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# Tracking N-terminal protein processing from the Golgi to the chromatophore of a rhizarian amoeba

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T.M. wrote the paper and performed the analysis.

The author responsible for distribution of materials integral to the findings presented in this article in accordance with the policy described in the Instructions for Authors (<https://academic.oup.com/plphys/pages/General-Instructions>) is: T. Meinzel ([thierry.meinzel@i2bc.paris-saclay.fr](mailto:thierry.meinzel@i2bc.paris-saclay.fr)).

Dear Editor,

Sample analysis from various ecosystems has substantially enriched recent knowledge about the diversity of life (Karsenti et al., 2011). One major aim of these efforts—facilitated by rapid advances in single and meta-genomic DNA sequencing—has been to explore the diversity of “invisible” or “uncultivable” organisms. While prokaryotes and viruses are by far the most abundant and biodiverse organisms on Earth (Lynch et al., 2014), the breadth of eukaryotic diversity is now better appreciated by complete genome sequences of model organisms in each class, facilitating and reinforcing classification efforts (Supplemental Figure S1). Several complete and well-annotated nuclear and organelle genomes contributing to eukaryotic diversity are now available through unique portals (Grigoriev et al., 2021).

The current representation of the eukaryotic tree of life is based on the dichotomy between Unikonta and Bikonta, both groups containing unicellular and multicellular organisms (Supplemental Figure S1). Unlike Unikonta, which are basically heterotrophic organisms, Bikonta often photosynthesize via a unique organelle, the plastid. There are three major phylogenetic branches of Bikonta: Archaeplastida, Chromalveolata, and Excavata (Supplemental Figure S1). Of these, the Archaeplastida include plants, green algae, and red algae, all of which are characterized by a common ancient primary endosymbiosis of a photosynthetic  $\beta$ -cyanobacterium ~1.6 billion years ago. This ancestral inclusion led to primitive photosynthetic organisms. Red and green algae (Supplemental Figure S1) are among the closest

representative fossils of this primordial event, since most other photosynthetic subgroups—Euglenozoa, Stramenopiles (diatoms), and Apicomplexa (Supplemental Figure S1)—acquired a secondary inclusion of a photosynthetic eukaryote (green or red algae) via phagocytosis into another non-photosynthetic eukaryote much more recently, ~190 million years ago (Falcatore et al., 2020). Other organelles, like in dinoflagellates, have a tertiary origin (Yoon et al., 2005).

Primary endosymbiosis of photosynthetic bacteria represents not only a remarkably ancient event but also a unique event creating sophisticated interdependence between the plastid and nucleus: genes migrated from the plastid to the nucleus and proteins were retargeted from the nuclear genome to the plastid. Exactly how proteins traffic to the plastid has been studied for decades in Archaeplastida. Most proteins display an N-terminal transit peptide (TP) that allows them to cross the plastid membranes, and this leader peptide is usually cleaved off once the protein enters the plastid (Christian et al., 2020). In the stroma, the proteins usually undergo further modifications including N-terminal acetylation (NTA) of the new N-terminus (Giglione and Meinzel, 2021) and some are retargeted to the thylakoid for further processing. Secondary endosymbiosis also gave rise to complex import mechanisms (Agrawal and Striepen, 2010; Hopkins et al., 2012).

As primary endosymbiosis is such a rare event, knowledge of at least one other independent similar event and how genes and proteins exchange between two genomes is key to a better understanding of eukaryogenesis, its trajectory,

and evolutionary events. *Paulinellidae*, which include the freshwater *Paulinella chromatophore* and *Paulinella micropora* and the marine *Paulinella longichromatophora* (Lhee et al., 2019), are euglyphid testate amoebae of the Rhizaria supergroup (Figure 1) displaying a siliceous shell and a photosynthetic organelle, like diatoms. Unlike diatoms, however, these organisms are unique as their plastid—the chromatophore—originates from the much more recent primary endosymbiosis of an  $\alpha$ -cyanobacterium only  $\sim 100$  million years ago and independent of Archaeplastida (Stephens et al., 2021). Despite difficulties in analyzing these organisms due to partial gene annotation (see, e.g. the 1 Gbp *P. micropora* [http://cyanophora.rutgers.edu/P\\_micropora/](http://cyanophora.rutgers.edu/P_micropora/) [Lhee et al., 2021]; the *P. chromatophora* genome is 10-fold larger) and lack of genetic manipulation tools, *Paulinella* spp. are useful models of primary endosymbiosis, a major event in eukaryogenesis that also involves the mitochondrion (Szathmáry, 2015).

