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The *Chlamydomonas* Genome Reveals the Evolution of Key Animal and Plant Functions

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Chlamydomonas reinhardtii is a unicellular green alga whose lineage diverged from land plants over 1 billion years ago. It is a model system for studying chloroplast-based photosynthesis, as well as the structure, assembly, and function of eukaryotic flagella (cilia), which were inherited from the common ancestor of plants and animals, but lost in land plants. We sequenced the ~120-megabase nuclear genome of *Chlamydomonas* and performed comparative phylogenomic analyses, identifying genes encoding uncharacterized proteins that are likely associated with the function and biogenesis of chloroplasts or eukaryotic flagella. Analyses of the *Chlamydomonas* genome advance our understanding of the ancestral eukaryotic cell, reveal previously unknown genes associated with photosynthetic and flagellar functions, and establish links between ciliopathy and the composition and function of flagella.

Chlamydomonas reinhardtii is a ~10-μm, unicellular, soil-dwelling green alga with multiple mitochondria, two anterior flagella for motility and mating, and a chloroplast that houses the photosynthetic apparatus and critical metabolic pathways (Fig. 1 and fig. S1) (*1*). *Chlamydomonas* is used to study eukaryotic photosynthesis because, unlike angiosperms (flowering plants), it grows in the dark on an organic carbon source while maintaining a functional photosynthetic apparatus (*2*). It also is a model for elucidating eukaryotic flagella and basal body functions and the pathological effects of their dysfunction (*3*, *4*). More recently, *Chlamydomonas* research has been developed for bioremediation purposes and the generation of biofuels (*5*, *6*).

The Chlorophytes (green algae, including *Chlamydomonas* and *Ostreococcus*) diverged from the Streptophytes (land plants and their close relatives) (Fig. 2) over a billion years ago. These lineages are part of the green plant lineage (Viridiplantae), which previously diverged from opisthokonts (animals, fungi, and Choanozoa) (*7*). Many *Chlamydomonas* genes can be traced to the green plant or plant-animal common ancestor by comparative genomic analyses. Specifically, many *Chlamydomonas* and angiosperm genes are derived from ancestral green plant genes, including those associated with photosynthesis and plastid function; these are also present in *Ostreococcus* spp. and the moss *Physcomitrella patens* (Fig. 2). Genes shared by *Chlamydomonas* and animals are derived from the last plant-animal common ancestor and many of these have been lost in angiosperms, notably those encoding proteins of the eukaryotic flagellum (or cilium) and the associated basal body (or centriole) (*8*). *Chlamydomonas* also displays extensive metabolic flexibility under the control of regulatory genes that allow it to inhabit distinct environmental niches and to survive fluctuations in nutrient availability (*9*).

Genome sequencing and assembly. The 121-megabase (Mb) draft sequence (*10*) of the *Chlamydomonas* nuclear genome was generated at 13× coverage by whole-genome, shotgun end-sequencing of plasmid and fosmid libraries, followed by assembly into ~1500 scaffolds (*1*). Half of the assembled genome is contained in 25 scaffolds, each longer than 1.63 Mb. The genome is unusually GC-rich (64%) (Table 1), which required modification of standard sequencing protocols. Alignments of expressed sequence tags (ESTs) to the genome suggest that the draft assembly is 95% complete (*1*).

The *Chlamydomonas* nuclear genome comprises 17 linkage groups (figs. S2 to S18) presumably corresponding to 17 chromosomes, consistent with electron microscopy of meiotic synaptonemal complexes (*11*). Seventy-four scaffolds, representing 78% of the draft genome, have been aligned with linkage groups (Fig. 3 and figs. S2 to S18). Sequenced ESTs from a field isolate (*1*) of *Chlamydomonas*, fertile with the standard laboratory strain, identified 8775 polymorphisms, result-

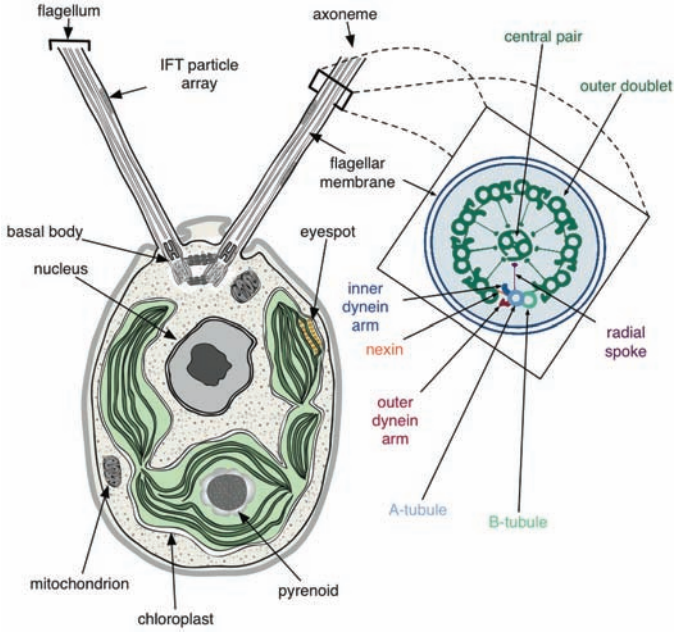

Fig. 1. A schematic of a *Chlamydomonas* cell (from transmission electron micrographs) showing the anterior flagella rooted in basal bodies, with intraflagellar transport (IFT) particle arrays between the axoneme and flagellar membrane, the basal cup-shaped chloroplast, central nucleus and other organelles. An expanded cross section of the flagellar axoneme, as redrawn from (*48*), shows the nine outer doublets and the central pair (9+2) microtubules; axoneme substructures are color-coded and labeled (see inset).

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ing in a marker density of 1 per 13 kb (*12*, *13*). By comparing physical marker locations on scaffolds with genetic recombination distances, we estimated 100 kb per centimorgan (cM) on average.

The *Chlamydomonas* genome has approximately uniform densities of genes, simple sequence repeats, and transposable elements. Several AT-rich islands coincide with gene- and transposable element–poor regions (figs. S2 to S18). As in most eukaryotes, the ribosomal RNA (rRNA) genes are arranged in tandem arrays. They are located on linkage groups I, VII, and XV, although assembly has only been completed on the outermost copies. We identified 259 transfer RNAs (tRNAs) (*1*) (table S1), 61 classes of simple repeats, ~100 families of transposable elements (*1*), and 64 tRNA-related short interspersed elements (SINEs) (tables S2 and S3), which is unusual for a microorganism. We also identified tRNAs clusters and a number of recent tRNA duplications (fig. S19), as well as clusters of genes associated with specific biological functions (fig. S20). Few chloroplast and mitochondrial genome fragments were detected in the nuclear genome ("cp" and "mito" in Fig. 3, and figs. S2 to S18).

Protein coding genes and structure. Ab initio and homology-based gene prediction, integrated with EST evidence, was used to create a reference set of 15,143 protein-coding gene predictions (*1*) (tables S4, S5, and S6). More than 300,000 ESTs were generated from diverse environmental conditions; 8631 gene models (56%) are supported by mRNA or EST evidence (*14*), and 35% have been edited for gene structure and/or annotated by manual curation, as of June 2007. Protein-coding genes have, on average, 8.3 exons per gene and are intron-rich relative to other unicellular eukaryotes and land plants (*15*) (fig. S21); only 8% lack introns (Table 1) (*1*). The average *Chlamydomonas* intron is longer (373 bp) than that of many eukaryotes (*16*), and the average intron number and size are more similar to those of multicellular organisms than those of protists (fig. S21) (*1*, *17*). Only 1.5% of the introns are short (<100 bp), and we did not observe the bimodal intron size distribution typical of most eukaryotes (fig. S21A). Furthermore, 30% of the intron length is due to repeat sequences (*1*), which suggests that *Chlamydomonas* introns are subject to creation or invasion by transposable elements.

Gene families. We identified 1226 gene families in *Chlamydomonas* encoding two or more proteins (*1*); of these, 26 families have 10 or more members (table S7). The genes of 317 of the 798 two-gene families are arranged in tandem, which suggests extensive tandem gene duplications. Gene families contain similar proportions of the total gene complement of *Chlamydomonas*, human, and *Arabidopsis*. As in *Arabidopsis*, *Chlamydomonas* has large families of kinases and cytochrome P-450s, but the largest one is the class III guanylyl and adenylyl cyclase family. With 51 members, the *Chlamydomonas* family is larger than that in any other organism (*18*). Although these cyclases are not found in plants, in animals they catalyze the synthesis of cGMP and cAMP (*18*), which serve as second messengers in various signal transduction pathways. Cyclic nucleotides are critical for mating processes, as well as flagellar function and regulation in *Chlamydomonas* (*19*–*21*), and may be vital for acclimation to changing nutrient conditions (*22*, *23*). *Chlamydomonas* also encodes diverse families of proteins critical for nutrient acquisition (*23*, *24*).

Transporters. The transporter complement in *Chlamydomonas* suggests that it has retained the diversity present in the common plant-animal ancestor. *Chlamydomonas* is predicted to have 486 membrane transporters (figs. S22 and S23) (*1*) that fall into the broad classes of 61 ion channels, 124 primary (active) adenosine triphosphate (ATP)–dependent transporters and 293 secondary transporters; eight are unclassified. The 69-member ATP-binding cassette (ABC) and 26-member P-type adenosine triphosphatase (ATPase) families are large, as in *Arabidopsis*, and overall, the complement of transporters in *Chlamydomonas* resembles that of both *Ostreococcus* spp. and land plants (fig. S22). Furthermore, a number of plant transporters not found in animals are encoded on the *Chlamydomonas* genome (fig. S22 and table S8).

We also found copies of genes encoding animal-associated transporter classes, including some with activities related to flagellar function (e.g., the voltage-gated ion channel superfamily) (*25*) (fig. S22 and table S8). A number of these transporters redistribute intracellular Ca^{2+} in response to environmental signals such as light. Changing Ca^{2+} levels may modulate the activity of the flagella, which are structures found in animals but not in vascular plants (see below).

The *Chlamydomonas* genome also encodes a diversity of substrate-specific transporters that are important for acclimation of the organism to the fluctuating, often nutrient-poor, conditions of soil environments (*24*). Of the eight sulfate transporters, four are in the H^+/SO_4^{2-} family (characteristic of the plant lineage), three are in the Na^+/SO_4^{2-} family (not found in plants but present in opisthokonts), and one is a bacterial ABC-type SO_4^{2-} transporter (associated with the plastid envelope). The 12-

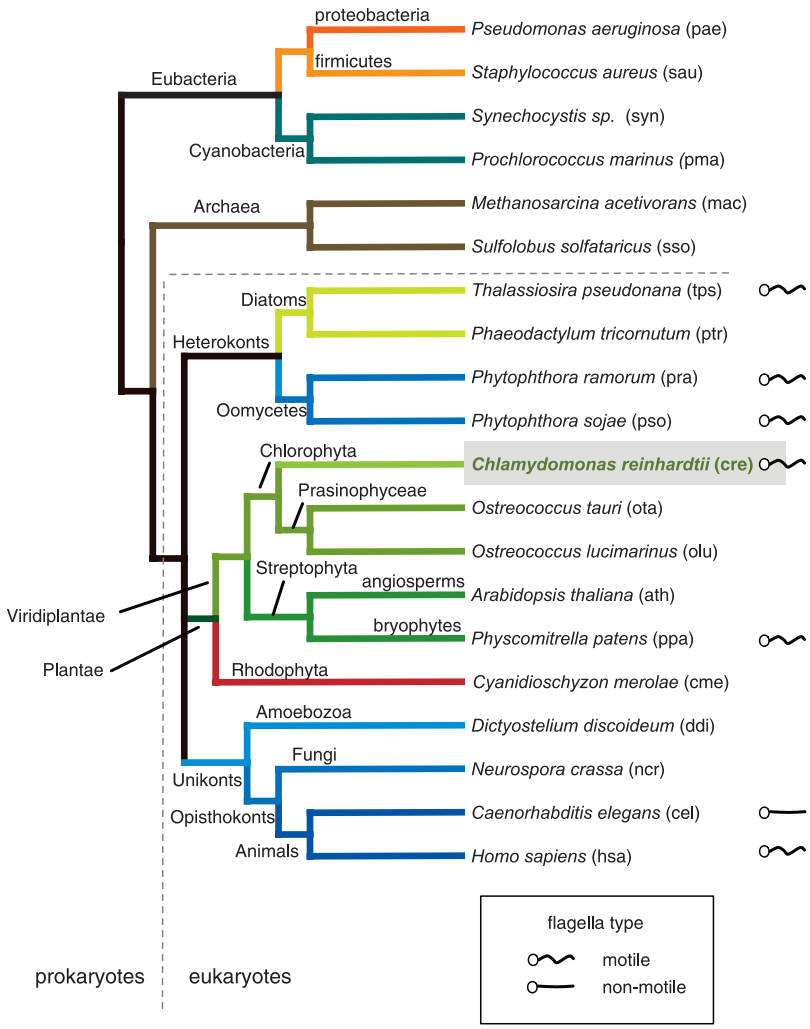


Fig. 2. Evolutionary relationships of 20 species with sequenced genomes (*54*, *55*) used for the comparative analyses in this study include cyanobacteria and nonphotosynthetic eubacteria, Archaea and eukaryotes from the oomycetes, diatoms, rhodophytes, plants, amoebae and opisthokonts. Endosymbiosis of a cyanobacterium by a eukaryotic protist gave rise to the green (green branches) and red (red branches) plant lineages, respectively. The presence of motile or nonmotile flagella is indicated at the right of the cladogram.

member PiT phosphate transporter and 6-member KUP potassium channel families are larger than in other unicellular eukaryotes, and the former underwent a lineage-specific expansion. *Chlamydomonas* has 11 AMT ammonium transporters, which is only surpassed by the number in rice.

Phylogenomics and the origins of *Chlamydomonas* genes. To explore the evolutionary history of *Chlamydomonas*, we initially compared the *Chlamydomonas* proteome to a representative animal (human) and angiosperm (*Arabidopsis*) proteome (*1*). We plotted the best matches, calculated on the basis of BLASTP (Basic Local Alignment Search Tool for searching protein collections) scores, of every *Chlamydomonas* protein to the *Arabidopsis* and human proteomes (Fig. 4A). Most *Chlamydomonas* proteins exhibit slightly more similarity to *Arabidopsis* than to human proteins. Many *Chlamydomonas* proteins with greater similarity to animal homologs are present in the flagellar and basal body proteomes (Fig. 4A and below). This is consistent with the maintenance of flagella and basal bodies as cilia and centrioles, respectively, in animals (*8*), and their loss in angiosperms.

A mutual best-hit analysis of *Chlamydomonas* proteins against proteins from organisms across the tree of life (*1*) identified 6968 protein families of orthologs, co-orthologs (in the case of recent gene duplications), and paralogs (*1*). Of the *Chlamydomonas* proteins, 2489 were homologous to proteins from both *Arabidopsis* and humans (Fig. 4B). *Chlamydomonas* and humans shared 706 protein families (774 and 806 proteins, respectively), but these were not shared with *Arabidopsis*. These genes were either lost or diverged beyond recognition in green plants (table S9), and are enriched for sequences encoding cilia and centriole proteins (*8*, *26*). Conversely, 1879 protein families are found in both *Chlamydomonas* and *Arabidopsis* (1968 and 2396 proteins, respectively), but lack human homologs. *Chlamydomonas* proteins with homology to plant, but not animal, proteins were either (i) present in the common plant-animal ancestor and retained in *Chlamydomonas* and angiosperms, but lost or diverged in animals; (ii) horizontally transferred into *Chlamydomonas*; or (iii) arose in the plant lineage after divergence of animals (but before the divergence of *Chlamydomonas*). This set is enriched for proteins that function in chloroplasts (table S9 and below).

The plastid and plant lineages. The plastids of green plants and red algae are primary plastids, i.e., direct descendants from the primary cyanobacterial endosymbiont (*27*). Diatoms, brown algae, and chlorophyll a– and c–containing algae are also photosynthetic, but their photosynthetic organelles were acquired via a secondary endo-

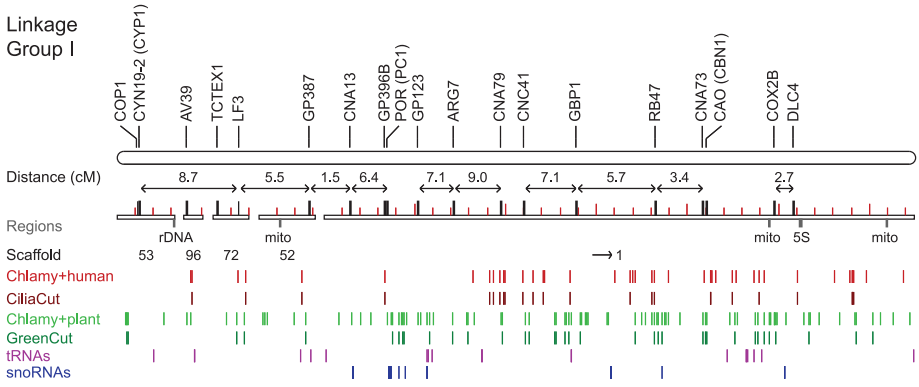


Fig. 3. Linkage group I depicted as a long horizontal rod, with genetically mapped scaffolds shown as open rectangles below (the scaffold number is under each scaffold, and arrows indicate the orientation of the scaffold where it is known; other scaffolds were placed in their most likely orientation on the basis of genetic map distances. The scale of each map is determined by molecular lengths of the mapped scaffolds. Short and long red ticks are drawn on scaffolds every 0.2 Mb and 1.0 Mb, respectively. We assumed small 50 kb gaps between scaffolds. Genetic distances between markers (centimorgans), where they are known, are shown by two-headed arrows above the scaffold, with the gene symbol and any synonyms in parentheses shown at the top. Genomic regions are labeled below the scaffolds: 5*S*, rDNA, mito (insertion of mitochondrial DNA). *Chlamydomonas* genes with homologs in other organisms/lineages ("Cuts" as defined in the text and Fig. 5) are shown as tracks of vertical bars: light red, genes shared between *Chlamydomonas* and humans, but not occurring in nonciliated organisms; dark red, genes in CiliaCut; light green, genes shared between *Chlamydomonas* and *Arabidopsis*, but not in nonphotosynthetic organisms; dark green, genes in GreenCut; magenta, predicted tRNAs, including those that represent SINE sequences; dark blue, small nucleolar RNAs (snoRNAs).

Table 1. Comparison of *Chlamydomonas* genome statistics to those of selected sequenced genomes. nd, Not determined. [Source for all but *Chlamydomonas* (*1*)]

	Chlamydomonas	*Ostreococcus tauri*	*Cyanidioschyzon*	*Arabidopsis*	Human
Assembly length (Mb)	121	12.6	16.5	140.1	2,851
Coverage	13×	6.7×	11×	nd	~8×
Chromosomes	17	20	20	5	23
G+C (%)	64	58	55	36	41
G+C (%) coding sequence	68	59	57	44	52
Gene number	15,143	8,166	5,331	26,341	~23,000
Genes with EST support (%)	63	36	86	60	nd
Gene density (per kb)	0.125	0.648	0.323	0.190	~0.0008
Average bp per gene	4312	nd	1553	2232	27,000
Average bp per transcript	1580	1257	1552	nd	nd
Average number of amino acids per polypeptide	444	387	518	413	491
Average number of exons per gene	8.33	1.57	1.005	5.2	8.8
Average exon length	190	750	1540	251	282*
Genes with introns (%)	92	39	0.5	79	85†
Mean length of intron	373	103	248	164	3,365
Coding sequence (%)	16.7	81.6	44.9	33.0	~1
Number of rDNA units (28*S*/18*S*/5.8*S* + 5*S*)	3 + 3	4 + 4	3 + 3‡	12 + 700	5 + nd
Number tRNAs	259§	nd	30	589	497
Selenocysteine (Sec) tRNAs	1	nd	nd	0	1

*National Center for Biotechnology Information (NIH) NCBI 36 from Ensembl build 38. †[Source (*56*)]. ‡Three regions contain 5*S* rDNA exclusively, and three regions contain 28*S*-18*S*-5.8*S* rDNAs exclusively. §65 tRNAs that were included in SINE elements were removed from the tRNA-scanSE predictions.

symbiosis (*28*, *29*). Because of shared ancestry, nucleus-encoded plastid-localized proteins derived from the cyanobacterial endosymbiont are closely related to each other and to cyanobacterial proteins.

We searched the 6968 families that contain *Chlamydomonas* proteins for those that also contained proteins from *Ostreococcus*, *Arabidopsis* and moss, but that did not contain proteins from nonphotosynthetic organisms. The search identified 349 families, which we named the GreenCut (Fig. 5A, table S10 and table SA); each of these families has a single *Chlamydomonas* protein. On the basis of manual curation of GreenCut proteins of known function (*1*) (table S11), we estimated ~5 to 8% false-positives and ~14% false-negatives (*1*). By comparing GreenCut proteins to those of the red alga *Cyanidioschyzon merolae*, which diverged before the split of green algae from land plants (Fig. 2), we identified the subset of proteins present across the plant kingdom; we named this subset the PlantCut (Fig. 5A, table S10 and table SA). GreenCut protein families that also included representatives from the diatoms *Thalassiosira pseudonana* (*30*) or *Phaeodactylum tricornutum* (*31*) were placed in the DiatomCut (Fig. 5A and table S10 and table SA). Given the phylogenetic position of diatoms and their secondary endosymbiosis-derived plastids, we hypothesize that protein families present in both the PlantCut and DiatomCut should contain only those GreenCut proteins associated with plastid function. This subset is referred to as the PlastidCut (Fig. 5A).

The GreenCut contains proteins of the photosynthetic apparatus, including those involved in plastid and thylakoid membrane biogenesis, photosynthetic electron transport, carbon fixation, antioxidant generation, and a range of other primary metabolic processes (table S11 and table SA). Although light-harvesting chlorophyll-binding proteins are poorly represented (*1*), we identified specialized chlorophyll-binding proteins, as well as a photosynthesis-specific kinase, involved in state transitions. Numerous GreenCut entries are enzymes of plastid-localized metabolic pathways (lipid, amino acid, starch, nucleotide, and pigment biosynthesis) or are unique to plants or highly divergent from animal counterparts. Although tRNA synthetases are conserved between kingdoms, those in the GreenCut represent organellar isoforms that are often targeted to both plastids and mitochondria in plants (*32*). GreenCut proteins that do not function in the plastids tend to be green lineage–specific or highly diverged from animal counterparts. For example, the *Chlamydomonas* GreenCut protein TOM20 (*1*), an outer mitochondrial membrane receptor involved in protein import, evolved convergently from a different ancestral protein in plants than in fungi and animals (*33*).

Of the 214 proteins in the GreenCut without known function, 101 have no motifs or homologies from which function can be inferred, and we can predict only a general function for the others (table S12). Given that 85% of the known proteins in the GreenCut are localized to chloroplasts (table S13), we predict that the set of unknowns contains many novel, conserved proteins that function in chloroplast metabolism and regulation.

The most reducing and oxidizing biological molecules are generated in chloroplasts via the activity of photosystem I and photosystem II, respectively. The flow of electrons through the photosystems causes damage to cellular constituents as a consequence of the accumulation of reactive oxygen species. Therefore, regulation of these molecules is important. Accordingly, plastids house more redox regulators than do mitochondria. Thioredoxins are critical redox-state regulators, and we identified novel thioredoxins in the GreenCut (table S12). These novel thioredoxins have noncanonical active sites or are fused to domains of inferred function (e.g., a vitamin K–binding domain) in plastid metabolism (fig. S1). These findings reveal the potential for identifying unique redox signaling pathways with selectivity and midpoint potentials associated with specific thioredoxin redox sensors (*1*).

Chlamydomonas has a structure called the eyespot (Fig. 1) which can sense light and trigger phototactic responses. The eyespot is composed of several layers of pigment granules, similar to plastoglobules in plants, and thylakoid membrane, which are directly apposed to the chloroplast envelope and a region of the plasma membrane carrying rhodopsin-family photoreceptors. The pigment granules or plastoglobules contain many proteins with unknown function, many of which are present in the GreenCut, and are likely critical to plastid metabolism; these include SOUL domain, AKC (see below), and PLAP (plastid- and lipid-associated protein) protein families (*34*–*36*). SOUL domain proteins of the GreenCut (SOUL4 and SOUL5) have homologs in the *Arabidopsis* plastoglobule proteome (*34*, *35*), and at least one (SOUL3) is associated with the eyespot. The SOUL domain, originally identified in proteins encoded by highly expressed genes in the retina and pineal gland, can bind heme (*37*, *38*). This domain may be important as a heme carrier and/or in maintaining heme in a bound, non-

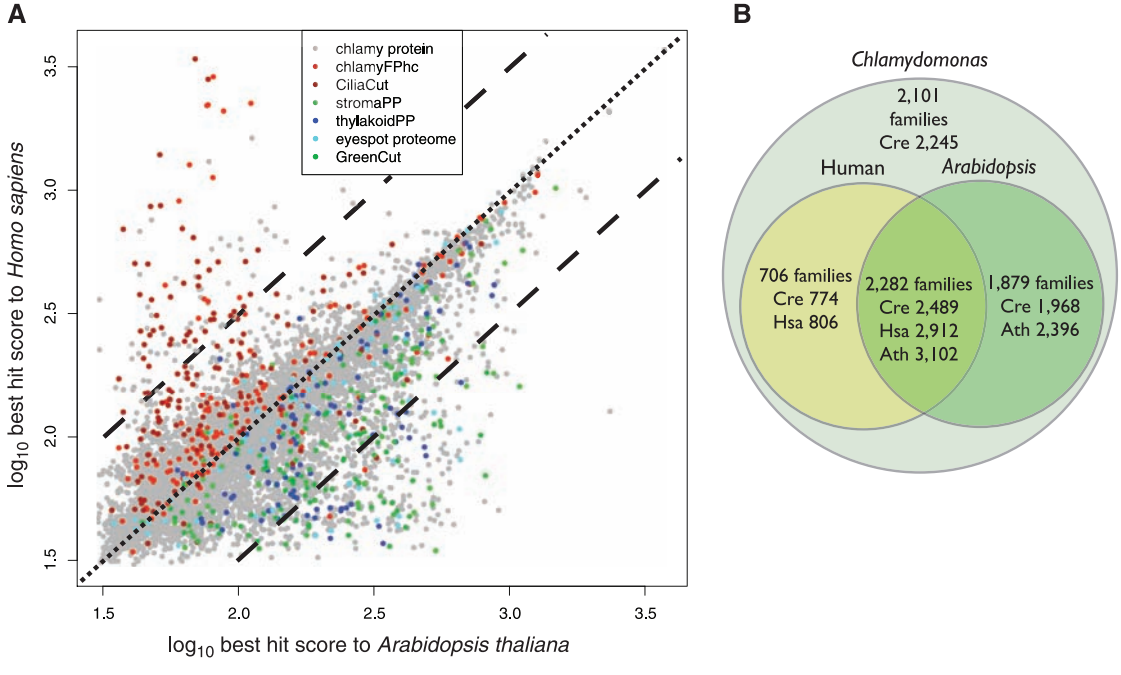


Fig. 4. (**A**) Scatter plot of best BLASTP hit score of *Chlamydomonas* proteins to *Arabidopsis* proteins versus best BLASTP hit score of *Chlamydomonas* proteins to human proteins. Functional or genomic groupings are colored [see inset key in (A)]: *Chlamydomonas* flagellar proteome (*42*) high confidence set (chlamyFPhc); CiliaCut; *Arabidopsis* stroma plastid proteome (stromaPP); *Arabidopsis* thylakoid plastid proteome (thylakoidPP); eyespot proteome; GreenCut; remaining proteins are gray. (**B**) *Chlamydomonas* protein paralogs were grouped into families together with their homologs from human and *Arabidopsis*. The outer circle represents the proteins in *Chlamydomonas*, 7476 (out of 15,143 total), that fall into 6968 families. Another 7937 proteins cannot be placed in families. Counts of families (and the numbers of proteins from each species in them) with proteins from *Chlamydomonas* and human only, *Chlamydomonas* and *Arabidopsis* only, and *Chlamydomonas* and human and *Arabidopsis*, are shown in the inner circles and the overlap between the two inner circles, respectively. Cre, *Chlamydomonas*; Hsa, human; Ath, *Arabidopsis*.

phototoxic form until it associates with proteins or may function in signaling circadian cues.

We also identified plant-specific AKCs (ABC1 kinase in the chloroplast, AKC1 to 4 in the GreenCut), one of which (designated EYE3) is required for eyespot assembly (*39*). These AKCs are distinct from the mitochondrial ABC1 kinase that regulates ubiquinone production (*40*). Protein phosphatases present in the GreenCut and plastoglobules may turn off signaling initiated by the AKCs.

The PLAPs (PLAP1 to 4 in the GreenCut), also called plastoglobulins, are also associated with the eyespot or plastoglobule. These proteins were originally identified by their abundance in carotenoid-rich fibrils and chromoplast plastoglobules and may be structural or organizational components of this plastid subcompartment. Other GreenCut proteins associated with plastoglobules (*34*, *36*) include short-chain dehydrogenases, an aldo-keto isomerase, various methyltransferases with unspecified substrates, esterases and lipases, and a protein with a pantothenate kinase motif.

In sum, the eyespot or plastoglobules contain proteins that likely function in the synthesis, degradation, trafficking, and integration of pigments and lipophilic cofactors into the metabolic machinery of the cell and, most notably, into the photosynthetic apparatus, where they are in high demand. The numerous proteins in the GreenCut associated with the eyespot/plastoglobules may reflect the diverse repertoire of compounds, such as quinones, tocopherols, carotenoids, and tetrapyrroles (fig. S1B), required by photosynthetic organisms.

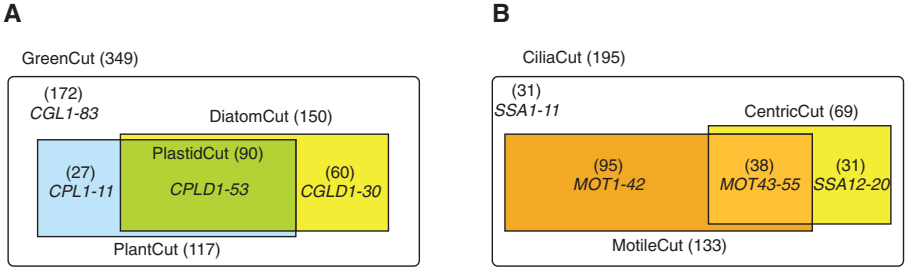
The 90 proteins in the PlastidCut (Fig. 5A) are likely to function in basic plastid processes because they are conserved in all plastid-containing eukaryotes. Sixty-one of these have unknown functions, with genes for most (except CPLD6 and CPLD29) expressed in chloroplast-containing cells, as assessed from EST representation in *Chlamydomonas* and *Physcomitrella*. For *Arabidopsis* homologs, expression (*41*) indicates that the genes represented in the PlastidCut tend to be expressed in leaves or all tissue, similar to genes that function in photosynthesis or primary chloroplast metabolism. Greater than 70% of previously unknown PlastidCut proteins have homologs in cyanobacteria, which suggests a critical, conserved, plastid-associated function.

Flagellar and basal body gene complement. *Chlamydomonas* uses a pair of anterior flagella to swim and sense environmental conditions (Fig. 1). Each flagellum is rooted in a basal body, which also functions as a centriole during cell division. The flagellar axoneme has the nine outer doublet microtubules plus a central pair (9+2) (Fig. 1) characteristic of motile cilia (cilia and eukaryotic flagella are essentially identical organelles). In addition to motile cilia, animals contain nonmotile cilia that function as a sensory organelle and typically lack outer and inner dynein arms, radial spokes, and central microtubules (Fig. 1), all of which are involved in the generation and regulation of motility. Both types of cilia have sensory functions and share conserved sensing and signaling components.

The loss of flagella in angiosperms, most fungi, and slime molds allowed us to identify cilia-specific genes through searches for proteins retained only in flagellate organisms (*8*, *26*). We searched the 6968 *Chlamydomonas* protein families (see above) for those that also contained proteins from human and a *Phytophthora* spp., but not from aciliates, and identified 186 protein families that we named the CiliaCut; these families contain 195 *Chlamydomonas* (Fig. 5B and table SB) and 194 human proteins. One hundred and sixteen of the *Chlamydomonas* proteins had been computationally identified (*8*, *26*), and 45 were identified in this study (*1*).

The *Chlamydomonas* CiliaCut proteins of unknown function that are missing from *Caenorhabditis*, which has only nonmotile sensory cilia (*26*), were designated MOT (motile flagella), whereas proteins of unknown function shared with *Caenorhabditis* were designated SSA (sensory, structural and assembly) (Fig. 5B). Thirty-five percent of CiliaCut proteins are in the *Chlamydomonas* flagellar proteome (*42*), double the number known from previous studies, and 27 of 101 previously identified flagellar proteins (*42*) are present in the CiliaCut. The CiliaCut contained δ-tubulin, which is required for basal body assembly (*43*), and a previously undescribed dynein light chain. Some flagellar proteins were not found by this analysis because they have orthologs in plants and fungi, whereas others are absent because they lack human orthologs. Most dynein heavy chains are missing, most likely due to the difficulty of identifying members of large gene families with a mutual best hit approach (*1*).

We manually curated 125 CiliaCut proteins (fig. S24) and identified large subsets as flagellar structural components (16%), mediating protein-protein interactions (26%), signaling (11%), GTP-binding (6%) and trafficking (6%). These results are consistent with proteomic



Cut	proteins in *Chlamydomonas* AND...	NOT IN...
GreenCut	*Ostreococcus*, *Arabidopsis* AND *Physcomitrella*	non-photosynthetic organisms
PlantCut	*Ostreococcus*, *Arabidopsis*, Physcomitrella AND *C. merolae*	non-photosynthetic organisms
DiatomCut	*Ostreococcus*, *Arabidopsis*, *Physcomitrella* AND a diatom	non-photosynthetic organisms

Cut	proteins in *Chlamydomonas* AND...	NOT IN...
CiliaCut	human AND a *Phytophthora*	non-ciliated organisms
MotileCut	human AND a *Phytophthora*	non-ciliated organisms OR *C. elegans*
CentricCut	human, a *Phytophthora* AND *T. pseudonana*	non-ciliated organisms

Fig. 5. Summary of genomic comparisons to photosynthetic and ciliated organisms. (**A**) GreenCut: The GreenCut comprises 349 *Chlamydomonas* proteins with homologs in representatives of the green lineage of the Plantae (*Chlamydomonas*, *Physcomitrella*, and *Ostreococcus tauri* and *O. lucimarinus*), but not in nonphotosynthetic organisms. Genes encoding proteins of unknown function that were not previously annotated were given names on the basis of their occurrence in various cuts. CGL refers to conserved only in the green lineage. The GreenCut protein families, which also include members from the red alga *Cyanidioschyzon* within the Plantae, were assigned to the PlantCut (blue plus green rectangles). CPL refers to conserved in the Plantae. GreenCut proteins also present in at least one diatom (*Thalassiosira* and *Phaeodactylum*) were assigned to the DiatomCut (yellow plus green rectangle). CGLD refers to conserved in the green lineage and diatoms. Proteins present in all of the eukaryotic plastid-containing organisms in this analysis were assigned to the PlastidCut (green rectangle). CPLD refers to conserved in the Plantae and diatoms. The criteria used for the groupings associated with the GreenCut are given in the lower table. (**B**) CiliaCut: The CiliaCut contains 195 *Chlamydomonas* proteins with homologs in human and species of *Phytophthora*, but not in nonciliated organisms. This group was subdivided on the basis of whether or not a homolog was present in *Caenorhabditis*, which has only nonmotile sensory cilia. The 133 CiliaCut proteins without homologs in *Caenorhabditis* were designated the MotileCut (orange rectangle). Unnamed proteins in this group were named MOT (motility). Proteins with homologs in *Caenorhabditis* are associated with nonmotile cilia (white and yellow areas). Proteins in this group that were not already named were named SSA. The CentricCut (yellow plus light orange box) is made up of 69 CiliaCut homologs present in the centric diatom *Thalassiosira*. These proteins can be divided into those also in the MotileCut (38 proteins; light orange box) or those not present in the MotileCut (31 proteins; yellow box).

analysis of the flagellum (*42*) and highlight the importance of signaling even in motile flagella.

