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Formation of cristae and crista junctions in mitochondria depends on antagonism between Fc1 and Su e/g

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Crista junctions (CJs) are important for mitochondrial organization and function, but the molecular basis of their formation and architecture is obscure. We have identified and characterized a mitochondrial membrane protein in yeast, Fc1 (formation of CJ protein 1), which is specifically enriched in CJs. Cells lacking Fc1 lack CJs, exhibit concentric stacks of inner membrane in the mitochondrial matrix, and show increased levels of F₁F₀-ATP synthase (F₁F₀) supercomplexes. Overexpression of Fc1 leads to increased CJ

formation, branching of cristae, enlargement of CJ diameter, and reduced levels of F₁F₀ supercomplexes. Impairment of F₁F₀ oligomer formation by deletion of its subunits e/g (Su e/g) causes CJ diameter enlargement and reduction of cristae tip numbers and promotes cristae branching. Fc1 and Su e/g genetically interact. We propose a model in which the antagonism between Fc1 and Su e/g locally modulates the F₁F₀ oligomeric state, thereby controlling membrane curvature of cristae to generate CJs and cristae tips.

Introduction

Mitochondria are ubiquitous organelles and fulfill a multitude of crucial functions in eukaryotic organisms. Mitochondria are made up by two membranes: the outer membrane (OM) and the inner membrane (IM). The IM is composed of two subdomains: the inner boundary membrane (IBM) and the cristae membrane (CM). The IBM is closely apposed to the OM, both forming a double-layered envelope of the organelle. CMs are invaginations of the IBM that protrude into the matrix space. Large variations exist in the morphology of CMs (Munn, 1974; Fawcett, 1981; for review see Zick et al., 2009), and aberrant mitochondrial structures have been described for numerous pathological

situations in humans (DiMauro et al., 1985; Wallace, 2005). Tubular-, lamellar-, and even triangle-shaped structures of the CM have been observed (Fawcett, 1981). Cristae are connected to the IBM by narrow tubular- or slotlike structures of varying length, so-called crista junctions (CJs), as revealed by EM of serial sections of mitochondria (Daems and Wisse, 1966) and by electron tomography (Mannella et al., 1994; Perkins et al., 1997, 1998; Nicastro et al., 2000; Frey et al., 2002; for reviews see Frey and Mannella, 2000; Mannella et al., 2001). The diameter of CJs was found to be rather small, ranging from 12 to 40 nm (Nicastro et al., 2000; Perkins et al., 2003; for review see Frey and Mannella, 2000). This led to the suggestion of CJs forming barriers for the movement of proteins and metabolites between the intracristal and the intermembrane space as well as

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Abbreviations used in this paper: BN-PAGE, blue native PAGE; CJ, crista junction; CM, cristae membrane; DIC, differential interference contrast; IBM, inner boundary membrane; IM, inner membrane; mtDNA, mitochondrial DNA; OM, outer membrane; PK, proteinase K; SEC, size exclusion chromatography; TAP, tandem affinity purification.

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between the CM and the IBM (Mannella et al., 1994). Such a role of CJs has been proposed to have important consequences for the regulation of oxidative phosphorylation, as a barrier of this kind might limit the diffusion of metabolites like ADP into the intracristal space and modulate the pH gradient across the IM (Perkins et al., 1997; Renken et al., 2002; for reviews see Mannella et al., 2001; Mannella, 2006a). Also, subcompartmentalization of the IM was suggested based on biochemical subfractionation of mitochondria (Werner and Neupert, 1972; Pon et al., 1989), localization of individual mitochondrial proteins by immuno-EM (Gilkerson et al., 2003), or fluorescence microscopy (Wurm and Jakobs, 2006). Recently, this subcompartmentalization was addressed by determining the distribution of 20 mitochondrial proteins using quantitative immuno-EM (Vogel et al., 2006). The IBM appears to be segregated from the CM by the CJ, yet proteins are able to dynamically redistribute between the two subcompartments of the IM depending on the physiological state of the cell (Vogel et al., 2006). Furthermore, CJs undergo remodeling during apoptosis; this was suggested to allow release of the intracristal pool of cytochrome *c* to the cytoplasm, thereby triggering programmed cell death (Scorrano et al., 2002; Cipolat et al., 2006; Frezza et al., 2006).

Despite the apparent importance of the structural organization of mitochondria, the components responsible for the morphology and biogenesis of cristae, and in particular of CJs, are largely unknown. Deletion of the dimer-specific subunit *e* (Su *e*) or Su *g* of the F₁F₀-ATP synthase (F₁F₀) leads to defective oligomerization of this complex (Arnold et al., 1998) and to altered cristae morphology with extended onion-like structures in yeast (Paumard et al., 2002). Furthermore, remodeling of CJs during apoptosis was reported to depend on the mitochondrial dynamin-like protein OPA1 (Frezza et al., 2006). Prohibitins were recently suggested to play a role in cristae morphogenesis by controlling OPA1 processing (Merkwirth et al., 2008). The yeast orthologue of OPA1, Mgm1, was proposed to be required for cristae maintenance in addition to its role in IM fusion (Wong et al., 2003; Meeusen et al., 2006). Another protein reported to determine cristae morphology in human cells is mitofilin. Down-regulation of this protein led to massive proliferation of the IM, resulting in multiple layers that appear to be interconnected accompanied by a disappearance of CJs (John et al., 2005). Mitofilin was initially named heart muscle protein, as it is highly expressed in rat heart tissue (Icho et al., 1994). Proteomic analysis of human heart mitochondria further indicated high abundance of this protein in mitochondria (Taylor et al., 2003). The name mitofilin was coined because of its colocalization with filament-like mitochondria in fibroblasts (Odgren et al., 1996).

In summary, a few proteins are known to influence cristae morphology, but, so far, no protein has been shown to be specifically localized to CJs or to be part of a CJ protein complex. In this study, we investigated the molecular basis governing the architecture of cristae and CJs in mitochondria of the yeast *Saccharomyces cerevisiae*. We describe ultrastructural alterations of the IM of yeast mitochondria lacking the open reading frame *AIM28* (*YKR016w/FMP13*) with hitherto unknown function. This gene was recently listed as a putative orthologue of mammalian mitofilin (Olichon et al., 2006). We provide both

morphological and biochemical evidence that this protein is directly involved in the formation and molecular structure of CJs and is specifically enriched at CJs. We propose to name it Fcj1 (formation of CJ protein 1). Fcj1 regulates the oligomeric state of the F₁F₀ supercomplex in an adverse manner, and, moreover, CJ formation and structure turn out to be dependent on F₁F₀ supercomplex formation itself. Furthermore, Su *e* and Su *g* of F₁F₀, which promote oligomerization of F₁F₀, are essential for the formation of endings of cristae, the tips, or rims. The F₁F₀ complex has been proposed to represent a key element in the determination of cristae structure (Paumard et al., 2002), a proposal which rests on its very high abundance in the mitochondrial IM. In this study, we present a working model that explains the control of cristae architecture by the antagonistic effects of Fcj1 and F₁F₀ Su *e* and Su *g* on the oligomeric state of F₁F₀.

Results

Deletion of Fcj1 leads to altered mitochondrial morphology and partial loss of mitochondrial DNA (mtDNA)

In a search for genes affecting mitochondrial structure, we analyzed a collection of yeast cells that showed deficiency in growth on nonfermentable carbon sources because mutants with this phenotype are often accompanied by alterations in mitochondrial structure. We focused on genes coding for putative mitochondrial proteins that, so far, do not have an assigned function. Possible candidate strains were checked for mitochondrial morphology by rhodamine staining and then by expression of matrix-targeted GFP. We identified a mutant that showed alterations in the typical mitochondrial network. In the strain lacking the open reading frame *FCJ1* (*AIM28/FMP13/YKR016w*), abnormally large spherical mitochondria were observed. These were mostly located in the periphery of the cell and formed atypical networks (Fig. 1 A). In addition, mitochondria occasionally appeared hollow. For further characterization, we determined the percentage of rho⁰/rho⁻ cells. Depending on the cell density and the growth medium used, this value varied from 15 to 40% for the mutant cells. The respiratory rates of isolated mitochondria from such cells in state III (+ADP) and state IV (-ADP) and the maximal uncoupled rates were reduced by 30% as compared with wild type (Table I). The membrane potential, Δψ, of the mutant mitochondria was also reduced compared with wild type, although to a lesser extent (Table I). The respiratory control ratio (state III/state IV rates) in the mutant was not altered, indicating that the respiratory chain and the ATP synthase were still well coupled. Taking all of these observations into account, compromised respiration in the mutant cells was rather caused by a secondary loss of mtDNA than by primary deficiencies of enzymatic activities involved in oxidative phosphorylation.

Fcj1 is located in the mitochondrial IM facing the intermembrane space with its large hydrophilic domain

The *FCJ1* (*AIM28/FMP13/YKR016w*) gene encodes a protein of 540 aa residues. In silico analysis by the MitoProt II program (Claros and Vincens, 1996) yielded a high probability for

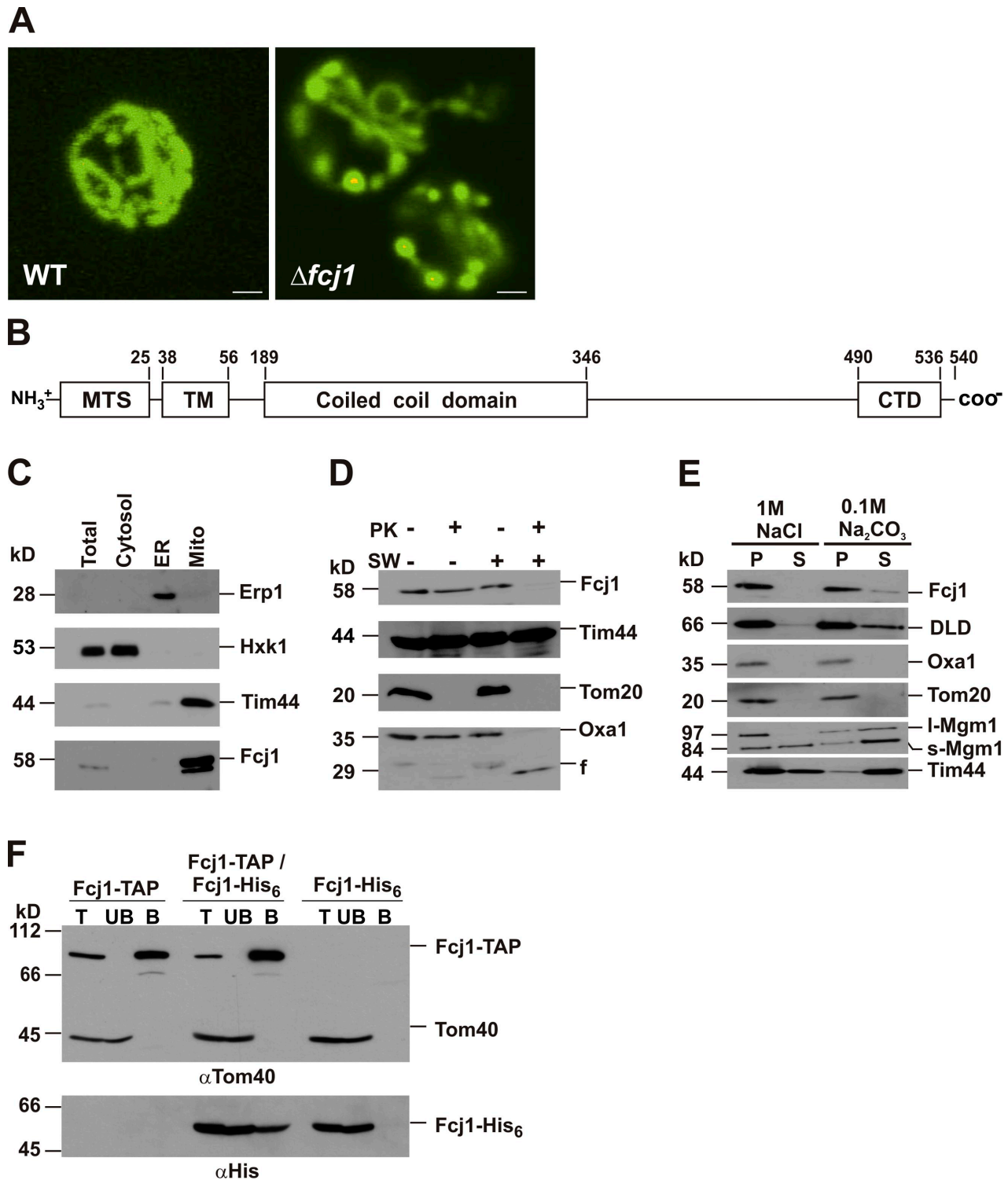


Figure 1. Fcj1 is required for normal mitochondrial morphology. (A) Wild-type (WT) and $\Delta fcj1$ cells expressing mitochondria-targeted GFP were grown on nonfermentable medium and visualized by fluorescence microscopy. (B) Domain structure of Fcj1 depicting the mitochondrial-targeting sequence (MTS), the transmembrane segment (TM), the coiled-coiled domain, and the conserved C-terminal domain (CTD) with corresponding positions of amino acid residues. (C) Subcellular fractionation of wild-type cells: mitochondria (Mito), microsomes (ER), and cytosol. Equal amounts of protein (50 μ g) were analyzed by Western blotting with the indicated marker proteins: Tim44 (Mito), Erp1 (ER), and Hxk1 (Cytosol). (D) Submitochondrial localization of Fcj1. Wild-type mitochondria and mitoplasts generated by hypotonic swelling (SW) were treated with PK. f, specific proteolytic fragment of Oxa1. (E) Membrane association of Fcj1. Wild-type mitochondria were extracted with NaCl or Na carbonate. Membrane-bound (P) and soluble (S) fractions were loaded and analyzed by Western blotting using the indicated marker proteins. DLD, D-lactate dehydrogenase. (F) Homotypic interaction of Fcj1. Mitochondria from cells expressing a His-tagged (Fcj1-His₆) or a TAP-tagged variant of Fcj1 (Fcj1-TAP) or both were subjected to TAP affinity chromatography. Total (T; 10%), bound (B; 100%), and unbound (UB; 10%) material was analyzed by Western blotting with the indicated antibodies. Tom40 and TAP-tagged Fcj1 were detected simultaneously using rabbit antibodies against Tom40. Bars, 1 μ m.

Table 1. Bioenergetic characterization of wild-type and $\Delta fcl1$ cells

Bioenergetic properties of isolated mitochondria	Wild type	$\Delta fcl1$
Uncoupled respiration (nmol oxygen/min/mg of protein)	248.5 \pm 41.27 (n = 3)	174.6 \pm 30.9 (n = 3) ^a
Respiratory control (state III/state IV)	3.01 \pm 0.23 (n = 3)	2.93 \pm 0.09 (n = 3)
Membrane potential $\Delta\Psi$ ($\Delta F/F$)	0.285 \pm 0.019 (n = 3)	0.221 \pm 0.032 (n = 3) ^b
ATPase activity (nmol P _i /min/mg of protein)		
–oligomycin	1,393 \pm 231 (n = 3)	1,264 \pm 121 (n = 3) ^c
+oligomycin	275 \pm 93 (n = 3)	256 \pm 102 (n = 3) ^d

F, fluorescence. For measuring bioenergetic properties, mitochondria were isolated from cells grown at 30°C on complete liquid media containing 2% (vol/vol) lactate. Mean values together with standard deviations (\pm) of *n* measurements are given.