A recent publication in *Plant Physiology* by Oberleitner et al. (2022) expands the current knowledge in this field. The authors extracted chromatophores from *P. chromatophora*. Using state-of-the-art mass spectrometry called “High-efficiency Undecanal-based N-Termini EnRichment” (HUNTER) (Weng et al., 2019), they assessed N-terminal protein processing in the organelle. HUNTER involves chemical labeling of free amino groups (i.e. N-termini and lysine chains) with dimethylation before trypsin-mediated protein hydrolysis into small fragments compatible with mass spectrometry. In this case, trypsin only cleaves after arginines because lysines are modified. After enrichment of N-terminal peptides, this approach allows easy recognition of protein N-termini, which are N-tagged due to chemical dimethylation or natural NTA, the most frequent N-terminal modification of eukaryotic proteins (Aksnes et al., 2019). Searches were carried out using > 60,000 open-reading frames (ORFs) translated from a *P. chromatophora* transcriptome and the



**Figure 1** Alignment of the full-length ORFs encoding the 60S ribosomal protein L12 (RPL12A) from various organisms. The ORFs encoding ribosomal protein L12 were aligned with ClustalX2.1. The ORF sequences derived from gene models from the Viridiplantae *Arabidopsis thaliana* (At2g37190, P508832), the rhizarian *Cercozoa* causative agent of cabbage clubroot, *Plasmodiophora brassicae* (P508832) as the closest relative of *Paulinellidae*, *Toxoplasma gondii* (A0A125YT49), *Homo sapiens* (P30050), *Saccharomyces cerevisiae* (P0CX53), *Trypanosoma brucei* (XP\_011775943.1), *Euglena gracilis* (QLA09605.1), *Dictyostelium discoideum* (Q54J50), the dinoflagellate *Polarella glacialis* (A0A813EJD3), and *Paulinella micropora* (g77960.t1). Both ORFs from *P. chromatophora* (m.87086 and m.108443) are derived from the transcriptome. None of the two corresponding L12 transcripts display the trans-splicing tag (CTTTCTG) and is therefore expected to correspond to a full-length transcript (nucleotide sequences of all *P. chromatophora* transcripts are available at <https://www.ncbi.nlm.nih.gov/bioproject/PRJNA311736>). The various in-frame start codons are colored in red. The numbering of the amino acid sequence is indicated on the right side in italics. The zone colored in yellow in *Paulinella* sequences shows significant divergence from the other otherwise very conserved sequences below. The N-terminal acetylated peptide sequence retrieved in this study is shown in green. NB PSI-BLAST of the N-terminal extension (137 residues) of *P. micropora* shows no homology with any known eukaryotic protein.

867 ORFs derived from the translated chromatophore genome (Singer et al., 2017), all available at <https://www.ebi.ac.uk/pride/archive/projects/PXD006531> and UniProtKB. The 28 ORFs from the mitochondrial genome—only available in *P. micropora* (<https://www.ncbi.nlm.nih.gov/bioproject/PRJNA587507>)—were not considered.