The 62 CiliaCut proteins that *Chlamydomonas* shares with *Caenorhabditis* are predicted to have structural, sensory, or assembly roles in the cilium. As expected, the 133 CiliaCut proteins missing from *Caenorhabditis* (Fig. 5B) (*1*), designated the MotileCut, include a number of proteins associated with motility (*42*) (table S14). This data set also contains 31 proteins of unknown function found in the flagellar and basal body proteomes, 36 known but uncharacterized proteins, and 55 novel proteins (designated MOT1 to MOT55); these flagellar proteins are all predicted to be involved specifically in motility.

A comparison of CiliaCut proteins with proteins encoded by the *Physcomitrella* genome indicates that *Physcomitrella* has lost five of the outer dynein arm proteins (Fig. 1, table S14). However, *Physcomitrella* contains inner dynein arm subunits IDA4 and DHC2, as well as subunits of the central microtubules, the radial spokes, and the dynein regulatory complex (table S14). From this we conclude that *Physcomitrella* sperm flagella have a "9+2" axoneme containing inner dynein arms, central microtubules, and radial spokes, but lack the outer dynein arms. Although the structure of the *Physcomitrella* sperm flagellum is not known, sperm flagella of the bryalean moss *Aulacomnium palustre* have just such an axoneme (*44*).

In contrast, the motile flagella of centric diatoms lack the central pair of microtubules (*45*, *46*). Orthologs of 69 of the 195 CiliaCut proteins (named CentricCut, Fig. 5B) were predicted to be present in the centric diatom *Thalassiosira*. As expected, *Thalassiosira* lacks all central pair proteins. However, it also lacks all radial spoke and inner dynein arm proteins, but has most of the outer dynein arm proteins. The contrasting patterns of loss of axonemal structures predicted for *Physcomitrella* and *Thalassiosira* suggest that the central pair and radial spokes function as a unit with the inner arms, but are dispensable for the generation of motility by the outer arms.

Intraflagellar transport (IFT), which is conserved in ciliated organisms except malaria parasites (*47*), is essential for flagellar growth (*48*). The IFT machinery consists of at least 16 proteins in two complexes (A and B) that are moved in anterograde and retrograde directions by the molecular motors kinesin-2 and cytoplasmic dynein 1b, respectively (Fig. 1). Our analysis of *Thalassiosira* reveals that it has components of the anterograde motor and complex B, but has lost the retrograde motor and complex A (table S14). This is intriguing, as retrograde IFT is essential for flagellar maintenance in *Chlamydomonas* (*49*) and is important for recycling IFT components (*50*). In addition, both *Physcomitrella* and *Thalassiosira* have lost the Bardet-Biedl syndrome (BBS) genes. BBS gene products are associated with the basal body in *Chlamydomonas* and mammals (*8*, *51*) and sensory cilia in *Caenorhabditis* (*52*), where they may be involved in IFT (*53*).

We searched the CiliaCut proteins for proteins shared with *Ostreococcus* spp., a green alga lacking a flagellate stage. The *Ostreococcus* spp. retain 46 (24%) of the 195 CiliaCut proteins but, consistent with loss of the flagellum, are missing genes encoding the IFT-particle proteins and motors, the inner and outer dynein arm proteins, the radial spoke and central pair proteins, and 32 out of 39 flagella-associated proteins (FAPs) (table S14). They have also lost many genes encoding basal body proteins, including all BBS proteins (table S14), which suggests that *Ostreococcus* also lack basal bodies. However, *Ostreococcus* spp. have retained many other CiliaCut proteins (table S14), which suggests either that they recently lost their flagella, or that they retained flagellar proteins for other cellular functions.

Conclusions. This analysis of the *Chlamydomonas* genome sheds light on the nature of the last common ancestor of plants and animals and identifies many cilia- and plastid-related genes. The gene complement also provides insights into life in the soil environment where extreme competition for nutrients likely drove expansion of transporter gene families, as well as sensory flagellar and eyespot functions (e.g., facilitating nutrient acquisition and optimization of the light environment). As more of the ecology and physiology of *Chlamydomonas* and other unicellular algae are explored, additional direct links between gene content and functions associated with the soil life-style will be unmasked with increased potential for biotechnological exploitation of these functions.

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Supporting Online Material

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Supporting Material for

The *Chlamydomonas* Genome Reveals the Evolution of Key Animal and Plant Functions

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CHLAMYDOMONAS GENOME: SUPPLEMENTAL MATERIAL

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1. MATERIALS AND METHODS

A. Strains: High quality genomic DNA was prepared from strain CC-503 *cw92 mt+*, a cell wall-deficient mutant isolated from strain 137c, which contains *nit1* and *nit2* mutations. A BAC library was prepared from the same strain (*1*). Most of the cDNA libraries were derived from wild-type strain CC-1690 21 gr *mt+* and most of the ESTs were sequenced at Stanford (*2, 3*). Strains CC-503 and CC-1690 were derived from the same original field isolate collected in Massachusetts in 1945, but their parent strains have been cultured separately since the mid-1950s. CC-2290 or S1D2 mt^-, which was used for generating some ESTs at the DOE - Joint Genome Institute (JGI) (see below), was collected in the 1980s in Minnesota (*4*). These strains are available from the *Chlamydomonas* Resource Center (*5*). ESTs from the Kazusa DNA Research Institute, Institute of Applied Microbiology, Tokyo, were from strain C-9. This strain also derives from the 1945 field isolate, and is listed in the *Chlamydomonas* Resource Center collection as strain CC-408 (*6*).

B. Whole genome shotgun sequencing and sequence assembly: The initial data set was derived from whole-genome shotgun sequencing (*7*) of 11 libraries supplemented with BAC end sequences. We used nine plasmid libraries, six with an insert size of 2-3 kb, three with an insert size of 6-8 kb and two fosmid libraries with an insert size of 35-40 kb. The reads from the different libraries were as follows: 2,153,471 reads from the 2-3 kb insert libraries comprising 1,683 Mb of raw sequence, 894,846 reads from the 6-8 kb insert libraries comprising 887 Mb of raw sequence, and 184,542 reads from the 35-40 kb insert libraries comprising 184 Mb of raw sequence (including BAC end sequence). The reads were screened for vector sequence with cross_match (*8*) and trimmed for vector and low quality sequences. Reads shorter than 100 bases after trimming were excluded from the assembly. This reduced the data set to 1,903,662 reads from the 2-3 kb insert libraries comprising 807 Mb of raw sequence, 830,326 reads from the 6-8 kb insert libraries comprising 544 Mb of raw sequence, and 153,719 reads from the 35-40 kb insert libraries comprising 49 Mb of raw sequence.

The high GC content of the *Chlamydomonas* genome caused reduced cloning efficiency and premature termination of sequencing reactions, resulting in uneven shotgun sequence coverage across the genome and reduced read lengths. To overcome

this difficulty, DMSO (5% final) was added to both the amplification and sequencing reactions. In addition, the RCA Finishing Kit (Amersham Biosciences, Piscataway, NJ) improved amplification of GC-rich sequences and reduced band compression and the formation of secondary structures that resulted in sequencing errors.

The trimmed read sequence data were assembled with release 1.0.3 of Jazz, a whole genome shotgun assembler developed at the DOE Joint Genome Institute (*9*). A word size of 14 was used for seeding alignments between reads, with a minimum of 15 shared words required before an alignment between two reads would be attempted. To reduce the number of collapsed repeats, words present in the sequence data in more than 65 copies were excluded from the set used to seed alignments. A mismatch penalty of −30.0 was used, which generally allows assembly of > 97% identical sequences. The genome size and sequence coverage were estimated to be 130 Mb and 13.0X, respectively. The initial assembly contained 125.5 Mb of scaffold sequence, of which 15.5 Mb (12.4%) represented gaps. There were 7,091 scaffolds, with a scaffold N/L50 of 26/1.63 Mb, and a contig N/L50 of 658/41.7 kb. Short scaffolds (<1 kb length) were removed.

The assembly was next filtered for redundant scaffolds that matched larger scaffolds (<5 kb length where >80% matched a scaffold of >5 kb length). Mitochondrion and chloroplast genome sequences, available prior to the nuclear assembly, were used to identify scaffolds comprising organelle sequence. Finally, scaffolds that showed homology to prokaryotic and non-cellular contaminants [i.e. viroids, viruses, other unclassified, top-level categories at NCBI (*10*)] were identified and removed. After filtering, 121.0 Mb of scaffold sequence remained, of which 15.3 Mb (12.7%) represented gaps.

The filtered assembly (v3.0) contained 1,557 scaffolds, with a scaffold N/L50 of 25/1.63 Mb, and a contig N/L50 of 608/44.5 kb. The sequence coverage was 12.8X ± 0.3X. To estimate the completeness of the assembly, a set of 168,110 ESTs was aligned with BLAT (*11*) to both the entire set of unassembled trimmed reads prior to running through Jazz (pre-assembled) and the assembled sequence; 159,136 ESTs (94.7%) were more than 80% covered by the unassembled data, 160,841 (95.7%) were more than 50% covered and 161,241 (95.9%) were more than 20% covered. By way of comparison,

159,084 ESTs (94.6%) matched the assembled sequence, showing that the assembly covers approximately 95% of the pre-assembled reads.

Whole genome alignment with WU-BLASTN (*12*) of the *Chlamydomonas* v3.0 assembly to the genome sequence of *Ralstonia eutropha* JMP134 (*13*) and *Populus trichocarpa* (*14*) revealed 299 *Chlamydomonas* scaffolds with regions identical to *Ralstonia* or *Populus* genomic sequence. 291 of these scaffolds (each ≤40 kb and assembled from ≤22 sequence reads, and together totaling 1.9 Mb of sequence) were manually removed. A new assembly with the remaining 1,226 scaffolds (assembly v3.1) was generated and is available for download on the JGI *Chlamydomonas* genome browser (*15*).

Of the 74 scaffolds that could be mapped to linkage groups only two show evidence of misassembly (i.e. contain segments that map to two different linkage groups). The approximate positions of the breakpoints are known: the segment of scaffold_6 with coordinates from 1 to ~1.23 Mb maps to LG V and the segment from ~1.44 Mb to 2.94 Mb maps to LG VII; the segment of scaffold_14 from 1 to ~0.874 Mb maps to LG III and the segment ~0.879 Mb to 2.12 Mb maps to LG XVIII.

The Stanford Human Genome Center has been finishing the genome of *Chlamydomonas* since April 2005 with the goal of releasing a finished reference sequence in 2007. The finishing process has been complicated by extreme variations in GC content, sequence hairpins and the presence of many small tandem repeats. Experiments performed to improve the quality of the genome sequence include: resequencing using dGTP chemistry, custom primer walks using a variety of different chemistries and conditions, transposon sequencing and the generation of small insert shatter libraries. In addition, a BAC library (with a mean insert size of 174 kb) provided by Andreas Gnirke from Exelixis (South San Francisco, CA, USA) has been end-sequenced; this library has been used to make further scaffold joins across the genome, reducing the scaffold number (>25 kb) from 168 to 91.

C. EST sequencing and sequence assembly: *E. coli* colonies harboring cDNA clones from *Chlamydomonas* strain S1D2 were plated onto solid agarose medium at a density of approximately 1,000 colonies per plate. The bacteria were grown at 37°C for 18 h and individual colonies were picked robotically and inoculated into LB medium with an

appropriate antibiotic in a 384 well plate format. Plasmid DNA was amplified by a rolling circle mechanism (Templiphi, GE Healthcare, Piscataway, NJ) and purified. The insert of each clone was sequenced from both ends with primers complementary to flanking vector sequences (Forward: 5'-ATTTAGGTGACACTATAGAA: Reverse: 5'-TAATACGACTCACTATAGGG) using Big Dye terminator chemistry; the products of the sequencing reactions were resolved by an ABI 3730 sequenator (ABI, Foster City, CA), yielding a total of 34,403 reads. Detailed sequencing protocols can be found in (*16, 17*).

The JGI EST Assembly Pipeline was run on a combined set of 196,594 sequences comprising the 34,403 S1D2 sequences together with ~160,000 sequences from NCBI mRNA and EST databases (*18*) and ~ 2,000 other sequences from various libraries. The pipeline began with the cleanup of 5'and 3' end reads from individual cDNA clones. The Phred program (*8, 19*) was used to call the bases and generate quality scores. Vector, linker, adapter, poly-A/T, and other artifact sequences were removed with the cross_match software, and an internally-developed algorithm that identifies short patterns. Low quality sequence reads were identified with internally-developed software, which masks regions with a combined quality score of less than 15. The longest high quality region of each read was used as an individual EST. ESTs shorter than 150 bases and those representing common contaminants, including *E. coli* genomic sequence, vector sequences, and sequencing standards are removed from the data set. EST clustering was performed ab initio, on the basis of alignments between pairs of trimmed, high quality ESTs. Pairwise EST alignments were generated with the Malign software (*20*), which is a modified version of the Smith-Waterman algorithm (*21*) that has been developed at the JGI for use in whole genome shotgun assembly. ESTs with 150 bp overlaps that align at ≥98% identity were assigned to the same cluster. These were relatively strict clustering cutoffs intended to avoid placing divergent members of gene families into the same cluster. However, this could separate splice variants into different clusters. Optionally, ESTs that do not share alignments were assigned to the same cluster if they were derived from the same cDNA clone. EST cluster consensus sequences were generated by running the Phrap program on the ESTs of each cluster. All alignments generated by Malign are required to extend to within a few bases of the ends of both

ESTs. Therefore, each cluster resembles a 'tiling path' across the gene that matches well with the genome-based assumptions underlying the Phrap algorithm. Additional improvements of the Phrap assemblies were achieved by using the 'forcelevel 4' option, which decreases the chances of generating multiple consensus sequences for a single cluster, where the differences in the consensus sequences may only represent sequencing errors. EST clustering generated 38,869 clusters containing 40,219 consensus sequences.

D. Generation of gene models and annotation: The genome assembly was annotated using the JGI Annotation Pipeline, which combines several gene prediction, annotation and analysis tools. First, the genome assembly was masked using RepeatMasker (*22*) and a custom repeat library (see below). Next, the EST (*3*) and full-length cDNAs were clustered into 32,960 consensus sequences (see above) and aligned to the scaffolds with BLAT (*11*). Model organism protein sequences from the non-redundant (NR) set of proteins from the National Center for Biotechnology Information (Genbank) (*18*) were aligned to the scaffolds with BLASTX (*23*). Gene models and associated transcripts/proteins were predicted or mapped using data from 5,476 putative full-length cDNAs derived from available mRNA, EST and ACEG sequences, and methods such as Genewise (*24*) and *ab initio* approaches such as Fgenesh and Fgenesh+ (*25*). Fgenesh was trained on 495 known genes and reliable homology-based models. The clustered ESTs/cDNAs were used to extend and correct predicted gene models where the exons overlapped and splice junctions were not consistent in comparing EST sequences to gene models. The use of EST information often added 5' and/or 3' UTRs to the models. With gene structure in place, function was assigned to models based on Smith-Waterman (*21*) homology to annotated genes from NR (*18*), KEGG (*26-28*) and KOG (*29*) databases. InterproScan (*30*) was used to identify predicted domains and the Gene Ontology (GO) (*31*) was used to identify function and/or subcellular location. Of the gene models present in the gene catalog (see below), 3,137 models from version 2 of the genome assembly (chlre.v2.0) were mapped forward (Table S4).

Although multiple models with overlapping sequences were generated for each locus, a single model was chosen for the gene catalog set. Model selection was based on maximizing protein sequence relationship and EST support for splice sites, ORFs and model completeness (i.e. inclusion of 5' methionine, 3' stop codon, and UTRs). After a

first automatic filtering, the catalog was refined by the annotators, including through generation of *ad hoc* gene models. The catalog was frozen on July 6, 2006, yielding 15,143 gene models, at 14,673 loci ("Frozen Gene Catalog"). All analyses discussed in this paper were carried out on this set. 9,461 (62%) predicted proteins from the Frozen Gene Catalog appear to be full-length, on the basis of the presence of start and stop codons. 4,369 (29%) also have both 5' and 3' UTRs. Furthermore, the majority of predicted genes are supported by EST (56%) or BLASTP (*23*) homology (63%) evidence (Table S5). Of the 6,298 predicted proteins without homology, 30% are *ab initio* fgenesh models with no apparent support and 59% have some support on the basis of EST or distant sequence relationships (E-value > 1E-5). Of the latter group 309 (4.9%) were annotated by users. An analysis based on Smith-Waterman alignments (E-value < 1E-5) (Table S6) yielded 9,435 (62%) gene models with homology to proteins in the COG database (*29, 32*) and/or with Gene Ontology annotations (*31*). Of the predicted gene models 35% have a manually assigned gene function. Furthermore, as of June 2007, 5,141 had been manually-annotated in an attempt to improve the gene set prior to submission to DDBJ/EMBL/GenBank (ABCN01000000). This resulted in an overall decrease in the number of gene models from 15,413 to 14,662. Annotation is on-going and data are available at the JGI genome portal (*15*). Periodic updates will be submitted to DDBJ/EMBL/GenBank (*33*).

E. Identification of transposons and simple sequence repeats: Censor (*34*) was used to identify occurrences of known transposon sequences. These sequences were clustered into families of transposons and retrotransposons and consensus sequences were manually curated. This process identified many new transposon families. The newly identified transposons were annotated and deposited in Repbase (*35*). The genome also contains an extensive range of simple sequence repeats that were identified with Censor (*34*). These have been compiled in a library (similar to the library associated with RepeatMasker).

F. Annotation of snoRNA genes: The snoRNA genes were identified using snoRMP (snoRNA Mining Platform), which is based on the SnoScan (*36*) and SnoGPS (*37*) algorithms, combined with secondary structure prediction and comparative genomic

analysis. These approaches predict snoRNA function and have been used successfully for snoRNA gene identification in yeast, plants, mammals and other genomes (*38, 39*).

G. Identification of membrane transporters: To identify membrane-associated transport systems, the complete, predicted proteome was searched against a curated database of transport proteins (*40*) using BLASTP (*23*). All query proteins with significant hits (E-value < 0.001) were collected and searched against the NCBI non-redundant protein and PFAM databases (*41*). Transmembrane protein topology was predicted by TMHMM (*42*) and a web-based interface was implemented to facilitate annotation processes, which incorporate (i) number of hits to the transporter database, (ii) the BLAST and HMM search E-value and score, (iii) the number of predicted transmembrane segments, and (iv) description of top hits to the non-redundant protein database. Detailed transporter profiles and abbreviations for transporter families can be found in (*40, 43*) and at the website TransportDB (*44*). The MPT and IISP transporter families were not included as complete data on these two families in all eukaryotes is not available.

H. Generation of paralogous gene families: We constructed *Chlamydomonas* gene families to investigate both the size and functions of proteins associated with these families. Protein sequences were compared by an all-against-all WU-BLASTP (*12*). The bit score was parsed from the BLAST output and used as the basis for Markov Clustering (MCL) (*45*) with an inflation index of 2.0. PFAM domains were assigned to members of families by RPSBLAST (*23*) (expect score < 1E-10). In the absence of PFAM domain homology, gene families were annotated with InterproScan (*29*). A correlation of >0.5 between nucleotides in the EST and nucleotides in the gene model was taken as evidence for expression of the gene. Sequences from each family were blasted to the NR data base (*18*) to determine homology. For comparison, the same analysis was performed for human, *Arabidopsis*, *Dictyostelium*, *Ostreococcus* spp., and *Neurospora crassa*. *Chlamydomonas* sequences with homology to transposable elements or which contain fragments from transposable elements, exhibit overlapping exonic regions, and do not have support for being expressed are unlikely to represent bonafide *Chlamydomonas* protein-coding genes and were not analyzed further.

In addition to the 51-member type III adenylyl/guanylyl cyclase domain-containing family, there is another family of three proteins with cyclase domains linked to heme NO-binding domains, as well as a pair of cyclases that is in a separate family type. This brings the total number of potential cyclases encoded on the *Chlamydomonas* genome to 56.

I. Best BLASTP score scatter plot of *Chlamydomonas* proteins against human and *Arabidopsis* proteins: The BLASTP scores of every *Chlamydomonas* protein against every human protein and Arabidopsis protein were taken from the BLAST analysis that we performed as part of the construction of homologous protein families (below). A scatter plot was generated with the coordinates of every point determined by the best blast score of the *Chlamydomonas* protein to *Arabidopsis* proteins on the x-axis and to human proteins on the y-axis.

J. Construction of families of homologous proteins: As a pre-requisite to comparing gene content of *Chlamydomonas* to other organisms at the whole-genome scale, we constructed families of homologous proteins from all sequences from *Chlamydomonas* and a wide phylogenetic range of prokaryotic and eukaryotic organisms (Fig. 2). Where several closely-related genome sequences were available, we chose manually- or well-annotated species to represent clades of interest. The shared ancestry (homology) of family members enabled us to infer shared function, allowing functional annotations to be transferred among family members. To create protein families, we first blasted [WU-BLASTP 2.0MP-WashU (20- Apr-2005) (macosx-10.3-g5-ILP32F64 2005-04-21T15:44:27)] (*12*) all protein sequences in *Chlamydomonas* to all protein sequences in the red alga (*Cyanidioschyzon,* strain 10D) (*46*), green algae *Ostreococcus tauri* (assembly v2.0) and *O. lucimarinus* (assembly v2.0) (*47-49*), the land plants *Arabidopsis thaliana* (*50*), and *Physcomitrella patens* (assembly v.1) (*51*), the cyanobacteria *Synechocystis* sp. strain PCC6803 (GenBank Accession: BA000022) and *Prochlorococcus marinus* strain MIT9313 (*52*), bacteria including *Pseudomonas aeruginosa* (strain PA01) (GenBank Accession: AE004091.1) and *Staphylococcus aureus* (subsp. aureus, strain N315) (GenBank Accessions: BA000018.1 AP003139.1), the Archaea *Methanosarcina acetivorans* strain C2A (*53*) and *Sulfolobus solfataricus* strain P2 (*54*), the oomycetes *Phytophthora ramorum* (v1) (*55*) and *P. soja*e (assembly v1) (*56*),

the diatoms *Thalassiosira pseudonana* (assembly v3.0) (*57*) and *Phaeodactylum tricornutum* (assembly v2.0) (*58*), the amoeba *Dictyostelium discoideum* (*59, 60*), the fungus *Neurospora crassa* (assembly v7.0; annotation v3.0) (*61*), and the metazoans human (*61-63*) and *Caenorhabditis elegans* (*62*). The blast score of each pair of proteins was extracted and used as a measure of evolutionary distance. Assignment of orthology was determined by mutual best hit between two proteins, using this metric. In creating individual protein families, we first generated all possible ortholog pairs consisting of one *Chlamydomonas* protein and a protein from another organism. Next, paralogs were added to each pair of proteins. A paralog from a given organism was added if its p-dist (defined as 1 – the fraction of identical aligning amino acids in the proteins) was less than a certain fraction of the p-dist between the two orthologs in the pair. The fractions were chosen to be 0.5 for pairs of organisms involving *Chlamydomonas* and a eukaryote and 0.1 for *Chlamydomonas* and a prokaryote. Two considerations led to the choice of these values. In order to assign function correctly, we wanted to include only 'in-paralogs' (paralogs that had duplicated after speciation) (*63*). Secondly, we determined empirically that higher (less stringent) values led to the generation of unwieldy protein families with >22,000 members that could not be analyzed further. In a last step, all pair-wise families of two orthologs plus paralogs were merged if they contained the same *Chlamydomonas* proteins. This created 6,968 families of homologous proteins. Each individual family consists of one or more *Chlamydomonas* paralog(s), mutual best hits to proteins of other species (orthologs) and any paralogs in each of those species. The set of protein families was used in subsequent 'cuts' for analysis of proteins associated with chloroplast or ciliary function (see below). To accomplish this, we built a software tool that allowed us to search for protein families containing any desired combination of species. We call the search results a 'cut' as it represents a phylogenetic slice through the collection of protein families.

The random nature of gene duplication and subsequent divergence and loss that leads to large gene families means that it is sometimes impossible to precisely assign orthology and paralogy between genes. As a result, mutual best hit relationships between sequences may not exist, preventing family construction, or may not be between correct proteins, leading to inclusion of non-homologous proteins in families. This problem was

particularly evident in the large family containing the Light Harvesting Complex Proteins (LHCP), for which only two members were included, and the axonemal dynein proteins, for which only two of 14 members in *Chlamydomonas* were included. Furthermore, a cytoplasmic dynein sequence from a diatom was included in the IDA4 inner dynein arm family, probably because the flagella-less diatom is missing genuine inner or outer dynein arms, and its cytoplasmic dynein therefore represents the mutual best hit.

K. Making the 'GreenCut': Having constructed families of homologous proteins, centered on *Chlamydomonas* proteins, we used our search tool (see above) to identify protein families in which all members were present in species in the green lineage of the Plantae, which includes *Chlamydomonas*, the prasinophyte algae *Ostreococcus* spp. (*47*) the angiosperm *Arabidopsis*, and the bryophyte *Physcomitrella* (*50, 51*), but not present in nonphotosynthetic organisms. We refer to this as the 'GreenCut' (Supplemental File 1).

Estimation of false negative frequency: The algorithm was designed to generate a conservative list of proteins, which might result in loss of some proteins that are specific to the green lineage or chloroplast function. We used the components of the photosynthetic apparatus to gauge the effectiveness of the method in recovering proteins expected to be unique to green chloroplasts. Since the cytochrome b_6f complex and the ATP synthase function are also in respiratory membranes in bacteria, we considered only the photosystems, their unique donors and acceptors (plastocyanin, ferredoxin, FNR) and Calvin Cycle enzymes that function only in photosynthetic carbon metabolism (Rubisco and phosphoribulokinase). Using only nucleus-encoded proteins, we generated an "expect inventory" of PsbO, P, Q, R, S, W, X, Y, PsaD, E, F, G, H, K, L, O, plastocyanin, ferredoxin, FNR, RbcS and phosphoribulokinase. Of these 21 proteins, 18 appear in the GreenCut, which gives a potential false negative frequency of ~14%.

Estimate of false positive frequency: There are 135 encoded proteins in the Knowns (K) and Known by Inference (KI) categories. Each of the K and KI proteins was assigned to a subcellular compartment based primarily on annotation of their *Arabidopsis* homologs (TAIR database), but also based on experimental evidence in the literature for *Chlamydomonas* or other photosynthetic organisms (tomato, spinach and tobacco) (**Table S13**). At least 85% (115/135) of the proteins were assigned to the chloroplast, with 9%

(12 out of 135) in other intracellular compartments and the remaining 8 proteins having an undetermined localization. The proteins we regard as false positives are RAD9/At3g05480, ERD2B/At1g19970 (KDEL receptor), SEC12/At5g50550, CYN23b/At1g26940 (ER cyclophilin), CGL28/At1g53650 (RNA binding protein), EFL1/At2g21340 and MER/At3g27730, which represent 5% of the total number of proteins. If CGL22/At2g03670, AMI2/At1g08980, SNE1/At5g28840 and CCD1/At3g63520 are included as false positives (some of these proteins appear to function in processes with plant specific peculiarities), the number increases to 8%. The high percentage of chloroplast localized proteins, as well as proteins that have functions unique to plants, gives an indication of the validity of the method, providing a basis for assessing functions of the unknown proteins. In fact, for one protein, PRMT3403/At3g12270, its presence in a cluster with moss and algae prompted a re-evaluation of the group and an assignment of function as the ribosomal protein arginine methyl transferase, resulting in the movement of the protein from the UP to the KI category. Phylogenetic analysis now places PRMT3403 and At3g12270 together in a green lineage-specific clade.

L. Making the 'CiliaCut': Having made families of putatively homologous proteins (see above), we searched the families for those in which all members were from ciliated organisms; the collection of proteins in these families is designated 'CiliaCut'. To make the CiliaCut, we searched the complete set of homologous protein families for families with members in human, *Chlamydomonas* and at least one *Phytophthora*, but not in the non-ciliated organisms *Arabidopsis*, *Neurospora*, *Cyanidioschyzon*, *Dictyostelium* or eubacteria and archaea. *Phytophthora* are ciliated protists that diverged from animals and plants a relatively short time before animals and plants diverged from each other. Despite this deep divergence, both the core motility machinery and signal transduction pathways are likely to be associated with *Phytophthora* flagella; *Phytophthora* spp. have motile flagellate zoospores that chemotax to their host plants (*64*), implying that their flagella also contain signal transduction components. Therefore, the proteins required for these core pathways should be present in the CiliaCut dataset, and their inclusion adds specificity to the CiliaCut.

There were fourteen *Chlamydomonas* genes in the CiliaCut families that appeared to contain transposons. These were removed from the analyses. The remaining CiliaCut proteins were classified based on the function of characterized orthologous family members, PFAM domain predictions, published information, protein domain searches, and previous comparative genomics (*65, 66*), proteomics (*67, 68*), tissue-specific gene expression studies (*69*), and the ciliome database (*70*).

Estimation of sensitivity and specificity in the 'CiliaCut': There is no simple way to assess how many of the genes in the CiliaCut are genuinely cilia-related and how many of the genuinely cilia-related genes are missing (analagous to the analysis performed for the GreenCut). Nonetheless, we made two attempts to address this issue. First, we compared the CiliaCut proteins to those in the *Chlamydomonas* Flagellar Proteome (chlamyFP) (*67*) and second, we compared the CiliaCut proteins to a curated list of proteins known to be involved in flagellar function.

We assumed that the high confidence proteins from the chlamyFP were very likely to be genuine. 35% (68 out of 195) of CiliaCut proteins are in the chlamyFP high confidence set, whereas only 15% (104 of 687) and 17% (32 of 187) of the proteins in the studies of Li (*66*) and Avidor-Reiss (*65*), respectively, are present in chlamyFP. This represents a greater than two-fold increase in specificity in the CiliaCut relative to previous work, presumably reflecting the inclusion of distantly related flagellate organisms as well as the inclusion of additional information based on the completion of genome sequences.

We also examined the known flagellar proteins identified prior to the generation of chlamyFP. We made a list of 13 randomly-chosen proteins known to be flagella-specific, including only one protein from each protein family; this avoids under-clustering of members of large gene families (see above). One of these genes (tektin) was not present in the CiliaCut, nor is it present in the genomes of 2 species of *Phytophthora*. Presence in at least one *Phytophthora* was required for inclusion in the CiliaCut. Of the remaining 12 proteins, 6 (50%) are in the CiliaCut. Similarly, 44% of the CiliaCut genes are upregulated following deflagellation (*71*) and 58% of these upregulated genes are in CiliaCut. These analyses suggest that the CiliaCut is 50-60% complete.

2. SUPPORTING TEXT

A. Transposons and simple sequence repeats: Known and novel families of transposons were identified and curated (see above). Most remarkable is the presence of SINEs (Tables S2 and S3), small interspersed transposable elements ancestrally related to tRNAs, which rely on LINEs (long interspersed transposable elements) for their propagation. There are 5 families (>200 copies) of SINEs, two of which have precisely kept the tRNA structure and intron position (see section B, immediately below). This is the first example of SINE families described in a unicellular organism.

The repeat landscape of the *Chlamydomonas* genome is dominated by GC-rich, simple sequence runs and transposons, totalling 2.1% and 8.9% of the genomic sequence respectively. The transposons include ~100 families of transposable elements represented by 147 consensus sequences (a unique transposon family is defined as less than 75% identical to transposons in other families). There are also many non-autonomous transposable elements that do not encode proteins. The most thoroughly studied transposon in *Chlamydomonas* is Gulliver (*GUL*) (*72*), whose pattern has been used as a feature of various *Chlamydomonas* field isolates to determine their ancestry. *GUL*, which is present at 14 positions on the genome, is scattered among different scaffolds. Genetic mapping of the *GUL* transposons is consistent with their locations on the physical map.

B. tRNA genes: Most of the 259 *Chlamydomonas* tRNAs (Table S1) are clustered on the genome and appear to result from recent gene duplications (Fig. S19A). The tRNA number in *Chlamydomonas* compares with 390 in *Dictyostelium discoideum*, 272 in *Saccharomyces cerevisiae*, 284 in *Drosophila melanogaster*, 496 in *Homo sapiens*, and 630 in *Arabidopsis thaliana*. However, prediction tools such as tRNAscan-SE (*73*) lead to an inflated number of tRNAs because of the highly conserved tRNA SINE retrotransposon elements (see above). SINE elements have evolved from tRNAs and can be abundant in eukaryotic genomes (*74*). The *Chlamydomonas* genome contains 40 SINEX-3 elements with 5 different anticodons that resemble 34 tRNA-Arg-CCG, 1 tRNA-Arg-ACG, 3 tRNA-Trp-CCA, 1 tRNA-Gly-CCC and 1 tRNA-Gln-CTG (Table S2). There are also 29 tRNA-related SINE elements that resemble 11 tRNA-Asp-ATC and 18 tRNA-Asp-GTC (Table S3). In all cases the SINE and authentic tRNA sequences are highly similar, and all SINE retrotransposon elements have an intron of 11-13

nucleotides between positions 37 and 38 of the tRNA sequence. Furthermore, many SINE-tRNA sequences end with a genome-encoded CCA, which is also present on some authentic *Chlamydomonas* tRNAs (see below). It is possible, as suggested for mammals, that these SINEs are important for transcriptional control, especially related to stress responses (*74, 75*).

There are a number of interesting features associated with *Chlamydomonas* tRNAs. A surprisingly large fraction (60%) of *Chlamydomonas* tRNAs contain introns as compared to human (7%), *Drosophila melanogaster* (5%) and *Saccharomyces cerevisiae* (22%). As in the SINE elements, the introns are located at position 37/38, but the size of the intron is extremely variable, ranging from 8-57 nucleotides. Seven of the tRNAs have the 3' terminal CCA encoded on the genome; a sequence normally added post-transcriptionally, after exonucleolytic trimming of the precursor tRNAs. The presence of a CCA in the genomic tRNA sequence is common in some bacteria and archaea but, to our knowledge, has rarely been described in eukaryotes (*76*). As in bacteria, the *Chlamydomonas* genome encodes RNAse PH and RNAse Z homologs, which in *Bacillus subtilis* are responsible for trimming CCA-containing and CCA-free tRNAs, respectively (*77*).