^a70% of wild type.

^b78% of wild type.

^c91% of wild type.

^d93% of wild type.

mitochondrial targeting (0.9967), including a predicted cleavage site of the mitochondrial processing peptidase between positions 16 and 17 and a possible site of the mitochondrial intermediate peptidase between residues 24 and 25 (Fig. 1 B). Furthermore, Fcjl1 contains a predicted single transmembrane segment close to the N terminus. Fcjl1 shares 13% sequence identity with human mitofilin and 12% with mouse mitofilin. At the C terminus, it contains a short segment of higher similarity (Fig. S1). Several structural features such as the position of the transmembrane segment, a coiled-coil domain, and the conserved C-terminal domain argue for Fcjl1 being a member of the mitofilin protein family.

Upon subcellular fractionation, Fcjl1 was completely recovered in the mitochondria (Fig. 1 C). Incubation of isolated mitochondria with proteinase K (PK) did not lead to degradation of Fcjl1; however, after selective opening of the OM, Fcjl1 was degraded (Fig. 1 D). Treatment of mitochondria with either high salt or Na carbonate did not result in extraction of the protein (Fig. 1 E). To test whether Fcjl1 can form oligomeric structures, two differently C-terminally tagged variants of Fcjl1, a His-tagged (Fcjl1-His₆) and a tandem affinity purification (TAP)-tagged variant (Fcjl1-TAP), were coexpressed in yeast. Upon TAP affinity chromatography, Fcjl1-His₆ was copurified with Fcjl1-TAP, indicating that Fcjl1 undergoes homotypic interactions (Fig. 1 F). Affinity purification of Fcjl1-His₁₂ from $\Delta fcl1 + Fcjl1-His_{12}$ mitochondria solubilized in Triton X-100 did not lead to copurification of proteins other than Fcjl1-His₁₂ (Fig. S2 A). The size of the Fcjl1-His₁₂ complex in Triton X-100-solubilized mitochondria was \sim 180 kD, as determined by size exclusion chromatography (SEC). Moreover, the complex retained its size after affinity purification, which is consistent with the formation of a stable homo-oligomeric complex (Fig. S2 B). The size would suggest a trimeric complex; however, a dimeric or tetrameric complex seems possible as well. We also analyzed the size of the Fcjl1 complex after TAP from mitochondria harboring Fcjl1-His₆ and Fcjl1-TAP. The size of the Fcjl1-TAP-Fcjl1-His₆ complex was shifted to a molecular mass of \sim 320 kD or \sim 220 kD depending on the detection of the TAP tag or the His tag, respectively (Fig. S2 C). This increase in molecular mass is consistent with the presence of one or more subunits of Fcjl1 containing the 20-kD TAP tag, further strengthening the view of a homo-oligomeric Fcjl1 complex. Collectively, Fcjl1 is a protein engaged

in homotypic interactions that is anchored to the mitochondrial IM with its major part exposed to the intermembrane space.

Fcjl1 is enriched at CJs

We determined the distribution of Fcjl1 over the various subdomains of the IM using quantitative immuno-EM. Cryosections of chemically fixed wild-type cells were immunodecorated with antibodies against Fcjl1 and visualized by immunogold. Low concentrations of antibodies were used to keep the level of unspecific immunodecoration low. The specificity of the Fcjl1 antibody was confirmed by Western blotting as well as by immunogold labeling of wild-type and $\Delta fcl1$ cells (Fig. S3, A and B). Gold particles present in a large number of sections of wild-type cells were projected on a model, as described previously (Vogel et al., 2006), representing OM, IBM, CM, and a CJ (Fig. 2, A and B). Fcjl1 was most prominently clustered in close proximity to the CJ (Fig. 2, A and B), a region which, so far, was rather characterized by a quite low density of numerous other mitochondrial proteins (for comparison see Vogel et al. [2006]). A region nearly lacking gold particles was observed in the CM adjacent to the CJ region (Fig. 2 B). This necklike region represents the narrow tubular segment of the CM, exhibiting a relatively high positive membrane curvature. The curvature here is opposite to the one at the base of CJs. Fcjl1 was present to some extent in the remainder of the CM, including the cristae tips and in the IBM distant from the CJ region. The term cristae tip refers to its appearance in electron micrographs of mitochondrial sections. Depending on the section plane, they represent the highly positively curved rims of lamellar-like CM sheets. To evaluate these findings in a quantitative way, a sliding window was moved along the IM of our model starting from the bottom, counting only gold particles that were located within 14-nm distance from the IM (Fig. 2, C and D). This was performed for Fcjl1 and as a control for Cox2, a subunit of cytochrome *c* oxidase, as well as for Su *e* and Su *g* of F₁F₀ (Fig. 2, C and D; raw data for Cox2 and Su *e* and Su *g* were taken from Vogel et al. [2006]). With Fcjl1, the highest number of gold particles was observed in the region corresponding to the CJ (Fig. 2, C and D, gray boxes). The number of gold particles representing Cox2 was very low in the CJ and IBM region and increased in the CM (Fig. 2 C; Vogel et al., 2006). Su *e* and Su *g* of F₁F₀ were not enriched in the CJ region

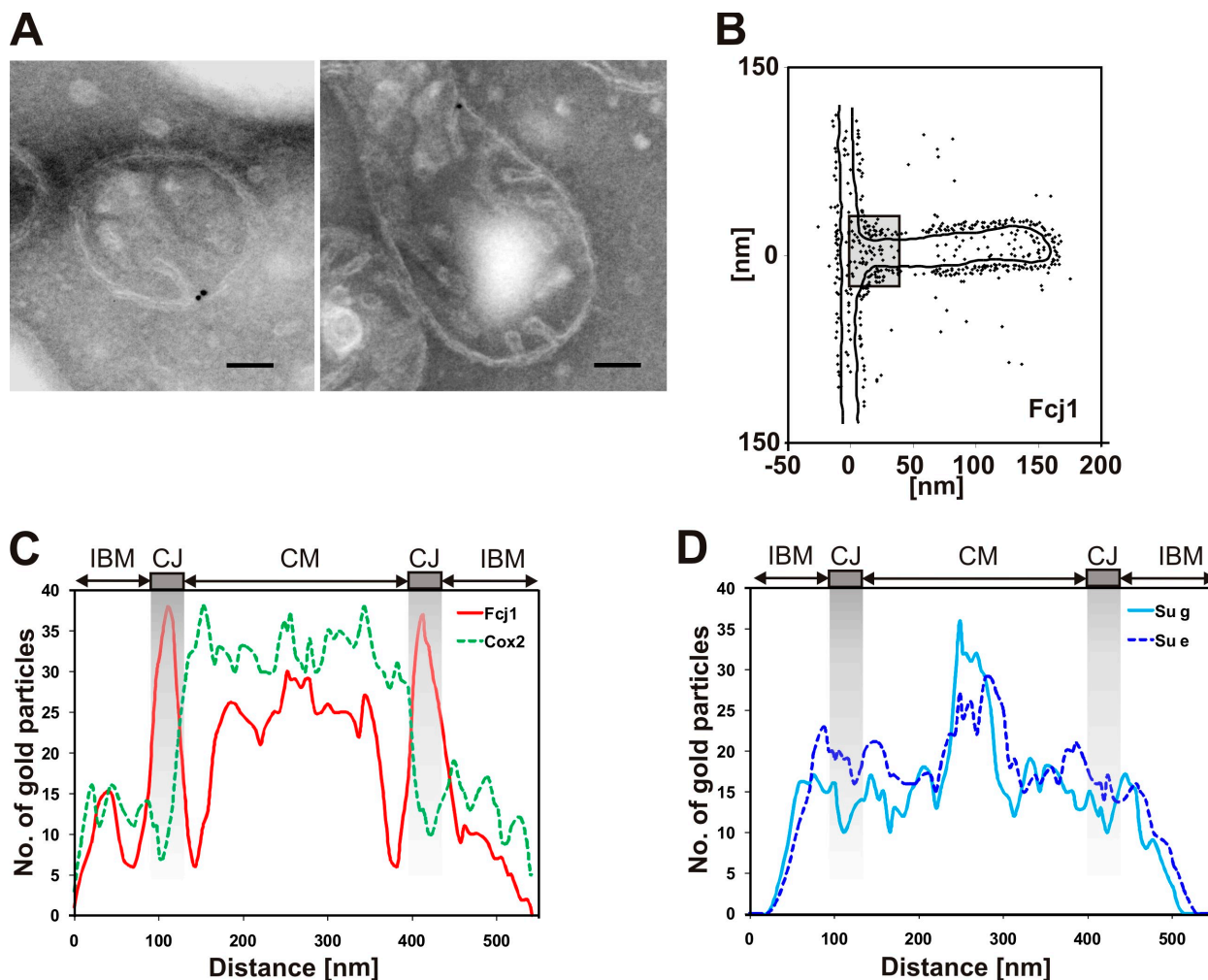


Figure 2. **Fcjl is enriched at CJs.** (A) Immunogold labeling of Fcjl in wild-type cells. (B) Representation of gold particles after immunogold labeling of Fcjl plotted on a model of CM, IBM, and OM (Vogel et al., 2006). (C and D) Quantification of protein densities in the IM. The number of gold particles occurring within a 14-nm distance from the IM was determined by moving a sliding window in silico along the IM in the model from bottom to top (raw data from C and from Vogel et al. [2006]). Gray boxes indicate the CJ region. The protein densities of Fcjl and Cox2 (C) and of Su g and Su e (D) are shown. Bars, 100 nm.

but rather appeared accumulated in the cristae tips (Fig. 2 D). Collectively, the enrichment of Fcjl at CJs was unique and not observed for any of the proteins analyzed previously (Fig. 2, C and D; Vogel et al., 2006). The presence of Fcjl in lower amounts in the planar parts of the cristae, where the levels of Su e and Su g are also intermediate, could reflect a balance of function in these regions; in part, they could also be caused by the topological complexity of bent cristae and perhaps, to a lesser degree, by background signals. Altogether, our data demonstrate that Fcjl is the only protein identified so far specifically enriched at CJs at sites where the amounts of Su e and Su g are relatively low.

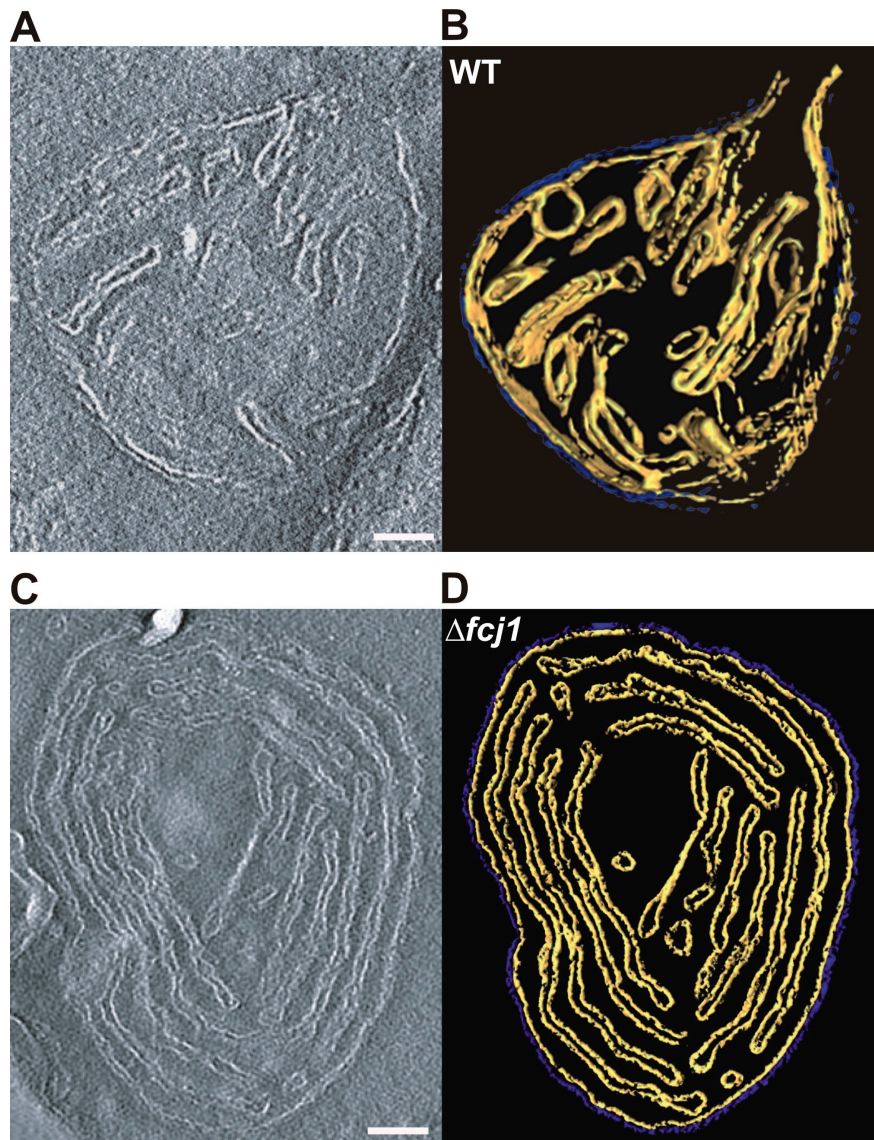
Deletion of Fcjl leads to loss of CJs and altered cristae morphology

The morphology of mitochondrial IMs was studied by EM and tomographic reconstruction from sections of yeast cells prepared by the Tokuyasu method (Tokuyasu, 1989). In wild-type cells, mitochondria showed a large number of CJs, and the IM

and OM were often in very close proximity at the bases of CJs (Fig. 3, A and B; Table II; and Video 1). In $\Delta fcjl$ cells, mitochondria were increased in size and contained numerous internal cristae with no apparent CJs (Fig. 3, C and D; Table II; and Video 2). These CMs were arranged mainly as parallel, concentric, stacked vesicular structures. The lumen of the vesicular structures was identified as intracristal space (Fig. S3 C). In a further approach, rapidly frozen hydrated spheroplasted yeast cells were subjected to cryosectioning and cryo-EM tomography. In addition, isolated mitochondria were analyzed by cryo-EM tomography. Both procedures led to high resolution structures of CJs in wild-type cells (Fig. 4, A and B; Fig. S4, A–C; and Video 3) and revealed the absence of CJs in $\Delta fcjl$ mitochondria (Fig. 4 C, Fig. S4 D, and Video 4).