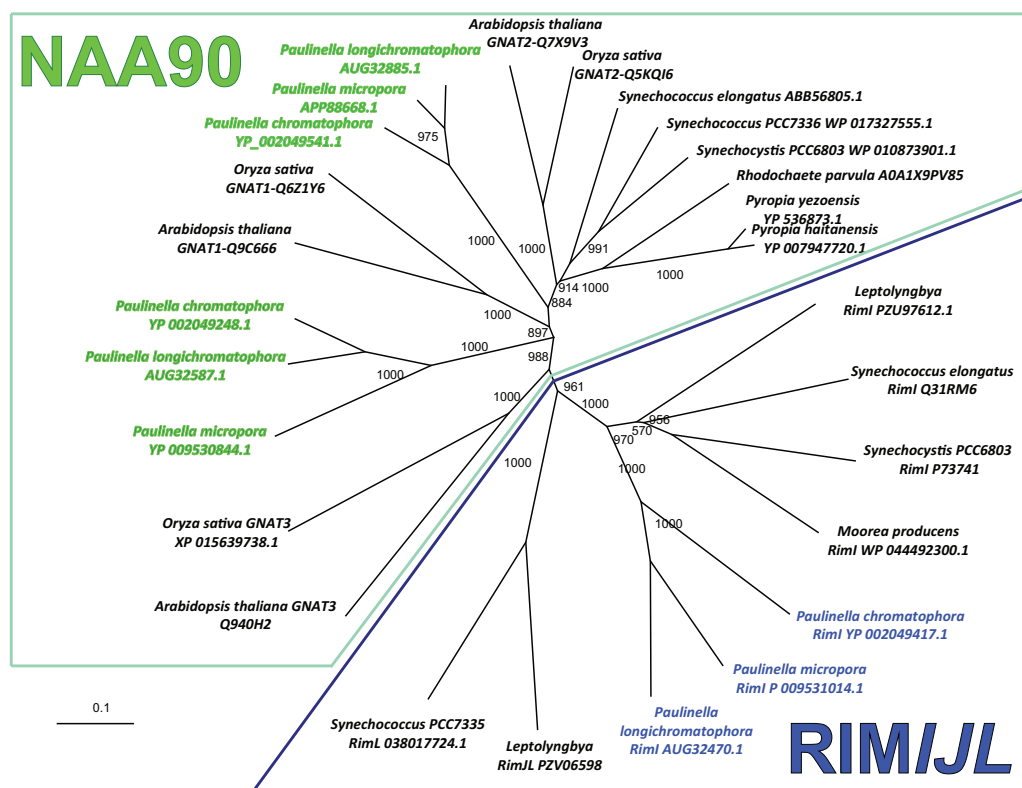
In their previous classical proteomic analysis, the same group had already suggested that chromatophores import two classes of nuclear-encoded proteins: one featuring N-terminal targeting signals (crTP) also functional in plants and the other devoid of these signals (Singer et al., 2017). The short imported proteins without crTP were long thought to traffic via the Golgi (Nowack and Grossman, 2012). Thanks to the cutting-edge mass spectrometry and sequence analysis performed in the current study, the authors make a decisive step forward in our understanding of proteins with a crTP. They show that this ~200-residue pre-sequence—which is longer than a chloroplast TP (< 100 residues)—is bipartite and only partially removed once in the chromatophore. The first part of the crTP contains a motif (Yxx $\phi$ , where  $\phi$  is a hydrophobic residue) believed to mediate protein trafficking via the Golgi through adaptor protein 1 complexes that bind to ADP-ribosylation factors (Park and Guo, 2014). The second domain is involved in translocation across the chromatophore inner membrane. Finally, the authors show that in addition to proteolytic cleavage of the crTP upon chromatophore import, many new N-termini also undergo NTA. This modification is also frequently found on cytosolic nuclear-encoded proteins and chromatophore-encoded proteins. The authors suggest that there is a dedicated cytosolic acetylation machinery similar to that seen in other eukaryotes (NatA/B/C/D/E). In addition, the authors suggest the existence of a dedicated NTA machinery in the chromatophore, but they failed to identify homologs with NatGs seen in all Archaeplastida (Giglione and Meinzel, 2021).