In some organisms, tRNAs are clustered on the genome. In *Dictyostelium* about 20% of the tRNA genes occur as pairs or triplets separated by 5-20 kb. *Arabidopsis* contains large families of tandemly arrayed tRNA that are on the same DNA strand (*78*). In *Chlamydomonas*, tRNA gene clustering is even more striking, with 160 tRNAs (approximately 60% of the total) associated on the same or opposite DNA strands, and separated by spacers that can be as short as 3-7 nt. As an example of clustered and duplicated tRNAs, we analyzed 12 tRNA-Val genes on scaffold 20 (Fig. S19A); 5 of these have an anticodon AAC and a genome-encoded CCA terminal-sequence while 7 have an anticodon CAC. These genes are grouped in two repeat units contained within a 35 kb genomic region. One of the repeat units contains 3 sets, each with 2 tRNAs; this represents duplications in which the tRNAs have remained within ~2 kb on the genome. The second repeat unit contains 2 sets, each with 3 tRNAs. These tRNA-Val sets are on opposite strands and separated on the genome by ~8.5 kb, but the positions and orientations of the genes within each set are essentially identical. Individual genes from

each of the putative gene pairs (genes 7 and 12, 8 and 11, 9 and 10 in Fig. S19A) have anticodons that are identical and introns that are identical, or nearly identical, suggesting a duplication of one entire set. The duplication is likely to have occurred recently on the basis of the near sequence identity between the analogous introns and the neighbor-joining tree made from the intron sequences (Fig. S19B).

C. snoRNA genes: The snoRNA genes are crucial to the biosynthesis of ribosomal RNAs, mediating important steps in folding, site-specific nucleotide modification and precursor cleavage via sequence-specific interactions. The box C/D and box H/ACA snoRNAs guide methylation and conversion of uridine to pseudouridine in their targets, respectively. The *Chlamydomonas* draft genome contains 315 snoRNA genes encoding 124 families, with 71 of the box C/D type and 53 of the box H/ACA type. The box C/D snoRNAs were predicted to guide methylation at 91 sites on rRNAs (31 on 18S, 1 on 5.8S, and 59 on 28S), and 3 sites on U6 snRNAs. Among the 91 rRNA methylation sites, there are 71 analogous sites in other organisms, although 20 are likely *Chlamydomonas* specific. Box H/ACA snoRNAs were predicted to guide pseudouridylation at 63 sites on rRNAs (28 on 18S and 35 on 28S), and 2 sites on U6 snRNA. Among the 63 rRNA pseudouridylation sites, there are 42 analogous sites in other organisms.

About 50% of the *Chlamydomonas* snoRNA genes are present as a single copy on the genome; the rest exist in families of 2 to13 paralogs. Most (71%) snoRNA genes are arranged on the genome in 70 gene clusters, each with 2-6 genes; 52 of these clusters are intron-encoded. Out of the 315 snoRNA genes, 94 were initially predicted to lie between protein-coding genes. After examination of EST and homology data, only 28 were confirmed as intergenic (13 loci). The remaining snoRNAs are found in introns. The polycistronic arrangement of snoRNAs in *Chlamydomonas* is similar to that of rice, although such an arrangement is not observed in vertebrates.

D. Introns and spliceosomal RNAs: Most eukaryotes have a characteristic population of introns with a mode size of between ~60 and 110 nucleotides, although longer introns are common in the human and other large genomes because of repetitive elements embedded in the introns. Surprisingly, the intron size for *Chlamydomonas* gene models, generated as described above, averages 373 nucleotides, which is considerably larger than that of many other eukaryotes (Fig. S21A). Furthermore, the peak intron size in the 60-110

nucleotide range, a feature of the typical bimodal distribution observed for many eukaryotes (Fig. S21A), is missing. These observations are not an annotation artifact as an almost identical peak value for intron length was obtained in the analysis of EST-derived ACEGs.

We calculated the proportion of nucleotides in introns that overlap predicted repeat sequence (see above). 30% of intron sequence consists of repeats, nearly three times the proportion for the whole genome of 11%. This suggests invasion by repeats as a possible mechanism of intron expansion.

Chlamydomonas introns show classical 3' and 5' splice site consensus sequences (CAG^ and G^GTG, respectively), but the classical sequence surrounding the branchpoint (CTNAY) is often difficult to recognize. This suggests that canonical base-pairing between the U2 snRNA and the branchpoint sequence contributes only marginally to the assembly of the spliceosome onto most pre-mRNAs. Similarly, the U1 consensus AAACUUACCU sequence that binds the 5' splice site of introns is not a perfect match to the consensus splice site in *Chlamydomonas* introns (ACG^**GU**GCG).

Altogether, 30 loci were identified that encode the 5 spliceosomal snRNAs. Two of the five U1 genes, four of the six U2 and one of the two U4 genes (all transcribed by Pol II) show EST coverage, with various degrees of truncation at the 5' end. In general, the snRNA-encoding sequences are found within introns of protein coding genes (supported by EST or homology-based analyses). An alternative transcription start gives rise to a transcript extending several hundred base pairs beyond the mature 3' end of the snRNA. The snRNAs are polyadenylated and spliced, using the same canonical exon/intron boundaries as the "host" gene. These observations are consistent with the highly unusual notion that *Chlamydomonas* snRNAs are transcribed as long precursors that are spliced and polyadenylated before maturation. Polyadenylation has been shown for *Dictyostelium* snRNAs (*79*) but splicing of a snRNA precursor has not been described.

E. Outlying proteins in scatter plot comparison of *Chlamydomonas* proteins to proteins in *Arabidopsis* and human: As expected, proteins from the high confidence *Chlamydomonas* Flagellar Proteome (chlamyFP set) (*67*) and CiliaCut (Fig. 4A, red and purple points, respectively) are shifted toward the human axis and conversely, many

proteins associated with thylakoid, stroma, eyespot proteomes, and GreenCut (dark blue, green, light blue and dark green points, respectively) lie closer to the *Arabidopsis* axis. Two high confidence chlamyFP points represent proteins with general enzymatic functions and activities that may not be strictly related to flagella function or biogenesis. There is one dark red point outlier from the CiliaCut which closely aligns with a homolog in *Arabidopsis*. There are also two outliers in the thylakoid proteome (Fig. 4A) that are more similar to human than to *Arabidopsis* proteins. In both proteomics sets, the outliers might represent contaminants present in the preparations used to generate the proteomic database.

In analogous analyses, we generated scatter plots of the best blast scores between *Chlamydomonas* proteins and proteins of other photosynthetic organisms (*Arabidopsis*, *Ostreococcus tauri* and *Thalassiosira pseudonana*) (Fig. S25). As expected, these plots show significantly fewer outlying proteins and reveal a closer overall similarity of *Chlamydomonas* proteins to those of *Arabidopsis* than to those of either *O. tauri* or *T. pseudonana*.

F. Transporters of the PlastidCut: Three transporters in the PlastidCut, CPLD21-CPLD23, are predicted to be sugar nucleotide transporters, consistent with the key role of plastids in sugar metabolism. More proteins, including exchangers/carriers that are involved in transporting the substrates and products of plastid metabolism such as phosphate, phosphate-esterified carbon compounds and organic acids, are conserved if we consider only the green lineage. A novel plastid transporter, TIM22B, was also identified in this analysis. This plastid-localized protein has evolved from the expansion of a family of mitochondrial pre-protein translocases (*80*) and is an interesting candidate for functional analysis because it may be involved in the movement of peptide substrates with bound ligands, such as FeS clusters or other minerals that are metabolized in the plastid.

3. SUPPORTING FIGURES

Fig. S1. Photosynthetic electron transport and isoprenoid metabolism: (**A**) 'Z' scheme of photosynthesis, showing photosystems (PS) II and I which are complexes of Psb and Psa polypeptides, respectively, and the cytochrome b_6f complex; Fd, ferredoxin; Trx, thioredoxin; redrawn from (*81*); (**B**) summary of isoprenoid metabolism with enzymes of the pathway mentioned in the text (purple), and end-products (orange); adapted from (*82*). The chloroplast is the site of synthesis of heme, chlorophyll, quinones (phylloquinone, plastoquinones), tocopherols (Vitamin E), and carotenoids, each derived from a common pool of isoprenoid pathway precursors and many having functions in light harvesting, photoprotection (e.g. antioxidants), and as cofactors for electron transfer reactions (*82, 83*). We noted many proteins in the UP categories of the GreenCut are predicted to function in isoprenoid metabolism based on their similarity to known enzymes in these pathways (see Table S12).

Supplemental Figure 1

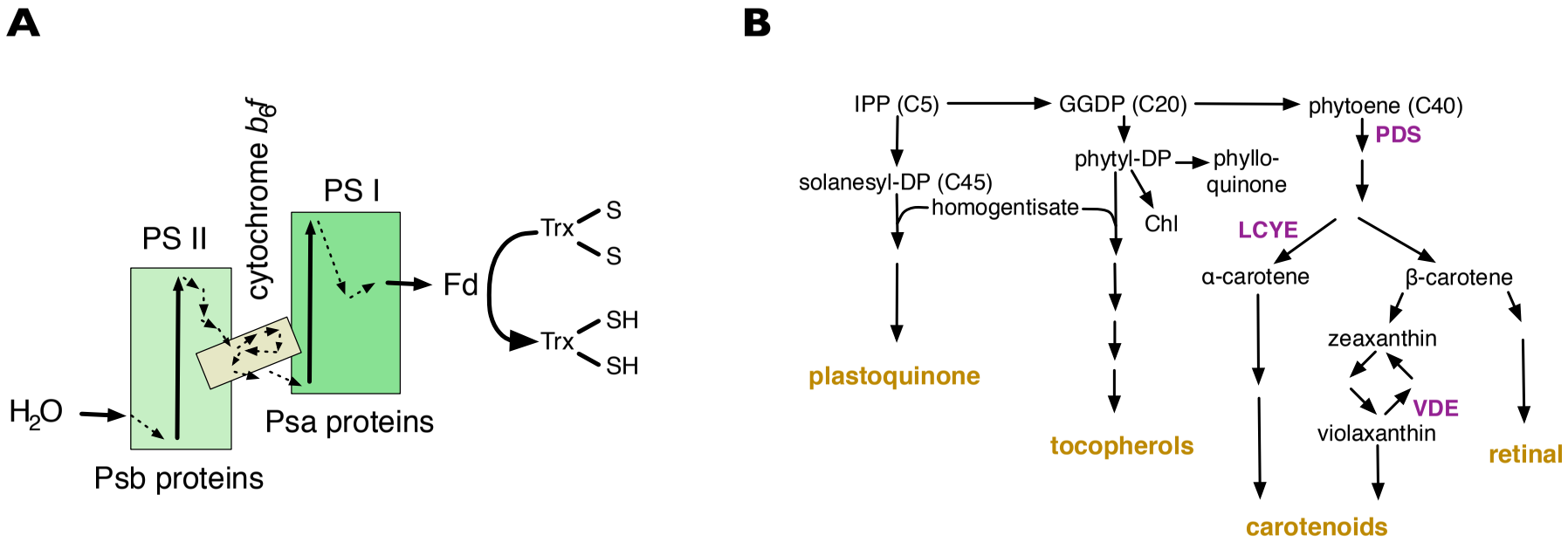


Fig. S2-S18. Features of genome organization: Each Linkage Group is depicted as a long horizontal rod, with genetically-mapped scaffolds shown as open rectangles below (the scaffold number is under each scaffold and arrows indicate orientation where determined; the reverse strand is assumed where orientation is not known). The scale of each map is determined by molecular lengths of the mapped scaffolds. Short and long red ticks are drawn on scaffolds every 0.2 Mb and 1.0 Mb, respectively. We assumed small 50 kb gaps between scaffolds, except where there is genetic evidence of a larger gap (e.g. see Linkage Group X). Genetic distances between markers (cM), where they are known, are shown by two-headed arrows above the scaffold. Genomic regions are labeled below the scaffolds: 5S, rDNA, mito (insertion of mitochondrial DNA), T (telomere), Cp (chloroplast DNA insertion). *Chlamydomonas* genes with homologs in other organisms/lineages ("Cuts" are defined in the text and Fig. 5) are shown as tracks of vertical bars: light red, genes shared between *Chlamydomonas* and humans, but not occurring in non-ciliated organisms; dark red, genes in "CiliaCut"; light green, genes shared between *Chlamydomonas* and *Arabidopsis*, but not in non-photosynthetic organisms; dark green, genes in "GreenCut" ; magenta, predicted tRNAs, including those that represent SINE sequences; dark blue, snoRNAs. Below, on separate axes, are features of the genomic sequence (in 25 kb windows): %GC (grey), gene density (red), transposable element (TE) density (blue), and simple repeat (Rep) density (teal). The %GC graph includes horizontal lines denoting 25, 50 and 75% GC. The other three graphs show a mean (solid horizontal line) and +/– SD (dashed horizontal line) for the scaffold, and are scaled to the densest region on any of the mapped scaffolds, which are as follows: gene density, 12 per 25 kb window; TE density, 44 per 25 kb window; repeat density, 46 per 25 kb window.

Fig S2. Overview of linkage group I

Fig S3. Overview of linkage group II.

Fig. S4. Overview of linkage group III.

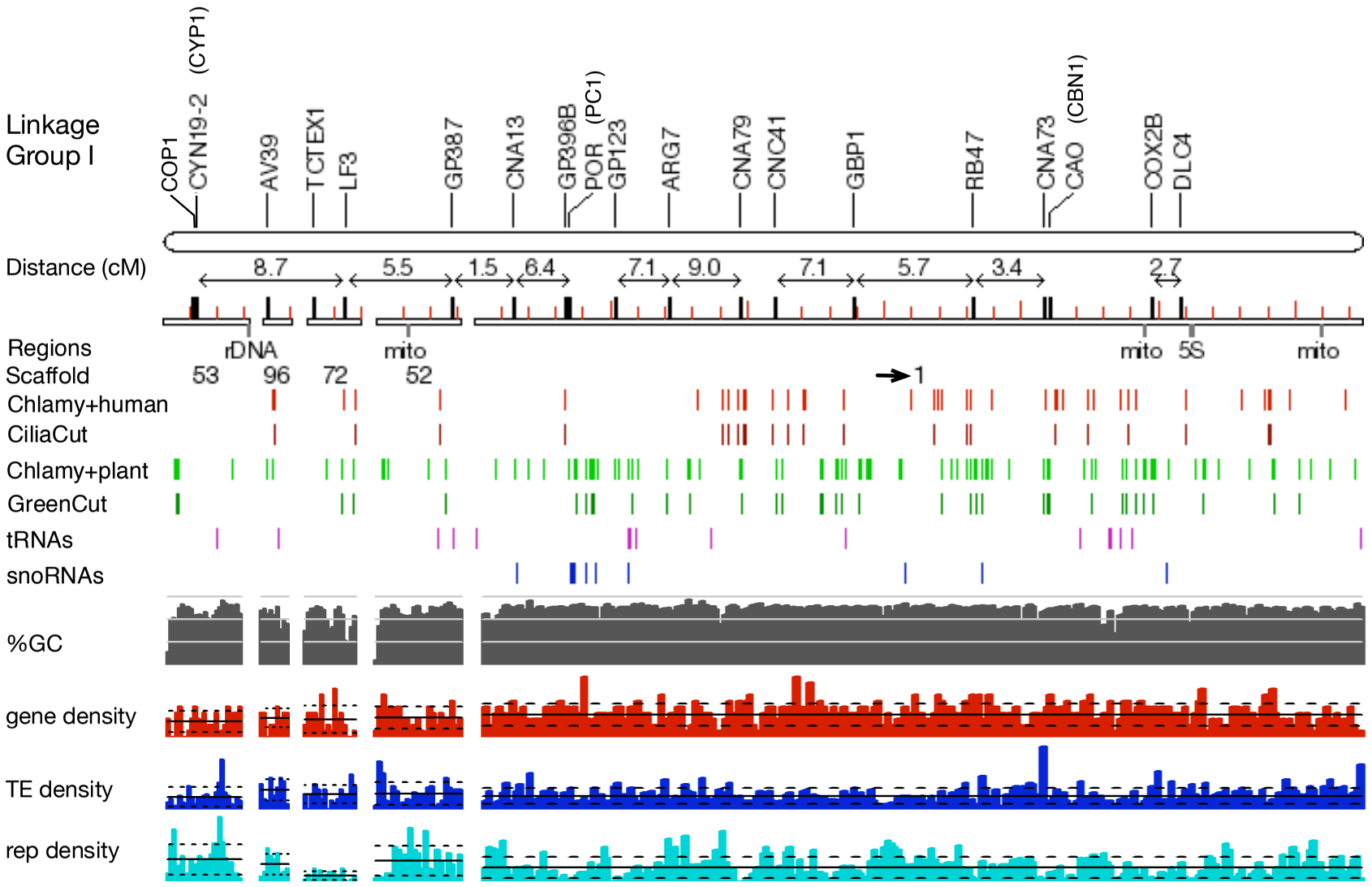
Fig. S5. Overview of linkage group IV.

Fig. S6. Overview of linkage group V.

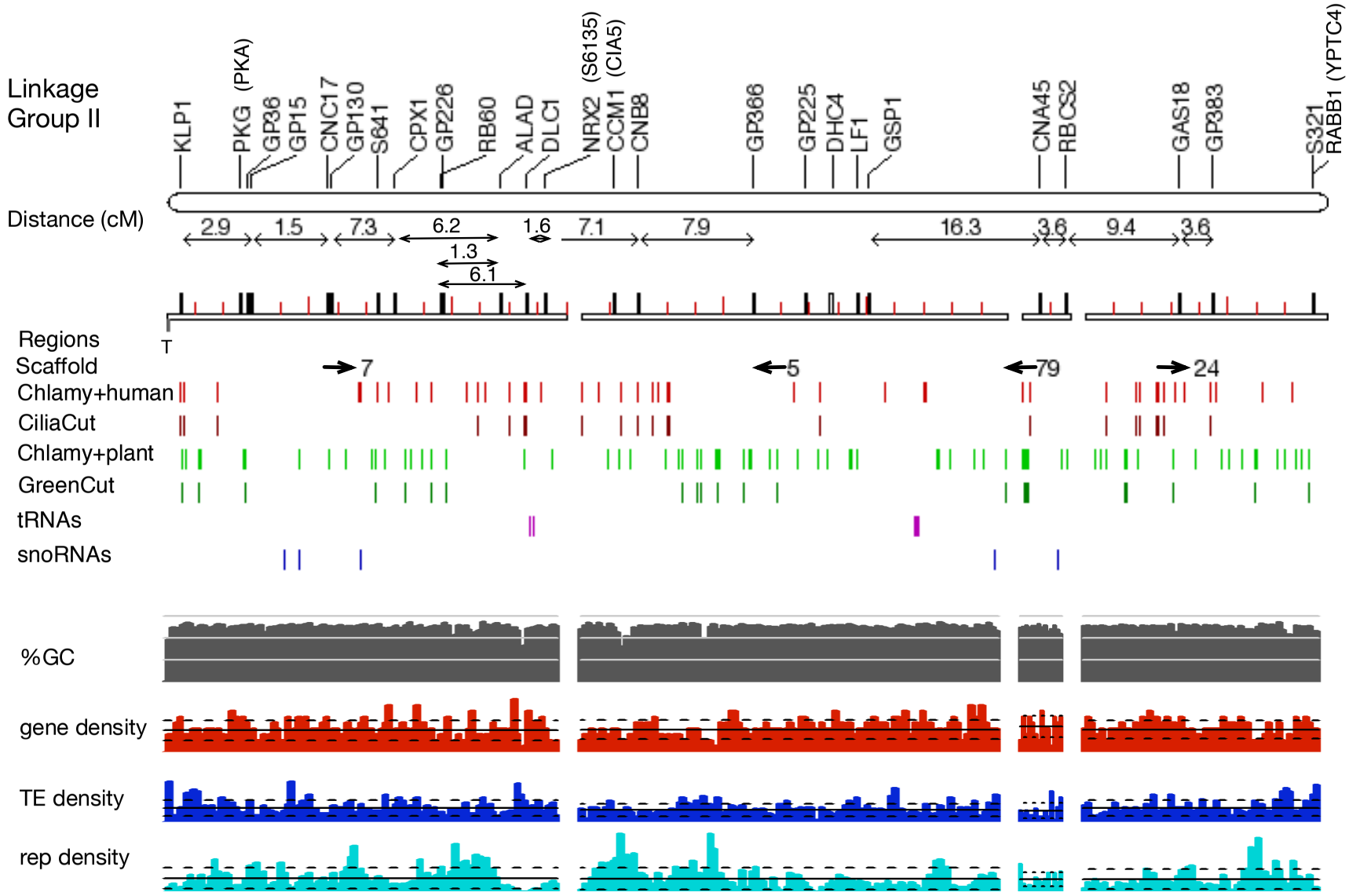
Fig. S7. Overview of linkage group VI.

Fig. S8. Overview of linkage group VII.

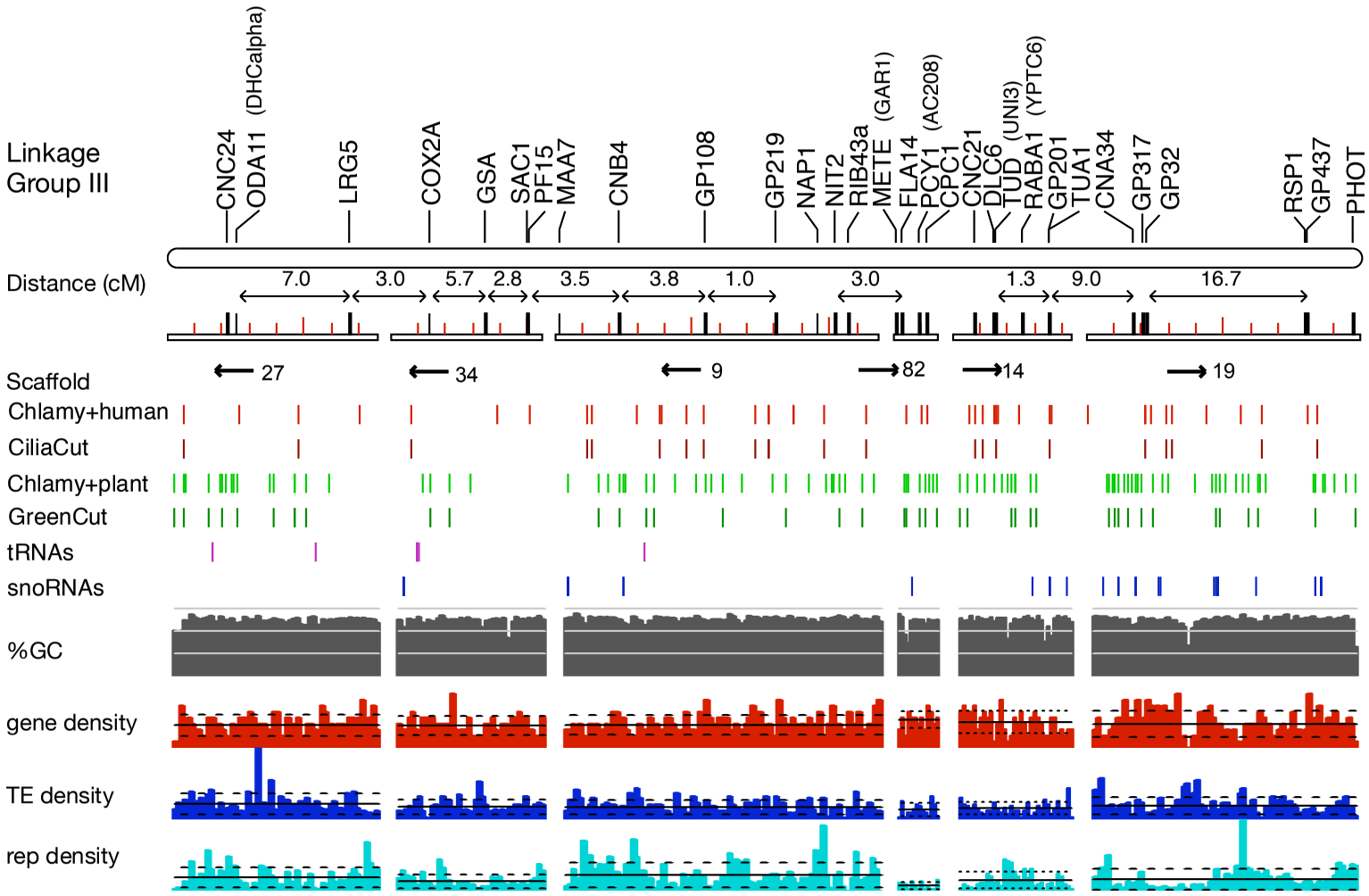
Supplemental Fig 2



Supplemental Fig 3



Supplemental Fig 4



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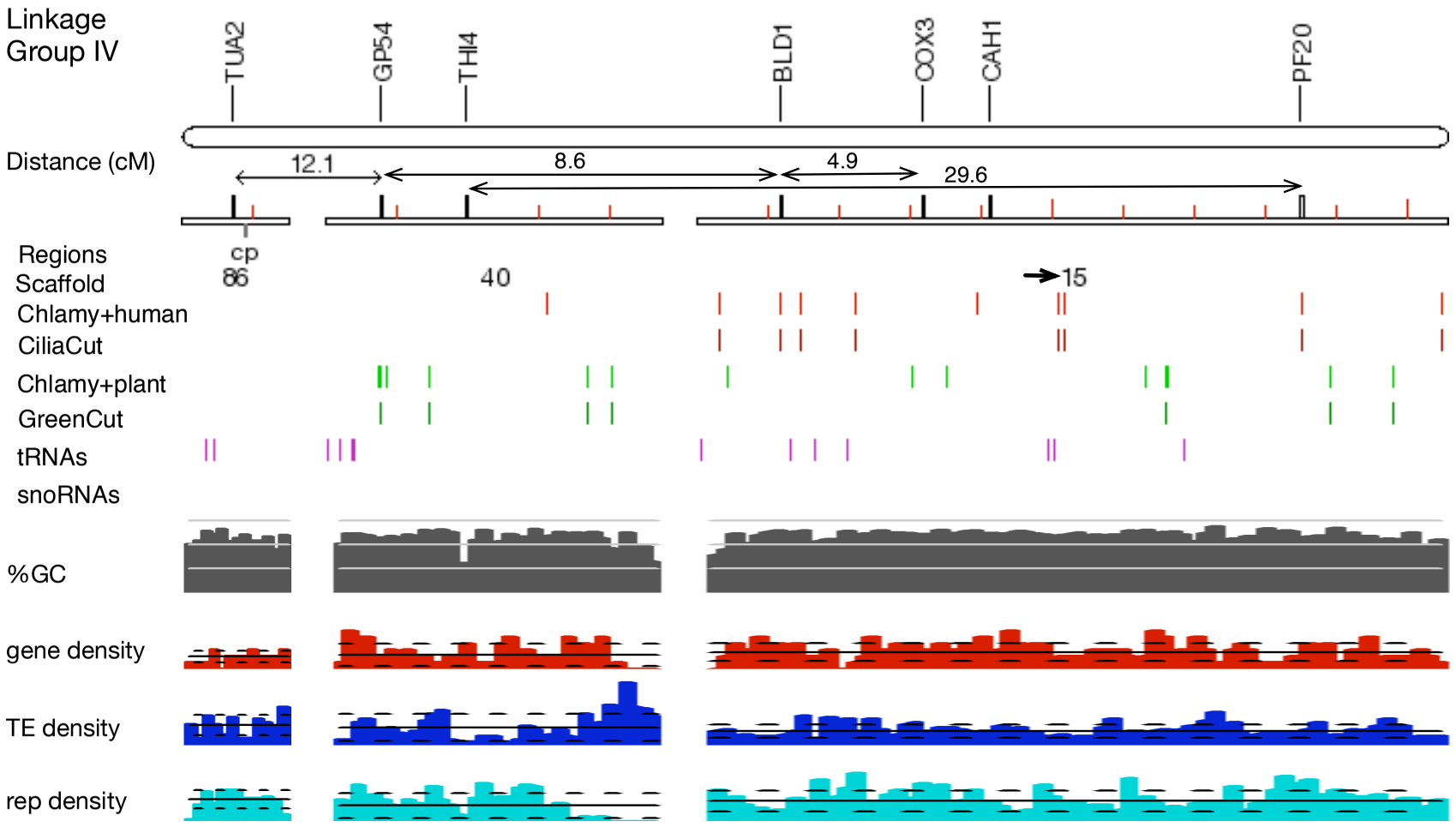
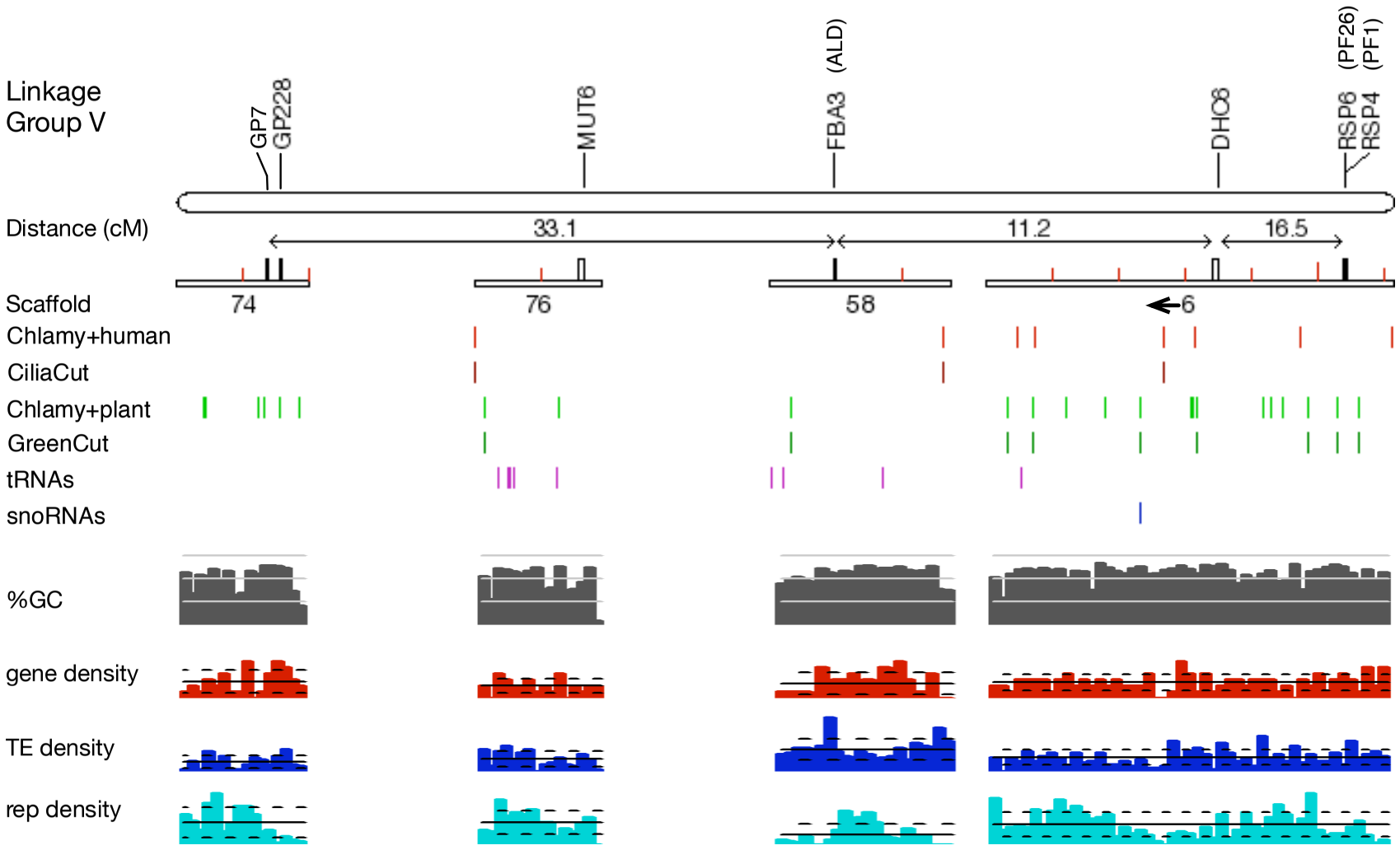
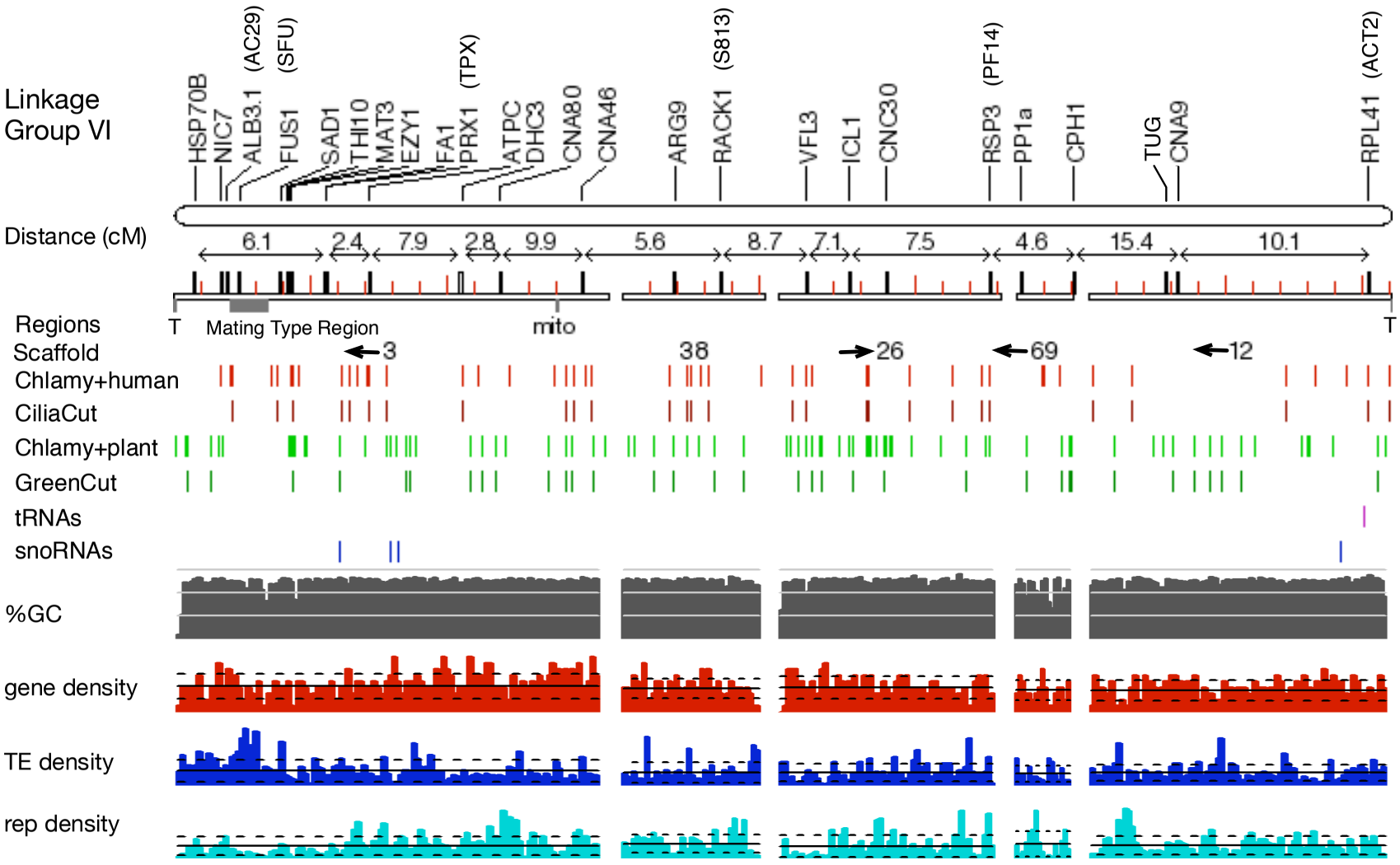