The narrow matrix space between adjacent CMs in mitochondria of $\Delta fcjl$ cells contained particles of uniform size and shape that were arranged in a zipperlike pattern (Fig. 4, D–G; and Videos 4 and 5). Top views reveal the distribution and organization of these complexes as short linear assemblies or as two

Figure 3. **Cristae morphology is altered, and CJs are absent in cells lacking Fcj1.** (A–D) EM tomograms of chemically fixed yeast cells. (A) Single slice of a tomogram of a mitochondrion in a wild-type cell. (B) Surface-rendered view of A. (C) Single slice of a tomogram of a mitochondrion from a $\Delta f cj 1$ cell. (D) Surface-rendered view of C. The OM (blue) and IM (yellow) are shown. WT, wild type. Bars, 100 nm.



parallel lines with a square-like or a hexagonal arrangement (Fig. 4, D–G). The dimensions and relative distances of these structures corresponded very well to those of the F_1 part of the F_1F_0 -ATP synthase ($\sim 10 \times 10 \times 10$ nm). These highly abundant structures have been assigned to dimers or oligomers of F_1F_0 in numerous other studies (Allen et al., 1989; Dudkina et al., 2005, 2006; Minauro-Sanmiguel et al., 2005; Buzhynskyy et al., 2007; Strauss et al., 2008).

To provide further evidence that the zipperlike structures represent oligomers of F_1F_0 , we deleted *Su e* or *Su g* of F_1F_0 . Loss of these subunits is known to impair dimerization/oligomerization of F_1F_0 (Arnold et al., 1998). Indeed, neither in $\Delta f cj 1/\Delta s u g$ (Fig. 5 A) nor in $\Delta f cj 1/\Delta s u e$ mitochondria (not depicted) were the putative F_1 particles arranged in a zipperlike manner. Rather, they were distributed randomly over the IM, which is in contrast to what was observed with $\Delta f cj 1$ mitochondria

Table II. **Analysis of electron micrographs of cells with disturbed IM organization upon deletion of Fcj1, Su e, or Su g**

Strain	Number of mitochondrial sections	Number of CJs	Number of CJs per mitochondrial section	Percentage of CJs connected to one or more CJs	Number of cristae tips	Number of cristae tips per mitochondrial section
WT	65	120	1.8	0	245	3.8
$\Delta f cj 1$	75	0	0	0	731	9.7
$\Delta s u e$	58	170	2.9	84.1	12	0.2
$\Delta s u g$	64	163	2.5	81.0	10	0.2

WT, wild type. The numbers of CJs and cristae tips per mitochondrial section were determined from electron micrographs of chemically fixed whole cells of the indicated strains grown at 30°C on complete liquid media containing 2% (vol/vol) lactate.

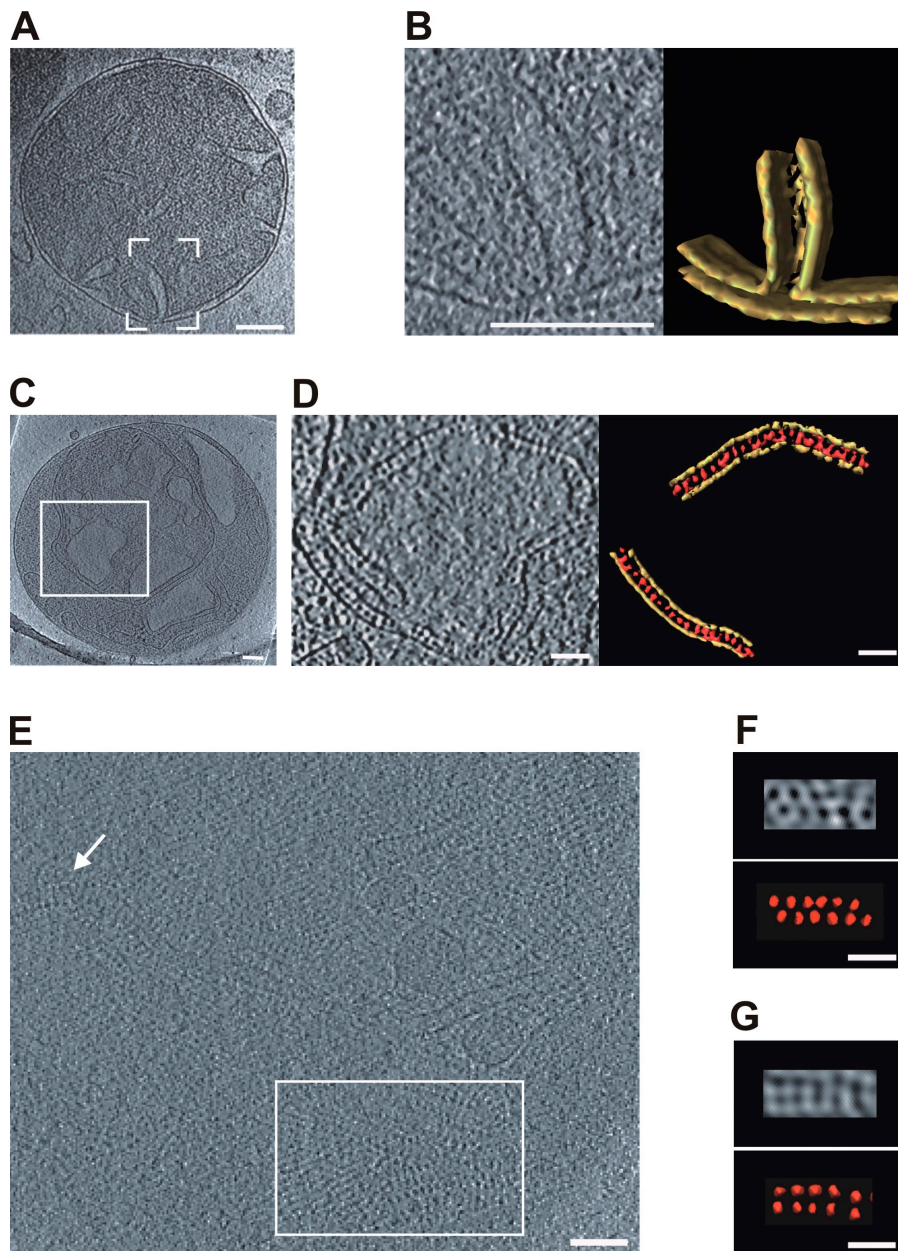


Figure 4. $\Delta fcj1$ mitochondria contain zipper-like structures typical of oligomers of the F_1F_0 -ATP synthase. Cryo-EM tomograms of isolated mitochondria. (A) Slice through a tomogram of a vitrified wild-type mitochondrion. (B) A CJ magnified (left) and surface rendered (right) corresponding to the boxed area in A is shown. Panels A and B are reprinted from Zick et al. (2009) with permission from *Biochimica et Biophysica Acta*. (C–G) $\Delta fcj1$ mitochondria. (C) Slice through a tomogram of a vitrified $\Delta fcj1$ mitochondrion. (D) Magnified view of boxed area in C (left) and surface-rendered representation (right) showing a zipperlike, regular arrangement with characteristics of the F_1 parts of F_1F_0 -ATP synthase. The IM is shown in yellow, and F_1F_0 -ATP synthases are shown in red. (E) A 10-nm thick slice of a tomogram cutting through particles typical of the F_1 parts of F_1F_0 -ATP synthase in side views (arrow) and top views (boxed area). (F and G) Sections through the tomogram and volume-rendered top views of putative F_1F_0 -ATP synthases arranged in double-row hexagonal stripes (F) and double-row square stripes (G) are shown. Bars: (A–C and E) 100 nm; (D, F, and G) 50 nm.

as a comparison (Fig. 5 B). This was confirmed by determining the distances between a large set of particles and their respective nearest neighbors. In $\Delta fcj1$ mitochondria, the vast majority of observed distances was between 14 and 16 nm, whereas in $\Delta fcj1/\Delta su g$, an apparently random distribution with a broad range of distances was detected (Fig. 5 C). Thus, the zipperlike structures in $\Delta fcj1$ mitochondria indeed represent oligomers of F_1F_0 that can be disrupted by deletion of *Su e* or *Su g* of F_1F_0 .

In intact wild-type mitochondria, F_1F_0 supercomplexes were neither observed in this study nor in other studies, probably as the result of the high protein density of the matrix and lack of contrast (Nicastro et al., 2000). The increased volume of mitochondria lacking *Fcjl* (Table S1) probably caused a reduced protein density in the matrix and thereby facilitated the visualization of the supramolecular organization of F_1F_0 in intact mitochondria. Only in mitochondrial membrane fragments

from wild-type yeast was an oligomeric arrangement of this complex previously observed (Buzhynskyy et al., 2007; Strauss et al., 2008). Collectively, the lack of *Fcjl* correlates, on the one hand, with the absence of CJs and, on the other hand, with the appearance of regular arrangements of large protein complexes characteristic of F_1F_0 supercomplexes.

***Fcjl* has a direct role in determining the number and the architecture of CJs**

To further study the role of *Fcjl* in the formation of CJs, we used overexpression and down-regulation of *Fcjl*. *Fcjl* was expressed from a plasmid in wild-type yeast cells under control of a doxycycline-repressible promoter. Wild-type cells containing the empty plasmid were also analyzed. Upon five- to 10-fold overexpression of *Fcjl*, the number of CJs per cell was increased two- to threefold as compared with control cells. The branching

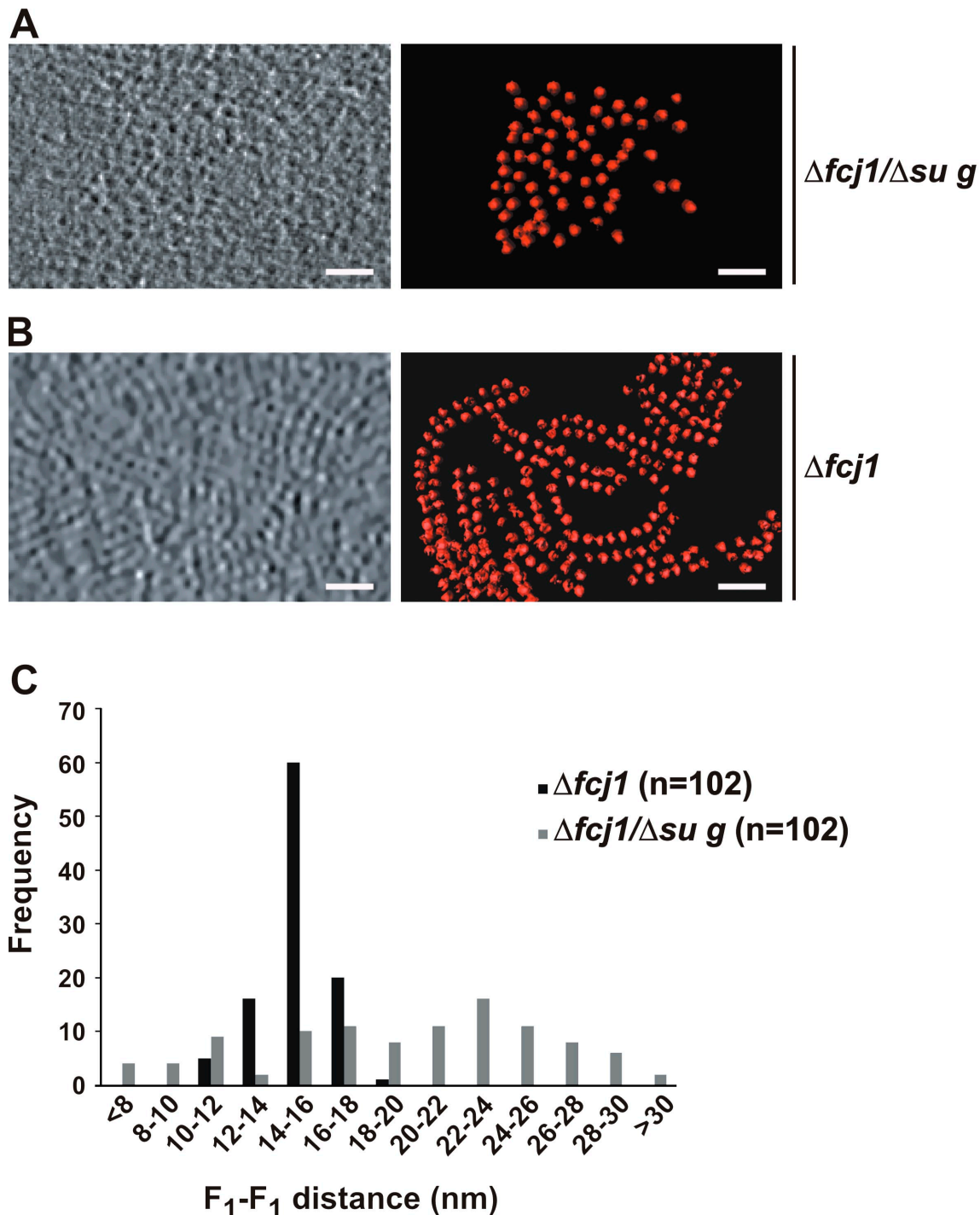


Figure 5. Zipperlike structures observed in $\Delta fcj1$ mitochondria depend on the presence of dimer-specific *Su e* and *Su g* of F_1F_0 -ATP synthase. Cryo-EM tomograms of isolated mitochondria of indicated strains. (A) Top views of putative F_1F_0 -ATP synthases in $\Delta fcj1/\Delta su g$ mitochondria (left) and a surface-rendered representation (right) are shown. (B) Top views of putative F_1F_0 -ATP synthases in $\Delta fcj1$ mitochondria (left; magnified view of boxed area in Fig. 4 E) and a surface-rendered representation (right) are shown. (C) Frequency distribution of F_1 - F_1 distances. The center to center distance of an F_1 particle to its nearest neighbor was determined for cryo-EM tomograms of isolated mitochondria of the indicated strains ($n = 102$ for both strains). Bars, 50 nm.

of cristae, which is very rarely observed in wild-type cells, was increased ~ 17 -fold in cells overexpressing *Fcj1* (Fig. 6 A). Moreover, the diameter of CJs appeared enlarged and showed a higher variation when *Fcj1* was overexpressed (Fig. 6, A and B). Thus, *Fcj1* overexpression promotes the formation of additional CJs and alters the molecular architecture of CJs.

Down-regulation of *Fcj1* led to a progressive decrease of the number of CJs as well as of cristae branches (Fig. 6, C and D). These alterations of cristae morphology occurred already at the early time points (13.5 and 23 h). Mitochondria with the characteristics of the $\Delta fcj1$ phenotype with continuous concentric stacks of IM appeared at later time points (37.5 and 47.5 h).

