996) Towards a mechanism of AMP-substrate inhibition in adenylate kinase from Escherichia coli. FEBS Lett 397: 273 – 276
Six E, Lagresle-Peyrou C, Susini S, De Chappedelaine C, Sigrist N, Sadek H, Chouette M, Cagnard N, Fontenay M, Hermine O et al (2015) AK2 deficiency compromises the mitochondrial energy metabolism required for differentiation of human neutrophil and lymphoid lineages. Cell Death Dis 6: e1856
Soboll S, Scholz R, Heldt HW (1978) Subcellular metabolite concentrations. Dependence of mitochondrial and cytosolic ATP systems on the metabolic state of perfused rat liver. Eur J Biochem 87: 377 – 390
Soboll S, Akerboom TP, Schwenke WD, Haase R, Sies H (2010) Mitochondrial and cytosolic ATP/ADP ratios in isolated hepatocytes. A comparison of the digitoxin method and the non-aqueous fractionation procedure. Biochim J 192: 951 – 954
Stojanovski D, Bragoszewski P, Chacinska A (2012) The Mia pathway: a tight bond between protein transport and oxidative folding in mitochondria. Biochim Biophys Acta 1823: 1142 – 1150
Strobel G, Zollner A, Angermann M, Bandlow W (2002) Competition of spontaneous protein folding and mitochondrial import causes dual subcellular location of major adenylate kinase. Mol Biol Cell 13: 1439 – 1448
Suzuki Y, Ali M, Fischer M, Riemer J (2013) Human copper chaperone for superoxide dismutase 1 mediates its own oxidation-dependent import into mitochondria. Nat Commun 4: 2430
Topf U, Wrobel L, Chacinska A (2016) Chatty mitochondria: keeping balance in cellular protein homeostasis. Trends Cell Biol 26: 577 – 586
Tyanova S, Temu T, Sinitsyn P, Carlson A, Hein MY, Geiger T, Mann M, Cox J (2016) The Perseus computational platform for comprehensive analysis of (pro)teomics data. Nat Methods 13: 731 – 740
Ueda E, Tamura Y, Sakaue H, Kawano S, Kakuta C, Matsumoto S, Endo T (2019) Myristoyl-group-aided protein import into the mitochondrial intermembrane space. Sci Rep 9: 1185
Varshavsky A (2017) The ubiquitin system, autophagy, and regulated protein degradation. Annu Rev Biochem 86: 123 – 128
Vongsamphanh R, Fortier PK, Ramotar D (2001) Pir1p mediates translocation of the yeast Apn1p endonuclease into the mitochondria to maintain genomic stability. Mol Cell Biol 21: 1647 – 1655
Wai T, Langer T (2016) Mitochondrial dynamics and metabolic regulation. Trends Endocrinol Metab 27: 105 – 117
Waterhouse AM, Procter JB, Martin DM, Clamp M, Barton GJ (2009) Jalview Version 2—a multiple sequence alignment editor and analysis workbench. *Bioinformatics* 25: 1189–1191

Wilson CH, Indarto D, Doucet A, Pogson LD, Pitman MR, McNicholas K, Menz RI, Overall CM, Abbott CA (2013) Identifying natural substrates for dipeptidyl peptidases 8 and 9 using terminal amine isotopic labeling of substrates (TAILS) reveals in vivo roles in cellular homeostasis and energy metabolism. *J Biol Chem* 288: 13936–13949

Wilson CH, Zhang HE, Gorrell MD, Abbott CA (2016) Dipeptidyl peptidase 9 substrates and their discovery: current progress and the application of mass spectrometry-based approaches. *Biol Chem* 397: 837–856

Wu JJ, Tang HK, Yeh TK, Chen CM, Shy HS, Chu YR, Chien CH, Tsai TY, Huang YC, Huang YL et al (2009) Biochemistry, pharmacokinetics, and toxicology of a potent and selective DPP8/9 inhibitor. *Biochem Pharmacol* 78: 203–210

Yogevo O, Pines O (2011) Dual targeting of mitochondrial proteins: mechanism, regulation and function. *Biochim Biophys Acta* 1808: 1012–1020

Yu DM, Ajami K, Gall MG, Park J, Lee CS, Evans KA, Mclaughlin EA, Pitman MR, Abbott CA, McCaughan GW et al (2009) The in vivo expression of dipeptidyl peptidases 8 and 9. *J Histochem Cytochem* 57: 1025–1040

Zhang H, Maqsudi S, Rainczuk A, Duffield N, Lawrence J, Keane FM, Justa-Schuch D, Geiss-Friedlander R, Gorrell MD, Stephens AN (2015) Identification of novel dipeptidyl peptidase 9 substrates by two-dimensional differential in-gel electrophoresis. *FEBS J* 282: 3737–3757

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