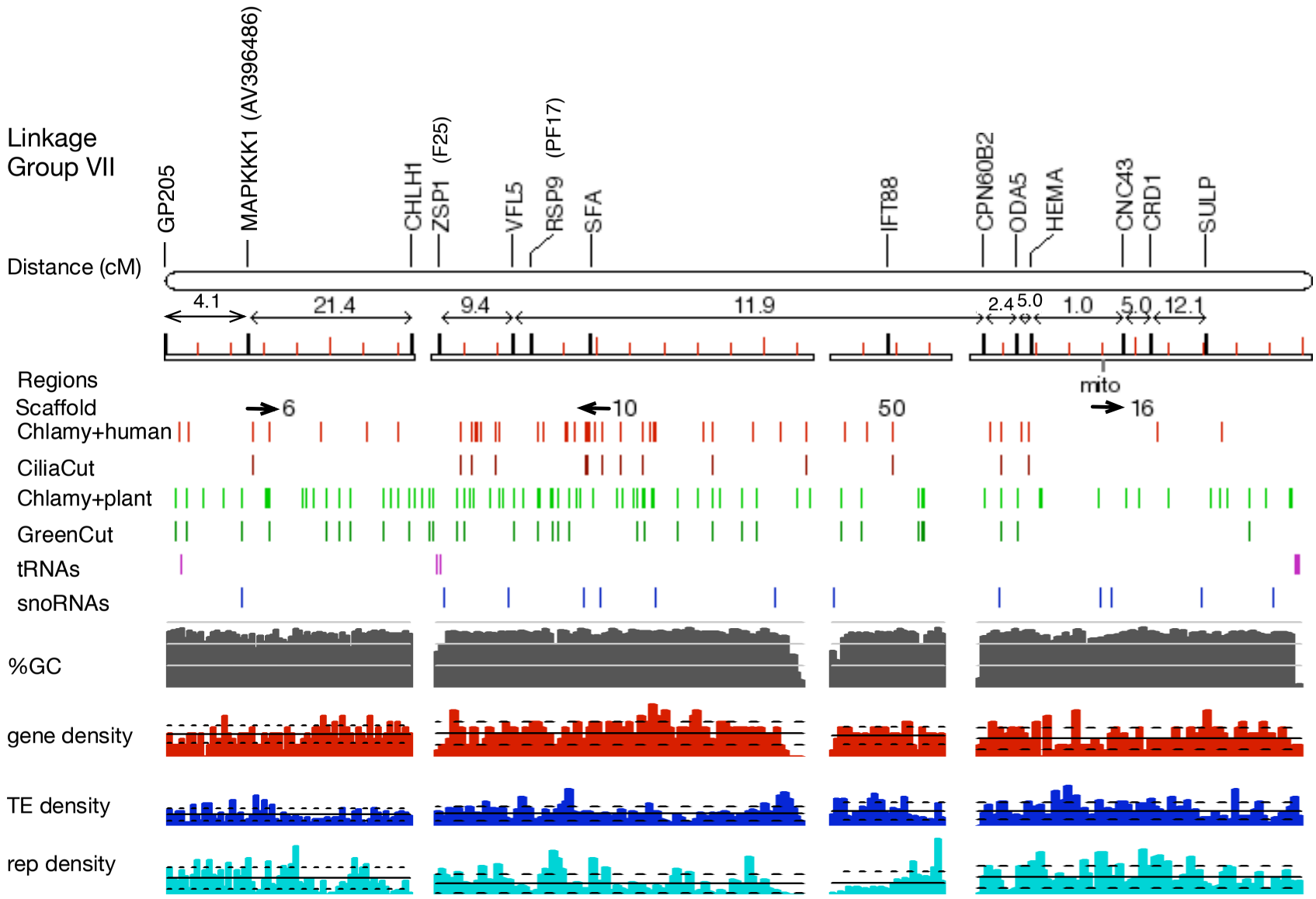
Supplemental Fig 6



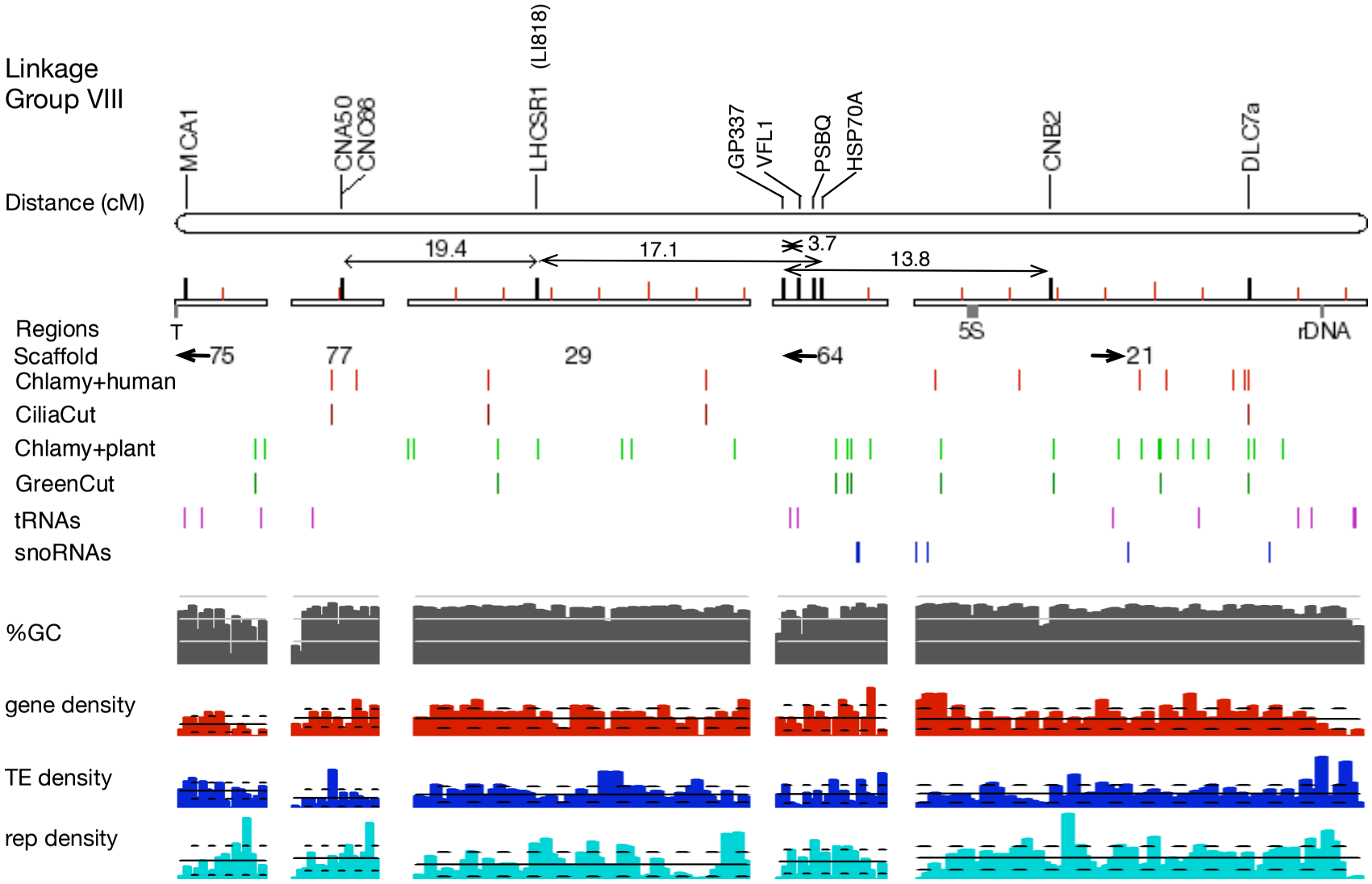
Supplemental Fig 7



Supplemental Fig 8



Supplemental Fig 9



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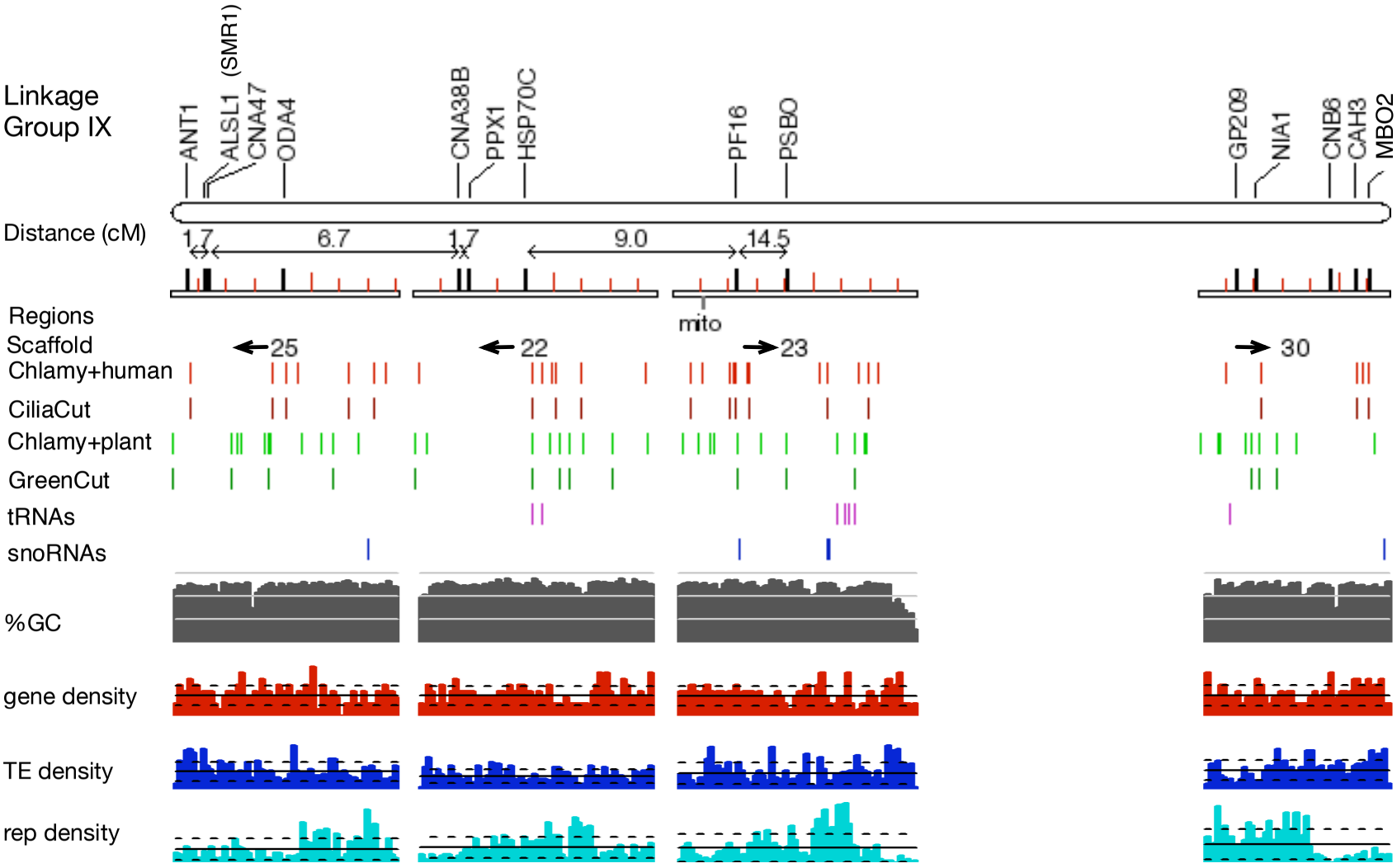

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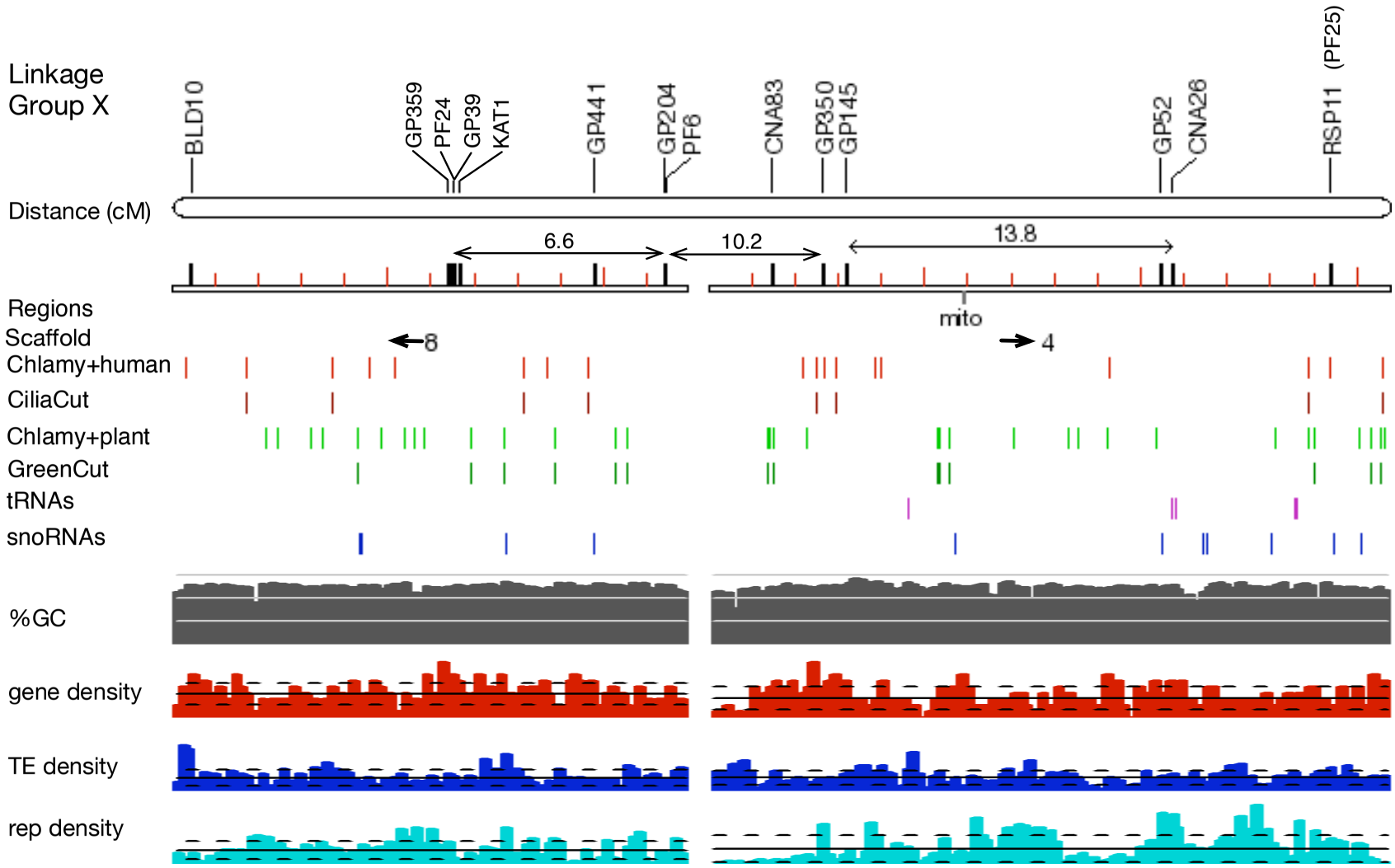
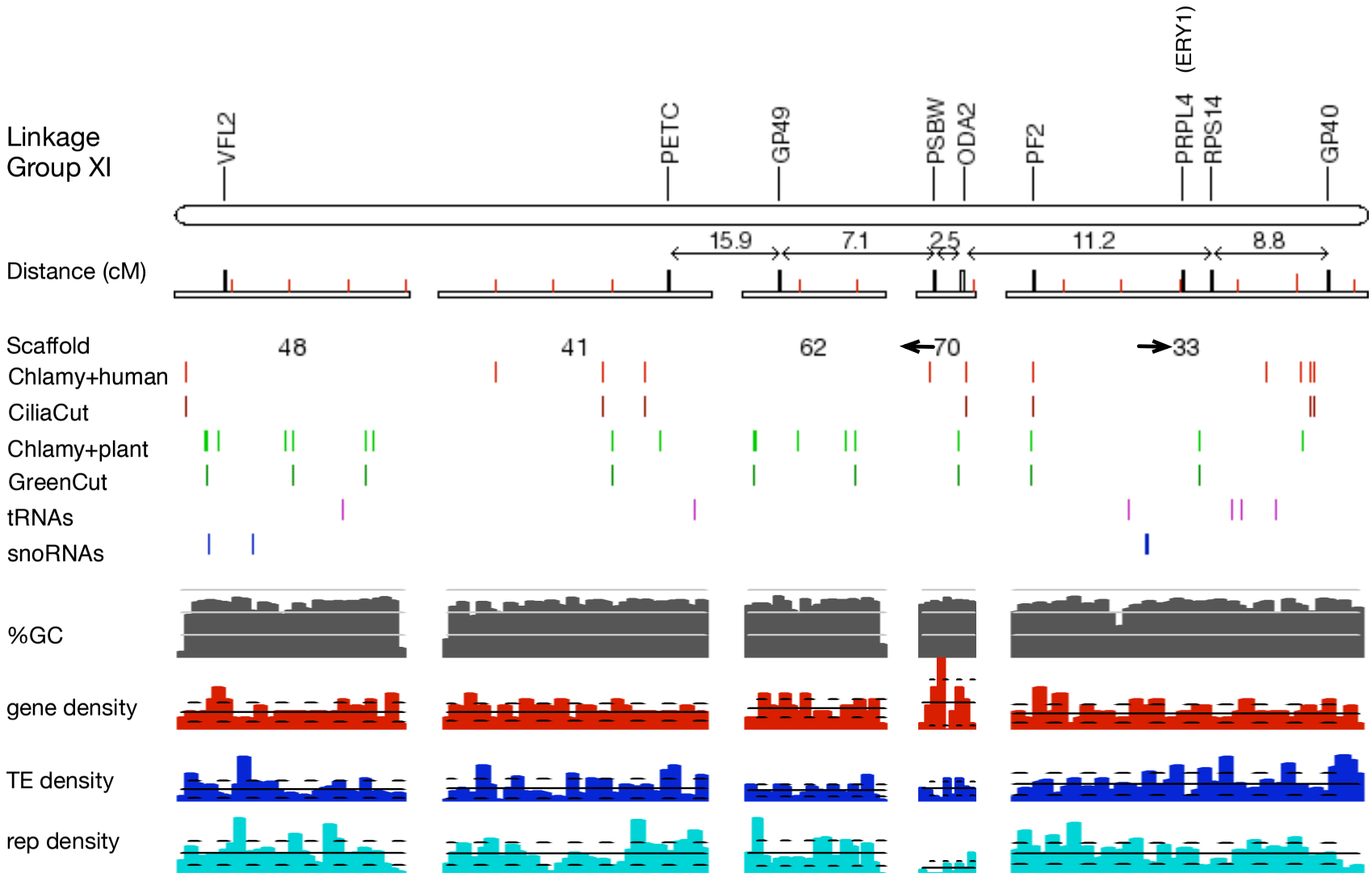
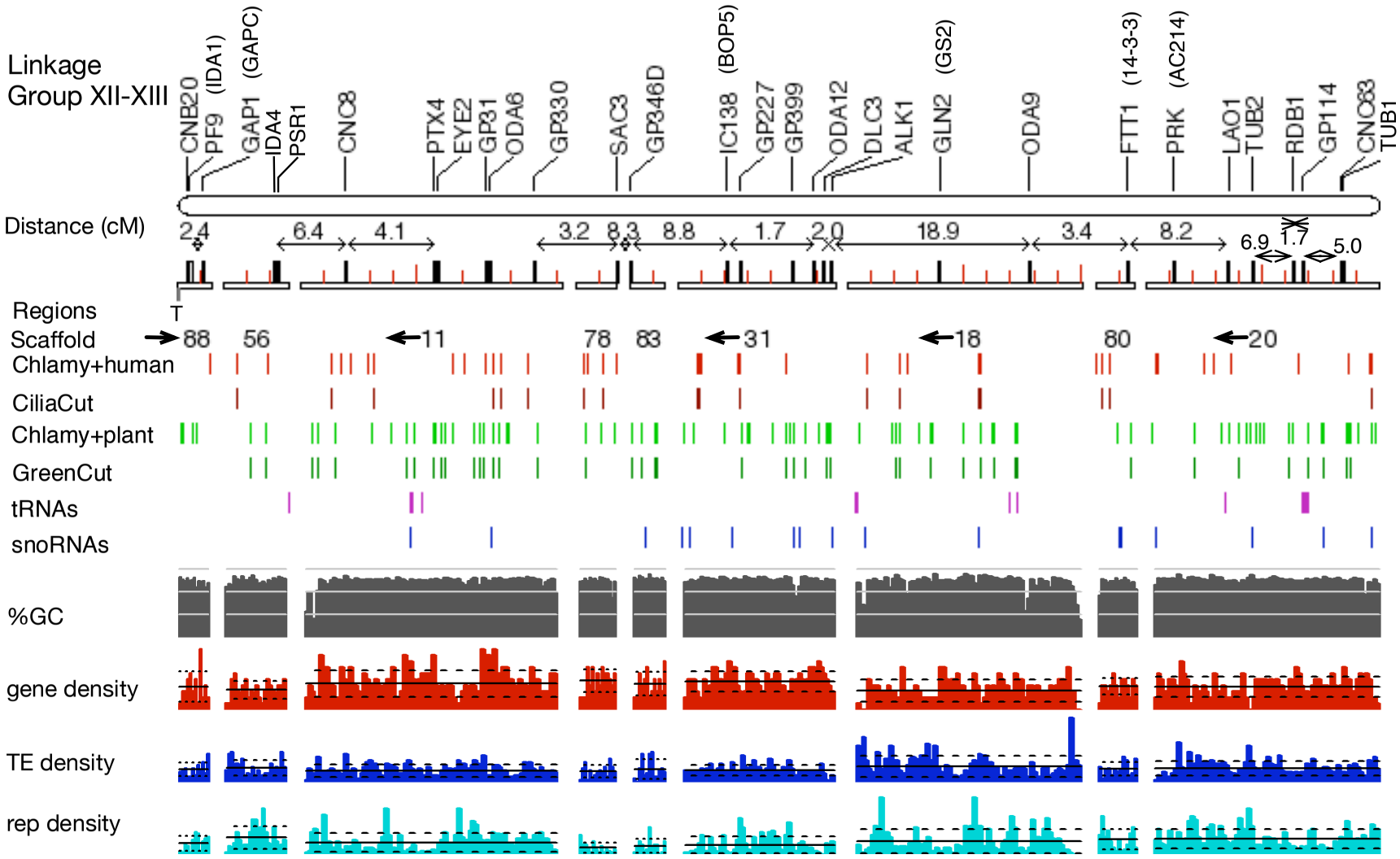