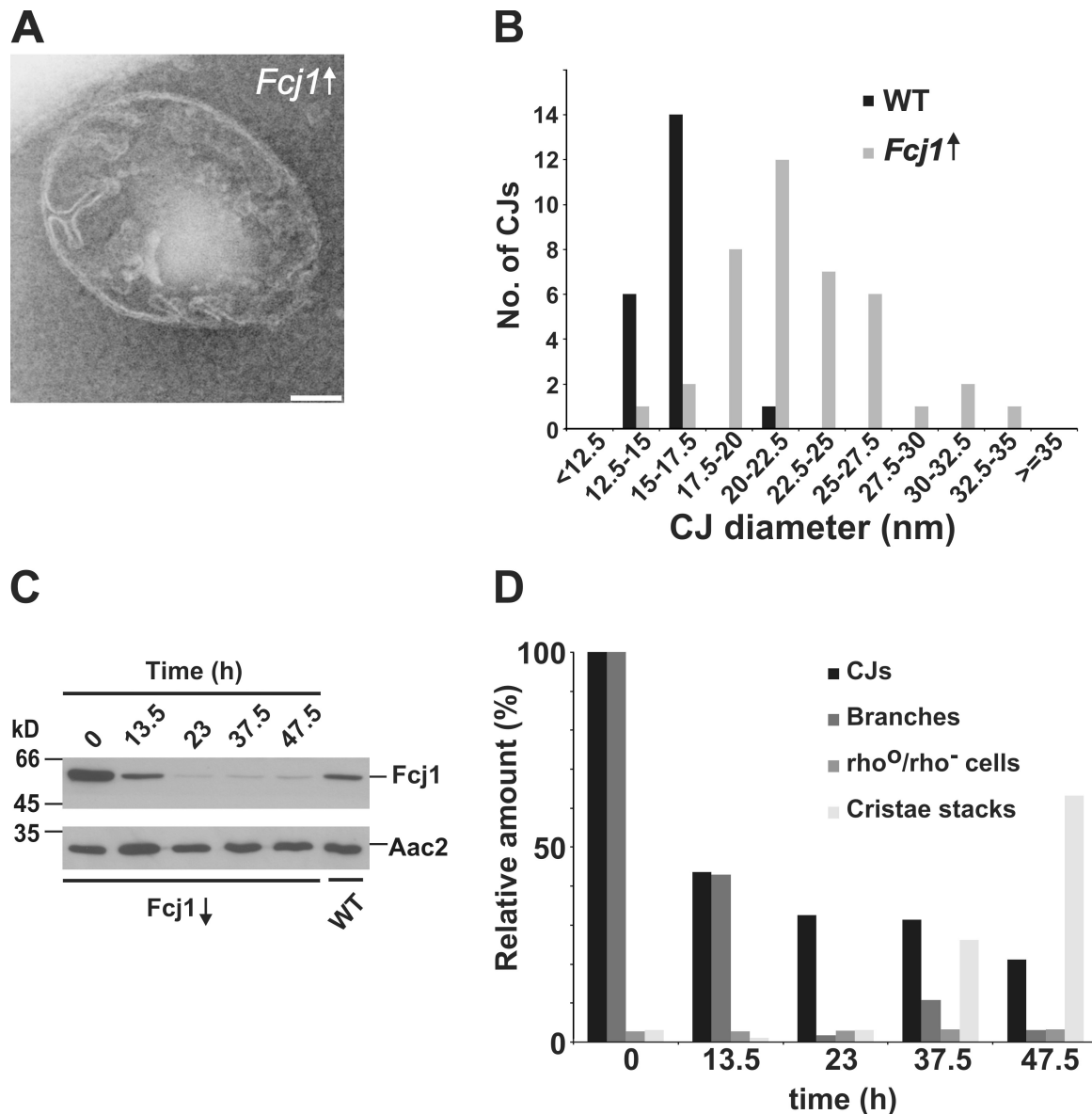


Figure 6. Fcj1 is directly involved in determining the number and the architecture of CJs. (A) Electron micrograph of a mitochondrion in a section of chemically fixed cells overexpressing Fcj1. (B) Distribution of diameters of CJs in wild-type (WT) control strain (W303) containing empty pCM189 plasmid ($n = 21$) and Fcj1-overexpressing strain (W303) containing pCM189-Fcj1 plasmid (Fcj1[↑]; $n = 40$). Cells were grown on nonfermentable, selective minimal media. A histogram of the number of diameters within the indicated ranges was plotted for both strains. (C and D) Fcj1 was down-regulated in a $\Delta fcj1$ strain harboring the pCM189-Fcj1 at different times after doxycycline addition. Wild-type control as in B was used. (C) Expression levels of Fcj1 were monitored by Western blot analysis. (D) Phenotypic analysis of down-regulation of Fcj1. The number of CJs and branches per mitochondrial section ($m_{0h} = 26$; $m_{13.5h} = 69$; $m_{23h} = 49$; $m_{37.5h} = 66$; $m_{47.5h} = 57$) was determined from electron micrographs of chemically fixed whole cells after the indicated time periods of down-regulation ($m =$ number of mitochondrial sections). The number of CJs and cristae branches per mitochondrial section before down-regulation of Fcj1 (0 h) was defined as 100%. The number of ρ^0/ρ^- cells and of cells containing cristae stacks is related to the number of total cells at each time point.

The incomplete disappearance of CJs is attributed to the incomplete down-regulation of Fcj1 (Fig. 6 C). The formation of ρ^0/ρ^- cells was not observed in the course of the experiment, excluding the possibility that our observations are caused by a secondary loss of mtDNA. In summary, these observations strongly suggest a direct involvement of Fcj1 in CJ formation.

Unchanged levels of Su e or Su g of the F_1F_0 -ATP synthase in $\Delta fcj1$ cells

Mutant strains lacking Su e or Su g of the F_1F_0 -ATP synthase were reported to have altered cristae morphology (Paumard et al., 2002).

Therefore, we asked whether protein levels of Su e or Su g were altered in $\Delta fcj1$ cells and also whether levels of Fcj1 were altered in $\Delta su e$ or $\Delta su g$ cells. In the absence of Fcj1, similar levels of Su e or Su g were observed (Fig. 7 A). Thus, altered cristae morphology in $\Delta fcj1$ mutant cells was not caused by a change in the levels of Su e or Su g. Likewise, Fcj1 levels were unaffected in $\Delta su e$ and $\Delta su g$ mutant cells, suggesting that Fcj1 is not responsible for the altered cristae morphology described for these mutants (Paumard et al., 2002). In addition, deletion of Fcj1 had no effect on the levels of several nuclear as well as mitochondrial encoded proteins that are involved in mitochondrial

functions such as mitochondrial protein import, mitochondrial fusion and fission machineries, and oxidative phosphorylation (Fig. 7 A). Notably, levels of proteins involved in determining mitochondrial morphology such as Mgm1, Fzo1, Fis1, and Dnm1 were not affected. Consistent with this, mitochondria in $\Delta fcl1$ cells formed a tubular, albeit altered, network. The formation of the short isoform of Mgm1 (s-Mgm1) was also not impaired. This further supports our conclusion that mitochondrial energy production and protein import are functional in cells lacking Fcjl as both processes are required for the formation of s-Mgm1 (Herlan et al., 2004).

Fcjl1 impairs oligomerization of the F_1F_0 -ATP synthase

In view of the presence of a zipperlike arrangement of F_1 particles in $\Delta fcl1$ mitochondria, we asked whether Fcjl affects oligomerization of F_1F_0 . First, we determined whether solubilization of F_1F_0 with a mild detergent depends on the level of Fcjl. Mitochondria isolated from wild-type cells, cells overexpressing Fcjl, and cells lacking Fcjl were solubilized with digitonin. The proportion of nonsolubilized F_1F_0 -ATP synthase subunits (F1 β and Atp4) was highest in mitochondria lacking Fcjl, lower in wild type, and lowest in cells overexpressing Fcjl (Fig. 7 B). The solubilization of other IM proteins such as Dld1 and Aac2 did not show such a dependence on the level of Fcjl. The nonsolubilized fraction of the F_1F_0 -ATP synthase could subsequently be solubilized with Triton X-100. Apparently, F_1F_0 forms a high molecular mass complex that is less easily dissociated by digitonin in mitochondria lacking Fcjl.

We then examined the supramolecular organization of F_1F_0 by blue native PAGE (BN-PAGE) and in-gel visualization of its ATPase activity. Formation of monomers, dimers, and putative tetramers of F_1F_0 was observed in mitochondria of all strains (Fig. 7 C). Higher order oligomers were observed in $\Delta fcl1$ mitochondria even at high digitonin concentrations, only at very low concentrations in wild-type mitochondria, and not when Fcjl was overexpressed. These high molecular mass complexes of F_1F_0 represent a small subpopulation of the soluble fraction. They were very stable even at high digitonin to protein ratios when Fcjl was lacking but not when Fcjl was present. Overexpression of Fcjl impaired the formation of these higher oligomers markedly and also decreased the amount of the tetrameric forms (Fig. 7 C and Fig. S5 A). The amount of dimers and monomers was not affected significantly. Likewise, the abundance of higher oligomers of F_1F_0 when determined by SEC was dependent on the presence of Fcjl (Fig. 7 D). The majority of Fcjl was recovered in the fractions corresponding to the higher oligomeric forms of F_1F_0 (>2 MD), showing that Fcjl is present in a high molecular mass complex. Thus, the size of F_1F_0 oligomers was considerably larger upon solubilization with digitonin as compared with solubilization with Triton X-100 (Fig. S2). Collectively, various lines of evidence, including biochemical data as well as EM tomography, show that Fcjl affects the stability of F_1F_0 oligomers in an adverse manner but obviously does not influence that of the dimers of this complex.

Fcjl1 acts in an antagonistic manner to Su e and Su g of the F_1F_0 -ATP synthase

The observed effect of Fcjl on the stability of oligomeric forms of F_1F_0 suggested a functional link between these proteins. Su e and Su g are known to be required for dimerization/oligomerization of F_1F_0 and are required for normal cristae morphology. To determine whether there exists a genetic interaction between these subunits and Fcjl, we determined the doubling times on a nonfermentable carbon source of each single and corresponding double deletion strain. The strain lacking Fcjl was most severely impaired in growth on a nonfermentable carbon source, showing a ~ 1.75 -fold longer doubling time as compared with wild type (Fig. 8 A). Additional deletion of either Su e or Su g of F_1F_0 led to an efficient suppression of the growth defect of the $\Delta fcl1$ strain. These double deletion strains had significantly shorter doubling times than the $\Delta fcl1$ strain. This was further confirmed by drop dilution growth tests (Fig. 8 B). In conclusion, these data demonstrate a genetic interaction of Fcjl with Su e/Su g of F_1F_0 , placing all proteins in the same pathway. The latter subunits are promoting the assembly of F_1F_0 -ATP synthase oligomers, whereas Fcjl has the opposite effect. In the double deletion strains ($\Delta fcl1/\Delta su e$ and $\Delta fcl1/\Delta su g$), the formation of F_1F_0 -ATP synthase oligomers was not observed (Fig. S5 B). Dimers were present to a very low extent, resembling the phenotype of single deletions of Su e or Su g. This is consistent with the view that dimers are the building blocks for oligomerization of F_1F_0 , as previously suggested (Paumard et al., 2002; Wittig et al., 2008). Collectively, Fcjl interacts functionally with Su e and Su g in an antagonistic mode that appears crucial for the formation of CJs.

Deletion of Su e and Su g leads to alterations of the IM structure of mitochondria

To relate the oligomeric states of F_1F_0 to the structure of cristae, we determined the number of cristae tips and the number of CJs in the deletion mutants of Su e and Su g and compared them with the numbers in wild-type and $\Delta fcl1$ cells. Cristae tips per mitochondrial section were ~ 2.5 -fold more abundant in sections of $\Delta fcl1$ cells than in wild type (Table II).

Mitochondria of strains lacking Su e or Su g were reported to contain onion-like structures and extended CMs (Paumard et al., 2002), but it was not investigated whether formation of CJs is affected. Therefore, we performed a quantitative analysis of CJs in these strains. Next to onion-like structures, we observed CJs in these strains, and the numbers of CJs per mitochondrial section were modestly increased compared with those in the corresponding wild type (Table II). Thus, Su e and Su g are not essential for CJ formation. However, CJs often appeared in pairs and were connected via CMs crossing a mitochondrial section, which is a structure rarely observed in wild type (Fig. 8, C and D; and Table II). In line with this observation, the number of cristae tips per mitochondrial section observed in $\Delta su e$ and $\Delta su g$ mitochondria was extremely low, ~ 20 times lower than in wild type (Table II). Moreover, branching of cristae was found in 27% and 9% of mitochondrial sections of $\Delta su e$ and $\Delta su g$ strains, respectively, but not in sections of the corresponding wild-type strain (Fig. 8, C and D). Furthermore, the mean diameters of CJs in $\Delta su e$ and $\Delta su g$ cells were increased, although to a smaller extent