The study suggests that the recent acquisition of a novel organelle such as the plastid involves the Golgi as the initial route for protein targeting. This mechanism has already been suggested in *Oryza sativa* (rice, a land plant) for a couple of proteins both featuring cleavage of a 20–30 amino acid signal peptide and N-terminal glycosylation (Kitajima et al., 2009; Kaneko et al., 2016). Whether other Bikonta display such a mechanism is unknown and how Archaeplastida have progressively gained direct transfer from the cytosol to the plastid for a majority of the imported ORFs is a conundrum. Identifying other organisms with intermediary endosymbiosis is critical to resolving this problem.

While the added value of the biological model is very appealing, the HUNTER technology is limited in its capacity to further expand the data, as it relies on the quality of proteome annotation and sample preparation, as with any other mass spectrometry-based approach. As the authors mention, proteins may degrade during sample preparation or as a natural cellular process, and both natural and unnatural proteolytic species can accumulate. The authors indicate that

NTA is one marker of natural proteolysis. Due to high instability of the acetyl donor, acetyl-CoA, NTA is unlikely to occur once the organelle is disrupted. For instance, they identified various N-termini of the large subunit of Rubisco encoded in the chromatophore, none of which corresponded to the full length active subunit and several of them showing NTA. Rubisco is highly abundant and it is indeed likely that degradation intermediates may have accumulated. It is striking that these intermediates may be further post-translationally N-acetylated. One cannot exclude that—during fractionation—the chromatophore undergoes proteolytic stress leading in organello to this accumulation. Another bias in the approach is one inherent to subcellular fractionation: in such approaches, there is often contamination with other, closely linked subcellular compartments, such as mitochondria in chloroplast preparations from plants. In this study, the identification of 60S ribosomal (L12) and cytoskeleton (fascin or actin) protein homologs is evidence of such contamination, as mentioned and cross-checked by the authors in their Supplemental Table S1 (column AA, sheet “nucleus-encoded”). It shows cytosolic contamination was < 5% and confirms high purity of the chromatophore preparations (see also Supplemental Figure S1 and Supplemental Table S2 in Singer et al., 2017). Among the contaminants, one actin homolog displaying NTA at Glu3 is peculiar, since NTA is a unique processing activity displayed by NAA80 (NatH) on cytosolic actin (Aksnes et al., 2019). Furthermore, this unambiguous characterization reveals that in addition to the aforementioned set of N-terminal acetyl transferases (Nats), there exists a nuclear-encoded NatH in Cercozoa (NatH homologs have not been identified in Archaeplastida to date; Giglione and Meinzel, 2021). This further reinforces the idea that the primary eukaryote engulfing the cyanobacteria at the origin of the chromatophore is distinct. A third limitation arises due to the partial nuclear genomic information, its extreme complexity so far only addressed through 60,000 sequences from the transcriptomic analysis and no gene model, except by comparison with those derived from the *P. micropora* draft genome (Lhee et al., 2021). Many ORFs are only partial and the N-terminus is the most uncertain part, with a majority of ORFs lacking the canonical methionine start codon. As a result, the position of the N-terminus in many identifications is uncertain and it is difficult to conclude whether or not a signal peptide is cleaved or whether the cleavage is a priori canonical (e.g. peptides 631/635). It may also be true that the first Met codon is not the start codon and that another alternative Met (or an unusual non-AUG codon; Kearse and Wilusz, 2017; Na et al., 2018) is used (e.g. peptides 66/693/291 with Met6/32/37 and cleavages at positions 45/46/50). To reinforce their conclusions, the authors filter their search to only transcripts displaying the 5'-trans-splicing tag, demonstrating that the full-length cDNA was retrieved (see their Supplemental Table S1, column W). This corresponds to 26.1% of the nuclear encoded unique transcripts of the transcriptome (Supplemental Figure S6





**Figure 2** Two gene homologs with strong homologies to cyanobacterial and chloroplast *N*-acetyltransferases NatG occur in the chromatophore genome of *Paulinella* spp. A *Synechococcus elongatus* GNAT2 homolog was used to challenge BLAST *Paulinella* protein diversity. In total, 14 sequences were selected, and the Nats of several cyanobacteria, including RimiJL, were added to NAA90 members from rice (*O. sativa*) and *Arabidopsis* (*A. thaliana*) and red algae plastids. In total, 27 sequences were aligned and the bootstrap tree was constructed (Grzela et al., 2017). Internal values labeled on each node record the stability of the branch over the 1,000 bootstrap replicates. Together with NAA70, NAA90 is one of two subgroups of NatGs and made of GNAT1/2/3.