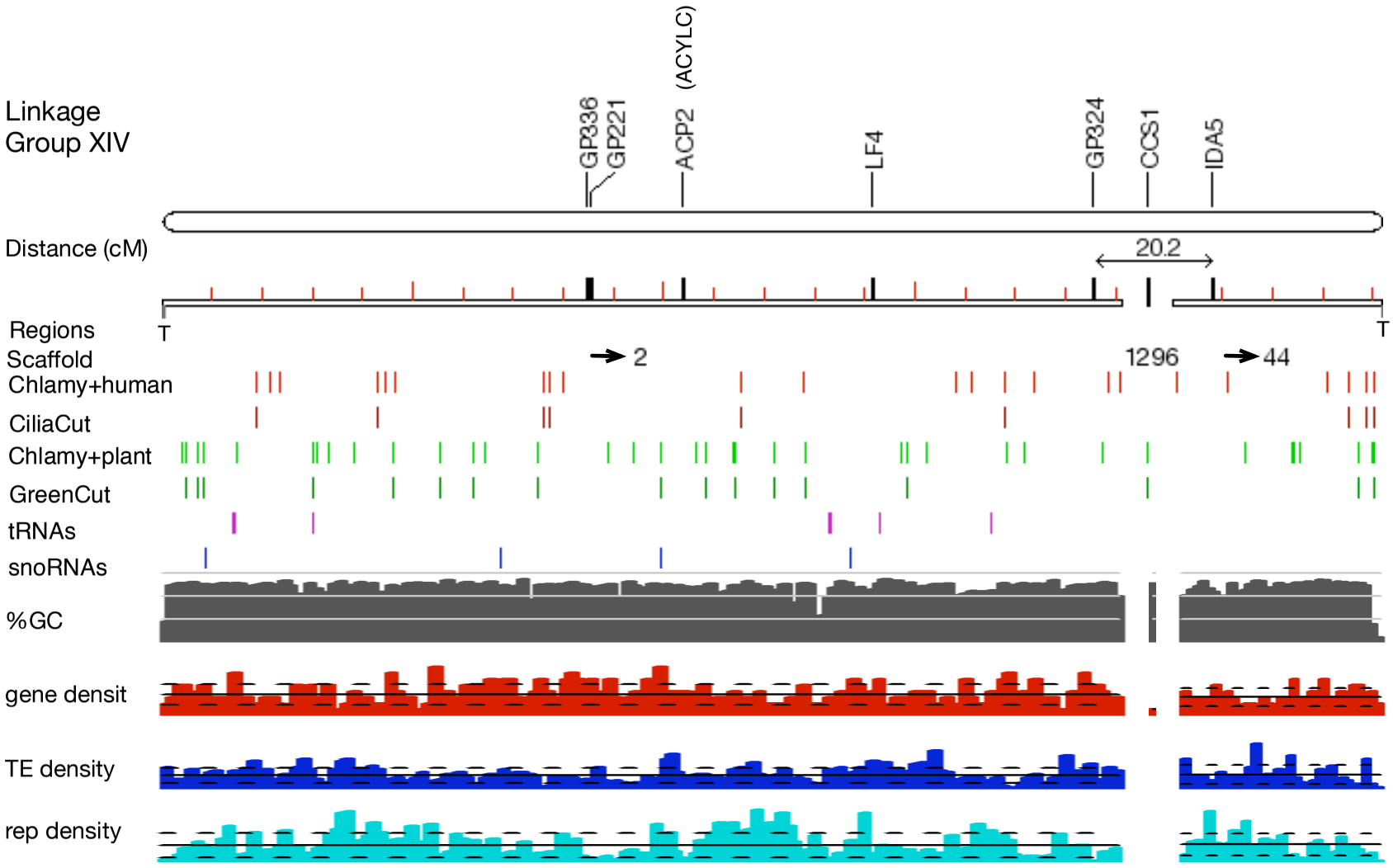
Supplemental Fig 12



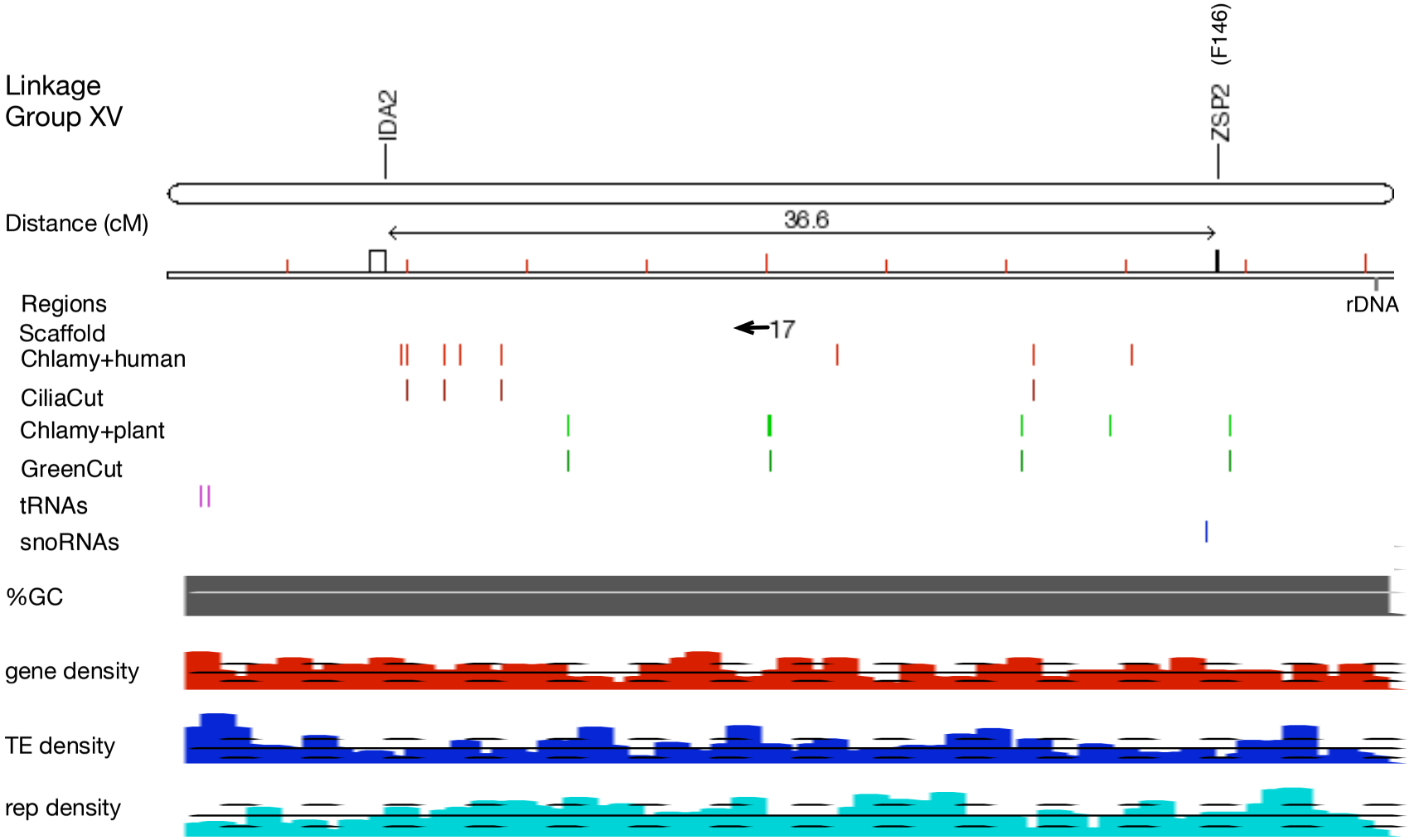
Supplemental Fig 13



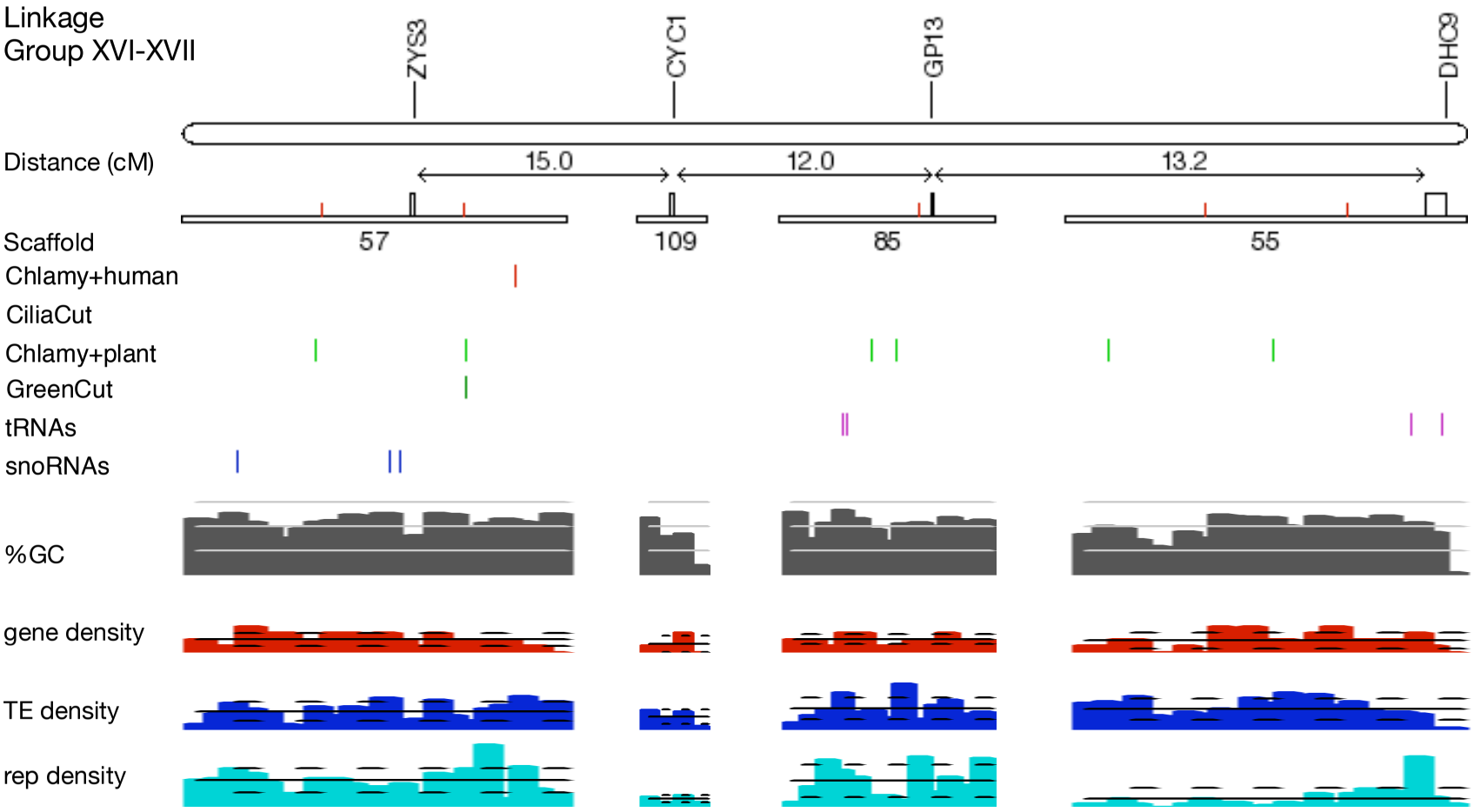
Supplemental Fig 14



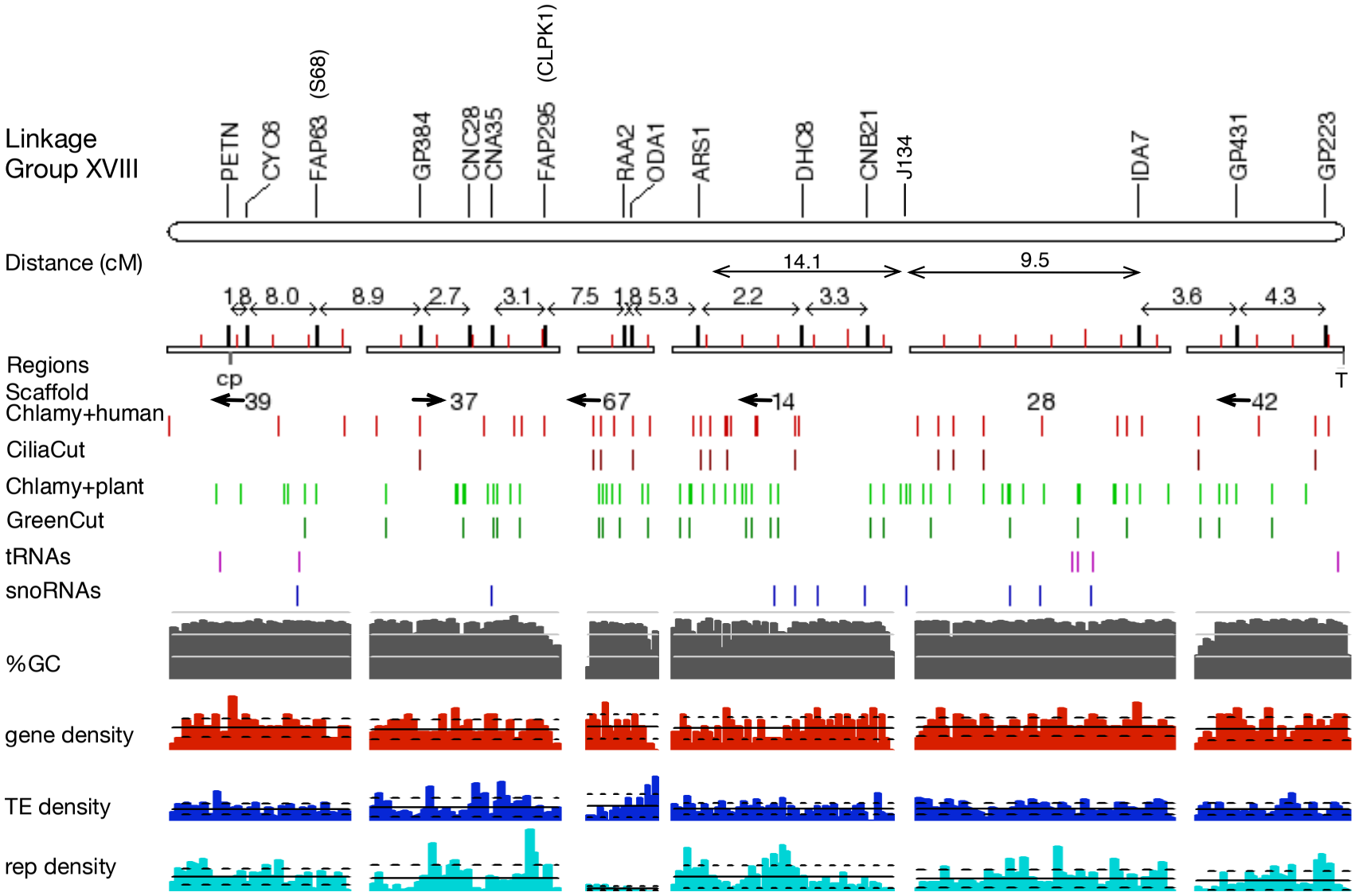
Supplemental Fig 15



Supplemental Fig 16



Supplemental Fig 17



Supplemental Fig 18

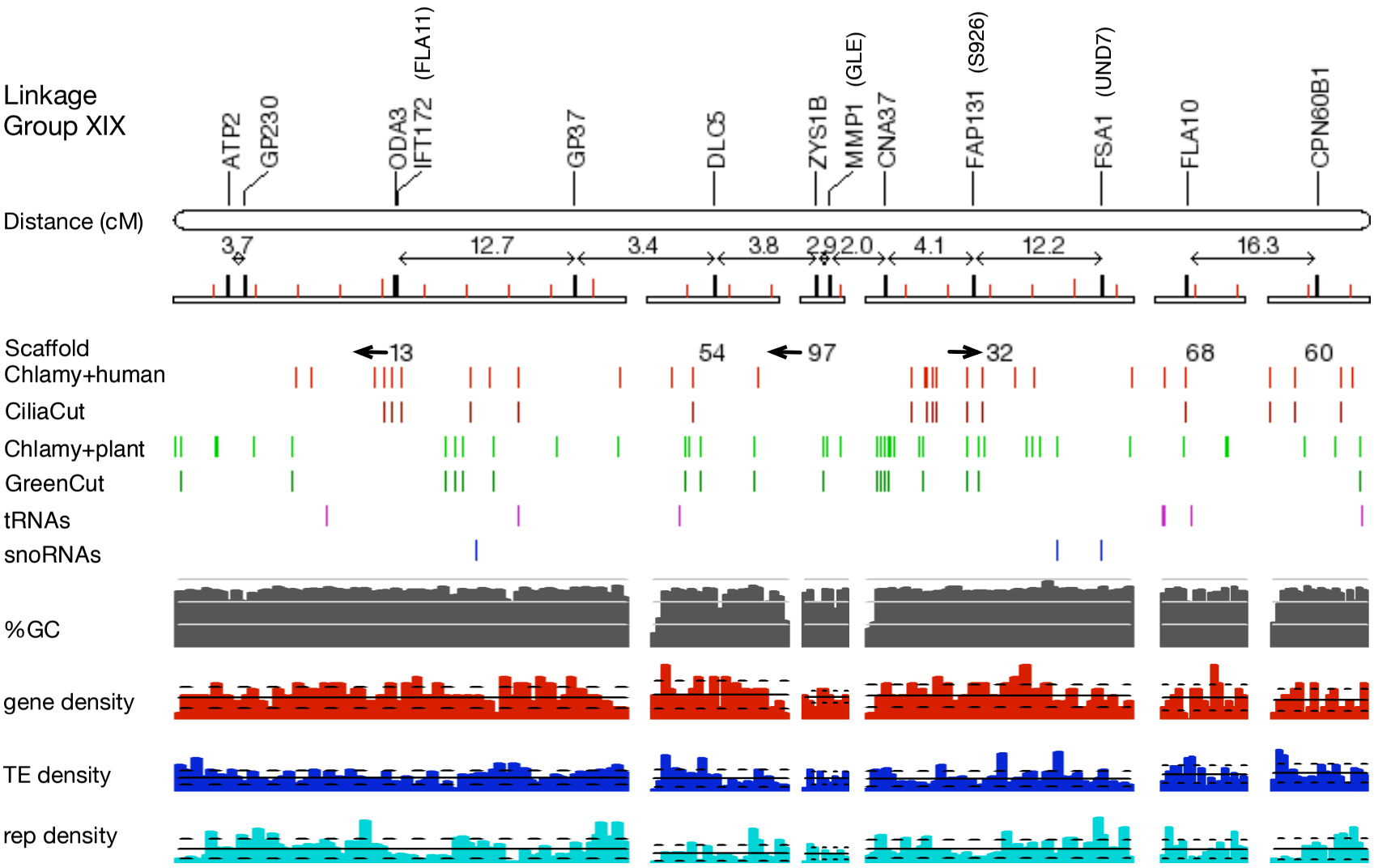
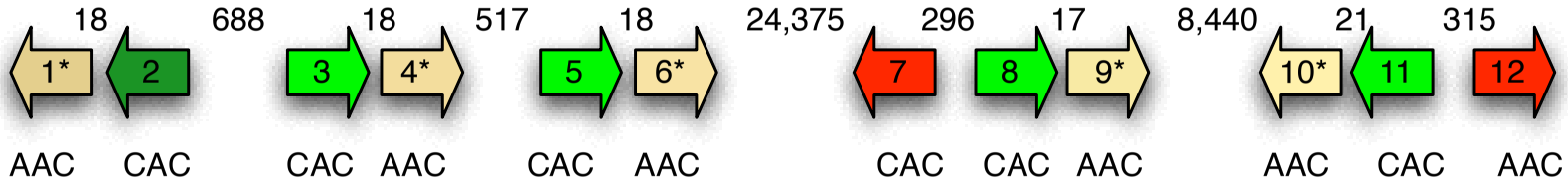


Fig. S19. Intron evolution in tRNA-Val cluster: (**A**) The 12 tRNAs, numbered consecutively, on scaffold 20:1350500-1386900 (LG XII-XIII) are depicted as arrows that indicate orientation on the chromosome, and color indicating those tRNAs that share sequence similarity (especially in the introns; see Fig. S19B). The spacing in bp between the tRNAs is indicated by the numbers above the intergenic regions. The anticodon is shown below each gene, and the asterisk within the arrow indicates that the tRNA has a genome-encoded CCA. (**B**) A neighbor-joining tree of the tRNA intron sequences with sequence differences between introns of the paired genes highlighted in bold black.

Supplemental Fig 19

A



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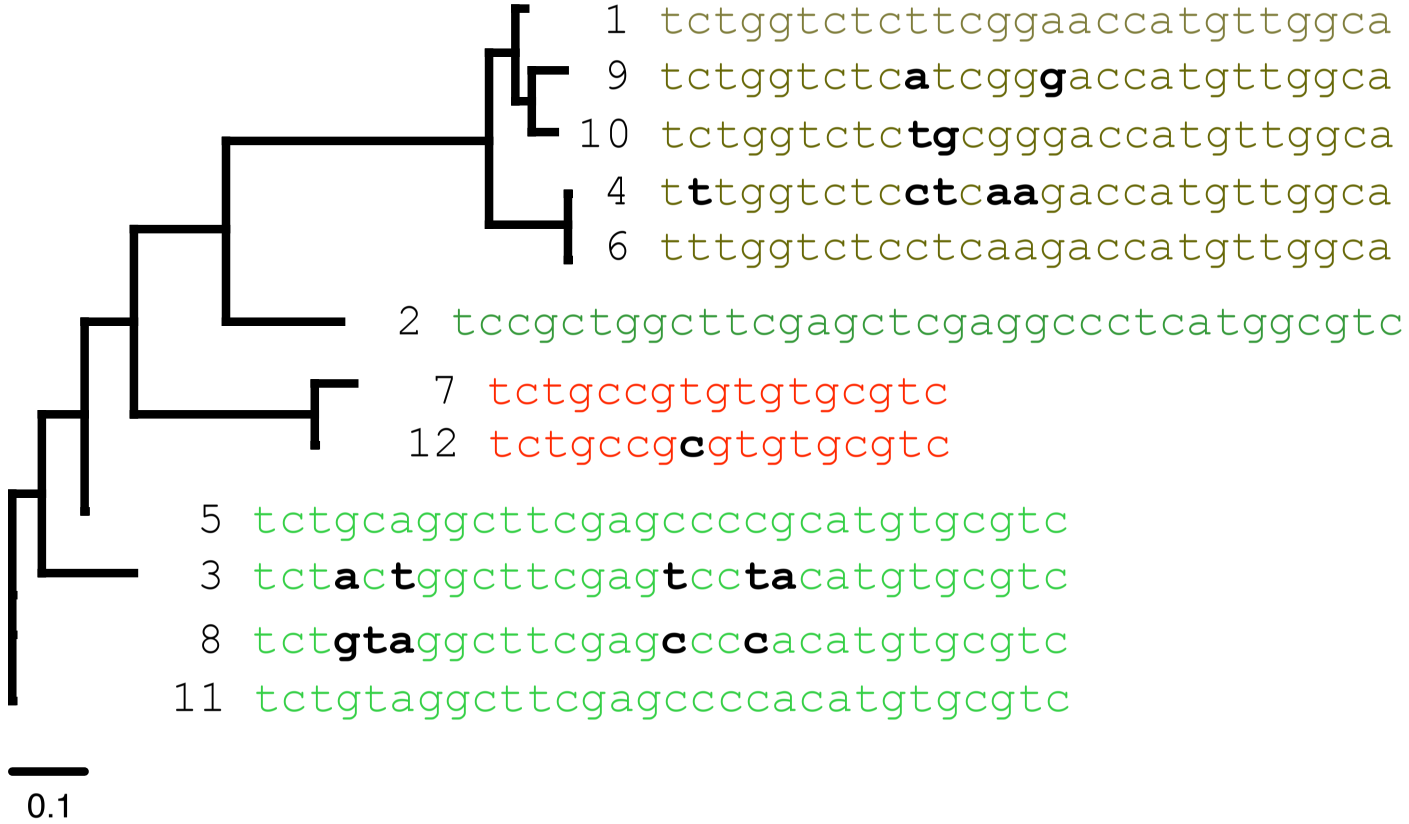


Fig. S20. The carbon concentrating mechanism region: The ~100 kb region of the genome (scaffold 15) that contains several genes associated with the carbon concentrating mechanism (CCM). Arrows are used to depict the different genes and their lengths and orientations and each gene is labeled with a JGI Chlre.v3.0 protein ID and gene name (where one has been assigned). Coordinates (bp) on scaffold 15 are shown along the line at the top. The red arrows depict the six CCM genes (*CCP2*, *LCID*, *CAH2*, *CAH1*, *LCIE* and *CCP1*), which were identified from both sequence and experimental data. The arrangement of the genes suggests three recent duplications. Neighboring and intervening genes are shown as open arrows. On the lower portion, red dashed lines connect the duplicated CCM sequences, with % nucleotide identity shown in boxes. One additional gene pair of unknown function in this region shows significant paralogy (black dashed lines connecting 170976 & 189424).

Supplemental Fig 20

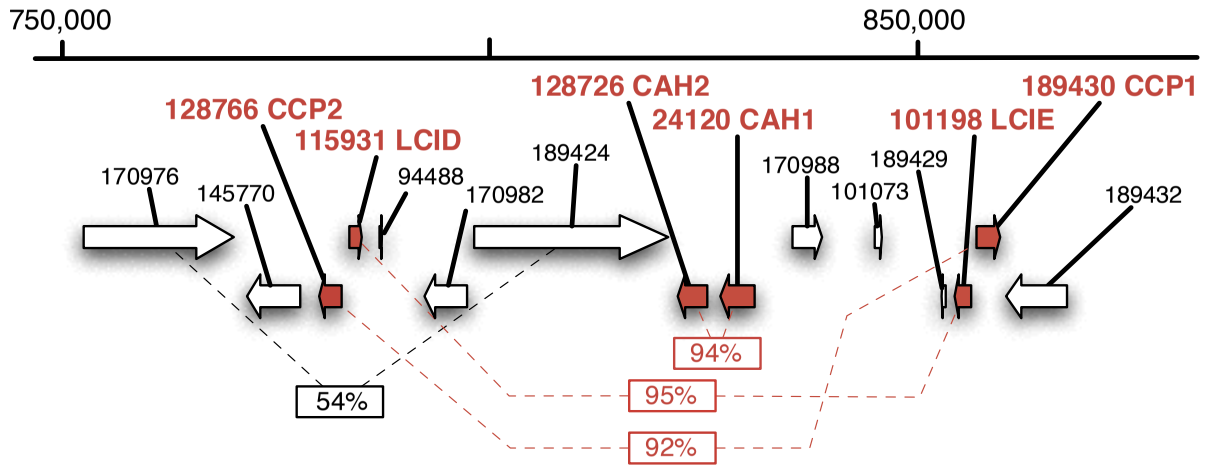
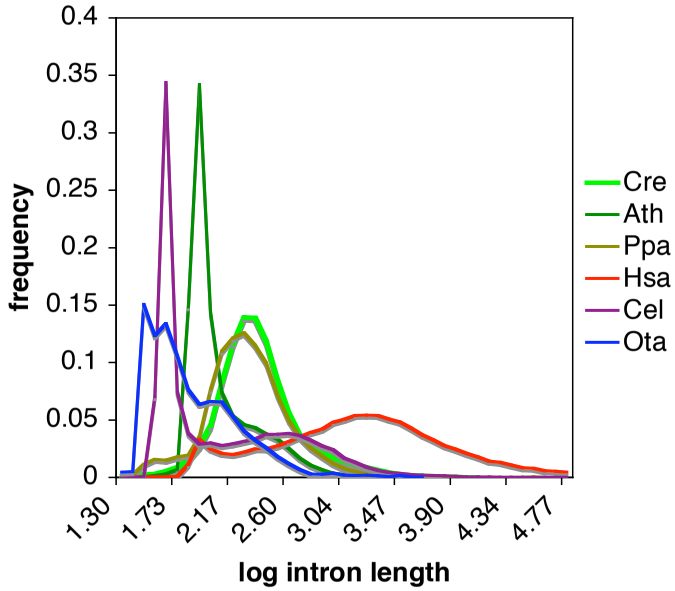


Fig. S21. Comparison of *Chlamydomonas* intron characteristics to those of other eukaryotes: Introns were collected from the genomes of the organisms listed (see Fig. 2), and graphs were plotted of (**A**) the log lengths of the introns against frequency in the genome, or (**B**) the average length for introns in each of the organisms against the average number of introns.

Supplemental Fig 21

A



B

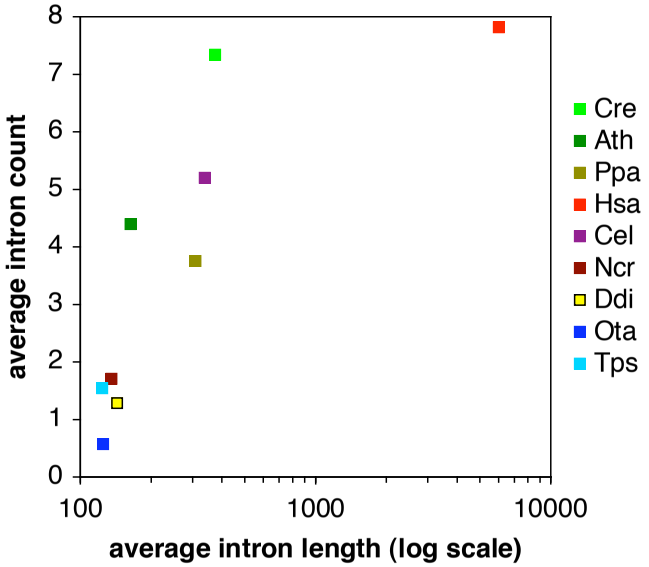


Fig. S22. Summary of transporter families: Transporter families (described along the top of the figure; the abbreviations can be found at (*44*)) that are present in organisms or groups of organisms listed on the left are colored with a red box. The criterion used for identification of the transporters is described in the **MATERIALS AND METHODS** section of this text. Families of transporters present in *Chlamydomonas* are highlighted with a horizontal green bar. Transporter families and organisms were automatically clustered hierarchically to generate the order in which they are displayed, and then grouped by coarse phylogenetic (vertical) and transporter superfamily (horizontal) membership. The analysis has been performed for transporter families present in animals (*H. sapiens*, *C. elegans*, *D. melanogaster*, *A. gambiae*), various single cell eukaryotes (sing euk: *E. histolytica* HM1:IMSS, *C. parvum* genotype 2 isolate, *E. cuniculi*, *T. parva*, *P. falciparum* 3D7, *P. vivax*, *T. thermophila* SB210, *T. brucei* TREU927/4 GUTat10.1, *L. major Friedlin*, *T. cruzi* CL Brener TC3, *T. whippelii* TW08/27, *T. whipplei* Twist), fungi (*S. pombe*, *A. oryzae*, *C. posadasii* C735, *A. nidulans* FGSC-A26, *A. fumigatus* Af293, *N. crassa* 74-OR23-IVA, *C. neoformans*, *S. cerevisiae* S288C), amoeba (*D. discoideum*), land plants (*O. sativa*, *A. thaliana*), *Ostreococcus* spp. (ostreo), *Chlamydomonas* (chlamy), the red alga *C. merolae 10D* and the diatom *Thalassiosira* (red alg+diat), and 220 bacteria. The color shows the proportion of species within the group that have genes for members of the indicated transporter family: black (family absent in all species); bright red (family present in all species); intermediate red color (family present in some species).

Supplemental Fig 22

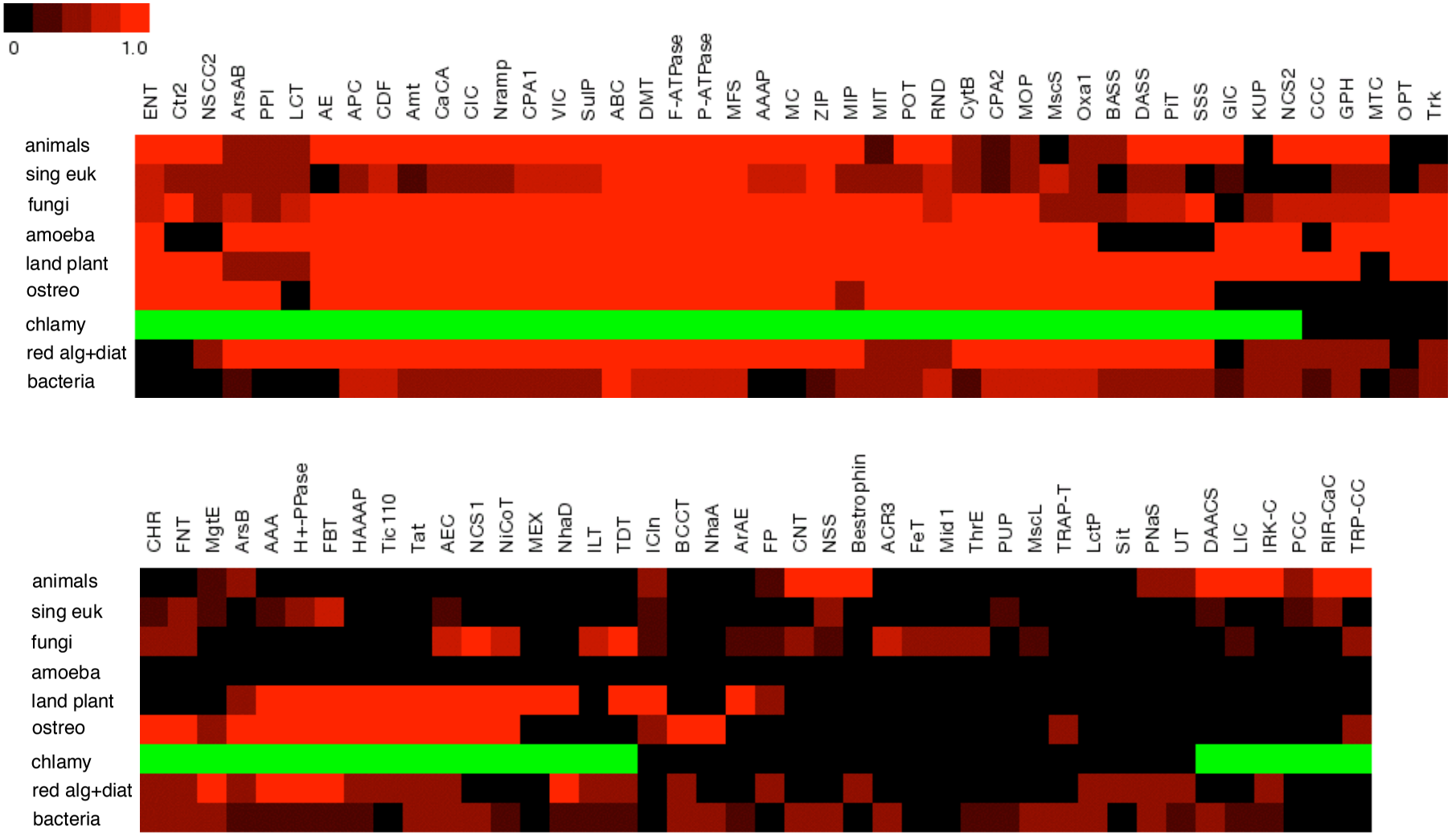


Fig. S23. Complete repertoire of transporter families: Details of clustering of transporter families across bacteria and eukaryotes are shown (summarized in Fig S23). Organisms are in rows; transporter families in columns. Euclidean distance clustering was performed in both dimensions. Red indicates presence of a transporter family; black, absence.

Supplemental Fig 23

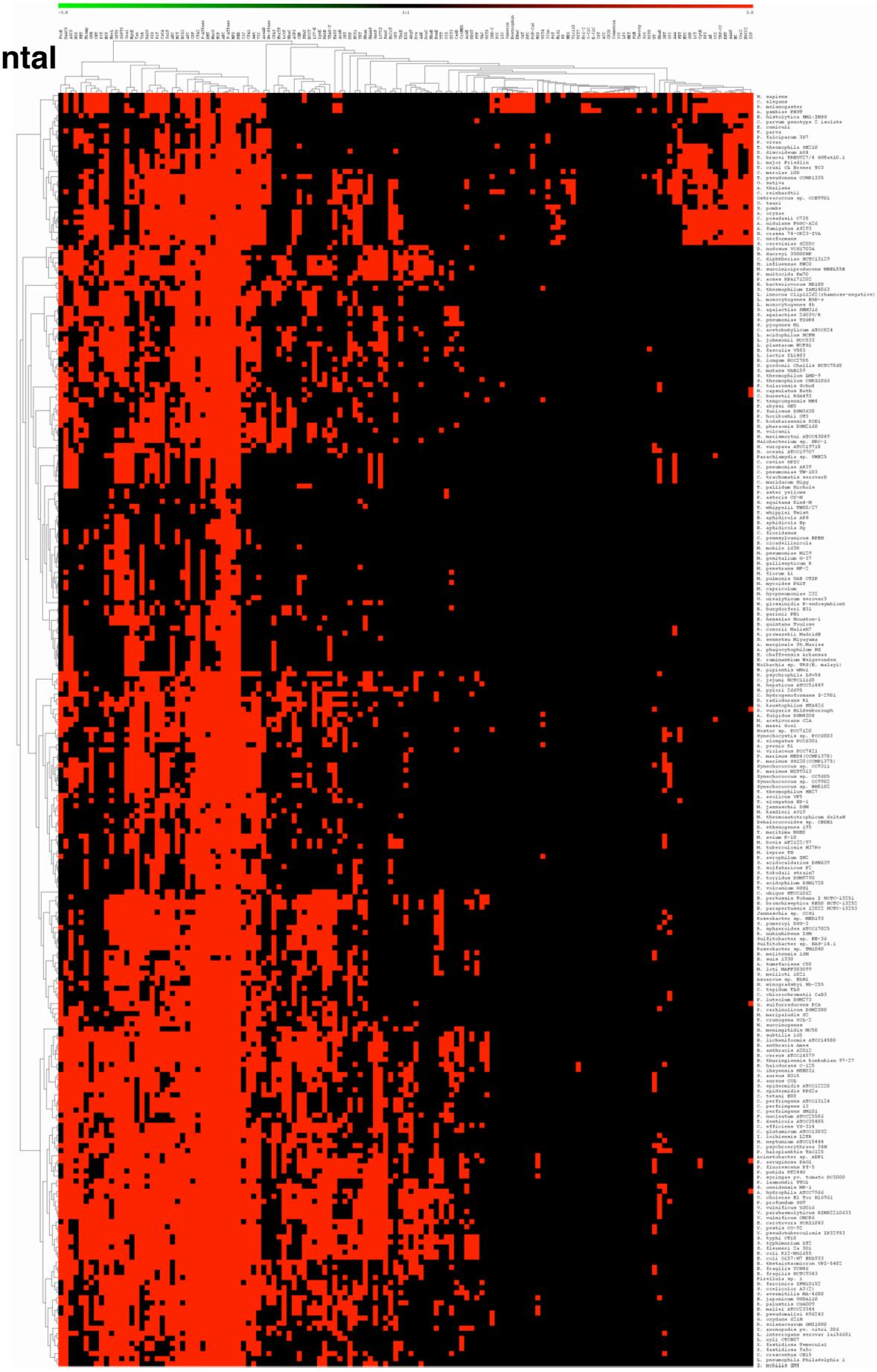


Fig. S24. Classification of CiliaCut proteins: Functional classification of CiliaCut proteins by manual annotation. Classification was based on the published function of characterized protein family members (if any), and/or the molecular function of predicted PFAM domains. 125 (67%) of the CiliaCut proteins were successfully classified; the remaining 80 either were not associated with functional information or the functional information available was ambiguous and is not included.

Supplemental Fig 24

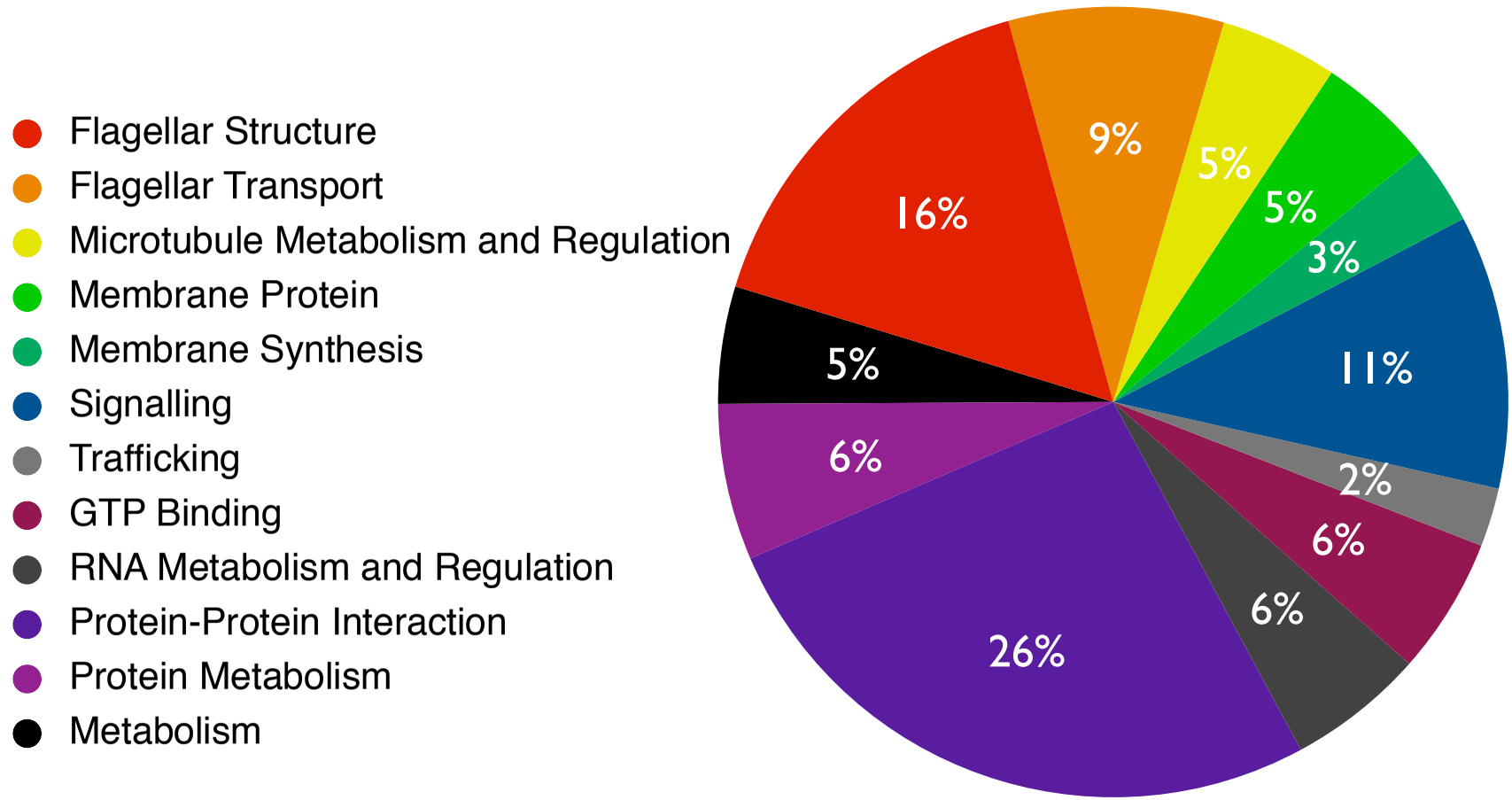
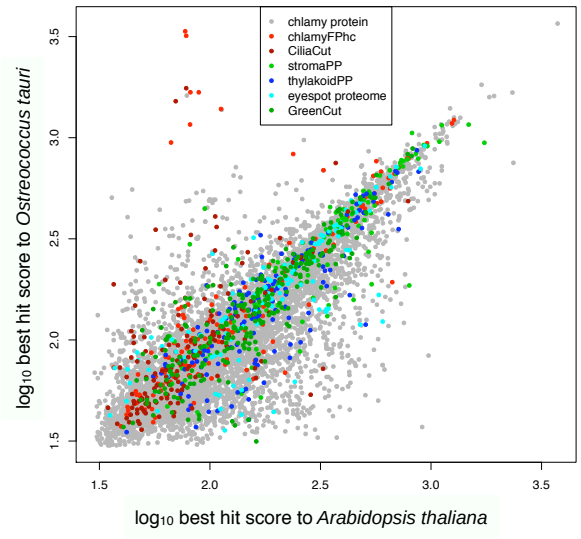


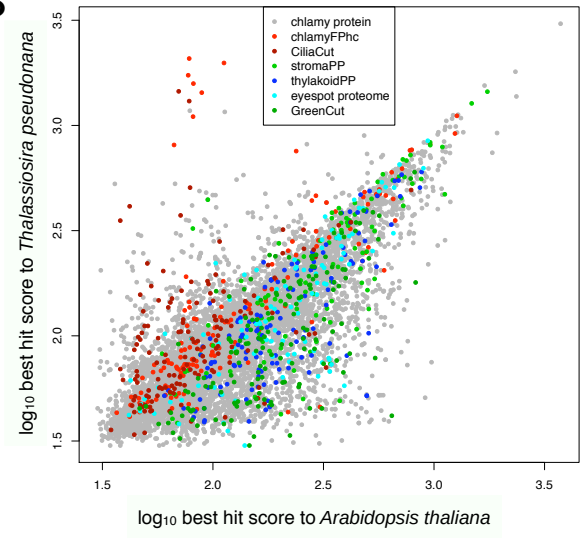
Fig. S25. Best hit scatter plots: Each *Chlamydomonas* protein is plotted by $\log_{10}$ of its best blast hit score to (**A**) *Arabidopsis*, *Ostreococcus tauri*; (**B**) *Arabidopsis*, *Thalassiosira*; (**C**) *Thalassiosira*, *Ostreococcus tauri*. Proteins are grey or colored by membership of functional or comparative genomic grouping: *Chlamydomonas* Flagellar Proteome (*67*) high confidence set (ChlamyFP, red); Stroma Plastid Proteome (stromaPP, green); Thylakoid Plastid Proteome (thylakoidPP, blue); *Chlamydomonas* PS cut7 (cyan); *Chlamydomonas* eyespot proteome (yellow).

Supplemental Fig 25

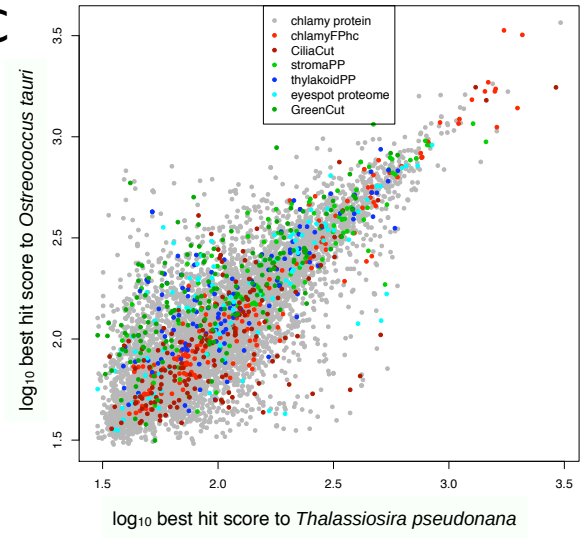
A



B



C



4. SUPPORTING TABLES

Four anticodon amino acids

amino acid	anticodon				total
Ala	AGC	GGC	CGC	TGC	
	13		10	5	28
Gly	ACC	GCC	CCC	TCC	
		17	1	1	19
Pro	AGG	GGG	CGG	TGG	
	13		6	1	20
Thr	AGT	GGT	CGT	TGT	
	6		3	2	11
Val	AAC	GAC	CAC	TAC	
	7		10	1	18

Six anticodon amino acids

amino acid	anticodon						total
Ser	AGA	GGA	CGA	TGA	ACT	GCT	
	5		5	1		8	19
Arg	ACG	GCG	CCG	TCG	TCT	CCT	
	11		3	1	1	2	18
Leu	AAG	GAG	CAG	TAG	TAA	CAA	
	3		10	1	1	2	17

Two anticodon amino acids

amino acid	anticodon		total
Phe	AAA	GAA	
		9	9
Asn	ATT	GTT	
		7	7
Lys	CTT	TTT	
	11	1	12
Asp	GTC	ATC	
	11		11
Tyr	ATA	GTA	
		8	8

Cys	ACA	GCA	
		7	7
Glu	CTC	TTC	
	13	1	14
His	ATG	GTG	
		5	5
Gln	CTG	TTG	
	6	1	7

Other amino acids

amino acid	anticodon			total
Meti	CAT			
	8			8
Mete	CAT			
	6			
Ile	AAT	GAT	TAT	
	7	1	1	9
SeC	TCA			
	1			1
Trp	CCA			
	5			5

Table S1. Summary of tRNA complement of *Chlamydomonas*: The 259 tRNAs encoded on the *Chlamydomonas* genome are grouped according to how many anticodons encode each amino acid, with total numbers for each amino acid and each anticodon indicated.