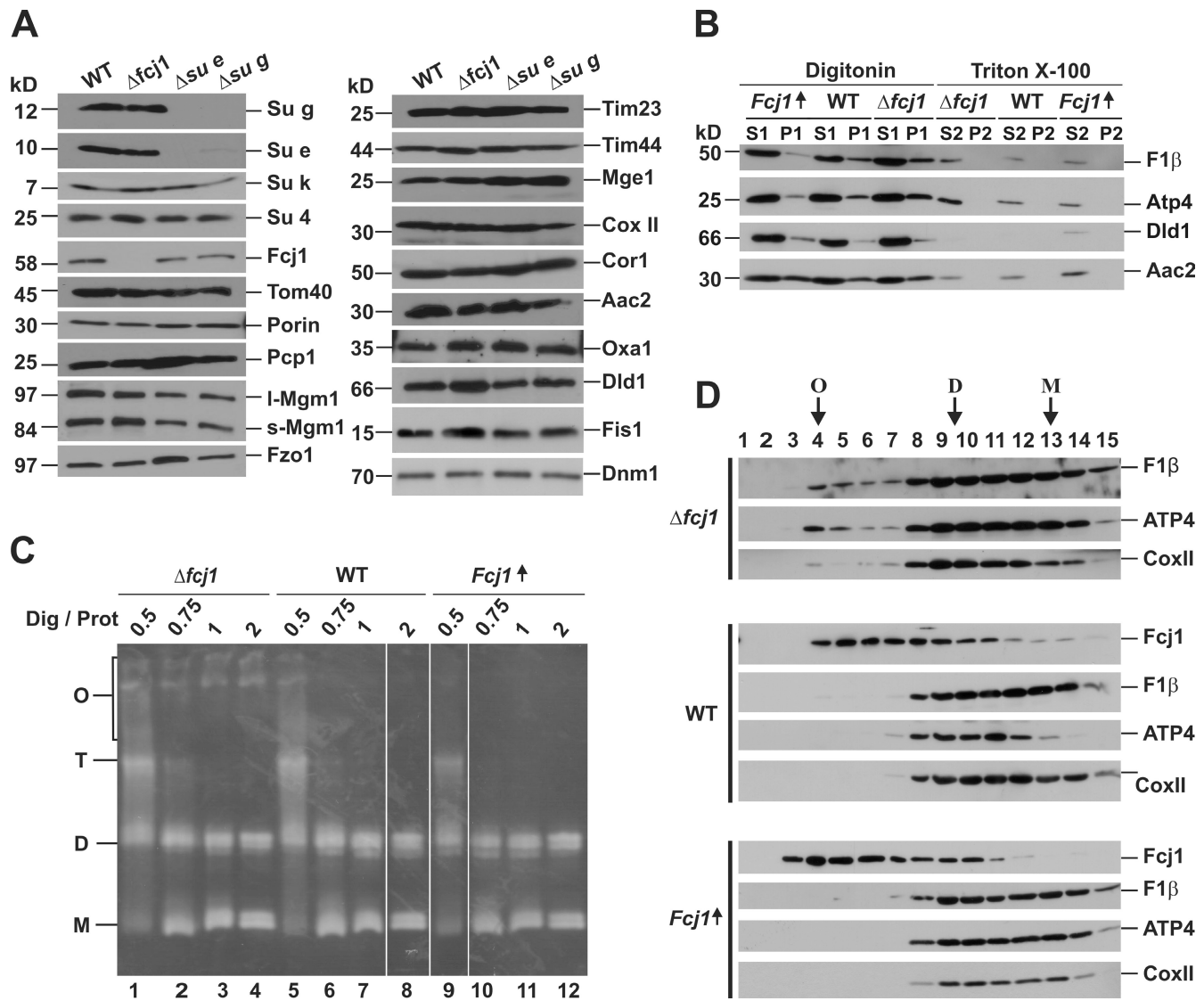


Figure 7. Fcj1 reduces the stability of F₁F_o-ATP synthase oligomers. (A) Protein levels in wild-type (WT), $\Delta fcj1$, $\Delta su e$, and $\Delta su g$ mitochondria. Aliquots were subjected to SDS-PAGE and decorated with the indicated antibodies. (B) Solubility of F₁F_o-ATP synthase in detergents. Isolated mitochondria of Fcj1-overexpressing (Fcj1 \uparrow), wild-type, and $\Delta fcj1$ strains were solubilized in digitonin and centrifuged to obtain supernatant (S1) and pellet (P1) fractions. Nonsolubilized material (P1) was treated with Triton X-100 and centrifuged to obtain supernatant (S2) and pellet (P2) fractions. Aliquots were analyzed by Western blotting with the indicated antibodies. (C) BN-PAGE analysis. Wild-type, $\Delta fcj1$, and Fcj1-overexpressing mitochondria (400 μ g) were solubilized at the indicated ratios of digitonin/protein (Dig/Prot; wt/wt) and subjected to BN-PAGE and to analysis in-gel of F₁F_o-ATPase activity. All lanes were originated from a single gel at identical image settings, but lanes 8 and 9 were cut and pasted in the appropriate order for clarity. (D) SEC. Wild-type, $\Delta fcj1$, and Fcj1-overexpressing mitochondria were solubilized at a digitonin/protein ratio of 1 (wt/wt) and fractionated by Superose 6 gel filtration chromatography, and aliquots were analyzed by Western blotting. The oligomeric forms of the F₁F_o-ATP synthase (O, oligomers; T, tetramers; D, dimers; M, monomers) are indicated as size markers.

than in cells with overexpressed Fcj1, and showed a high variation (Fig. 8 E and not depicted). In wild-type cells, a slightly lower mean CJ diameter was measured when compared with the wild-type control shown in Fig. 6 B. We attribute this to different growth media (selective minimal vs. complete media) and strain backgrounds (W303 vs. BY4742; Table S2). We also investigated the ultrastructure of mitochondria of the double deletion strains $\Delta fcj1/\Delta su e$ and $\Delta fcj1/\Delta su g$. These mitochondria were characterized by the presence of internal, concentric or vesicle-like CMs and the virtual absence of CJs (Fig. S5, C and D). In conclusion, Su e and Su g, on the one hand, promote the formation of cristae tips, and Fcj1, on the other hand, is necessary for formation

of CJs. These two types of mitochondrial membrane proteins modulate the oligomeric state of F₁F_o in an antagonistic manner. This suggests that the architecture of the cristae with its elements of the cristae endings, connections with the IBM, and perhaps also the planar arrangement of cristae sheets are controlled by the local concentrations of these two types of antagonistic proteins.

Discussion

This study addresses the question of how the complex structure of cristae in mitochondria is determined on a molecular basis. In this study, we report on proteins that, in conjunction with

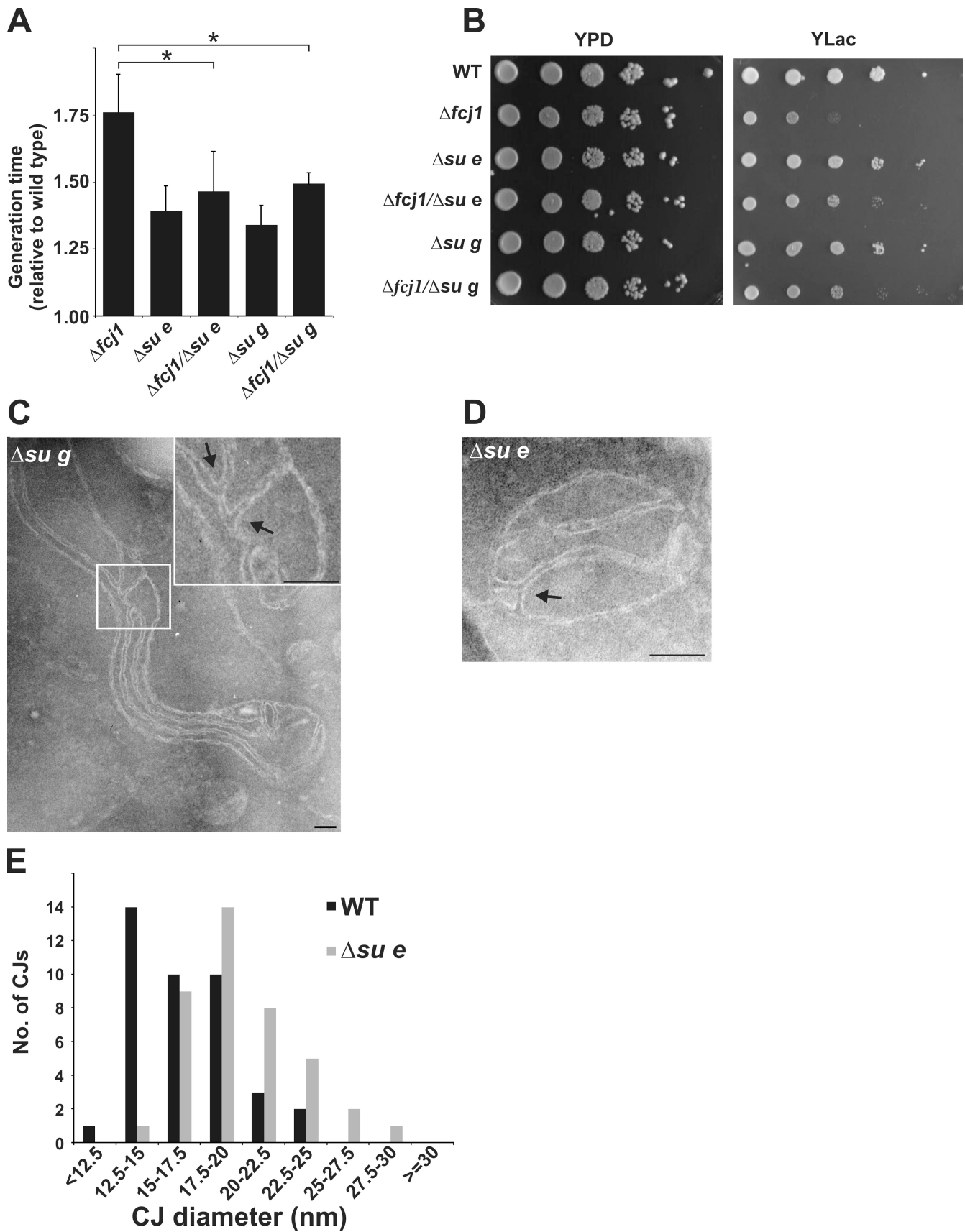


Figure 8. **Functional link between Fcj1 and the F₁F₀-ATP synthase.** (A and B) Genetic interaction of Fcj1 with Su e and Su g. (A) Generation times of the indicated strains (BY4742) grown in complete liquid lactate media during exponential growth at 30°C (n = 4). Error bars show standard deviations. Statistically significant differences (*, P < 0.05) according to a *t* test are indicated. (B) Growth of indicated strains tested by drop dilution in 1:10 steps on

F_1F_0 -ATP synthase, determine cristae architecture. The IM protein Fcj1 plays an essential role in the formation of CJs and cristae shape. *Su e* and *Su g* of F_1F_0 , whose depletion was previously shown to lead to onion-like cristae (Paumard et al., 2002), have an influence on cristae structure, which is opposing that of Fcj1. We present a model of how these components determine the structure of the various parts of cristae, which is based on our results as discussed in this study (Fig. 9, A and B).

Fcj1 is anchored to the IM by a single N-terminal transmembrane segment, exposing the major part of the protein to the intermembrane space. It undergoes homotypic interactions and is quite abundant with an estimated number of 5,730 molecules per cell (Ghaemmaghami et al., 2003). Yet, it is much less abundant than the major constituent of the IM, the F_1F_0 -ATP synthase with an estimated number of 41,500 and 164,000 molecules for the $F_1\alpha$ and $F_1\beta$ subunits, respectively (Ghaemmaghami et al., 2003). The mammalian protein mitofilin shares very low sequence similarity with Fcj1; however, it is likely to represent an orthologue on the basis of its depletion phenotype (John et al., 2005). Fcj1 is present at high levels at the base of CJs but is also present in other parts of cristae, although to a lesser extent. Deletion of Fcj1 leads to a loss of CJs. Overexpression results in an increase of the number of CJs and to internal branching of cristae, suggesting a direct role of Fcj1 in the formation of junctions. Fcj1 has a regulatory influence on the oligomeric state of F_1F_0 . Deletion of Fcj1 leads to an increase in the level of F_1F_0 supercomplexes. This effect is also apparent upon morphological analysis of mitochondria lacking Fcj1. Cryo-EM revealed the formation of extensive F_1F_0 supercomplex structures in the matrix space between cristae of this strain. Consistent with these observations, overexpression of Fcj1 leads to a reduction of the level of F_1F_0 supercomplexes.

Manipulation of the levels of the F_1F_0 *Su e* and *Su g* yielded equally intriguing results. As previously documented, depletion of these subunits leads to accumulation of concentric cristae structures (Paumard et al., 2002). Deletion of either *Su e* or *Su g* has been reported to strongly favor the dissociation of the F_1F_0 into monomers (Arnold et al., 1998). Remarkably, this has no effect on the enzymatic activity of the ATP synthase. In this study, we show that these mitochondria contain CJs but exhibit a virtual absence of cristae tips. This goes along with a strongly increased number of bridgelike cristae that extend across whole sections of mitochondria, ending on both sides in CJs. Moreover, branching of cristae is observed in these mitochondria, much like in Fcj1-overexpressing cells. Thus, branching of cristae may represent internal CJs that emerge within a planar region of the cristae, a region in which CJs are normally absent.

In summary, our data show opposing functions of Fcj1 in relation to F_1F_0 *Su e* and *Su g*. Fcj1 is obviously critical for formation of CJs, whereas formation of cristae tips requires *Su e* and *Su g*. This is in line with the specific enrichment of these

components within the respective regions of the IM, as revealed by quantitative immunogold labeling. It appears to be their relative distribution that determines cristae structure in mitochondria.

Building on our results, we propose a working hypothesis for the molecular basis of the generation of CJs and cristae tips. In the mitochondria of most species and organs, the following structural elements can be discriminated (Fig. 9, A and B). The cristae sheets are made up of two leaves of IM arranged in close apposition, leaving a narrow intermembrane space in between. These sheets are delimited by tips or rims, in which the lipid bilayer is bending over, showing a strong positive curvature. At their base, the cristae are connected to the IBM, which forms a kind of second envelope tightly apposed to the OM. These connections, the CJs, have a rather complex 3D architecture. At the base where they open out into the IBM, the membrane exhibits negative curvature. It is followed by a narrow tubular neck region with highly positive curvature. This structure widens out into the planes of the cristae, a region in which both negative and positive curvatures are present.

According to our model, Fcj1 directly or indirectly interferes with the formation of higher oligomers and thereby favors negative membrane bending. By this mechanism, the presence of Fcj1 affects the building of the bases of CJs. However, in the regions where the formation of higher oligomeric forms of F_1F_0 is not disturbed, these oligomers impose a positive curvature on the membrane and thereby favor the formation of cristae tips. The idea of F_1F_0 oligomers introducing a positive curvature to the IM and leading to tip formation has also been put forward in a recent study (Strauss et al., 2008). Our findings attribute this effect to the action of *Su e* and *Su g*. As we show in this study, they are both enriched in the rim region, and their absence leads to a virtual absence of cristae tips. However, their lower abundance in CJs supports negative curvature. Interestingly, Fcj1 is present in very low amounts in cristae necks, the IM region connecting the base of CJ with the cristae sheets. This membrane has an extremely high positive curvature. Thus, we suggest that the ratios of Fcj1 to *Su e*/*Su g* are decisive in shaping cristae. The sheets of cristae appear to contain both Fcj1 and *Su e*/*Su g*, and this may determine their planar shape. A balanced distribution of these antagonistic components may also be necessary for the dynamics and fission/fusion of cristae. A dynamic nature of cristae was proposed previously (for reviews see Mannella et al., 2001; Mannella, 2006b) based on EM tomography studies of isolated mitochondria showing that cristae morphology apparently is reversibly changing under various physiological conditions (for reviews see Mannella et al., 2001; Mannella, 2006b; Zick et al., 2009).

Altogether, our hypothesis can explain all of the findings presented in this study and in other studies (Paumard et al., 2002; Strauss et al., 2008). It does not exclude that other components such as prohibitins or OPA1 or others yet to be identified also contribute to CJ and cristae tip formation (for reviews

fermentable (yeast peptone dextrose [YPD]) and respiratory (lactate medium [YLac]) carbon sources. (C and D) Electron micrographs of mitochondria of chemically fixed $\Delta su g$ (C) and $\Delta su e$ (D) cells. Branches are indicated by arrows. Enlargement of the indicated box is shown as an inset in C. (E) Diameters of CJs in electron micrographs of mitochondrial sections of wild-type (WT; $m = 40$) and $\Delta su e$ strains ($m = 40$; $m =$ number of mitochondrial sections). Histograms of the number of observed diameters within the indicated ranges are shown. Bars, 100 nm.

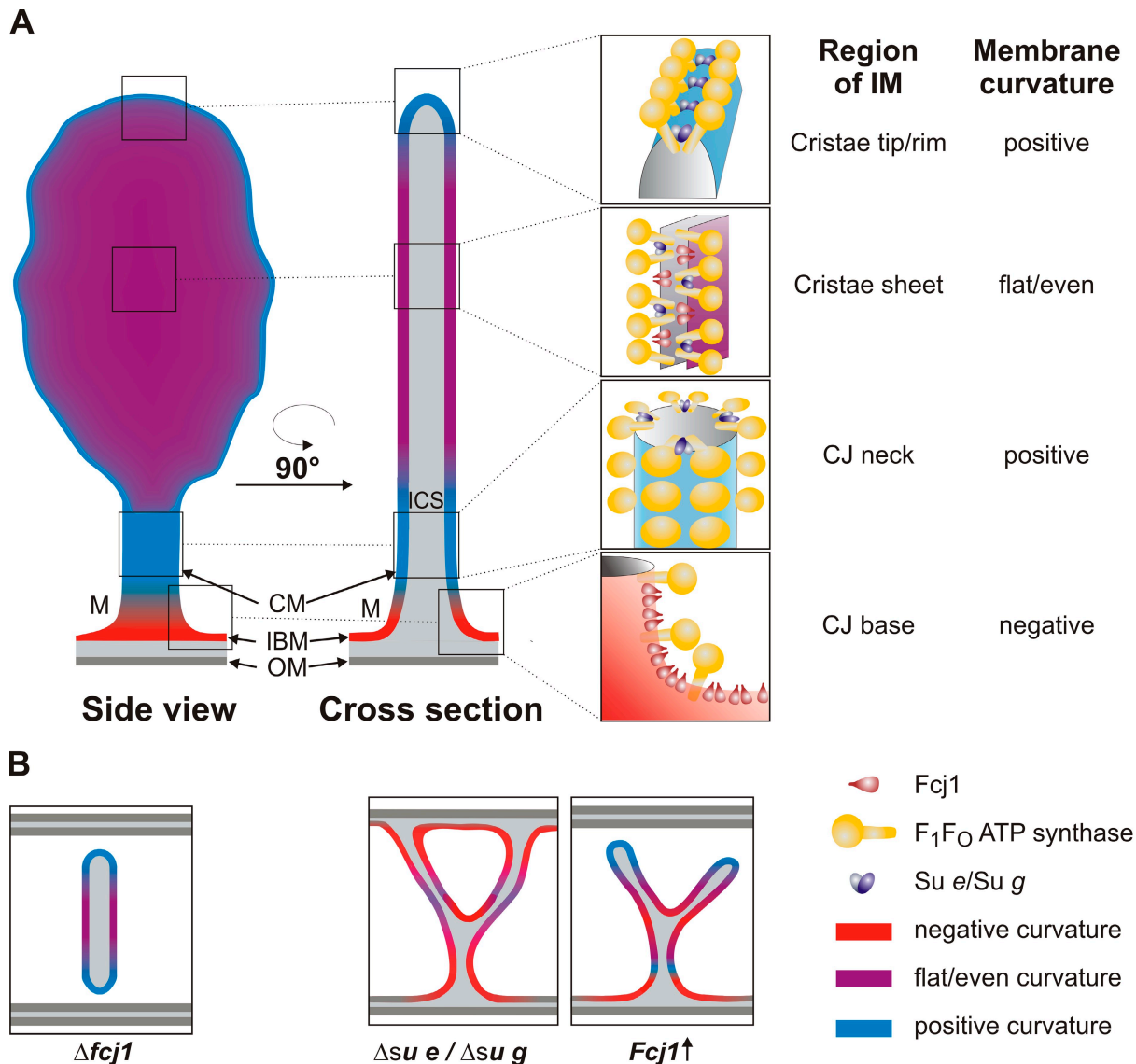


Figure 9. Working model of the formation of cristae and CJs in mitochondria. (A) Schematic representation of membrane curvatures at distinct regions of the CM and of the submitochondrial localization of Fc1 and Su e and Su g in wild-type mitochondria. Positive membrane curvature is indicated in blue, negative curvature in red, and regions with both or no apparent curvature are colored in purple. An overview of a representative cristae sheet with one CJ is shown in side view (left) or as a cross section after 90° rotation (middle). Enlargements of boxed areas in 3D view show the proposed arrangement of Fc1, Su e/Su g, and of F₁F₀ in the respective regions of the CM and their influence on membrane curvature. (B) Schematic representation of membrane curvature in CM structures observed in the indicated mutant mitochondria. Upon depletion of Fc1, positive curvatures of the CM are predominating, whereas membrane structures that require negative bending of the membrane are absent. The lack of Su e/Su g leads to an apparent reduction in positive membrane curvatures, which results in an enlargement of CJ diameter, branching of cristae, and a strong reduction in the number of apparent cristae tips. Increased levels of Fc1 produce a similar phenotype. However, the formation of cristae tips is not affected in this mutant. ICS, intracristal space; M, matrix space.

see Merkwirth and Langer, 2009; Zick et al., 2009). Also, it raises several intriguing questions such as how Fc1 is recruited to the base of the CJ. Fc1 itself or a possible reaction partner may interact with the OM and thereby anchor CJs to the OM. Of particular interest are the physical details of Fc1 and Su e/Su g interaction with F₁F₀ and how these interactions regulate membrane curvature. Finally, an intriguing aspect is the obvious problem of how cristae stacks are formed when CJs are lacking and how protein import and complex assembly in the CM occurs in this situation. One possibility would be that there is a rapid fusion/fission mechanism of cristae to and from the IBM. The life time of these connections might be too short lived to be

seen by EM but long enough for diffusion of newly inserted proteins and lipids. Several other explanations may seem equally possible. In any case, it will be a great challenge to find out how mitochondria can survive the virtual complete absence of CJs and still maintain mitochondrial function and inheritance.

Materials and methods

Strains and growth conditions

The strains and plasmids used are listed in Tables S2 and S3. Culturing of yeast strains was performed using standard methods (Sherman et al., 1986) at 30°C on complete liquid media containing 2% (vol/vol) lactate. Strains containing plasmids (Table S2) were grown on selective liquid

media containing 2% (vol/vol) lactate supplemented with 0.1% (wt/vol) glucose. For overexpression of Fcj1, the wild type and the $\Delta f c j 1$ strain containing the plasmid pCM189-Fcj1 were grown in the absence of doxycycline. For down-regulation of Fcj1, the same plasmid was used in a $\Delta f c j 1$ strain, and 20 $\mu\text{g}/\text{ml}$ doxycycline was added to the medium. Drop dilution growth tests were performed with 1:10 dilution steps and incubation on yeast peptone dextrose and lactate medium plates for 2–4 d at 24°C. $\text{Rho}^0/\text{rho}^-$ cell formation was determined on complete glycerol plates supplemented with 0.1% (wt/vol) glucose.

Fractionation of yeast cells

Yeast total cell extracts were prepared by alkaline lysis as previously described (Herlan et al., 2003). Cellular subfractionation of yeast cells was performed as previously described (Rowley et al., 1994). To extract peripherally bound membrane proteins, 10 mg/ml of mitochondria was swollen by dilution to a final concentration of 1 mg/ml in 20 mM Hepes/KOH, pH 7.4. After the addition of an equal volume of freshly prepared 0.2 M Na carbonate solution or 2 M NaCl solution, samples were incubated for 30 min at 4°C and centrifuged at 92,000 g for 30 min at 4°C. Equal fractions of membrane-associated and soluble proteins were analyzed by SDS-PAGE and immunoblotting. Spheroplasting of yeast cells, preparation of mitochondria, generation of mitoplasts, and PK treatment were performed according to standard procedures (Sirrenberg et al., 1996).

Recombinant DNA techniques

The yeast strain $\Delta f c j 1$ (W303 background) was obtained by genomic integration of the PCR product synthesized from the plasmid pFA6aHIS3Mx6 (Wach et al., 1997) using the primers Fcj1-HIS3 upstream and Fcj1-HIS3 reverse. The deletion of Fcj1 was verified by PCR using the primers Fcj1 upstream and Fcj1 downstream. pCM189-Fcj1 was generated by amplifying Fcj1 (YKRO16w) by PCR from yeast genomic DNA using *Pyrococcus furiosus* polymerase (Agilent Technologies) and Fcj1 forward and overexpression Fcj1 reverse primers (Table S4) and cloning into the pCM189 yeast overexpression vector using BamHI–NotI restriction sites. For the generation of pYX242–Fcj1-His₆ or pYX242–Fcj1-His₁₂, the primers Fcj1 upstream and Fcj1-His₆ reverse or Fcj1 upstream and Fcj1-His₁₂ reverse, respectively, were used, and the PCR products were cloned into pYX242 using BamHI–XhoI restriction sites. Constructs or empty plasmids (Table S3) were transformed into the wild-type or the $\Delta f c j 1$ strain to allow growth of all strains in the same selective media.

Antibody production

Antisera against Fcj1 were raised in rabbits against a synthetic peptide corresponding to amino acid residues 214–226 of Fcj1 (CNTQYEN-SKREFEKN) coupled to activated ovalbumin (Pineda Antikörperservice).

Protein purification

Protein purification by TAP was adapted from a previous study (Puig et al., 2001). In brief, 25 μl of IgG Sepharose 6 Fast Flow (GE Healthcare) bead slurry was washed three times with TBS, once with 0.1 M Gly, pH 2.5, again three times with TBS, and once with solubilization buffer (10 mM Tris/HCl, pH 8.0, 150 mM NaCl, 1% [vol/vol] Triton X-100, and protease inhibitors [complete protease inhibitor cocktail; Roche]), 5 mM EDTA, 1 mM PMSF, 0.5 mM 1,10-phenanthroline, 2 $\mu\text{g}/\text{ml}$ aprotinin, and 1 $\mu\text{g}/\text{ml}$ pepstatin. 1 mg of isolated mitochondria (Fcj1-TAP, Fcj1-His, or Fcj1-TAP–Fcj1-His; Table S2) was solubilized for 30 min on ice in 100 μl of solubilization buffer. The samples were centrifuged for 20 min at 57,000 g, and the supernatants were loaded onto the prepared IgG Sepharose beads. After incubation for 1 h at 4°C, the beads were washed with solubilization buffer containing 0.1% (vol/vol) Triton X-100, and bound proteins were eluted with 2x Laemmli sample buffer for 3 min at 95°C and analyzed by SDS-PAGE and Western blotting.

Purification by nickel nitrilotriacetic acid binding of His-tagged proteins was performed as described previously (Mokranjac et al., 2005). In brief, 3 mg of mitochondria prepared from cells expressing tagged versions of Fcj1 (Fcj1-His₁₂, Fcj1-TAP–Fcj1-His₆, or Fcj1 full length as a control; Table S2) were solubilized in solubilization buffer for 30 min on ice containing 5 mM imidazol without EDTA. After a clarifying spin, the supernatants were incubated for 1 h at 4°C with 30 μl of washed nickel nitrilotriacetic acid bead slurry (EMD). The beads were washed with solubilization buffer containing 0.1% (vol/vol) Triton X-100 and 30 mM imidazol. Bound proteins were eluted with Laemmli sample buffer at 95°C or with 300 mM imidazol at 4°C depending on the subsequent treatment of the samples. Bound material was analyzed either by Coomassie staining or subjected to SEC.

Analysis of bioenergetic parameters of mitochondria

Wild-type and $\Delta f c j 1$ cells were grown on a nonfermentable carbon source (see Strains and growth conditions), and enzymatic activities of oxidative phosphorylation of isolated mitochondria were assayed essentially as described previously (Velours et al., 2001). Oxygen consumption rates were measured with an Oxygraph (Hansatech Instruments) and analyzed by Oxyg32 version 2.25 software (Hansatech Instruments). 500 μg of freshly isolated mitochondria was resuspended in 2 ml of respiration buffer (250 mM sucrose, 5 mM KPi, pH 7.4, 1 mM EDTA, 6 mM MgCl₂, and 20 mM Hepes/KOH, pH 7.4). After adding 4 mM NADH, state IV respiration was measured. Further addition of 200 μM ADP allowed determination of state III respiration. To uncouple respiration from ATP synthesis, 3 μM carbonyl cyanide *m*-chlorophenylhydrazone was added, leading to maximal respiration rate. ATPase activity was measured with or without the addition of oligomycin to a final concentration of 10 $\mu\text{g}/\text{ml}$, as previously described (Velours et al., 2001). The membrane potential ($\Delta\psi$) was evaluated by measurement of fluorescence quenching of rhodamine123 as previously described (Emaus et al., 1986).

Analysis of oligomeric F₁F₀-ATP synthase complexes

To determine the solubility of F₁F₀, isolated mitochondria were incubated on ice for 30 min in digitonin solution at a digitonin/protein (wt/wt) ratio of 1 (150 mM NaCl, 20 mM Tris/HCl, pH 8, 5 mM EDTA, 1 mM PMSF, complete protease inhibitor cocktail, 0.5 mM 1,10-phenanthroline, 100 $\mu\text{g}/\text{ml}$ α -macroglobulin, 2 $\mu\text{g}/\text{ml}$ aprotinin, and 1 $\mu\text{g}/\text{ml}$ pepstatin). After centrifugation at 4°C for 10 min at 13,000 g, nonsolubilized material was resolubilized with 1% (vol/vol) Triton X-100.

SEC

Isolated mitochondria were solubilized for 30 min on ice at a digitonin/protein ratio of 1 (wt/wt) or at a Triton X-100/protein ratio of 0.5 in 30 mM Hepes/HCl, pH 7.4, and 150 mM potassium acetate including protease inhibitors as indicated in the previous section and centrifuged for 10 min at 13,000 g before the cleared lysate was loaded on a Superose 6 size exclusion column (GE Healthcare; Meier et al., 2005).

BN-PAGE

500 μg of mitochondria was incubated for 30 min at 4°C in 50 μl of digitonin buffer (30 mM Hepes/HCl, pH 7.4, 150 mM potassium acetate, 10% glycerol, 1 mM EDTA, 2 mM *ε*-amino caproic acid, and 1 mM PMSF) at the indicated ratios of digitonin/protein (wt/wt) and centrifuged at 4°C for 20 min at 57,000 g. Analysis of supernatants by BN-PAGE and in-gel ATPase activity was performed essentially as previously described (Schagger, 2001; Bornhövd et al., 2006).

Fluorescence microscopy

For the visualization of mitochondria, yeast strains were transformed with plasmid pVT100U–mitochondrial GFP–expressing mitochondrial matrix-targeted GFP (Westermann and Neupert, 2000) and grown on selective lactate medium lacking Leu. Confocal images were taken with a confocal microscope (LSM 510; Carl Zeiss, Inc.) equipped with a 63x objective and a differential interference contrast (DIC) setting. For imaging, living cells were embedded in 1% low melting point agarose and observed at RT. 512 × 512-pixel images at 1- μm focal increments were recorded, simultaneously imaging mitochondrially targeted GFP and DIC. Images were processed with an LSM image browser (Carl Zeiss, Inc.). For analysis of mitochondrial volume, means of at least 10 cells per yeast strain and five focal increments per cell were scored for the ratio of mitochondrial GFP within the cell (DIC picture).

EM and electron tomography

EM on chemically fixed, cryosectioned yeast cells using the Tokuyasu method (Tokuyasu, 1989) and quantitative immuno-EM were performed as previously described (Vogel et al., 2006). Cells were immunogold labeled using the indicated antibodies and goat anti-rabbit IgG conjugated to 10-nm gold particles (Dianova). The location of gold particles in mitochondria with clearly resolvable CMs connected by CJs to the IBM was plotted onto a single empirically determined model. To quantitatively determine the relative density of a protein along the IM, the number of gold particles within a 14-nm distance for all points on the IM was determined by moving in silico along the IM in the model from the bottom to the top of the model. CJ diameters were measured in electron micrographs of whole yeast cells prepared by the Tokuyasu method (Tokuyasu, 1989). For this, the distances from center to center of IMs of a CJ opening at the top were measured in various CJs for each yeast strain.