Scaffold	Class	tRNA Type	Anti-codon	Intron Begin	Intron End	CCA	tRNA part of SINE elements
scaffold_7	SINE-Arg	Arg	CCG	2542253	2542265	P	AGGGGGGTCGTCTAAATGGTtA AGACACTCAAGCCGatttcgttaag gcTTCGAGAGAtCCTGGGTTCGA ATCCCGGTCACCCCA
scaffold_203	SINE-Arg	Arg	CCG	7724	7712	P	GGGGGGGTCATCTAAATGGTtA AGACACTCAAGCCGatttcgttaag gcTTCGAGAGAtCCTGGGTTCGA ATCCCGGTCACCCCA
scaffold_40	SINE-Arg	Arg	ACG	84541	84553	P	GGGGGGGTCGTCTAAATGGTtA AGACACTCAAGACGatttcgttaag gcTTCGAGAGAtCCTGGGTTCGA ATCCCGGTCACCCCA
scaffold_121	SINE-Arg	Arg	CCG	47226	47238	P	GGGGGGGTCGTCTAAATGGTtA AGACACTCAAGCCGatttcgttaag gcCTCGAGAGAtCCTGGGTTCGA ATCCCGGTCACCCCA
scaffold_124	SINE-Arg	Arg	CCG	4376	4364	P	GGGGGGGTCGTCTAAATGGTtA AGACACTCAAGCCGatttcgttaag gcTTCGAGAGAtCCTGGGTTCGA ATCCCGATCACCCCA
scaffold_958	SINE-Arg	Arg	CCG	363	351	P	GGGGGGGTCGTCTAAATGGTtA AGACACTCAAGCCGatttcgttaag gcTTCGAGAGAtCCTGGGTTCGA ATCCCGATCACCCCA
scaffold_21	SINE-Arg	Arg	CCG	1832790	1832802	P	GGGGGGGTCGTCTAAATGGTtA AGACACTCAAGCCGatttcgttaag gcTTCGAGAGAtCCTGGGTTCGA ATCCCGGTCACCCCA
scaffold_21	SINE-Arg	Arg	CCG	1828284	1828272	P	GGGGGGGTCGTCTAAATGGTtA AGACACTCAAGCCGatttcgttaag gcTTCGAGAGAtCCTGGGTTCGA ATCCCGGTCACCCCA
scaffold_40	SINE-Arg	Arg	CCG	41699	41687	P	GGGGGGGTCGTCTAAATGGTtA AGACACTCAAGCCGatttcgttaag gcTTCGAGAGAtCCTGGGTTCGA ATCCCGGTCACCCCA
scaffold_40	SINE-Arg	Arg	CCG	9927	9915	P	GGGGGGGTCGTCTAAATGGTtA AGACACTCAAGCCGatttcgttaag gcTTCGAGAGAtCCTGGGTTCGA

scaffold_73	SINE-Arg	Arg	CCG	191771	191783	P	ATCCCGGTCACCCCA GGGGGGGTCGTCTAAATGGTtA AGACACTCAAGCCGatttcgttaag gcTTCGAGAGAtCCTGGGTTCGA ATCCCGGTCACCCCA
scaffold_113	SINE-Arg	Arg	CCG	6095	6107	P	GGGGGGGTCGTCTAAATGGTtA AGACACTCAAGCCGatttcgttaag gcTTCGAGAGAtCCTGGGTTCGA ATCCCGGTCACCCCA
scaffold_125	SINE-Arg	Arg	CCG	26556	26544	P	GGGGGGGTCGTCTAAATGGTtA AGACACTCAAGCCGatttcgttaag gcTTCGAGAGAtCCTGGGTTCGA ATCCCGGTCACCCCA
scaffold_218	SINE-Arg	Arg	CCG	208	196	P	GGGGGGGTCGTCTAAATGGTtA AGACACTCAAGCCGatttcgttaag gcTTCGAGAGAtCCTGGGTTCGA ATCCCGGTCACCCCA
scaffold_217	SINE-Arg	Arg	CCG	8299	8311	P	GGGGGGGTCGTCTAAATGGTtA AGACACTCAAGCCGatttcgttaag gcTTCGAGAGAtCCTGGGTTCGA ATCCCGGTCACCCCA
scaffold_285	SINE-Arg	Arg	CCG	10820	10808	P	GGGGGGGTCGTCTAAATGGTtA AGACACTCAAGCCGatttcgttaag gcTTCGAGAGAtCCTGGGTTCGA ATCCCGGTCACCCCA
scaffold_545	SINE-Arg	Arg	CCG	8056	8044	P	GGGGGGGTCGTCTAAATGGTtA AGACACTCAAGCCGatttcgttaag gcTTCGAGAGAtCCTGGGTTCGA ATCCCGGTCACCCCA
scaffold_729	SINE-Arg	Arg	CCG	1631	1643	P	GGGGGGGTCGTCTAAATGGTtA AGACACTCAAGCCGatttcgttaag gcTTCGAGAGAtCCTGGGTTCGA ATCCCGGTCACCCCA
scaffold_729	SINE-Arg	Arg	CCG	3577	3589	P	GGGGGGGTCGTCTAAATGGTtA AGACACTCAAGCCGatttcgttaag gcTTCGAGAGAtCCTGGGTTCGA ATCCCGGTCACCCCA
scaffold_7	SINE-Arg	Arg	CCG	2572686	2572674	P	GGGGGGGTCGTCTAAATGGTtA AGACACTCAAGCCGatttcgttaag gcTTCGAGAGAtCCTGGGTTCGA ATCCCGGTCACCCCA
scaffold_58	SINE-Arg	Arg	CCG	344247	344235	P	GGGGGGGTCGTCTAAATGGTtA AGACACTCAAGCCGatttcgttaag gcTTCGAGAGAtCCTGGGTTTGA

							ATCCCGGTCACCCCA GGGGGGGTCGTCTAAATGGTtA AGACACTCAAGCCGatttcgttaag
scaffold_121	SINE-Arg	Arg	CCG	13939	13951	P	gcTTCGAGAGAtCCTGGGTTTGA ATCCCGGTCACCCCA GGGGGGGTCGTCTAAATGGTtA AGACACTCAAGCCGatttcgttaag
scaffold_40	SINE-Arg	Arg	CCG	79872	79884	P	gcTTTGAGAGAtCCTGGGTTCGA ATCCCGGTCACCCCA GGGGGGGTCGTCTAAATGGTtA AGACACTCAAGCCGatttcgttaag
scaffold_112	SINE-Arg	Arg	CCG	36173	36161	P	gcTTTGAGAGAtCCTGGGTTCGA ATCCCGGTCACCCCA GGGGGGGTCGTCTAAATGGTtA AGACACTCAAGCCGatttcgttaag
scaffold_1105	SINE-Arg	Arg	CCG	3248	3236	P	gcTTTGAGAGAtCCTGGGTTCGA ATCCCGGTCACCCCA GGGGGGGTCGTCTAAATGGTtA AGACACTCAAGCCGattttcgttaag
scaffold_110	SINE-Arg	Arg	CCG	57194	57181	P	gcTTTGAGAGAtCCTGGGTTCGA ATCCCGGTCACCCCA GGGGGGGTCGTCTAAATGGTtA AGACACTCAAGCCGattttgttaag
scaffold_112	SINE-Arg	Arg	CCG	40724	40712	P	gcTTTGAGAGAtCCTGGGTTCGA ATCCCGGTCACCCCA GGGGGGGTCGTCTAAATGGTtA AGACACTCGAGCCGatttcgttaag
scaffold_87	SINE-Arg	Arg	CCG	68634	68622	P	gcTTCGAGAGAtCCTGGGTTCGA ATCCCGGTCACCCCA GGGGGGGTCGTCTAAATGGTtG AGACACTCAAGCCGatttcgttaag
scaffold_124	SINE-Arg	Arg	CCG	8824	8836	P	gcTTCGAGAGAtCCTGGGTTCGA ATCCCGGTCACCCCA GGGGGGGTCGTCTAAATGGTtG AGACACTCAAGCCGatttcgttaag
scaffold_965	SINE-Arg	Arg	CCG	3317	3329	P	gcTTCGAGAGAtCCTGGGTTCGA ATCCCGGTCACCCCA GGGGGGGTTGTCTAAATGGTtA AGACACTCAAGCCGatttcgttaag
scaffold_60	SINE-Arg	Arg	CCG	451585	451573	P	gcTTCGAGAGAtCCTGGGTTCGA ATCCCGGTCACCCCA GGGGGGGTTGTCTAAATGGTtA AGACACTCAAGCCGatttcgttaag
scaffold_270	SINE-Arg	Arg	CCG	2897	2909	P	gcTTCGAGAGAtCCTGGGTTCGA

scaffold_52	SINE-Arg	Arg	CCG	566079	566067	P	ATCCCGGTCACCCCA TGGGGGGTCGTCTAAATGGTtA AGACACTCAAGCCGatttcgttaag gcTTCGAGAGAtCCTGGGTTCGA ATCCCGGTCACCCCA
scaffold_1295	SINE-Arg	Arg	CCG	914	902	A	GGGGTcGTCTAAATGGTtAAGAC ACTCAAGCCGatttcgtcaaggcTTT GAGAGAtCCTGGGTTCGAATCC CAGTCACCCCA
scaffold_18	SINE-Arg	Arg	CCG	96788	96776	A	GGGGTcGTCTAAATGGTtAAGAC ACTCAAGCCGatttcgtcaaggcTTT GAGAGAtCCTGGGTTCGAATCC CAGTCACCCCA
scaffold_136	SINE-Arg	Trp	CCA	36284	36296	A	GGGGGGGTCGTCTAAATGGTtA AGACACTCAAGCCAatttcgttaag gcTTCGAGAGAtCCTGGGTTCGA ATCCCGGTCGCCCCA
scaffold_258	SINE-Arg	Trp	CCA	2370	2358	P	GGGAGGGTCGTCTAAATGGTtA AGACACTCAAGCCAatttcgttaag gcTTCGAGAGAtCCTGGGTTCGA ATCCCGGTCACCCCA
scaffold_808	SINE-Arg	Trp	CCA	3681	3693	A	GGGGGGGTCGTCTAAATGGTtA AGACACTCAAGCCAatttcgttaag gcTTCGAGAGAtCCTGGGTTCGA ATCCCGGTCGCCCCA
scaffold_99	SINE-Arg	Gly	CCC	112238	112322	P	GGGGGGGTCGTCTAAATGGTtA AGACACTCAAgCCCAtttcgttaag gcTTCGAGAGAtCCTGGGTTCAA ATCCCGGTCACCCCA
scaffold_285	SINE-Arg	Gln	CTG	9029	8945	P	GGGGGGGTCGTCTAAATGGTtA AGACACTCAAgCTGAtttcgttaag gcTTCGAGAGAtCCTGGGTTCGA ATCCCGGTCACCCCA

Table S2. tRNA-related SINE-3 family elements: Details of the scaffold on which the tRNA-related SINE-3 sequence lies, the class, the amino acid of the tRNA and anticodon sequence, the begin and end coordinates of the intron, the presence (P) or absence (A) of a 3' CCA and the sequence of the tRNA-related portion of the SINE-3 element are shown.

Scaffold	Class	tRNA Type	Anti-codon	Intron Begin	Intron End	CCA	tRNA part of SINE elements
scaffold_808	SINE-Asp	Asp	ATC	2922	2972	A	GGGGGGGTAGCTCAGTAGGTaAGAGCACTTCCTTATCAccctgcggacccgggttcaaatctcgtattcggcccgtttcccggcggataAGGTTGAGGtCGTGGGTTCGGATCCCACCCCCCTCA
scaffold_136	SINE-Asp	Asp	ATC	35519	35569	A	GGGGGGGTAGCTCAGTAGGTaAGAGCACTTCCTTATCAccctgcggacccgggttcgaatctcgtattcggcccgtttcccggcagataAGGTTGAGGtCATGGGTTCGGATCCCACCCCCCTCA
scaffold_42	SINE-Asp	Asp	ATC	853996	853946	A	GGGGGGGTAGCTCAGTAGGTaAGAGCACTTCCTTATCAccctgcggacccgggttcgaatctcgtattcggcccgtttcccggcggataAGGTTGAGGtCGTGGGTTCGGATCCCACCCCCCTCA
scaffold_98	SINE-Asp	Asp	ATC	137522	137572	A	GGGGGGGTAGCTCAGTAGGTaAGAGCACTTCCTTATCAccctgcggacccgggttcgaatctcgtattcggcccgtttcccggcggataAGGTTGAGGtCGTGGGTTCGGATCCCACCCCCCTCA
scaffold_986	SINE-Asp	Asp	ATC	868	818	A	GGGGGGGTAGCTCAGTAGGTaAGAGCACTTCCTTATCAccctgcggacccgggttcgaatctcgtattcggcccgtttcccggcggataAGGTTGAGGtCGTGGGTTCGGATCCCACCCCCCTCA
scaffold_20	SINE-Asp	Asp	ATC	685237	685287	A	GGGGGGGTAGCTCAGTAGGTaAGAGCACTTCCTTATCAccctgcggacccgggttcgaatctcgtattcggcccgtttcccggcggataAGGTTGAGGtCGTGGGTTTGGATCCCACCCCCCTCA
scaffold_104	SINE-Asp	Asp	ATC	32583	32633	A	GGGGGGGTAGCTCAGTAGGTaAGAGCACTTCCTTATCAccctgcggacccgggttcgaatctcgtattcggcccgtttcctggcggataAGGTTGAGGtCGTGGGTTCGGATCCCACCCCCCTCA
scaffold_55	SINE-Asp	Asp	GTC	536020	535977	A	TCCCCGGTAGCTCAATTGGTAGAGCATGCCGCTGTCAcatggcagacccaggttcgaatcacggattcggccgggttgaggCTGACAAGTATAGaTGCAGGTTCGGATCCTGCCCG

							GGGAA TCCCCGGTAGCTCAATTGGTAGAGCAT GCCGCTGTCAcatggcagacccaggttcgaa tcgcagattcggccaggttgaggCTGACAAG
scaffold_56	SINE-Asp	Asp	GTC	563038	563081	P	TATAGaTGCAGGTTCGGATCCTGCCCG GGGAA TCCCCGGTAGCTCAATTGGTAGAGCAT GCCGCTGTCAcatggcagacccaggttcgaa tcgcagattcggccaggttgaggCTGACAAG
scaffold_99	SINE-Asp	Asp	GTC	9414	9457	P	TATAGaTGCAGGTTCGGATCCTGCCCG GGGAA TCCCCGGTAGCTCAATTGGTAGAGCAT GCCGCTGTCAcatggcagacccaggttcgaa tcgcagattcggccaggttgaggCTGACAAG
scaffold_388	SINE-Asp	Asp	GTC	552	595	P	TATAGaTGCAGGTTCGGATCCTGCCCG GGGAA TCCCCGGTAGCTCAATTGGTAGAGCAT GCCGCTGTCAcatggcagacccaggttcgaa tcgcggattcggccgggttaggCTGACAAGT
scaffold_2134	SINE-Asp	Asp	GTC	666	624	A	ATAGaTGCAGGTTCGGATCCTGCCCG GGGAA TCCCCGGTAGCTCAATTGGTAGAGCAT GCCGCTGTCAcatggcagacccaggttcgaa tcgcggattcggccgggttgaggCTGACAAG
scaffold_120	SINE-Asp	Asp	GTC	50480	50437	A	TATAGaTGCAGGTTCGGATCCTGCCCG GGGAA TCCCCGGTAGCTCAATTGGTAGAGCAT GCCGCTGTCAcatggcagacccaggttcgaa tcgcggattcggccgggttgaggCTGACAAG
scaffold_2077	SINE-Asp	Asp	GTC	718	761	A	TATAGaTGCAGGTTCGGATCCTGCCCG GGGAA TCCCCGGTAGCTCAATTGGTAGAGCAT GCCGCTGTCAcatggcagacccaggttcgaa tctccgattcggccaggttgaggCTGACAAGT
scaffold_51	SINE-Asp	Asp	GTC	33405	33448	A	ATAGaTGCAGGTTCGGATCCTGCCCG GGGAA TCCCCGGTAGCTCAATTGGTAGAGCAT GCCGCTGTCAcatggcagacccaggttcgaa tctccgattcggccaggttgaggCTGACAAGT
scaffold_18	SINE-Asp	Asp	GTC	75512	75555	A	ATAGaTGCAGGTTCGGATCCTGCCCG GGGAA TCCCCGGTAGCTCAATTGGTAGAGCAT
scaffold_58	SINE-Asp	Asp	GTC	9057	9100	A	GCCGCTGTCAcatggcagacccaggttcgat tcacggattcggccgggttgaggCTGACAAG

scaffold_58	SINE-Asp	Asp	ATC	44296	44346	A	TATAGaTGCAGGTTCGGATCCTGCCCGGGGAA
scaffold_73	SINE-Asp	Asp	ATC	149364	149314	A	GGGGGGGTAGCTCAGTAGGTaAGAGCACTTCCTTATCAccctgcggacccgggttcgaatctcgtattcggcccgtttcccggcggataAGGTTGAGGtCGTGGGTTCGGATCCCACCCCCCTCA
scaffold_55	SINE-Asp	Asp	GTC	491372	491329	P	GGGGGGGTAGCTCAGTAGGTaAGAGCACTTCCTTATCAccctgcggacccgggttcgaatctcgtattcggcccgtttcccggcggataAGGTTGAGGtCGTGGGTTCGGATCCCACCCCCCTCA
scaffold_59	SINE-Asp	Asp	GTC	308788	308831	A	TCCCCGGTAGCTCAATTGGTAGAGCATGCCGCTGTCAcatggcagacccaggttcgaatcgcagattcggccaggttgaggCTGACAAGTATAGaTGCAGGTTCGGATCCTGCCCGGGGAA
scaffold_110	SINE-Asp	Asp	GTC	47423	47380	P	TCCCCGGTAGCTCAATTGGTAGAGCATGCCGCTGTCAcatggcagacccaggttcgatcacggattcggccgggttgaggCTGACAAGTATAGaTGCAGGTTCGGATTCTGCCCGGGGAA
scaffold_58	SINE-Asp	Asp	ATC	46217	46267	A	GGGGGGGTAGCTCAGTAGGTaAGAGCACTTCCTTATCAccctgcggacccgtgttcgaatctcgtattcggcccgtttcccggcggataAGGTTGAGGtCGTGGGTTCGGATCCCACCCCCCTCA
scaffold_59	SINE-Asp	Asp	GTC	404475	404518	A	TCCCCGGTAGCTCAATTGGTAGAGCATGCCGCTGTCAcatggcagacccaggttcgaatcacggattcggccgggttgaggCTGACAAGTATAGaTGCAGGTTCGGATCCTGCCCGGGGAA

scaffold_18	SINE-Asp	Asp	GTC	1388379	1388336	A	TCCCCGGTAGCTCAATTGGTAGAGCATGCCGCTGTCAcatggcagacccaggttcgaatcgcagattcggccaggttgaggCTGACAAGTATAGaTGCAGGTTCGGATCCTGCCCGGGGAA
scaffold_59	SINE-Asp	Asp	GTC	406230	406273	A	TCCCCGGTAGCTCAATTGGTAGAGCATGCCGCTGTCAcatggcagacccaggttcgaatcacggattcggccgggttgaggCTGACAAGTATAGaTGCAGGTTCGGATCCTGCCCGGGGAA
scaffold_59	SINE-Asp	Asp	GTC	464906	464949	A	TCCCCGGTAGCTCAATTGGTAGAGCATGCCGCTGTCAcatggcagacccaggttcgaatcacggattcggccgggttgaggCTGACAAGTATAGaTGCAGGTTCGGATCCTGCCCGGGGAA
scaffold_1	SINE-Asp	Asp	ATC	6483869	6483819	A	GGGGGGGTAGCTCAGTAGGTaAGAGCACTTCCTTATCAccctgcggacccgggttcgaatctcatattcggcccgtttcccggcggataAGGTTGAGGtCGTGGGTTCGGATCCCACCCCCCTCA
scaffold_59	SINE-Asp	Asp	GTC	31219	31176	A	TCCCCGGTAGCTCAATTGGTAGAGCATGCCGCTGTCAcatggcagacccaggttcgaatcgcggattcggccgggttgaggCTGACAAGTATAGaTGCAGGTTCGGATCCTGCCCGGGGAA

Table S3. tRNA-related SINE family elements: Details of the scaffold on which the tRNA-related SINE sequence lies, the class, the amino acid of the tRNA and anticodon sequence, the begin and end coordinates of the intron, the presence (P) or absence (A) of a 3’ CCA and the sequence of the tRNA-related portion of the SINE-3 element are shown.

Models	Number	Percentage
Homology based models	3,022	20
ab initio prediction	6,619	44
Transfers (mapping) of models from chlamy portal version 2.0 to 3.0	3,137	21
ACEGs-based models	439	3
'Known' genes - mapped (not predicted) by fgenesh+	1,112	7
EST based models	201	1
User created models	613	4
Total	15,143	100

Table S4. Gene model generation: Gene models in the Frozen Gene Catalog are categorized with respect to the ways in which they were generated. Generation of the model was through homology, *ab initio* predictions, correspondence with ACEGs and ESTs, or mapping of previous models by fgenesh. Some models were generated by users or carried over from assembly v2.0 of the *Chlamydomonas* assembly.

Supporting Evidence	Number	Percentage
Clustered ESTs support	8,522	56
Swissprot homologs Evalue < 10^{-5}	9,558	63
NR homologs Evalue < 10^{-5}	8,845	58
Pfam domains	6,161	41
Ostreococcous best hits	2,223	15
Cyanidioschyzon best hits	275	2
Greenplants/algae best hits	5,335	35
Fungi/Metazoa best hits	1,729	11
Bacteria, mostly cyanobacteria best hits	1,156	8
ab initio models without support	1,843	12
Manually curated *ab initio* models without support	309	2
Manually assigned name	3914	26

Table S5. Support for gene model assignment: The table lists the various methods and tools that support the generation of gene models.

Functional assignment category	Number	Percentage	Distinct categories
Unique KOG assignments, E-value < 10^{-5}	9,435	62	3,158
Unique Gene Ontology (GO) assignments	6,733	44	3,165
Unique KEGG/EC assignments (60% ID 60% coverage)	2,780	18	798

Table S6. Functional assignment of gene models from KOG, GO and KEGG analyses

Rank	No. of members	Associated protein domain
1	51	PF00211: adenylyl and guanylyl cyclase catalytic domain
2	44	PF00125: core histone H2A/H2B/H3/H4
3	39	PF00125: core histone H2A/H2B/H3/H4
4	35	PF00125: core histone H2A/H2B/H3/H4
5	35	PF00125: core histone H2A/H2B/H3/H4
6	29	PF00069: protein kinase domain
		PF07714: protein tyrosine kinase
7	22	PF00233: 3'5'-cyclic nucleotide phosphodiesterase
8	20	PF00025: ADP-ribosylation factor family
9	15	PF00069: protein kinase domain
		PF07714: protein tyrosine kinase
10	14	PF03110: SBP domain
11	14	PF00069: protein kinase domain
		PF07714: protein tyrosine kinase
12	14	IPR002290: serine/threonine protein kinase
13	14	PF00071: Ras family
14	14	PF00179: ubiquitin-conjugating enzyme
15	13	PF00067: cytochrome P450
16	12	PF00160: cyclophilin type peptidyl-prolyl cis-trans isomerase
17	12	PF03171: 2OG-Fe(II) oxygenase superfamily
18	12	PF07714: protein tyrosine kinase
19	11	PF00651: BTB/POZ domain
20	11	PF00249: Myb-like DNA-binding domain

21	11	PF01384: phosphate transporter family
22	11	PF00226: DnaJ domain
23	10	PF03016: exostosin family
24	10	PF00240: ubiquitin family
25	10	PF00504: chlorophyll a/b binding protein
26	10	PF00168: C2 domain

Table S7. Large protein families: Families of paralogous proteins within each species were made with MCL I=2.0 (*45*); PFAM domains (*41*) were assigned to proteins achieving a score <1e–10 with RPSblast (*23*). Protein families were ranked by size. The table lists the top 20 families based on the number of members in each. Representative PFAM domains are given with PF numbers and descriptions.

Transporter relationship	Members
Plant-specific transporters	MEX (maltose exporter), Tic110 (translocon of the inner chloroplast membrane), AAA (ATP:ADP Antiporter), Tat (twin arginine translocase), HAAAP (Hydroxy/Aromatic Amino Acid Permease), FBT (Folate-Biopterin Transporter), H^+-PPase (H+-translocating Pyrophosphatase), NhaD (Na+:H+ Antiporter)
Transporters associated with animals	DAACS (dicarboxylate amino-acids cation- Na^+ or H^+ symporter), IRK-C (inward rectifier K^+ channel), TRP-CC (transient receptor potential Ca^{2+} channel), LIC (neurotransmitter receptor, cys loop, ligand-gated ion channel), RIR-CaC (ryanodine-inositol 1,4,5-triphosphate receptor Ca^{2+} channel) and PCC (polycystin cation channel, involved in regulating intracellular Ca^{2+} levels)

Table S8. Plant- and animal-associated transporters of *Chlamydomonas*.

PFAM description	PFAM or KOG ID	JGI v3.0 protein ID (gene name)	notes
Animal-associated proteins			
Tubulin-tyrosine ligase family	PF03133	100760, 146893, 118345, 119250, 126569	Likely associated with flagellar function
Kinesin-associated protein (KAP)	PF05804	182554 (KAP1)	Likely associated with flagellar function
Dynein heavy chain	PF03028	130324 (DHC2)	Associated with flagellar function
Ion transport protein	PF00520	179342, 189093, 192415, 144131, 180826, 144354, 170854, 194450, 194451	Voltage-gated Na^{+}/Ca^{2+} ion channels; 194450, 194451 are adjacent on the genome; possibly involved in flagellar signaling
Pyridoxal-dependent decarboxylase	PF00278, PF02784	206067 (ODC1), 206062 (ODC2)	
Vitamin B12 dependent methionine synthase; Homocysteine S methyltransferase	PF02965, PF02574	76715 (METH1)	Cobalamin-dependent methionine synthase (METH), which is not found in vascular plants (*84*)
Selenocysteine-specific elongation factor	KOG0461	112829	The selenocysteine specific elongation factor, which is not found in vascular plants
Adenylate and guanylate cyclase catalytic domain	PF00211	193525 (CYG41), 187517 (CYG12)	See text above
Plant-associated proteins			
Ammonium transporter family	PF00909	182688 (AMT1D), 192308 (AMT1A), 183975 (AMT1B)	Similar to ammonium transporter AMT1 in *Arabidopsis*
S1 RNA binding domain	PF00575	195616 (EFT1)	EF-Ts; Chloroplast small

UBA/TS-N domain Elongation factor TS	PF00627 PF00889		ribosomal subunit protein *PSRP-7* and elongation factor Ts are encoded in this single transcript

Table S9. *Chlamydomonas* protein families similar to those in human or *Arabidospis*: Selected proteins (from scatter plot of **Fig. 4A)**, with closer similarity to human (top half) or *Arabidopsis* (bottom half) polypeptides but that are not members of phylogenomic or experimental groupings. Also given are the PFAM descriptions, JGI protein IDs and notes related to their potential functions.

Description	derivation of gene number	Total	total U or K		
GreenCut		**349**	135	109	K
green lineage of the plantae				26	KI
			214	101	U
				113	UP
PlastidCut		**90**	29	25	K
Common to all photosynthetic				4	KI
eukaryotes			61	26	U
CPLD1-53				35	UP
DiatomCut - PlastidCut	150 - 90 =	**60**	18	15	K
only in green lineage + 1 or more				3	KI
diatoms			42	18	U
CGLD1-30				24	UP
PlantCut - PlastidCut	117 - 90 =	**27**	9	7	K
only in plantae				2	KI
			18	7	U
CPL1-11				11	UP
ViridiCut	349-90-27-60 =	**172**	79	62	K
only in green lineage of plantae				17	KI
not in *Cyanidioschyzon* or diatoms			93	50	U
CGL1-83				43	UP

Table S10. Proteins in the GreenCut and their division into subgroups: The 349 proteins of the GreenCut were selected based on phylogenetic analyses as described in the Main Text. These were classified as either known (K) or unknown (U) with respect to function. The designation was based on experimental work in the literature for either *Arabidopsis* or *Chlamydomonas* proteins. The modifier I for the K category indicates a

function that is known by "inference" (based on a strong sequence identity and full coverage along its length to a protein in a related organism whose function is known). The modifier P for the U category stands for "Predicted" where the gene product is predicted to have a particular enzymatic activity or the sequence contains a structural motif. The distinction between KI and UP may be occasionally blurred because the classifications were made subjectively based on evaluation of the body of literature. Restricting the GreenCut only to those proteins conserved in at least one diatom yielded the DiatomCut with 150 proteins. Restricting the GreenCut only to those proteins conserved in plants yielded the PlantCut with 117 proteins. Restricting the GreenCut only to those proteins conserved in photosynthetic eukaryotes, which include diatoms and plants, yielded the PlastidCut with 90 proteins. The corresponding genes were named according to these groupings unless they had been previously named during manual curation. The name designation *CPL* was given (for conserved in the plant lineage) to genes encoding proteins in the GreenCut that are conserved also in *Cyanidioschyzon* but not in the diatoms, *CPLD* (for conserved in the plant lineage and diatoms) to genes corresponding to proteins in the GreenCut that are conserved in *Cyanidioschyzon* and at least one diatom (PlastidCut), CGLD (for conserved in the green lineage and diatoms) for genes encoding proteins conserved in the GreenCut plus at least one diatom, and CGL (for conserved in the green lineage) for those in the GreenCut that are not present in either *Cyanidioschyzon* or a diatom. This grouping was also designated the ViridiCut. Also see **Fig. 5** and **Supplemental File 1**.

Function	Associated gene products
Regulation of photosynthesis	PGR5, STT7, RCA2, APE1
Thylakoid membrane biogenesis	CCS1, HCF164, CCB factors, SUFD, EGY1, TAB2, MCA1, CSP41a, THF1
Plastid biogenesis	TOCs, TIC110, TIC40, HSPs, CYNs, FKBPs, CLP subunits, PRORS1
Plastid division	MINE1
Lipid biosynthesis	FAB2, LPAAT, KAS1, DGD1, FAT1, PLSB1
Other carbon metabolism	DLA2, DLD2, TAL2, MDH5, RPI2
Amino acid, nucleotide biosynthesis	CGL37 (shikimate kinase), RPPK2, DPR1, DPA1
Starch biosynthesis	STA6, STA11, STA1, PWD1, SSS2, AMYB1
Pigment, cofactor biosynthesis	CTH1, GUN4, DVR, UROD1, HMOX1, LCYE, ADCL1, CHLD, CAO
Metabolite transporters	LCI20, CEM1, RCP1, TPT3
Anti-oxidant pathways	GSH1, APXs, CDSP32, TRXL/HCF164, SNE1

Table S11. Proteins of known function in the GreenCut: Selected chloroplast proteins of known function in the GreenCut are grouped by general function. We excluded proteins of the photosynthetic apparatus, which had been used to estimate the false negative fraction in the GreenCut (see above); these are listed in **Supplemental File 1**. The enzymes LL-diaminopimelate aminotranferase and TGD2 (involved in lipid transfer from the endoplasmic reticulum) are unique to plants, while RPPK2 (phosphoribosyl diphosphate synthase), TAL2 (transaldolase), DLA2, DLD2 (of the pyruvate dehydrogenase complex) and ADCL1 (aminodeoxychorismate lyase) represent plastid-specific isoforms (*85-88*).

Functional Group	***Chlamydomonas* Protein name**	**Description**
SOUL proteins	SOUL4 SOUL5	Related to chicken heme protein identified in retina and pineal gland (which contain light-cued circadian clocks) Also, SOUL3 is found in *Chlamydomonas* eyespot and in *Arabidopsis* plastoglobule
Redox active proteins	TRXL1	Thioredoxin-like protein, unusual active site WCNAC
	TRX10	Thioredoxin-like protein, unusual active site WCPKC
	CITRX	Cytoplasmic in tomato, but highly conserved in the green lineage and diatoms
	CPLD41	Protein disulfide isomerace-like motif + VitK epoxide reductase motif, conserved in cyanobacteria.
	GRX6	Glutaredoxin, CGFS type, probably chloroplastic
	CPLD26	related to pyridoxamine 5' phosphate oxidase
	CPLD32	FAD dependent oxidoreductase
	CPLD49	saccharopine dehydrogenase-like
	CPLD25	short-chain dehydrogenase/reductase
	TEF5	Rieske [2Fe-2S] domain
Isoprenoid pathway	CPLD35	flavin containing amine oxidase related to phytoene desaturase
	VDR1	violaxanthin de-epoxidase related
	CPLD27	coclaurine N-methyl transferase
	CGL2	ubiquinol methyl transferase
	CPLD34	ubiquinol methyl transferase
	AKC1 AKC2 AKC3 AKC4	ABC1 kinases. The mitochondrial homolog regulates UQ biosynthesis. A *Chlamydomonas* AKC is the product of the EYE3 locus, required for assembly of the carotenoid pigmented eyespot. ORFs in cyanobacteria with very strong sequence similarity.
	PLAP1 PLAP2	plastid lipid associated protein or Plastoglobulins, conserved in cyanobacteria

	PLAP3	
	PLAP4	
Transporters	CPLD21	sugar nucleotide transporters, solute carriers
	CPLD22	
	CPLD23	
	ARSA	anion transporter
	CGL51	plastid metabolite exchanger
	CGL7	plastid metabolite exchanger
	CGLD4	ABC transporter
	CGL15	major facilitator superfamily
	MITC4	mitochondrial carrier
	TIM22B	plastid homolog of TIM17/22/23 family
Various	CPLD3	aldo-keto isomerase
metabolic	SNE3	NAD-dependent epimerase/dehydratase
reactions	CGLD13	related to nucleoside diphosphate sugar epimerase, putative chloroplast targeted
	CGL2	methyltransferase
	CGL33A/B	methyl transferase
	CGL75	methyl transferase motif
	CGL77	methyl transferase
	CGLD2	thioesterase
	CGLD24	thioesterase
	CGLD7	esterase / lipase / thioesterase
	CGL69	lipase
	CPLD15	lipase
	CGLD15	related to triacylglycerol lipase
	CGL76	esterase, epoxide hydrolase
	CPLD2	hydrolase
	CGL53	related to carbohydrate hydrolase
	CPLD4	inositol monophosphatase-related
	CGL14	pantothenate kinase motif
	CGL79	carbohydrate kinase motif
	CGLD12	potential galactosyl transferase activity
	CGLD24	related to diacylglycerol acyl transferase
	RIBFL1	related to riboflavin biosynthesis protein RibF
	CGL48	related to lysine decarboxylase domain
biogenesis and	CPLD17	organelle-targeted protein, related to OTU-like cysteine

nucleic acid		protease family
transactions	CPLD6	metal-dependent CAAX amino terminal protease family
	HEP2	Hsp70 escorting protein 2
	CPLD43	YGGT family
	RNB2	3'-5' Exoribonuclease II
	CPLD16	organelle-targeted, RNA methyl transferase related
	CGL43	RNA binding protein with S1 domain
	CGL72	hemolysin motif and RNA methyltransferase motif
	TPR2	tetratricopeptide repeat protein, organelle-targeted
	CGL71	TPR repeat protein related to YCF37
	CPLD46	DEAD/DEAH-box helicase possibly plastid targeted
	CGLD3	DEAD/DEAH box helicase domain and proline rich domain
	CGLD5A	ethylene response element dna binding domain containing protein
	CGLD5B	AP2-domain transcription factor
	CPL2	transcription factor like protein
	CGLD30	SET domain containing protein, putative histone methyltransferase
	CGL31	pterin carbinolamine dehydratase domain
	CGL49	ARF/SAR superfamily small monomeric GTP binding protein
Regulation	PP2C4	related to protein phosphatase 2C
	PP2C5	related to protein phosphatase 2C
	PP2C6	related to protein phosphatase 2C
	CPL3	related to protein serine / threonine phosphatase
	MAPK2	Mitogen-Activated Protein Kinase Homolog 2
	STPK25	MUT9 related serine/threonine protein kinase
Photosynthesis	CPLD45	possible function in PSII and possible lumen location

Table S12. Proteins of unknown function in the GreenCut: Proteins of the GreenCut with unknown functions are tabulated with potential activities associated with these proteins based on annotations of the *Chlamydomonas* genome at (*15*) and the *Arabidopsis* genome (*89*). Note the striking representation of redox-active proteins, proteins that might function in isoprenoid metabolism and proteins from the plastoglobule/eyespot proteomes (see Fig. S1).

GreenCut			cp	mito	other	unknown
349	135	K+KI	115	3	9	8
	214	U+UP	113	36	19	46

Table S13. Subcellular localization of proteins in the GreenCut: The experimental or predicted localization of the proteins in each group (known K, unknown U, which also includes both known inferred, KI, and unknown predicted, UP) is indicated as follows: cp, chloroplast; mito, mitochondrion; other, all other compartments; not known, no data and no prediction. For the known group, the subcellular location is experiment-based for 73% of the proteins. For the unknown group the subcellular location is experiment-based for only 15% of the proteins.

Category	Members	Significance
Motililty-associated (MotileCut)	PF16, PF20, KLP1 and hydin	central pair proteins
	RSP3 and RSP9	radial spoke proteins
	DHC2, DHC6 (inner dynein arm components), ODA4, ODA6 (outer dynein arm components), ODA1 (the outer dynein arm docking complex protein), and PF2 (component of the dynein regulatory complex)	
Outer dynein arm proteins lost in moss *Physcomitrella*	ODA4, ODA6, ODA9, DLC1 and DLC4	
DiatomCut	anterograde motor (KAP) and complex B (IFT57, IFT74, IFT81, IFT88)	Intraflagellar transport proteins present in centric diatom *Thalassiosira*
	retrograde motor (represented by D1bLIC) and complex A (represented by IFT140)	Intraflagellar transport proteins lost in centric diatom *Thalassiosira*
Comparison to *Ostreococcus*	ODA1, ODA4, ODA6, ODA9, Tctex1 DHC2, DHC6, RSP3, RSP9, PF16, PF20, KLP1, hydin, KAP, D1bLIC, IFT20, IFT52, IFT57, IFT74, IFT80, IFT81, IFT88, IFT140, IFT172, RIB43a, PKD2, FAPs 9, 21, 22, 32, 36, 43, 46, 47, 50, 60, 61, 66, 69, 73, 74, 75, 81, 94, 100, 111, 116, 118, 122, 134, 146, 155, 156, 161, 184, 198, 240, 251, 253, 259, 263, 264, 247	Flagellar proteins lost in *Ostreococcus*
	MKS1, NPH4, BLD1, BLD2, UNI3, POC11, POC18, FBB5, 9, 11, 15, and all of the BBS proteins (BBS2, 3, 5, 7, 8, 9)	Basal body proteins lost in *Ostreococcus*
	RIB72, PF2, MBO2, DLC1, PACRG1, DIP13, FAPs 14, 44, 45, 52, 57, 59, 67, 82, 106, 250, 267, and POC1	Flagellar proteins retained in *Ostrecoccus*

Table S14. CiliaCut proteins: Protein designations, association with flagella, or a specific sub- structure of the flagella, basal body. intraflagellar transport and/or affiliations with specific organisms are given.

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Supplemental File 1: GreenCut proteins

frozen catalog model protein ID	JGI updated model protein ID (link to protein page)	protein family (cluster) ID	name	*Chlamydomonas* Defline	Arabidopsis locus (link to TAIR)	Chlamy ESTs	P predicted, X experimental	Location, B = membrane, C = chloroplast, D = endosome, ER = endoplasmic reticulum, G = Golgi, L = lumen, M = mitochondrion, P = plasma membrane, S = stroma, T = thylakoid membrane, U = nucleus, iV = inner envelope, oV = outer envelope, X = peroxisome, Y= cytoplasm, blank = not known, no prediction	Function U or K for unknown or known, P = motif, domain or activity predicted, I = inferred	Gene Name Post-4/07 freeze	identified by Mulkidjanian *et al*. in cyanobacterial genome core
187094		5707704	LCYE	lycopene epsilon cyclase, putative chloroplast	At5g57030	yes	X	C	K		
193086		5707594	RBCMT1	ribulose-1,5 bisphosphate carboxylase	At1g14030	yes	P	C	K		
105530	138922	5708627	PRORS1	organellar class II (G, H, P and S) aminoacyl tRNA synthetase	At5g52520	yes	X	C/M	K		
196673	187308	5706588	OHP1	low CO2 and stress-induced one-helix protein	At1g34000	yes	X	T	K		
105908	205760	5707055	DPR1	dihydropicolinate (DAP) reductase	At3g59890	no	P	C	K		
186597		5707074	APX1	ascorbate peroxidase	At1g77490	yes	X	T	K		
153656		5707261	PSBQ	Oxygen evolving enhancer 3,OEE3	At4g05180	yes	X	L	K		
189624		5708869	PTOX1	alternative oxidase, possibly chloroplast-localized	At4g22260	yes	X	T	K		
130292	205741	5709232	PLSB1	Glycerol-3-phosphate acyltransferase, chloroplast precursor	At1g32200	yes	X	S	K		
195947		5709844	HMOX1	Heme oxygenase	At1g69720	yes	P	C	K		
196775		5705132	SUFD	iron-sulfur cluster assembly protein	At1g32500	yes	X	C	K		
195556	195553	5705722	FNR1	Ferredoxin-NADP reductase, chloroplast	At4g05390	yes	X	S	K		
128415		5705802	TAB2	conserved chloroplast PsaB RNA binding protein	At3g08010	yes	X	S	K		yes
113617	182653	5705826	THF1	conserved expressed chloroplast-localized protein similar to Arabidopsis Thylakoid Formation1	At2g20890	yes	X	T	K		yes
129557		5706628	DPA1	LL-diaminopimelate aminotransferase, putative	At4g33680	yes	X	S	K		
130316		5706989	PSBO	Oxygen-evolving enhancer protein 1 of photosystem II, chloroplast precursor	At5g66570	yes	X	T/L	K		yes
196500		5707144	DLA2	dihydrolipoamide acetyltransferase, possibly plastidic	At1g34430	yes	P	C	K		
194448		5707908	APE1	homolog of Arabidopsis APE1 that is required for acclimation of photosynthesis to various light intensity	At5g38660	yes	X	T	K		yes
196518	205763	5708531	DLD2	dihydrolipoamide dehydrogenase, plastid precursor, putative	At4g16155	no	P	C	K		yes

Supplemental File 1: GreenCut proteins

frozen catalog model protein ID	JGI updated model protein ID (link to protein page)	protein family (cluster) ID	name	*Chlamydomonas* Defline	Arabidopsis locus (link to TAIR)	Chlamy ESTs	P predicted, X experimental	Location, B = membrane, C = chloroplast, D = endosome, ER = endoplasmic reticulum, G = Golgi, L = lumen, M = mitochondrion, P = plasma membrane, S = stroma, T = thylakoid membrane, U = nucleus, iV = inner envelope, oV = outer envelope, X = peroxisome, Y= cytoplasm, blank = not known, no prediction	Function U or K for unknown or known, P = motif, domain or activity predicted, I = inferred	Gene Name Post-4/07 freeze	identified by Mulkidjanian *et al* . in cyanobacterial genome core
134594		5709272	CHLD	Magnesium chelatase subunit D, chloroplast precursor	At1g08520	yes	X	C	K		
195910		5709595	PRK	Phosphoribulokinase, chloroplast precursor	At1g32060	yes	X	T	K		
132194		5709835	UROD1	Uroporphyrinogen-III decarboxylase, chloroplast precursor	At2g40490	yes	P	C	K		
55336		5705127	CLPP5	active subunit of the chloroplast ClpP complex	At1g02560	yes	X	T/S	K		
78983	196952	5706992	SNE1	sugar nucleotide epimerase	At5g28840	yes	P		K		
127079		5709570	PGR5	thylakoid membrane protein	At2g05620	yes	X	T	K		
79446		5707319	FKB16-3	peptidyl-prolyl cis-trans isomerase, FKBP-type	At2g43560	yes	X	L	KI		
32852		5707923	FKB19	peptidyl-prolyl cis-trans isomerase, FKBP-type	At5g13410	yes	X	L	KI		
191582	205877	5708642	PSBP2	lumenal PsbP-like protein	At2g28605	yes	P	L	KI		
196558		5706467	CYN38	Peptidyl-prolyl cis-trans isomerase, cyclophilin-type	At3g01480	yes	X	L	KI		
141254		5707081	CPLD14	conserved protein	At5g52540	no	P	C	U		
108052	154041	5705756	CPLD13	conserved expressed protein	At5g40500	yes	P	C	U		
154010	185795	5707800	CPLD9	conserved expressed protein	At2g38695	yes	P	B	U		
121991	205876	5709810	CPLD20	conserved expressed protein	At5g47860	no	P	C	U		yes
122132	185598	5707211	CPLD31	conserved expressed protein	At5g52970	yes	X	L	U		yes
127973	182934	5707415	CPLD24	conserved expressed protein, perhaps chloroplast targetted	At1g16080	yes	P	C	U		
121199	193550	5707522	CPLD52	conserved expressed protein	At2g39080	yes	P	C	U		
115563	183275	5707541	CPLD28	conserved expressed protein	At1g73070	yes	P		U		
183448		5708382	TEF3	unknown function, chloroplast location proposed	At4g11960	yes	X	T	U		
137516	205570	5709921	CPLD18	conserved expressed protein	At2g21960	yes	P	C	U		
184621		5708622	CPLD33	conserved expressed protein	At2g48070	yes	P	C	U		
105761	151387	5705796	CPLD7	conserved expressed protein of unknown function	At2g45990	yes	P		U		

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184614		5710646	CPLD1	hypothetical protein with unknown function	At5g50100	yes	P	M	U		
118100	191167	5705021	CPLD36	conserved expressed protein	At1g64355	yes	P	C	U		yes
169505		5706200	CPLD47	conserved expressed membrane protein	At4g19100	yes	P		U		yes
193790	152648	5706548	CPLD48	conserved expressed protein	At3g60810	yes	P	M	U		
160068	206091	5707024	CPLD42	conserved expressed membrane protein	At1g54520	yes	P	C	U		
121963	185542	5707745	CPLD38	conserved expressed protein	At3g17930	yes	X	T	U		yes
101763	183668	5707799	CPLD5	probably chloroplast targeted conserved expressed protein	At2g47840	yes	P	C	U		yes
118702	184411	5710330	CPLD39	conserved expressed protein	At2g43945	yes	P	C	U		yes
120574	185128	5709264	CPLD51	putative plastid protein	At3g26710	yes	P	C	U	CCB1	yes
582	151721	5705112	CPLD3	conserved expressed protein	At5g53580	yes	P		U		
188978		5705849	CPLD12	conserved expressed protein of unknown function	At5g27560	yes	P	C	U		yes
121745	185467	5706036	CPLD11	conserved expressed protein	At3g19900	yes	P	C	U		yes
105340	150826	5709235	TEF9	conserved expressed protein, possible chloroplast localization	At3g61870	yes	X	iV	U		
178204		5705630	CPLD50	conserved protein	At5g03900	no	P	C	U		
146442		5705373	CPLD15	conserved organelle protein with lipase active site	At5g17670	yes	P	M	UP		
117277	196953	5707780	CPLD2	conserved expressed protein with hydrolase motif	At3g48420	yes	P	C	UP		
140668	205488	5707142	CPLD35	conserved expressed flavin containing amine oxidase domain	At3g09580	yes	P	C	UP		
166701		5707567	CPLD32	conserved FAD dependent oxidoreductase	At2g22650	no	P	M	UP		
134003		5708525	CPLD26	conserved expressed protein related to pyridoxamine 5' phosphate oxidase	At2g46580	yes	P	M	UP		
119132		5708535	CPLD27	conserved expressed protein related to putative coclaurine N-methyltransferase	At4g33110	yes	P		UP		
190093	146838	5709676	CPLD17	conserved expressed organelle-targeted protein, related to OTU-like cystein protease family	At3g57810	yes	P	C/M	UP		

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193755		5709825	CPLD21	conserved expressed protein of the solute carrier family	At1g76340	yes	P		UP		
144907		5709825	CPLD22	conserved expressed membrane protein, related to putative transporters	At1g76340	yes	P		UP		
149680		5709825	CPLD23	conserved, expressed membrane protein related to putative transporters	At1g76340	yes	P		UP		
116421	189893	5706016	PP2C4	conserved expressed protein in flagellar proteome related to protein phosphatase 2C	At1g79630	yes	P		UP		
113913	205878	5706840	CPLD10	conserved expressed protein	At1g16720	yes	P	C	UP		
116679		5708577	CPLD46	DEAD/DEAH-box helicase possibly plastid targeted	At1g70070	no	P	C	UP		
119827	205880	5705776	CPLD4	conserved expressed protein inositol monophosphatase-related	At4g39120	yes	P	C	UP		
107783	205604	5705143	CPLD49	saccharopine dehydrogenase-like protein	At1g50450	yes	P	M	UP		
122880	132949	5705312	ARSA	putative arsenite translocating ATPase-like protein	At3g10350	yes	P		UP		
185063		5708249	CPLD34	conserved expressed protein related to ubiquinone / menaquinone biosynthesis methyltransferase	At4g29590	yes	P	C	UP		
10730	205779	5710032	AKC2	conserved protein related to ABC1/COQ8 putative ser/thr kinase	At4g24810	no	P		UP		
105237	205572	5705895	CPLD25	conserved expressed protein of the short-chain dehydrogenase/reductase family	At4g13250	yes	P		UP	NYC1	
164377	205755	5706772	CPLD16	conserved expressed protein, organelle-targeted, RNA methyl transferase related	At1g54310	yes	P	M	UP		
183051		5707972	CPLD30	Putative protein of unknown function, similar to hypothetical rice polypeptide	At5g17170	yes	X	T	UP		
153915		5708800	CPLD53	conserved expressed protein	At2g47970	no	P	M	UP		
108495	154497	5709254	CPLD8	conserved expressed protein	At5g21920	yes	P	C	UP		
100330	188875	5709546	SOUL4	conserved expressed protein with SOUL heme binding motif	At5g20140	yes	X	T	UP		

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103242	205756	5710349	CPLD19	conserved expressed protein in the zing finger family	At3g60800	no	P	M	UP		
186610		5709991	CPLD29	conserved expressed protein	At1g71480	no	X	T	UP		
187371		5706348	CPLD45	conserved expressed protein with possible function in PSII and possible lumen location	At1g03600	yes	X	L	UP		yes
107288	178821	5706610	CPLD41	conserved expressed membrane protein	At4g35760	yes	P	C	UP		
101647	205830	5708018	CPLD43	YGGT family, conserved hypothetical integral membrane protein	At5g36120	yes	P	C	UP	CCB3	
139226	205743	5708201	AKC3	conserved expressed protein related to ABC1/COQ8 mitochondrial putative ser/thr kinase	At3g24190	yes	P	C	UP		
122683	205750	5710477	CPLD37	conserved expressed integral membrane protein	At1g78620	yes	X	iV	UP		
184818		5706622	CPLD44	conserved expressd thylakoid lumenal protein-like	At1g12250	yes	X	T	UP		
145444	205881	5706400	PLAP2	conserved expressed protein related to plastid lipid associated protein PAP	At3g26070	yes	X	T	UP		
107013	205882	5709050	CPLD6	conserved protein related to metal-dependent CAAX amino terminal protease family	At5g60750	no	P	C	UP		
113685		5706443	AKC4	conserved expressed ABC-1 like kinase	At5g64940	yes	P	C	UP		
147520		5707116	CLPR4	inactive subunit of chloroplast ClpP complex	At4g17040	yes	X	S	K		
195952		5707642	DVR	3,8-divinyl protochlorophyllide a 8-vinyl reductase, chloroplast precursor	At5g18660	yes	X	C	K		
39090	205922	5708078	CCD1	related to carotenoid 9,10-9',10' cleavage dioxygenase	At3g63520	yes	X	Y	K		
196553		5708985	DGD1	galactolipid galactosyltransferase	At3g11670	yes	X	C/M	K		
142479		5709516	MSH1	DNA mismatch repair MutS protein	At3g24320	no	X	C/M	K		
143831		5709793	EFL1	related to EDS5, enhanced disease susceptibility gene	At2g21340	yes	P		K		
184810		5707565	LHCB4	chlorophyll a-b binding protein of photosystem II	At2g40100	yes	X	T	K		

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155150		5709062	PSBW	Photosystem II subunit W, chloroplast precursor	At2g30570	yes	P	C	K		yes
24036	168073	5709237	LHCA4	light-harvesting protein of photosystem I	At3g47470	yes	X	T	K		
14590	184171	5710150	RPPK2	ribose-phosphate pyrophosphokinase (RPPK)	At1g10700	yes	P	C	K		
135886	205753	5710858	FAB2	plastid acyl-ACP desaturase	At2g43710	no	P	C	K		
99751	182896	5706454	PSB28	Photosystem II subunit 28, chloroplast precursor	At4g28660	yes	X	T	K		yes
11164		5705291	HCF164	Thioredoxin-like protein similar to Arabidopsis HCF164	At4g37200	yes	X	T	K		
196222	205768	5709052	GUN4	Tetrapyrrole-binding protein, chloroplast precursor (Genomes uncoupled 4) (GUN4)	At3g59400	yes	X	C	K		
193583	205771	5708540	EGY1	conserved expressed protein related to membrane associated metalloprotease required for chloroplast development	At5g35220	yes	X	C	K		
188387	205568	5709566	CSP41a	conserved expressed chloroplast RNA binding protein	At3g63140	yes	X	C	KI		
37663		5708031	CYN28	Peptidyl-prolyl cis-trans isomerase, cyclophilin-type	At5g35100	yes	P	C	KI		
185878	205934	5710428	MCA1	maturation/stability factor for petA mRNA	At5g02860	yes	P	C	KI		
141399	205883	5709710	CGLD1	putative plastid protein	At1g64150	yes	P	C	U		
195705		5709241	REX1B	Conserved Protein of Unknown Function	At5g04910	yes	P		U		
190046		5705203	CGLD6	conserved expressed protein	At3g50685	yes	P	C	U		
143294		5708371	CGLD8	Conserved protein, arabidopsis homolog is related to arabidopsis cyclin delta-3	At2g23370	no	X	M	U		
179251		5708881	CGLD11	conserved protein	At2g21385	no	P	C	U		
167973	205923	5709806	CGLD14	conserved protein in diatoms and the green lineage	At1g76450	yes	X	L	U		
158544	163712	5705622	CGLD16	conserved expressed protein	At2g05310	yes	P	C	U		
117443	205927	5708850	CGLD19	expressed protein conserved in green lineage and diatoms	At2g35610	yes	P	M	U		

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79776	205928	5710059	CGLD20	conserved protein with COG5222 RING zinc finger domain	At5g47430	yes	P		U		
169161	205932	5710265	CGLD21	conserved protein	At1g27510	yes	P	C	U		
102133	146879	5708767	CGLD22	conserved expressed protein	At2g31040	yes	P	C	U		yes
118123	205823	5710906	CGLD23	conserved expressed protein	At1g59840	no	P	M	U	CCB4	yes
122264	82483	5705809	CGLD25	conserved expressed protein	At2g04360	yes	P	C	U		
148682		5707240	CGLD26	conserved expressed protein	At4g24090	yes	P	C	U		
44653	184984	5709072	CGLD27	conserved expressed protein	At5g67370	yes	P	M	U		yes
187252		5709857	CGLD28	conserved expressed protein	At1g67080	yes	X	V	U		
107169		5708107	CGLD29	conserved protein of unknown function	At5g27290	no	P	M	U		
187910		5708434	CGLD9	conserved expressed protein	At1g44920	yes	P	C	U		
171897		5706717	CGLD2	conserved protein, related to thioesterase family	At5g48370	no	P	C	UP		
144028		5709127	CGLD3	conserved protein with DEAD/DEAH box helicase domain and proline rich domain	At1g59990	no	P	C	UP		
182736		5709334	CGLD4	conserved expressed protein with ABC transporter motifs	At1g03905	yes	P		UP		
195890		5707643	CITRX	Thioredoxin CITRX	At3g06730	yes	P	C	UP		
120927	131867	5707654	CGLD7	conserved, expressed esterase / lipase / thioesterase family protein	At5g38360	yes	P	C	UP		
196129	205754	5708809	TRXL1	conserved expressd thioredoxin-like protein, unusual active site WCNAC	At4g26160	yes	P	C	UP		
189909		5709030	CGLD12	protein with potential galactosyl transferase activity	At4g37690	yes	P	D	UP		
123134	205577	5709095	CGLD13	conserved protein related to nucleoside diphosphate sugar epimerase, putative chloroplast targeted	At4g31530	yes	P	C	UP		
190433	205924	5710340	CGLD15	conserved expressed protein related to triacylglycerol lipase	At3g62590	yes	P		UP		
142644		5707225	MITC4	putative mitochondrial carrier protein	At2g35800	yes	P	M	UP		
188559		5708306	RNB2	3'-5' Exoribonuclease II	At5g02250	yes	P	C	UP		

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190539		5705744	CGLD24	conserved expressed protein related to diacylglycerol acyl transferase	At3g51520	yes	P		UP		
107982	205919	5709335	CGLD5A	conserved ethylene response element dna binding domain containing protein	At2g33710	no	X	U	UP		
75247	205920	5709335	CGLD5B	AP2-domain transcription factor	At2g33710	yes	X	U	UP		
100030	205921	5707266	TPR2	conserved tetratricopeptide repeat protein, organelle-targetted	At3g05625	no	P	C	UP		
139332		5708358	FAP173	Conserved Uncharacterized Flagellar Associated Protein/ band 7 domain protein	At5g62740	yes	P		UP		
178960	153371	5708635	CGLD30	SET domain containing protein, putative histone methyltransferase	At4g15180	no	P	U	UP		
142077		5708874	CGLD10	conserved protein	At1g26760	yes	P	C	UP		
195571		5709344	VDR1	violaxanthin de-epoxidase related, chloroplast	At2g21860	yes	P	C	UP		
120332		5709408	AKC1	ABC1 family ser/thr kinase	At5g05200	yes	P		UP		
183765	205926	5705505	PLAP3	conserved expressed protein related to plastid lipid associated protein PAP	At4g00030	yes	P	C	UP		
154399		5705666	CGLD17	conserved protein related to Arabidopsis protein with Toprim domain	At1g30680	no	P	M	UP		
159133		5705768	CGLD18	conserved B-box zinc finger protein	At3g02380	yes	P		UP		
179586	21100	5708126	SOUL5	expressed protein conserved in photosynthetic organisms	At2g46100	yes	P	C	UP		
112947	205870	5705953	GWD1	R1 Protein, alpha-glucan water dikinase	At1g10760	yes	P	M	K		
30187	206003	5706169	TIC110	110 kDa translocon at the inner membrane of chloroplasts	At1g06950	yes	X	C	K		
174358		5707301	LPAAT1	related to plastidic lysophosphatidic acid acyltransferase (LPAAT)	At4g30580	no	X	C	K		
193847		5708454	PSAO	Photosystem I subunit O	At1g08380	yes	X	C	K		
115079	183141	5709118	AMYB1	beta-amylase	At3g23920	yes	X	C	K		
194013	205874	5710135	CPL7	conserved expressed protein related to GIF3	At4g00850	yes	P	U	K		
196703		5709211	FDX6	Apoferredoxin, chloroplast precursor	At1g32550	yes	P	C	K		
195403		5705996	VAMP72	R-SNARE, VAMP72-family	At1g04750	yes	X	D/P	KI		

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59842		5710052	CDKB1	plant specific cyclin dependent kinase	At2g38620	yes	P		KI		
154342		5706493	CPL1	conserved expressed protein	At5g08450	yes	P		U		
143849		5708976	CPL4	hypothetical protein	At1g16870	no	P	M	U		
104223	205873	5709922	CPL6	conserved expressed protein	At5g61670	yes	P	C	U		
191988	149843	5710225	TEF10	conserved expressed protein	At1g67700	yes	P	M	U		
159383	205606	5705087	CPL9	conserved expressed protein of unknown function	At1g02470	yes	P	C	U		
183430	205879	5710154	CPL10	Conserved protein of unknown function	At1g28140	yes	P	M	U		
189732		5705310	CPL11	conserved expressed organellar protein involved in translation	At3g01920	yes	P	M	U		
137528		5706710	MAPK2	Mitogen-Activated Protein Kinase Homolog 2	At1g73670	yes	P	M	UP		
181068		5707367	CPL3	conserved protein related to protein serine / threonine phosphatase	At1g07010	no	P	C	UP		
153736		5709002	STPK25	MUT9 related kinase, serine/threonine protein kinase	At3g13670	yes	P	M	UP		
177997		5709568	CPL5	conserved peptidase M16 family protein	At5g56730	yes	P		UP		
121874	152921	5711046	PP2C5	protein phosphatase 2C-like	At2g40860	yes	P		UP		
151947		5706325	RTB1	related to reticulon	At2g46170	yes	P	ER	UP		
152228		5707250	RIBFL1	conserved protein related to riboflavin biosynthesis protein RibF	At5g08340	no	P		UP		
188447	205871	5707326	CPL2	conserved, expressed, transcription factor like protein	At4g32890	yes	P	U	UP		
190008		5709319	PLAP1	conserved expressed protein related to lipid associated plastid protein, PAP	At2g46910	yes	P	C	UP		
105401	205875	5710136	CPL8	conserved expressed protein	At3g21140	yes	P	C	UP		
116298	205993	5711272	TEF30	conserved expressed protein, SHOOT1 homolog	At1g55480	yes	X	T	UP		
182996		5709782	CLPP4	active subunit of chloroplast ClpP peptidase	At5g45390	yes	X	T/S	K		
162226	183277	5710856	SSS2	soluble starch synthase II, ADP-glucose alpha-1,4 glucane alpha-4-glucanotransferase	At3g01180	yes	P	C	K		
145657		5711029	CYC4	cytochrome c, chloroplast precursor	At5g45040	no	P	C	K		

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128745		5704999	RCA1	rubisco activase	At2g39730	yes	X	S	K		
174723	184471	5705036	LHCA1	light-harvesting protein of photosystem I	At3g54890	yes	X	T	K		
196431		5706145	TAL2	Transaldolase, plastid form, putative	At1g12230	yes	P	C	K		
181975		5706166	GSH1	Gamma-glutamylcysteine synthetase	At4g23100	yes	X	C	K		
157805	205886	5706384	CDSP32	CDSP32, plastidic thioredoxin-like protein	At1g76080	yes	X	C	K		
132552	205887	5706711	KAS1	3-ketoacyl-CoA-synthase component of plastidic multimeric fatty acid synthase	At5g46290	yes	X	C	K		
153678		5706791	LHCA3	light-harvesting chlorophyll-a/b protein of photosystem I (Type III)	At1g61520	yes	X	T	K		
13382		5707640	TOC159	Similar to 159 kDa translocon at the outer membrane of chloroplasts	At2g16640	yes	X	oV	K		
165416		5708030	PSAG	photosystem I reaction center subunit V, chloroplast precursor	At1g55670	yes	X	T	K		
190221	183767	5708244	CLPR1	inactive subunit of chloroplast ClpP complex	At1g49970	yes	X	T	K		
169838	205893	5708248	MEX1	conserved expressed protein related to maltose exporter, RCP1	At5g17520	yes	X	iV	K		
113331		5708789	OKL1	conserved ser/thr protein kinase related to OST1 of Arabidopsis	At4g33950	no	P		K		
99594	144221	5708877	CGL28	conserved expressed putative RNA binding protein	At1g53650	yes	P		K		
192398		5709063	PWD1	phosphoglucan water dikinase	At5g26570	yes	X	C	K		
196599	205899	5709096	AMI2	Amidase	At1g08980	yes	X	Y	K		
169967		5709219	HY5	putative bZIP transcription factor	At5g11260	no	X	U	K		
176977	205943	5709310	CGL37	conserved protein related to shikimate kinase	At2g35500	yes	X	C	K		
80866	192083	5709407	MDH5	NADP-Malate Dehydrogenase	At5g58330	yes	X	C	K		
140500	154398	5709441	HSP90C	Heat shock protein 90C	At2g04030	yes	X	C	K		
186790	205903	5709538	LHCBP1	chlorophyll a-b binding protein, chloroplast precursor	At1g76570	yes	P	T	K		
172432	205655	5709628	TOM20	Mitochondrial translocase of outer membrane, 20 kDa	At3g27080	yes	X	M	K		

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142899		5709700	MER	conserved protein with DEAD/DEAH box helicase domain related to MER3	At3g27730	no	P		K		
113634	30522	5709746	ADCL1	aminotransferase related to 4-amino-4-deoxychorismate lyase	At5g57850	yes	P	C	K		
183027		5709759	TIC40	related to chloroplast protein translocon component Tic40 precursor	At5g16620	yes	X	iV	K		
187332		5709925	CGL55	conserved protein related to cleavage and polyadenylation factor 6	At2g33540	yes	X	U	K		
123555	186089	5709932	RCA2	Similar to RuBisCO activase (RCA)	At1g73110	yes	X	S	K		
182959		5710113	PSAH	Subunit H of photosystem I	At3g16140	yes	X	T	K		
118898	205910	5710220	APX2	putative L-ascorbate peroxidase	At4g32320	yes	P	C	K		
111002	205530	5710290	PSBY2	ycf32-related polyprotein of photosystem II	At1g67740	yes	X	T	K		
168074	182560	5710290	PSBY1	Ycf32-related subunit of photosystem II	At1g67740	yes	X	T	K		
193552	185309	5710305	LHL3	low molecular mass early light-induced protein	At4g17600	yes	X	T	K		
196341		5710467	PSBS1	chloroplast Photosystem II-associated 22 kDa protein	At1g44575	yes	X	T	K		
116665	171516	5710467	PSBS2	chloroplast Photosystem II-associated 22 kDa protein	At1g44575	no			K		
167738	205912	5710475	RPI2	ribose-5 phosphate isomerase-related protein	At5g44520	yes	P	S	K		
148916		5710509	ELI3	Early light-inducible protein	At4g14690	yes	X	T	K		
195512		5710545	TOC75	75 kDa translocon at the outer envelope membrane of chloroplasts	At3g46740	yes	X	oV	K		
116544	205633	5710571	TPT3	triose phosphate translocator	At5g46110	yes	X	iV	K		
196283		5710596	FAT1	acyl carrier protein thioesterase, putative	At3g25110	yes	X	S	K		
120159	205637	5710633	LSD1	zinc-finger protein Lsd1	At4g20380	yes	X	Y	K		
187891	205913	5710702	STA1	ADP-glucose pyrophosphorylase large subunit	At5g19220	yes	X	C	K		
194793		5710812	STT7	Chloroplast protein kinase required for state transitions	At1g68830	yes	X	C	K		
187188	205915	5711123	LCI20	Putative 2-oxoglutarate/malate translocator	At5g64290	yes	P	C	K		

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136037		5705101	STA6	ADP-glucose pyrophosphorylase small subunit	At5g48300	yes	X	C	K		
82986		5705899	RBCS1	ribulose-1,5-bisphosphate carboxylase/oxygenase small subunit 1, chloroplast precursor	At1g67090	yes	X	S	K		
60238		5705899	RBCS2	ribulose-1,5-bisphosphate carboxylase/oxygenase small subunit 2,chloroplast precursor	At1g67090	yes	X	S	K		
120177	184971	5705906	PSAD	Photosystem I reaction center subunit II, 20 kDa	At1g03130	yes	X	T	K		yes
130914		5706419	PSAF	photosystem I reaction center subunit III, chloroplast precursor	At1g31330	yes	X	T	K		yes
58334		5706509	FTSH2	membrane AAA-metalloprotease, chloroplast	At2g30950	yes	X	T	K		
147787		5707952	PETF	Apoferredoxin, chloroplast precursor	At1g60950	yes	P	C	K		
99956	205935	5708618	PSAL	Photosystem I reaction center subunit XI	At4g12800	yes	X	T	K		yes
128002	205856	5708969	CTH1	Copper target homolog 1, chloroplast precursor, functional variant	At3g56940	yes	X	T/iV	K		
140452		5708981	STA11	4-alpha-glucanotransferase	At5g64860	yes	X	C	K		
76146		5710097	PSAE	photosystem I 8.1 kDa reaction center subunit IV	At2g20260	yes	X	T	K		yes
195343		5710878	CCS1	c-type cytochrome synthesis 1	At1g49380	yes	X	T	K		
58407	205944	5705194	MINE1	chloroplast division site-determinant MinE	At1g69390	yes	X	C	K		
196011	205938	5705280	CTP2	heavy metal transporting ATPase (HMA), P-type ATPase superfamily, membrane protein	At5g21930	no	X	T	K		
192478		5706435	PSAK	photosystem I reaction center subunit psaK, chloroplast precursor	At1g30380	yes	X	T	K		yes
178067		5710773	TGD2	related to a permease-like component of an ABC transporter involved in lipid transfer from ER to chloroplast	At3g20320	no	X	iV	K		
195951		5709624	CAO	Chlorophyll a oxygenase, chloroplast precursor	At1g44446	yes	X	T	K		

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115641		5709014	PRMT3	Protein arginine N-methyltransferase, ChromDB PRMT3403	At3g12270	no	P		KI		
139925		5705554	FKB17-2	peptidyl-prolyl cis-trans isomerase, FKBP-type	At1g18170	yes	X	T	KI		
148057	33411	5705619	PSBP1	Oxygen Evolution Enhancer 2 of photosystem II	At1g06680	yes	X	L	KI		
170734		5706157	RAD9	DNA damage checkpoint protein	At3g05480	no	P	M	KI		
37152	205890	5707139	CYN23b	conserved expressed protein cyclophilin type, alternatively spliced form B	At1g26940	yes	P	D	KI		
127879		5707181	PSBP6	conserved expressed lumen targeted protein related to OEE2 protein	At5g11450	yes	X	L	KI		
149307	184451	5708551	PSBP9	PsbP-like protein of PSII	At3g56650	yes	X	L	KI		
193859		5708792	FKB16-4	peptidyl-prolyl cis-trans isomerase, FKBP-type	At3g10060	yes	X	L	KI		
100415	205898	5708843	CYN37	Peptidyl-prolyl cis-trans isomerase, cyclophilin-type	At3g15520	yes	X	T/L	KI		
156074		5709051	FKB18	peptidyl-prolyl cis-trans isomerase, FKBP-type	At1g20810	yes	X	L	KI		
104731	205916	5709488	PSBP4	lumenal PsbP-like protein	At4g15510	no	X	L	KI		
196705		5709748	FDX4	Apoferredoxin, chloroplast precursor	At4g14890	yes	P	C	KI		
188000		5710057	FKB20-2	peptidyl-prolyl cis-trans isomerase, FKBP-type	At3g60370	yes	X	L	KI		
176		5710400	ERD2B	KDEL Receptor B	At1g19970	no	P	ER	KI		
30719		5710576	CYN26	Peptidyl-prolyl cis-trans isomerase, cyclophilin-type	At1g74070	yes	P	L	KI		
142309		5707918	CEM1	putative proton extrusion protein cemA, chloroplastic	At4g31040	yes	P	C	KI		
115135		5708183	SYK1	putative tRNA synthetase class II (D, K and N) family protein	At3g13490	no	X	C/M	KI		
183308	162260	5707776	CGL1	conserved expressed protein of unknown function	At2g20920	yes	P	C	U		