Cryopreparation of isolated mitochondria

Cryopreparation was performed essentially as previously described (Dubochet et al., 1988). Freshly prepared mitochondria were resuspended in isotonic Hepes buffer (10 mM Hepes, pH 7.4, 1 mM EDTA, and 600 mM sorbitol) at a protein concentration of 10 mg/ml. 3–5 μ l of this mitochondrial suspension was applied onto an EM grid (R2/2; Quantifoil; or custom made) and mounted in a plunger equipped with a custom-made humidifying device (Cyrklaff et al., 1990). After adding 3–5 μ l of fiducial marker solution (3-nm protein A–gold [Sigma-Aldrich] in isotonic Hepes buffer), excess liquid was removed, and the grids were rapidly frozen in liquid ethane slush and stored in liquid nitrogen.

Cryo-EM of vitreous sections

Sample vitrification, cryosectioning, and imaging were performed as described previously (Al-Amoudi et al., 2004). In brief, spheroplasted yeast cells mixed with fiducial markers in isotonic buffer were soaked in 20% dextran (~40 kD), sucked in 300- μ m diameter copper tubes and high pressure frozen at >2,000 bars using EMPACT1 (Leica). Vitreous samples were cryosectioned in the microtome (Ultracut UCT; Leica) with nominal thickness of sections of 120 nm and a final thickness of ~200 nm as the result of specimen compression using either a 35° cryodiamond knife (Diatome) or a 45° glass knife with a clearance angle of 6°.

Electron tomography

Cryohydrated samples were mounted in a 70° tilt cryospecimen holder (model 626; Gatan GmbH) and examined in a cryoelectron microscope (CM 300; Philips) equipped with a field emission gun, a postcolumn energy filter (GIF 2002; Gatan GmbH), and a slow-scan charge-coupled device camera (Gatan GmbH) with 2,048 \times 2,048 pixels. Low electron-dose series (4,000–5,000 electrons/nm²) of typically 60–70 images were recorded using the Digital Micrograph package (Gatan GmbH) in tilt ranges between $\pm 60^\circ$ and $\pm 70^\circ$, with 2° tilt intervals, at nominal magnifications of 43,000 (0.82 nm/pixel) or 52,000 (0.68 nm/pixel), and with an objective lens defocus of 6–10 μ m. Chemically preserved material was imaged under similar conditions, but the areas selected for recording of tomographic tilt series were preirradiated with electron doses of >10,000 electrons/nm². Images of tilt series were aligned using fiducial markers and merged in 3D reconstructions by weighted back-projection using the EM image processing package (Hegerl, 1996) as well as the translocase of the mitochondrial OM package (Nickell et al., 2005) for postprocessing the volumes. The final thickness of EM tomograms was between 300 and 500 nm. The AMIRA visualization package (Mercury Computer Systems) was used for surface rendering of membranes and molecules in mitochondria in the original reconstructions as well as for displaying the results in figures.

Online supplemental material

Fig. S1 shows that *FCJ1* (*AIM28/FMP13/YKRO16W*) has very low sequence similarity to mitofilin. Fig. S2 shows that oligomeric protein complexes containing Fcj1 are the result of Fcj1 self-interaction. Fig. S3 shows the specificity of Fcj1-directed antibodies and immuno-EM in Δ *fcj1* cells. Fig. S4 shows that the deletion of Fcj1 leads to altered cristae morphology. Fig. S5 shows the analysis of double deletion strains Δ *fcj1*/ Δ *su e* and Δ *fcj1*/ Δ *su g* and of a strain overexpressing Fcj1. Videos 1 and 2 show EM tomograms of chemically fixed wild-type and Δ *fcj1* *S. cerevisiae* cells, respectively. Video 3 shows a cryo-EM tomogram of isolated wild type *S. cerevisiae* mitochondrion. Videos 4 and 5 show cryo-EM tomograms of isolated Δ *fcj1* *S. cerevisiae* mitochondrion. Table S1 shows the quantitative analysis of wild-type and Δ *fcj1* cells. Tables S2–S4 show the strains, plasmids, and oligonucleotides, respectively, used in this study. Online supplemental material is available at <http://www.jcb.org/cgi/content/full/jcb.200811099/DC1>.

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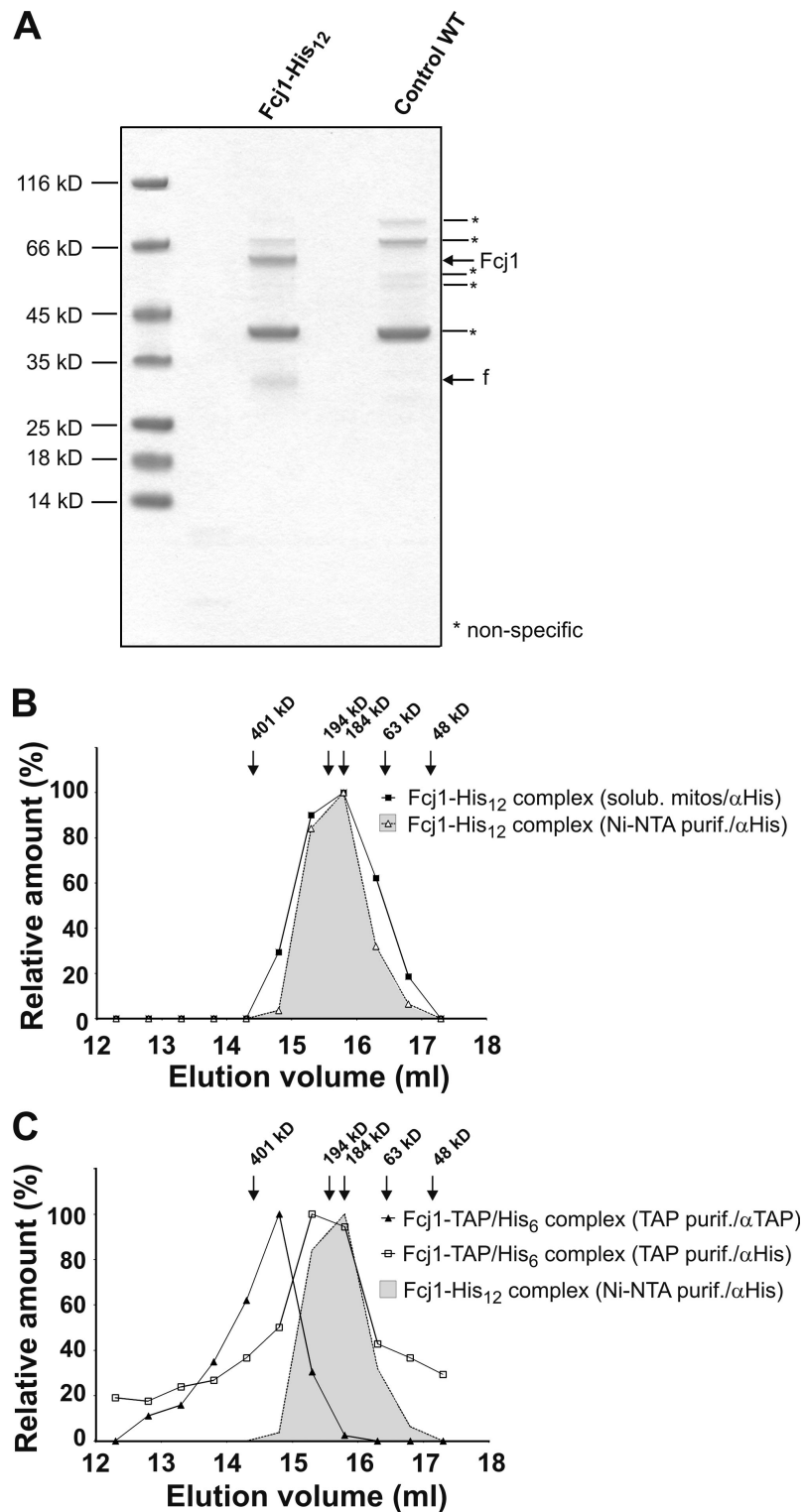


Figure S2. **Oligomeric protein complexes containing Fcj1 are caused by Fcj1 self-interaction.** (A) Nickel nitrilotriacetic acid pull-down of Fcj1 from solubilized mitochondria. A His-tagged variant of Fcj1 (Fcj1-His₁₂) and wild-type (WT) control were subjected to nickel nitrilotriacetic acid affinity purification. Bound material was analyzed by SDS-PAGE and Coomassie staining. Protein bands corresponding to Fcj1 and a degradation fragment of Fcj1 (f) were identified by immunoblotting (not depicted). Nonspecific protein bands are indicated by an asterisk. (B and C) Homo-oligomeric nature of the Fcj1 complex. (B) Mitochondria from cells expressing Fcj1-His₁₂ were isolated and either directly subjected to SEC on a Superose 6 column (closed squares) or nickel nitrilotriacetic acid (Ni-NTA) pull-down was performed, and the bound material was subjected to SEC (open triangles; gray). (C) Mitochondria from cells expressing both Fcj1 and Fcj1-TAP were solubilized and subjected to TAP affinity purification. Bound material was subjected to SEC. Analysis of SEC fractions was performed by SDS-PAGE and Western blotting with the indicated antibodies both against the His-tagged (open squares) and the TAP-tagged (closed triangles) versions of Fcj1. Bands were quantified by densitometry, setting the peak fraction to 100%. For comparison, the size distribution of the Fcj1 complex determined in B is included as a reference (gray). The peak fractions of protein size markers are indicated by arrows.

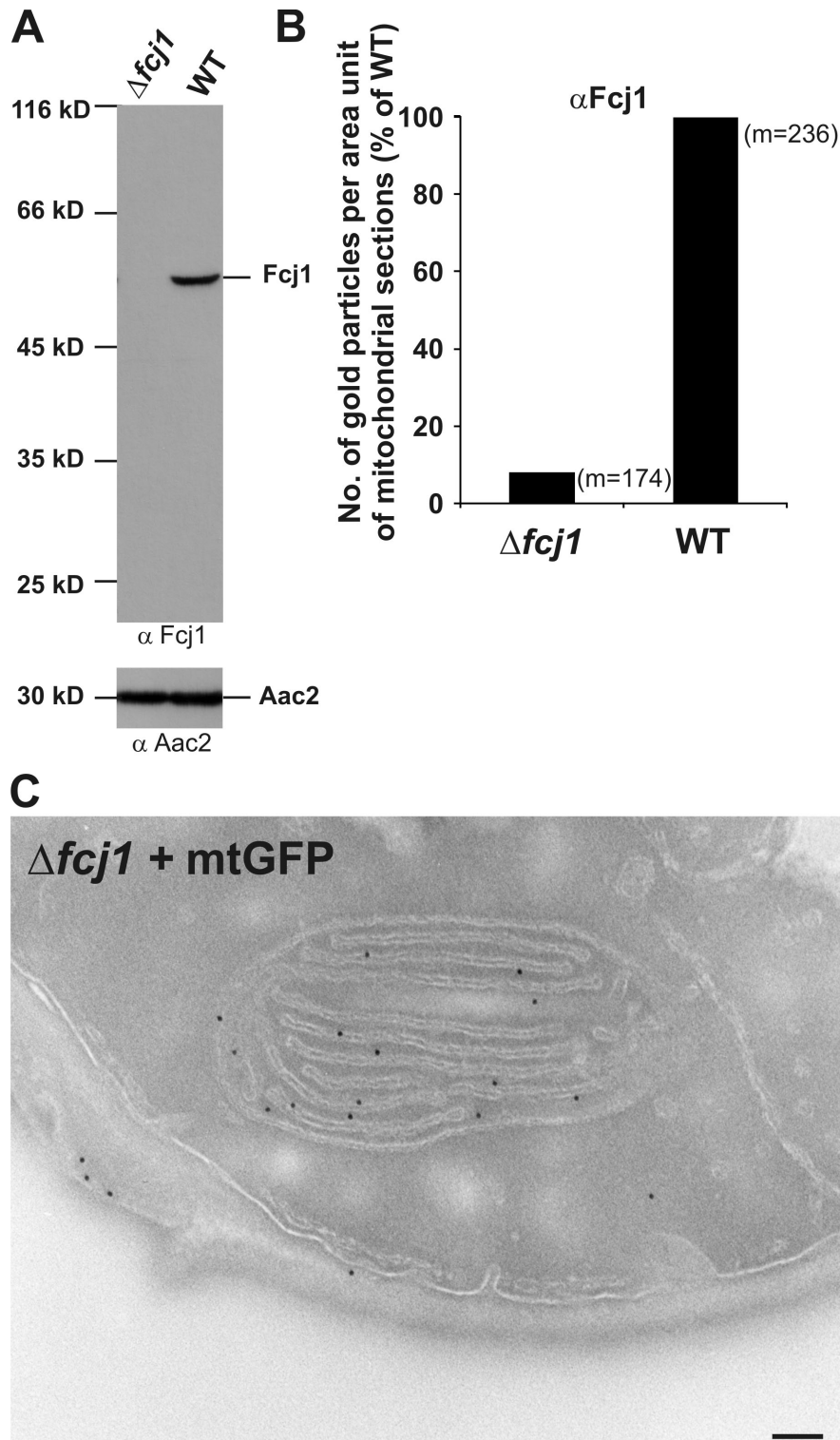


Figure S3. **Specificity of Fcj1-directed antibodies and immuno-EM in $\Delta fcj1$ cells.** (A) Western blotting. Equal amounts of wild-type (WT) and $\Delta fcj1$ mitochondria (30 μ g) were analyzed by Western blotting with antibodies against Fcj1 and Aac2. (B) Quantification of immuno-EM. After immunogold labeling of Fcj1, wild-type and $\Delta fcj1$ cells were screened for the presence of gold particles. The number of gold particles per mitochondrial area unit in wild-type sections was set as 100%. m, number of mitochondrial sections. (C) Matrix-targeted GFP (mtFGP) was expressed in $\Delta fcj1$ cells and labeled with immunogold using antibodies against GFP to distinguish between the matrix and intermembrane space. The majority of gold particles appears outside and between cristae stacks, indicating that these structures enclose the intracristal space rather than the matrix space. Bar, 100 nm.