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97676	142225	5709778	CGL81	conserved expressed protein of unknown function	At5g04440	yes	P	C	U		
187386	182434	5705069	CGL3	conserved expressed protein	At2g17972	yes	X	T	U		
178040	205579	5705095	CGL4	conserved expressed protein	At3g58800	yes	P	M	U		
131754	205884	5705147	CGL5	conserved expressed protein	At1g64680	yes	P	M	U		
166032	205885	5705652	CGL6	unknown function	At5g11960	yes	P		U		
155494		5706349	CGL9	conserved expressed protein	At1g19360	yes	P	M	U		
162817		5706744	CGL10	conserved expressed protein	At5g05360	yes	P	C	U		
193961	206051	5706767	CGL11	conserved expressed protein of unknown function	At4g24930	yes	X	L	U		
163196	184127	5706820	CGL12	conserved expressed protein	At5g39790	yes	P	M	U		
189798		5706832	CGL13	conserved expressed protein	At1g08030	yes	P	M	U		
145947		5708027	CGL16	conserved protein	At1g07040	no	P	C	U		
185270		5708074	CGL17	conserved expressed protein	At1g50020	yes	X	T	U		
191999	205653	5708231	CGL18	conserved expressed protein of unknown function	At5g55570	yes	P	C	U		
193846		5708436	CGL20	conserved expressed protein of unknown function	At2g17240	yes	P	C	U		
174967	206053	5708582	CGL21	conserved protein of unknown function	At5g08540	yes	X	T/V	U		
154373		5708649	CGL23	conserved expressed protein	At1g74530	yes	P	M	U		
172656		5708652	CGL24	conserved protein of unknown function	At1g28100	no	P		U		
101500	205896	5708811	CGL25	conserved hypothetical protein	At5g13500	yes	P	D	U		
175106	205897	5708813	CGL26	conserved expressed protein	At4g29520	yes	P	D	U		
168688		5708844	CGL27	conserved expressed protein of unknown function	At1g62780	yes	P	C	U		
167685		5709055	CGL29	conserved protein of unknown function	At3g55760	no	P	C	U		
157731	205900	5709109	CGL30	conserved expressed protein	At1g77090	yes	X	L	U		
186351	160683	5709179	CGL32	conserved expressed protein of unknown function	At3g57280	yes	X	iV	U		
117853	184125	5709207	CGL34	conserved expressed protein of unknown function	At5g24690	yes	X	iV	U		

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187792	182630	5709309	CGL36	conserved expressed protein of unknown function	At3g10405	yes	P	C	U		
117798	190875	5709352	CGL38	conserved expressed protein of unknown function	At1g50900	yes	P	C	U		
148272	205902	5709379	CGL39	conserved expressed protein of unknown function	At5g27710	yes	P	D	U		
186576		5709522	CGL40	conserved expressed protein of unidentified function	At1g49975	yes	P	C	U		
162607		5709610	CGL41	conserved expressed protein of unknown function	At4g04330	yes	P	C	U		
151791	205904	5709615	CGL42	conserved expressed protein of unknown function	At5g48790	yes	P	C	U		
176794		5709684	CGL46	conserved expressed protein of unknown function	At5g65440	no	P		U		
190282		5709895	CGL52	conserved expressed protein of unknown function	At1g65230	yes	P	C	U		
155280	205907	5709903	CGL54	conserved expressed protein of unknown function	At1g05385	yes	P	C	U		
182361		5710094	TEF14	putative thylakoid lumenal protein	At4g02530	yes	X	L	U		
191642	158401	5710131	CGL59	conserved expressed protein	At5g44650	yes	X	T	U		
93364	205909	5710143	CGL60	conserved expressed protein of unknown function	At4g25660	yes	P		U		
144728	205911	5710298	CGL61	conserved expressed protein of unknown function	At4g22920	no	P	C	U		
144132		5710389	CGL63	conserved expressed protein of unknown function	At4g26410	yes	P		U		
196478	196477	5710502	LPB1	LPB1 Low Photochemical Bleaching 1 protein	At3g56040	yes	P	C	U		
187487		5710717	CGL64	conserved expressed protein	At3g19340	yes	P		U		
111993	182181	5711454	CGL68	Acid phosphatase/vanadium-dependent haloperoxidase related, DUF212	At1g67600	yes	P	D	U		

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196285		5706534	CGL70	conserved expressed protein of unknown function	At5g08400	yes	P	C	U		yes
172512	205936	5709382	CGL83	conserved expressed protein of unknown function	At3g61770	yes	P	C	U		
94227	205937	5705110	CGL73	expressed protein conserved in the green lineage	At3g51140	yes	X	iV	U		
180279		5705322	CGL74	conserved expressed tpr repeat protein	At1g78915	no	X	T	U		
114879	162021	5708641	CGL78	conserved expressed protein of unknown function	At5g58250	yes	P	C	U		
166730		5710953	CGL80	conserved protein of unknown function	At1g08530	no	P	C	U		
120035	205894	5708383	CGL19	conserved expressed protein with Dof type zinc finger	At2g34140	yes	P		UP		
195615		5710501	GRX6	Glutaredoxin 6, CGFS type, probably chloroplastic	At2g38270	yes	P	C	UP		
172865		5706882	PP2C6	protein phosphatase type 2C	At1g68410	no	P		UP		
157262		5709435	TRX10	Thioredoxin-like protein, unusual active site WCTKC	At3g53220	yes	P		UP		
191951		5709625	CGL43	Putative RNA binding protein with S1 domain	At3g23700	yes	P	C	UP		
78189		5709734	CGL49	ARF/SAR superfamily small monomeric GTP binding protein	At5g17060	yes	P		UP		
99082		5709854	EFG4	GTP-binding elongation factor-like protein, similar to yeast Hbs1p	At5g10630	no	P		UP		
122546	153782	5709880	CGL51	conserved expressed protein related to sugar phosphate/phosphate translocator	At5g17630	yes	P	C	UP		
116397	205906	5709898	CGL53	conserved protein related to carbohydrate hydrolase	At2g20680	no	P	D	UP		
167017		5711558	CGL69	conserved protein with lipase motif	At3g07400	yes	P	M	UP		
178217		5706431	CGL75	conserved protein with methyltransferase motif	At5g64150	no	P	C	UP		
157632	205939	5706797	CGL76	conserved expressed protein with epoxide hydrolase / esterase motif	At1g52510	yes	P	C	UP		

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104299	205940	5707941	SNE3	NAD-dependent epimerase/dehydratase	At4g20460	yes	P	D	UP		
190827		5710587	CGL2	putative methyltransferase	At3g01660	yes	P	M	UP		
120386		5705774	CGL7	conserved protein of unknown function related to chloroplast P translocator	At1g21070	no	P	B	UP		
166663	205638	5706161	CGL8	SPX containing protein	At5g20150	yes	P		UP		
77597	205888	5707134	CGL14	conserved expressed protein related to pantothenate kinase family	At4g35360	yes	P		UP		
117924	205891	5707556	CGL15	conserved expressed permease of the major facilitator superfamily	At5g20380	yes	P	B	UP		
195501		5707906	SEC12	regulator of COP-II vesicle coat	At5g50550	yes			UP		
145283		5708634	CGL22	cdc48-like protein	At2g03670	no	P		UP		
183721	205901	5709154	CGL31	conserved expressed protein with pterin carbinolamine dehydratase domain	At5g51110	yes	P	C	UP		
142662		5709193	CGL33A	conserved methyltransferase of unknown function	At5g63100	no	P	M	UP		
142678		5709193	CGL33B	conserved methyltransferase of unknown function	At5g63100	no	P	M	UP		
144101	205630	5709217	CGL35	conserved expressed protein of unknown function	At4g17760	yes	P	U	UP		
132449		5709240	CGL82	Conserved, expressed protein with meprin and TRAF homology domain	At5g43560	yes	P		UP		
170340	205640	5709603	HEP2	Hsp70 escorting protein 2	At5g27280	yes	P	C	UP		
172469		5709663	CGL44	RabGAP/TBC Domain Protein	At5g53570	no	P	C	UP		
144926		5709668	CGL45	expressed protein conserved in the green lineage	At1g11800	yes	P	M	UP		
171590		5709693	CGL47	conserved protein with F-box domain	At5g45360	no	P		UP		
116762		5709718	CGL48	conserved protein related to lysine decarboxylase domain	At1g50575	no	P	C	UP		
144505	205639	5709721	PLAP4	conserved expressed protein related to plastid lipid associated protein PAP	At5g19940	yes	X	iV	UP		

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161769		5709745	CGL50	conserved expressed protein of unknown function	At4g09620	yes	P	C	UP		
167673		5709941	CGL56	conserved expressed protein	At3g59780	yes	P		UP		
178070		5710013	CGL57	conserved expressed protein	At2g31140	no	P	M	UP		
149808	205908	5710021	CGL58	conserved expressed protein	At2g01810	yes	P		UP		
194876		5710111	TIM22B	translocase of inner mitochondrial membrane 22 homolog	At5g24650	yes	X	M/C	UP		
172486		5710341	CGL62	putative cell cycle associated protein	At1g67270	no	P		UP		
174855	205914	5710771	CGL65	conserved expressed protein	At1g69210	no	P	M	UP		
148832	205917	5711124	CGL66	conserved expressed protein	At4g38640	yes	P		UP		
189296	145554	5711125	CGL67	Mpv17/PMP22 family protein with unknown function	At2g42770	yes	P	X	UP		
192099		5711472	TEF5	Rieske [2Fe-2S] domain, putative, chloroplast location proposed	At1g71500	yes	X	T	UP		
184916		5707937	CGL71	conserved TPR repeat protein related to YCF37	At1g22700	yes	X	T	UP		yes
96690	140949	5710543	CGL72	conserved expressed protein with hemolysin motif and RNA methyltransferase motif	At3g25470	no	P	M	UP		
148091		5707836	CGL77	conserved protein with unknown function	At1g60990	no	P	Y	UP		
169453		5709802	CGL79	conserved protein with carbohydrate kinase motif	At1g19600	yes	P		UP		

CPLD goes to 53 and stands for Conserved in the Plant Lineage and Diatoms = in green + Cme + 1 diatom

CGL goes to 83 and stands for Conserved in the Green Lineage = in only green

CPL goes to 11 and stands for Conserved in the Plant Lineage = in green + Cme

CGLD goes to 30 and stands for Conserved in the Green Lineage and Diatoms = in green + 1 diatom

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Proteins in CiliaCut, but neither CentricCut nor MotileCut																
24475	5705258		(ARL6a , BBS3B) Most similar to mammalian ARL6, causative gene for Bardet-Biedl syndrome 3, member of the ARF/Sar1 GTPase family. The C. elegans ARL6 undergoes intraflagellar transport. Two ARL6 paralogs in Chlamydomonas (see ARL6b).	GTP-Binding	BBS	Y		Y	Y					Y		
189076	5706101	FAP47		No data		Y		Y		Y	Y			Y		
140113	5706980	BBS8	Tetratricopeptide repeat protein 8 (Bardet-Bield syndrome 8) similarity	Protein-protein interraction	BBS	Y		Y	Y	Y						
130394	5708472	D1bLIC	Cytoplasmic dynein 1b light intermediate chain (homologue of mammalian D2LIC/LIC3), the retrograde motor for intraflagellar transport.	Flagellar transport	BBS	Y		Y		Y	Y			Y		
190054	5709103	BBS7	Simlar to Bardet-Biedl syndrome 7	No data	BBS	Y		Y		Y						
101137	5709290	BBS9	Bardet-Biedl syndrome 9	No data	BBS					Y						
182299	5709609	BBS5	Similar to Bardet-Biedl syndrome 5	No data	BBS, Duane retraction syndrome 2	Y		Y	Y	Y				Y		
185788	5709716	UNC119	Signal transduction protein	Trafficking	Bone mineral density variability 3		Y	Y	Y	Y						
98915	5709897	FBB17		Protein metabolism					Y	Y						
126758	5711323	BBS2		No data	BBS, C8 deficiency, type I	Y		Y		Y						
192205	5705047	IFT140	Intraflagellar transport particle protein IFT140	Flagellar transport	1p36 deletion syndrome; Bone mineral density variability 3	Y		Y		Y	Y			Y		
182072	5705822	IFT20	Intraflagellar transport particle protein 20	Flagellar transport	Bone mineral density variability 3	Y		Y	Y		Y			Y		
143468	5706244	FAP66		Protein-protein interraction		Y		Y	Y	Y	Y			Y		

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183240	5708965	IFT172	Intraflagellar transport protein 172	Flagellar transport		Y		Y	Y	Y	Y	Y		Y		
195385	5709351	FAP118	(FBB1) In the flagellar basal body proteome as FBB1	Protein-protein interaction		Y		Y	Y	Y	Y			Y		
24171	5709540	IFT80	Intraflagellar transport particle protein 80	Flagellar transport		Y		Y	Y	Y	Y	Y		Y		
81760	5709994	DYF13	(FBB2) Homologous to protein required for ciliogenesis in C. elegans.	Protein-protein interaction	Thromboxane synthase deficiency	Y		Y	Y	Y		Y		Y		
126867	5710863	FAP60		Protein-protein interaction		Y		Y		Y	Y			Y		
195877	5708672	FAP9	Ortholog RABL5 in human, member of the Ras superfamily of GTPases but the GTP-specificity motif abrogated (ATPase?).	GTP-Binding		Y				Y		Y			Y	Y
194946	5710084	PTP1	Putative Protein Tyrosine Phosphatase 1; Dephosphorylates phosphotyrosine residues	Signalling				Y		Y					Y	Y
102300	5706231	SSA1		Signalling		Y					Y			Y		
150669	5706961	SSA2		Protein metabolism						Y						
150490	5709110	SSA3	Contains an engulfement and cell motility, ELM, domain (IPR006816) found in a number of eukaryotic proteins involved in the cytoskeletal rearrangements required for phagocytosis of apoptotic cells and cell motility.	Unclear						Y						
107835	5709188	SSA4		No data	Deafness, autosomal dominant 2				Y	Y					Y	Y
176942	5709323	SSA5		Signalling					new gene model in v3							
176788	5709889	SSA6		No data					new gene model in v3							
180447	5709065	SSA7		Protein-protein interaction					new gene model in v3							
95290	5709079	SSA8		Metabolism						Y						
172167	5707709	SSA9		RNA metabolism					new gene model in v3							

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172866	5709515	SSA10	Similar to copper type II, ascorbate-dependent monooxygenase , similar to dopamine beta-monooxygenase	Metabolism	Amyotrophic lateral sclerosis-4, juvenile dominant				Y	Y						
118345	5708916	SSA11		Microtubule Regulation and Metabolism					Y	Y						
Proteins in MotileCut but not CentricCut																
169142	5705315	MOT1		Protein-protein interaction					new gene model in v3							
171647	5706202	RTN1		No data					new gene model in v3							
149708	5705337	FAP44		Protein-protein interaction		Y		Y	Y	Y	Y					
194338	5709355	FAP57	Hypothetical protein contains WD40 repeats	Protein-protein interaction	Deafness, autosomal dominant 2	Y		Y		Y	Y			Y		
112249	5710479	POC1	Found in basal body proteome [PMID: 15964273].	Protein-protein interaction			Y		Y						Y	
169983	5710231	MOT2		Unclear					new gene model in v3							
189109	5706889	FAP59		No data		Y					Y			Y		
148926	5705281	MOT3		No data					Y	Y						
144011	5705309	FAP61	(FAP61)	No data		Y		Y		Y	Y		Y	Y		
134599	5705986	DHC6	Dynein heavy chain 6 (putative flagellar inner arm dynein heavy chain)	Flagellar Structure		Y			Y	Y	Y	Y	Y			
167096	5706093	FAP74		No data	1p36 deletion syndrome; Bone mineral density variability 3	Y				Y	Y					
176821	5706137	MOT4		Unclear					new gene model in v3						Y	Y
154904	5706505	FAP263		No data	C8 deficiency, type I	Y		Y		Y				Y		
173632	5707289	MOT5	Alanine rich novel protein	No data		Y										
145396	5707456	FAP251		Protein-protein interaction		Y		Y		Y	Y			Y		
102649	5708715	MOT6		Signalling					Y	Y						

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146683	5708925	MOT7		No data										Y		
142494	5709113	FAP75	Flagellar Associated P-Loop Containing Protein	No data		Y							Y			
194218	5709378	MOT8		No data				Y	new gene model in v3							
194403	5709602	MOT9		No data					new gene model in v3							
190077	5709733	FAP155		Protein-protein interaction		Y		Y		Y	Y			Y		
165974	5710024	MOT10		Membrane Protein					new gene model in v3							
126569	5710065	MOT11		Microtubule Regulation and Metabolism				Y		Y						
13542	5710242	POC11	Found in basal body proteome as POC11 [PMID: 15964273].	No data	1p36 deletion syndrome; Bone mineral density variability 3		Y			Y						
195529	5710730	ARLP1	(ARL13) Expressed Protein. ARF-like 13, a member of the ARF/Sar1 family of Ras-like GTPases. C. elegans ortholog specifically expressed in flagellated cells	GTP-Binding		Y		Y	new gene model in v3							
188246	5710879	FAP69		Protein-protein interaction		Y		Y			Y			Y	Y	Y
121332	5711016	MOT12		No data						Y						
21780	5711546	ARM1	contains armadillo (Arm) repeat	Protein-protein interaction					new gene model in v3							
146448	5711707	FAP122		Signalling		Y				Y	Y			Y		
190653	5706159	FAP94		No data		Y		Y			Y			Y		
103782	5706246	PF16	Central pair associated protein	Flagellar Structure	Bone mineral density variability 3	Y					Y	Y	Y	Y	Y	Y
177591	5706388	FAP50		Protein-protein interaction		Y					Y				Y	
192295	5706575	FAP184		No data		Y		Y		Y	Y		Y	Y		
105624	5708696	FBB9		No data						Y				Y		
130542	5709331	VFL3		Flagellar Structure	C8 deficiency, type I			Y		Y						

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119090	5709342	FAP198	Presence of a cyt-b5 -like domain in the N terminal part of the protein	Unclear	Blood group, Radin antigen; Blood group, Scianna system; Bone mineral density variability 3	Y			Y	Y	Y			Y		
138013	5709708	SPEF1		Unclear		Y		Y		Y		Y	Y			
116240	5709840	HY3	(HYD3) Similar to mouse hydrocephaly protein hydin HY3	Unclear	Mouse: Hydrocephaly	Y				Y	Y		Y	Y		
141685	5710115	MOT15		Signalling				Y	new gene model in v3							
101210	5710416	PF20	WD-repeat containing protein PF20 of the central pair of the flagella. Associates with the intermicrotubule bridge.	Flagellar Structure		Y				Y	Y			Y		
193672	5710456	MOT16		Unclear				Y	new gene model in v3					Y	Y	Y
188960	5710498	FAP81		Unclear		Y		Y			Y					
193355	5710580	FAP43		Protein-protein interaction		Y		Y		Y	Y			Y		
195517	5710628	RAB23		GTP-Binding				Y	Y							
130324	5710955	DHC2	Dynein heavy chain 2 (putative flagellar inner arm dynein heavy chain)	Flagellar Structure	Aicardi-Goutieres syndrome 1	Y		Y		Y			Y			
138046	5710963	RSP3	Flagellar radial spoke protein 3 (RSP3). axonemal A-kinase anchoring protein KAP [PMID: 11309423; PMID: 16571668; PMID: 16267272; GI:134041]. Gene originally termed PF14 [PMID: 7204490; PMID: 2745550; PMID: 2377611]	Flagellar Structure		Y		Y	Y	Y	Y	Y	Y	Y	Y	Y
192915	5711490	MOT17		Signalling	Rippling muscle disease-1			Y						Y		
172483	5711560	FAP134		Protein-protein interaction	Bone mineral density variability 3	Y				Y	Y			Y		

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182960	5711716	RSP9	Flagellar radial spoke protein 9; A subunit in the radial spoke head; Gene originally termed pf17 [PMID: 7204490, PMID: 16507594; GI:83284713]	Flagellar Structure		Y		Y		Y	Y	Y	Y	Y	Y	Y
177575	5708309	MOT18		RNA metabolism					new gene model in v3						Y	
173947	5710187	MOT19		Membrane Protein					new gene model in v3							
192150	5709694	MOT20		Membrane Protein					new gene model in v3							
135100	5709694	MOT21	putative phosphate/phosphoenolpyruvate translocator protein	Membrane Protein				Y	new gene model in v3							
93765	5708783	FAP240		No data		Y					Y			Y		
177375	5709293	MOT22		Signalling					new gene model in v3							
173608	5710430	MOT23		Signalling					new gene model in v3							
184899	5710794	MOT24		Flagellar Structure		Y		Y	Y					Y		
192442	5711802	MOT25		Unclear				Y		Y				Y		
196807	5711862	ELG34	exostosin-like glycosyltransferase	Metabolism				Y	new gene model in v3							
191232	5711862	EGL12	exostosin-like glycosyltransferase	Metabolism				Y	new gene model in v3							
150998	5711866	TEX9	(FBB15)	No data	C8 deficiency, type I			Y		Y				Y		
186414	5709707	KLP1	Kinesin-like protein 1; kinesin associated with one of the central pair microtubules of the flagellar axoneme	Flagellar Structure	Aicardi-Goutieres syndrome 1	Y		Y			Y			Y		
189194	5711344	FAP146		No data	Deafness, autosomal dominant 2	Y		Y		Y	Y	Y		Y		
136082	5711457	UNI3	Delta tubulin (TUD)[gi:7441381] Required for assembly of the basal body/centriole and localizes to the basal body	Flagellar Structure			Y		new gene model in v3							

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117499	5709647	MOT26		No data	Myasthenia gravis, neonatal transient					Y						
168675	5707372	MOT27		No data					new gene model in v3					Y		
150732	5710991	SSMT	SET domain-containing methyltransferase; catalyzes methylation of the N-terminal alpha-amino group of the processed form of RuBisCO small subunit prior to holoenzyme assembly	Protein-protein interraction				Y	new gene model in v3							
140873	5711553	SMP10	(PRP1) Predicted snRNP core protein; SMP10 name replaces previous PRP1 name	RNA metabolism					new gene model in v3							
177784	5705352	MOT28		No data					new gene model in v3						Y	
181739	5709937	MOT29		No data					new gene model in v3							
179771	5709937	MOT30		No data					new gene model in v3							
151105	5710089	MOT31		Protein-protein interaction					new gene model in v3							
109243	5710241	MOT32		No data												
106614	5710241	MOT33		No data					new gene model in v3							
107462	5710241	MOT34		No data	1p36 deletion syndrome; Bone mineral density variability 3											
141109	5710324	FAP46		No data		Y					Y					
191879	5710410	MOT35		No data												
187854	5710441	FAP161		No data		Y		Y			Y		Y		Y	Y
190937	5710532	FBB5		No data				Y		Y				Y		
188180	5710541	FAP111		Protein-protein interaction		Y		Y		Y	Y			Y	Y	

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6466	5710586	POC18	Found in basal body proteome [PMID: 15964273]	Protein-protein interaction	Epilepsy, myoclonic, benign adult familial; Macular dystrophy, atypical vitelliform		Y		Y	Y						
180607	5710673	MOT36		No data												
146778	5711548	MOT37		Protein-protein interaction	1p36 deletion syndrome; Bone mineral density variability 3					Y						
151348	5711741	MOT38		No data						Y						
173581	5710018	MOT39		Protein turnover	Triphalangeal thumb-polysyndactyly syndrome					Y					Y	Y
188195	5710277	BLD2	Epsilon tubulin (TUE) [gi:20514387, PMID: 12429830]	Flagellar Structure			Y	Y	new gene model in v3							
189500	5710403	MOT40		No data				Y	new gene model in v3					Y		
142470	5710516	MOT41	This gene is in the location of Probe 2 used in PMID: 11805055.	No data				Y		Y				Y		
152883	5705024	MOT42		Protein turnover				Y	new gene model in v3							
179158	5710249	FAD5b	Fatty acid desaturase like, similar to Arabidopsis putative FAD5	Membrane synthesis/differentiation					new gene model in v3							
122385	5710249	FAD5D		Membrane synthesis/differentiation					new gene model in v3							
153533	5710249	FAD5C		Membrane synthesis/differentiation					new gene model in v3							
94516	5710642	MOT13		DNA Binding					new gene model in v3							
43319	5705234	MOT14		Unclear	1p36 deletion syndrome; Bone mineral density variability 3									Y		

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134605	5711269	CDJ2		Chaperone				Y	new gene model in v3							
Proteins in both CentricCut and MotileCut																
107408	5707103	MOT47		Protein-protein interraction	Epilepsy, myoclonic, benign adult familial; Macular dystrophy, atypical vitelliform					Y				Y		
196793	5708194	FAP45	Identified in Chlamydomonas basal body proteome as BUG28 [Keller et al., 2005; PMID: 15964273]. Weakly similar to nasopharyngeal epithelium-specific protein 1	No data	Leukemia, acute pre-B-cell; Atherosclerosis, susceptibility to	Y		Y	Y			Y				
129295	5708315	KIF6		Trafficking				Y						Y		
180221	5708949	NDK7	(BUG5) in basal body proteome as BUG5 [PMID: 15964273].	Metabolism		Y					Y	Y	Y	Y		
126286	5709885	Rib72	novel component of the ribbon compartment of flagellar microtubules.	Flagellar Structure		Y		Y		Y	Y	Y	Y	Y		
143267	5710023	MOT48		No data	C8 deficiency, type I			Y		Y						
182403	5709730	C1	RNA-binding protein with three KH domains and a protein-protein interaction domain (WW) at the C-terminus Subunit of the circadian RNA-binding protein CHLAMY 1 (Zhao et al., Euk. Cell, in press)	RNA metabolism				Y		Y						
186669	5707578	DLC1	Flagellar outer dynein arm light chain 1	Flagellar Structure		Y		Y	Y	Y	Y			Y		
115671	5706503	ECH1		Membrane synthesis/differentiation					new gene model in v3							
107386	5706608	MOT50		Protein metabolism												

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192170	5706578	MB02	Move backward only mutant defective in the production of the ciliary waveform. Mbo2p is a novel coiled-coil protein.	Protein-protein interaction	Aneurysm, intracranial berry	Y		Y		Y	Y		Y			
106450	5709417	FAP250		No data		Y				Y	Y			Y		
169559	5709521	FBB15		Signalling		Y				Y	Y			Y		
137793	5710049	NKRN1	(FAP106)	Signalling	Bone mineral density variability 3	Y		Y		Y	Y			Y		
168908	5708554	MOT51		Unclear						Y						
135463	5705051	FBB4		Signalling	Retinitis pigmentosa-10	Y		Y		Y				Y		
186878	5705821	FAP100		No data	Renal cell carcinoma; Glaucoma 1C, primary open angle	Y		Y		Y	Y			Y		
121413	5709547	FAP73		No data	Blood group, Radin antigen; Blood group, Scianna system; Bone mineral density variability 3	Y				Y	Y			Y		
132719	5709873	ODA1	Flagellar outer dynein arm-docking complex subunit 2 (ODA-DC 2)	Flagellar Structure		Y		Y	Y	Y	Y		Y	Y	Y	
132143	5710434	CTO59	(FBB5) Similar to C21orf59 (CTO59)	No data				Y	Y	Y				Y	Y	Y
192441	5705837	RABL2A	Expressed Protein. Distantly similar to a class of Rab-like proteins from mammals.	GTP-Binding				Y						Y		
3686	5705975	RJL1	Expressed Protein. Member of the RJL family in the Ras superfamily of GTPases (Nepomuceno-Silva et al. 2004, Gene 327:221-32)	GTP-Binding				Y								
77703	5708336	RIB43a	Coiled-coil protein associated with protofilament ribbons of flagellar microtubules (PMID 10637302).	Flagellar Structure		Y		Y	Y	Y	Y	Y		Y		
192763	5709569	MOT52		Microtubule Regulation and Metabolism	Bone mineral density variability 3			Y		Y						

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24252	5709882	ODA4	Flagellar outer dynein arm heavy chain beta	Flagellar Structure	Blood group, Radin antigen; Blood group, Scianna system; Bone mineral density variability 3	Y		Y		Y	Y	Y		Y		
188612	5709886	ODA6	Flagellar outer dynein arm intermediate chain 2, IC2, ODA-IC2, IC69, IC70	Flagellar Structure		Y		Y	Y	Y	Y	Y	Y	Y		
149002	5710454	MOT53		Protein-protein interaction					new gene model in v3							
188421	5710529	MOT54		Protein-protein interaction		Y								Y		
144241	5710529	FAP264		Protein-protein interaction				Y						Y		
145799	5711409	MOT43		Unclear					new gene model in v3							
189445	5711409	FAP147		Unclear				Y	new gene model in v3		Y					
142227	5708384	MOT44		Membrane Protein	C8 deficiency, type I					Y						
160148	5705956	FBB11		No data				Y		Y						
175396	5711057	MOT45		No data					new gene model in v3							
195180	5706594	MOT46		RNA metabolism	Hypotrichosis, Marie Unna type; Schizophrenia			Y								
129193	5709938	FAP156	Expressed Protein. Rab-type GTPase distantly related to Rab-like proteins from mammals.	GTP-Binding		Y		Y			Y	Y		Y	Y	
116664	5709999	MOT49		Protein-protein interaction					Y					Y	Y	Y
151144	5706167	PF2	Component of dynein regulatory complex (DRC) of flagellar axoneme; has similarity to mammalian growth-arrest specific gene product (Gas11/Gas8), trypanin related, PMID: 10969087 PMID: 11864997	Flagellar Structure		Y		Y	Y	Y	Y	Y	Y	Y		

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Proteins in CentricCut but not MotileCut																
128114	5706542	FAP52	(BUG14) in basal body proteome as BUG14 [PMID: 15964273].	Protein-protein interaction	Blood group, Radin antigen; Blood group, Scianna system; Bone mineral density variability 3	Y		Y	Y		Y			Y		
131284	5708587	DIP13	Similar to Sjogren's syndrome nuclear autoantigen 1.	Microtubule Regulation and Metabolism	Amyotrophic lateral sclerosis-4, juvenile dominant; Leukemia, T-cell acute lymphoblastic	Y	Y	Y		Y	Y			Y		Y
195496	5709491	SSA12		Unclear				Y	Y							
100760	5710578	FAP267		Microtubule Regulation and Metabolism		Y				Y						
127720	5706974	SSA13		Signalling	Epilepsy, myoclonic, benign adult familial; Macular dystrophy, atypical vitelliform			Y	Y	Y						
171688	5709575	SSA14	Some similarities with flavoprotein monooxygenases	Metabolism					new gene model in v3							
3897	5709171	DPY30	(FBB12) Desc Chromatin modifying protein complex member, identified by mutations in C. elegans defective in male sensory behavior.	Protein-protein interaction		Y		Y		Y		Y	Y			
97201	5705256	PACRG1	(BUG21) in basal body proteome as BUG21 [PMID: 15964273]. Homologous to mammalian PACRG parkin co-regulated gene.	No data		Y	Y		Y	Y	Y		Y	Y	Y	Y
191923	5709909	SSA15		Protein-protein interaction				Y	new gene model in v3							

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192420	5705900	FAP22	Similar to D. rerio cystic kidney disease gene qilin	No data	1p36 deletion syndrome; Bone mineral density variability 3	Y		Y	Y	Y	Y	Y		Y		
128761	5708192	ARLC2	(ARL3) Expressed Protein. Similar to the ARLC-type GTPases. , ARF-like 3, a member of the ARF/Sar1 GTPase family. Experimental evidence and presence only in organisms with flagella suggest a function in the flagellum/basal body.	Microtubule Regulation and Metabolism		Y		Y	Y		Y					
132451	5709415	IDA4	Flagellar inner arm dynein light chain p28	Flagellar Structure	Deafness, autosomal dominant 2	Y		Y	Y	Y	Y	Y	Y	Y		Y
24116	5709496	BLD1	(IFT52) Intraflagellar transport protein IFT52(Curr Biol. 2001. 11(20):1591-4. The C. elegans homologue is osm-6.	Flagellar transport		Y		Y	Y	Y	Y	Y		Y		
185392	5709600	FAP116	Similar to Microtubule Interacting TNF Receptor-Associated Factor 3 Interacting Protein 1	Trafficking		Y		Y	new gene model in v3		Y			Y		
108954	5709902	FAP32		No data		Y			Y	Y	Y	Y		Y		
98642	5710979	IFT57	Intraflagellar transport particle protein 57	Flagellar transport		Y			Y	Y	Y	Y		Y		
128801	5711212	TPR5	(FAP259) Desc TPR protein with similarity to human FLJ30990. similar to dyf-1 (C. elegans)	Protein-protein interraction	Duane retraction syndrome 2	Y		Y	Y	Y		Y		Y		
24421	5705296	IFT88	Intraflagellar transport particle protein 88	Flagellar transport	Bone mineral density variability 3; Mouse model: PKD, RP	Y		Y	Y	Y	Y	Y		Y		
147682	5706286	PP11		No data	Deafness, autosomal dominant 2			Y								
182554	5707722	KAP	Kinesin-associated protein; probable non-motor subunit of kinesin-II, the anterograde motor for intraflagellar transport.	Flagellar transport		Y		Y	Y	Y				Y		
138649	5707942	IFT81	Desc Intraflagellar Transport Protein 81	Flagellar transport		Y		Y		Y	Y	Y		Y		

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136521	5708482	IFT74/72	Intraflagellar transport particle protein 74/72	Flagellar transport		Y		Y		Y	Y			Y		
32880	5708502	NPH4	Found in basal body proteome as POC10 [PMID: 15964273]. Mammalian homolog is NPHP-4, also known as nephroretinin, gene mutated in Senior-Loken syndrome.	No data	Senior-Loken syndrome		Y	Y		Y						
130473	5709311	MKS1	Ortholog of the human Meckel Syndrome 1 gene	No data	Meckel Syndrome, Bone mineral density variability 3		Y	Y	Y	Y					Y	
169948	5706569	IRK1	putative inward rectifier K+ channel TC# 1.A.2	Membrane Protein	Bone mineral density variability 3					Y						
192430	5709753	SSA16		Signalling	Bone mineral density variability 3			Y		Y				Y		
129433	5710308	ODA9	Flagellar outer dynein arm intermediate chain 1, IC1, ODA-IC1, IC78	Flagellar Structure		Y		Y	Y	Y	Y	Y	Y	Y	Y	Y
169222	5706998	SSA17		Protein-protein interaction					new gene model in v3							
111541	5707143	SSA18		RNA metabolism						Y						
147671	5705982	SSA19		Protein metabolism						Y						
143218	5709875	SSA20		Protein metabolism						Y					Y	
TOTAL NUMBERS IN EACH COLUMN						88	10	102	47	106	68	28	21	85	24	17
TOTAL GENES IN DATASET THAT MAP TO v3:									187	687	331	138	73	146	318	126