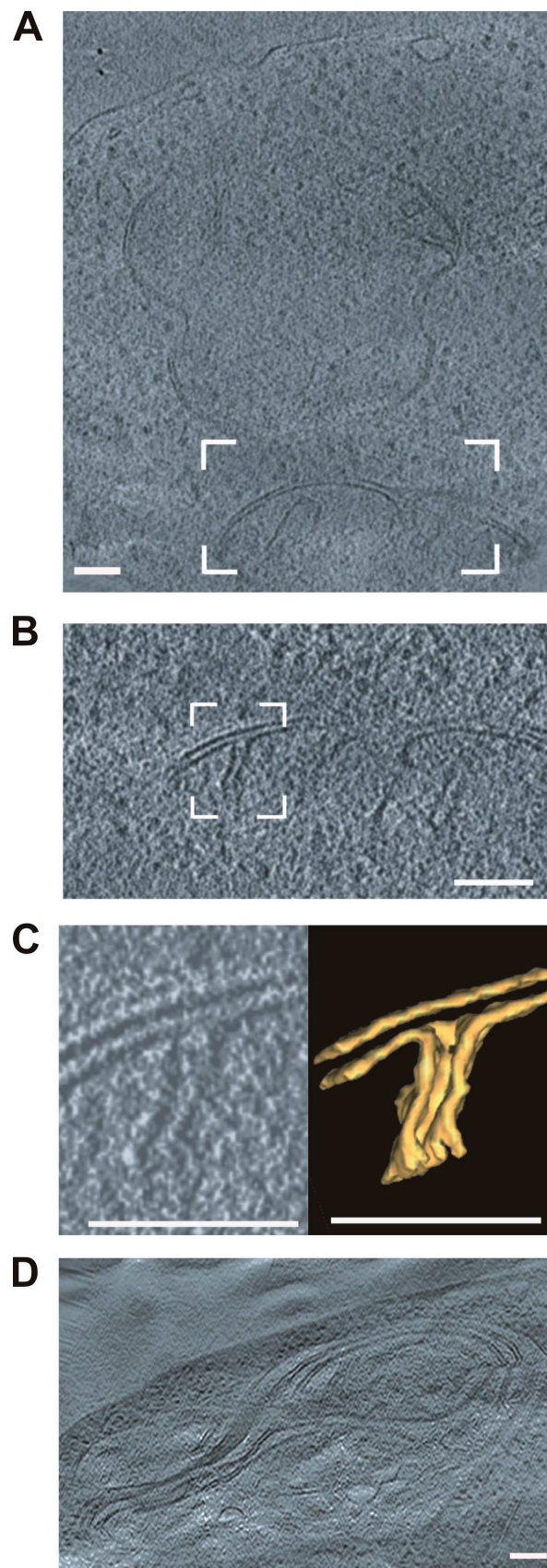


Figure S4. **Deletion of *Fcj1* leads to altered cristae morphology.** (A–D) Cryo-EM tomograms of cryohydrated sections of spheroplasted yeast cells. (A) Wild-type yeast cell with two cryosectioned mitochondria. (B) CJ of wild-type mitochondrion magnified from the boxed area in A. (C) An enlargement of the boxed area in B (left) and a surface-rendered view (right) are shown. (D) $\Delta fcj1$ cell with cryosectioned mitochondrion showing cristae stacks. Bars, 100 nm.

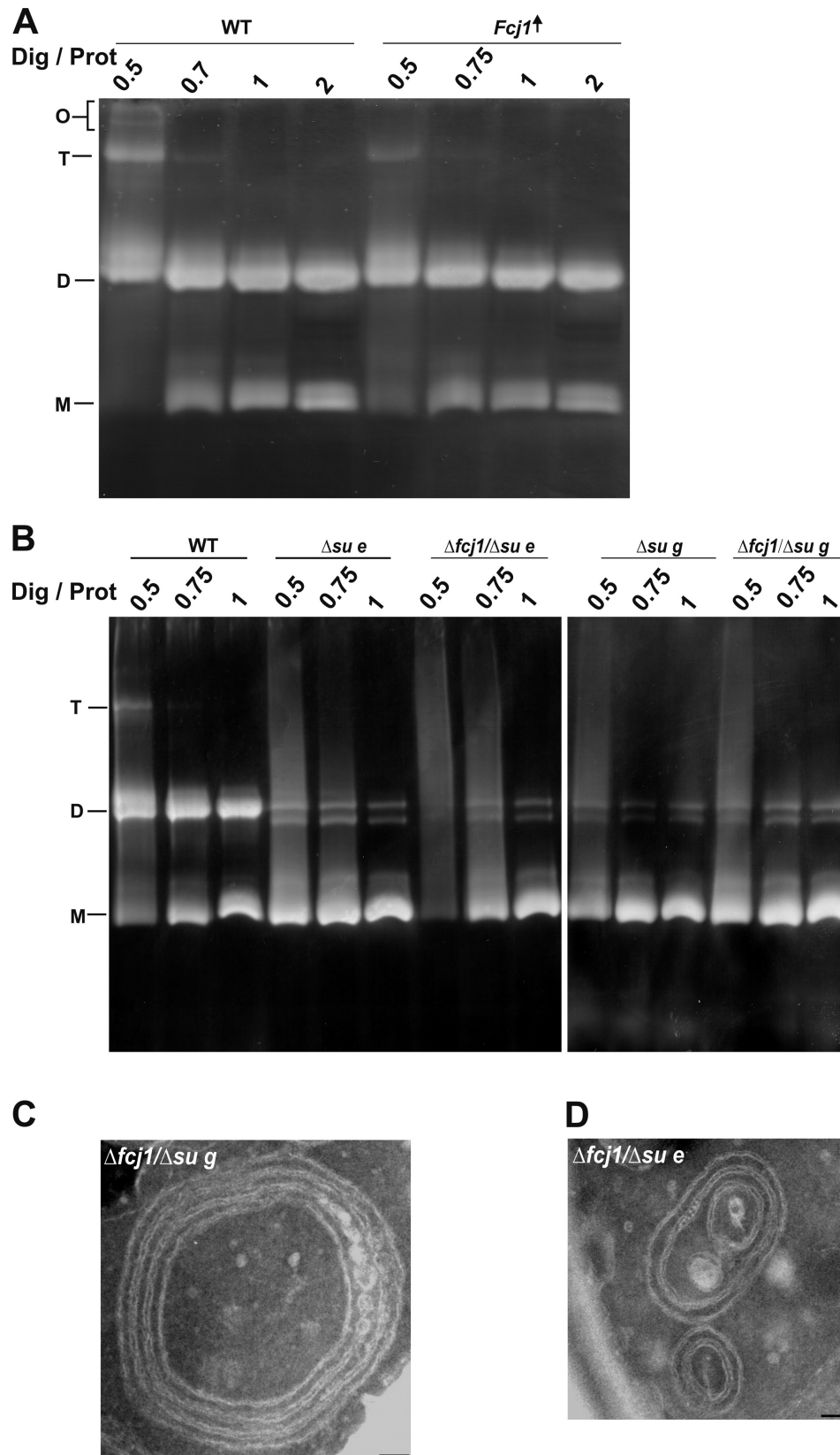


Figure S5. **Analysis of double deletion strains $\Delta fcj1/\Delta su e$ and $\Delta fcj1/\Delta su g$ and of a strain overexpressing *Fcj1*.** (A and B) Equal amounts (400 μ g) of wild-type (WT) and *Fcj1*-overexpressing mitochondria (*Fcj1*[↑]; A) or of wild-type, $\Delta su e$, $\Delta fcj1/\Delta su e$, $\Delta su g$, and $\Delta fcj1/\Delta su g$ mitochondria (B) were solubilized at the indicated ratios of digitonin/protein (Dig/Prot; g/g) and analyzed by BN-PAGE and by in-gel F_1F_0 -ATPase activity. (C and D) Electron micrographs of $\Delta fcj1/\Delta su g$ (C) and $\Delta fcj1/\Delta su e$ (D) mitochondria in chemically fixed yeast cells. Cells were cultivated on a nonfermentable carbon source before fixation. Bars, 100 nm.

Table S1. **Quantitative analysis of wild-type and $\Delta fcj1$ cells**

Stereological parameter used for comparison	Wild type	$\Delta fcj1$
Ratio of membrane length IM/OM	1.57 ± 0.25 (m = 5)	3.29 ± 0.64 (m = 12)
Ratio of mitochondrial volume/cell volume	0.21 ± 0.06 (m = 16)	0.26 ± 0.08 (m = 17)
Ratio of mitochondrial area per mitochondrion relative to wild type	1 (m = 38)	2.0 (m = 31)

m, number of mitochondria or mitochondrial sections. Electron micrographs of the indicated cells after chemical fixation (Tokuyasu method) were analyzed by standard stereological tools (Griffiths, G. 1993. *Fine Structure Immunocytochemistry*. Springer-Verlag, Berlin. 459 pp.) for membrane length and the mitochondrial area per section. The assessment of mitochondrial volumes was performed by analyzing images obtained after confocal fluorescence microscopy of strains expressing mitochondrial GFP (see Materials and methods). Mean values ± standard deviations are given.

Table S2. **Strains**

Name	Description	Background/genotype	Plasmids	Source
BY4741	WT	BY4741	NA	EUROSCARF
$\Delta fcj1$	$\Delta fcj1$	BY4741	NA	EUROSCARF
$\Delta su e$	$\Delta su e$	BY4742	NA	Open Biosystems
$\Delta su g$	$\Delta su g$	BY4742	NA	Open Biosystems
$\Delta fcj1/\Delta su e$	$\Delta fcj1/\Delta su e$	BY4742	NA	This study
$\Delta fcj1/\Delta su g$	$\Delta fcj1/\Delta su g$	BY4742	NA	This study
<i>Fcj1-TAP</i>	<i>Fcj1-TAP</i>	BY4741	NA	Open Biosystems
<i>Fcj1-His₆</i>	<i>Fcj1-His₆</i>	W303	pYX242- <i>Fcj1-His₆</i>	This study
<i>Fcj1-His₁₂</i>	<i>Fcj1-His₁₂</i>	W303	pYX242- <i>Fcj1-His₁₂</i>	This study
<i>Fcj1-TAP/Fcj1-His</i>	<i>Fcj1-TAP/Fcj1-His₆</i>	BY4741	pYX242- <i>Fcj1-His₆</i>	This study
$\Delta Fcj1 + pCM189-Fcj1$	<i>Fcj1</i> ↓ used for down-regulation experiments	W303	pCM189- <i>Fcj1</i>	This study
WT + pCM189	WT control strain for overexpression/down-regulation experiments	W303	pCM189	This study
WT + pCM189- <i>Fcj1</i>	<i>Fcj1</i> ↑ overexpression experiments	W303	pCM189- <i>Fcj1</i>	This study

EUROSCARF, European *Saccharomyces Cerevisiae* Archive for Functional Analysis; NA, not applicable; WT, wild type.

Table S3. **Plasmids**

Name	Description	Vector	Source
pYX242	Yeast expression vector containing constitutive <i>TPI</i> promoter	NA	EMD
pCM189	Yeast expression vector containing doxycycline-regulatable promoter system	NA	Garí et al., 1997
pVT100U-mtGFP	Expression of mitochondrial matrix-targeted GFP	NA	Westermann and Neupert, 2000
pCM189- <i>Fcj1</i>	Expression of <i>Fcj1</i> under a doxycycline-repressible promoter	pCM189	This study
pYX242- <i>Fcj1-His₆</i>	Expression of C-terminally His-tagged <i>Fcj1</i>	pYX242	This study
pYX242- <i>Fcj1-His₁₂</i>	Expression of C-terminally His-tagged <i>Fcj1</i>	pYX242	This study

mtGFP, mitochondrial GFP; NA, not applicable.

References

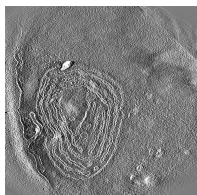
- Garí, E., L. Piedrafitra, M. Aldea, and E. Herrero. 1997. A set of vectors with a tetracycline-regulatable promoter system for modulated gene expression in *Saccharomyces cerevisiae*. *Yeast*. 13:837–848.
- Westermann, B., and W. Neupert. 2000. Mitochondria-targeted green fluorescent proteins: convenient tools for the study of organelle biogenesis in *Saccharomyces cerevisiae*. *Yeast*. 16:1421–1427.

Table S4. **Oligonucleotides**

Name	Sequence
<i>Fcj1</i> forward	5'-CCCCCATGGCAATGCGAAGAACTACTGCCTCACG-3'
Overexpression <i>Fcj1</i> reverse	5'-CCCCGCGCCGCTCACAACGTCTTATTCACAGTCTTC-3'
<i>Fcj1-His₆</i> reverse	5'-CCCCCTCGAGTCAGTGATGGTGATGGTGATGCAACGTCCTTATTCACAGTCTTC-3'
<i>Fcj1-His₁₂</i> reverse	5'-CCCCCTCGAGTCAGTGATGGTGATGGTGATGGTGATGGTGATGCAACGTCCTTATTCACAGTCTTC-3'
<i>Fcj1-HIS3</i> upstream	5'-AAAGGCATAAGAACGCAATTGAAAAGTCTAAAAACTAATATTCGTCGTACGCTGCAGGTGAC-3'
<i>Fcj1-HIS3</i> reverse	5'-TTGAGGTGTAATGACGTACATCTCTTTCTCTTTGATTATCTTATCGATGAATTCGAGCTC-3'
<i>Fcj1</i> upstream	5'-GGTAATTTCTTGCAATTC-3'
<i>Fcj1</i> downstream	5'-TCTTCTTATTAAGCCTCC-3'



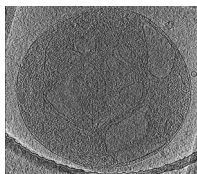
Video 1. **EM tomogram of chemically fixed wild-type *S. cerevisiae* cell.** Successive xy slices through the tomogram are shown moving along the tomographic z axis. The top right mitochondrion was used for surface rendering and is shown in Fig. 3 (A and B). The video is shown at 30 frames/s.



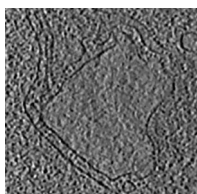
Video 2. **EM tomogram of chemically fixed $\Delta fcj1$ *S. cerevisiae* cell.** Successive xy slices through the tomogram are shown moving along the tomographic z axis. A single slice of mitochondrion was used for surface rendering and is shown in Fig. 3 (C and D). The video is shown at 30 frames/s.



Video 3. **Cryo-EM tomogram of isolated wild-type *S. cerevisiae* mitochondrion.** Successive xy slices through a tomogram represented in Fig. 4 A are shown moving along the tomographic z axis. The video is shown at 30 frames/s.



Video 4. **Cryo-EM tomogram of isolated $\Delta fcj1$ *S. cerevisiae* mitochondrion.** Successive xy slices through a tomogram represented in Fig. 4 C are shown moving along the tomographic z axis. Details are shown in Figs. 4 (D–G) and 5 B. See also Video 5. Zipperlike structures typical of the F_1 parts of F_1F_0 -ATP synthases are visible. The video is shown at 30 frames/s.



Video 5. **Cryo-EM tomogram of isolated $\Delta fcj1$ *S. cerevisiae* mitochondrion.** Successive xy slices through a tomogram represented in Fig. 4 D are shown moving along the tomographic z axis. This represents an enlargement slightly below the boxed area in Fig. 4 C. Zipperlike structures typical of the F_1 parts of F_1F_0 -ATP synthases are visible. The video is shown at 30 frames/s.