in Nowack et al., 2016). An interesting example is ribosomal L12, which corresponds to a strongly conserved protein over the entire length of its sequence—including the N-terminus—in Eukaryotes (Figure 1). In *P. chromatophora*, the L12 protein homolog is encoded by two transcripts (scaffold13378/m.87086 and scaffold19310/m.108443) only differing at their 5'-end, one long and the other much shorter. The retrieved peptide matches codon 35 after the fourth methionine of the long transcript and the other matches codon 2 after the first methionine of the short transcript (Figure 1). While the occurrence of NTA suggests that the peptide is present, the origin of the primary substrate, and therefore the exact N-terminal sequence, therefore needs further analysis. The authors select the short L12 transcript over the long transcript and reject this protein as a cytosolic contaminant. What the long transcript—likely even longer if not containing the trans-splicing tag—corresponds to is unknown. In the *P. micropora* draft genome annotation, the L12 protein can be retrieved (g77960.t1, 88% identity over the 162 residues of the short isoform; Figure 1), which indeed encodes a 140-residue extension. This extension does not match that of the long isoform. While this validates the interpretation of the current analysis, it raises questions about

protein synthesis and/or processing-related mechanisms in Chromalveolata (see for instance Gabr et al., 2022).

An important conclusion of this study relates to the occurrence of protein NTA in the chromatophore. The authors mention the similarity with observations in Archaeplastida, namely NTA of proteins of both origins, the “ragging” of proteins devoid of crTP starting on very close residues and undergoing partial trimming after initial cleavage, as observed in plants (Bienvenut et al., 2012; Rowland et al., 2015). A large panel of N-terminal residues may undergo NTA, either through a family of dedicated enzymes like those seen in the cytosol of all eukaryotes or through Nats with relaxed specificity, like in the chloroplast (Bienvenut et al., 2020; Giglione and Meinel, 2021). It is therefore important to identify the associated Nat(s), but the authors were unsuccessful. In a recent study of Nats in photosynthetic organisms, a family of eight nuclear-encoded and plastid-located Nats (NatG) with relaxed specificity was shown to perform plastid NTA in all Archaeplastida. NatGs are very similar to *N*-acetyltransferases from cyanobacteria like *Synechococcus* or *Leptolyngbya*, all featuring NTA of four photosystem 2 components (Yang et al., 2014; Giglione and Meinel, 2021). These Nats are specific to cyanobacteria and add to the classical RIMI/JL-specific set of Nats found in

bacteria. In addition, in the fossil red algae *Pyropia yezoensis* and *Rhodochaete parvula*, a NatG (GNAT2) homolog is “still” encoded in the plastid genome, suggesting that the gene progressively moved to the nucleus. Likewise, using cyanobacteria-specific Nats, one can identify NatG homologs in all three chromatophore genomes (Figure 2). These GNATs are also close homologs to the plant NAA90s, one of the two NatG subgroups (Giglione and Meinzel, 2021). In the absence of reverse genetics tools for Paulinellidae, alternative approaches to characterize and confirm *Paulinella* NatG activities would be interesting to implement (Asensio et al., 2022). One conclusion therefore is that NTA in *Paulinella* spp. arises through the same mechanism seen in Archaeplastida or Cyanobacteria. In this context, it would be interesting to explore the role of NTA in the chromatophore in photosynthesis adaptation, like in plants or other processes involving protein stability (Koskela et al., 2018). Finally, whether GNAT2 displays similar NTA and lysine acetyltransferase activity in *Paulinella* as it does in plants is also unknown.

## Supplemental data

The following materials are available in the online version of this article.

**Supplemental Figure S1.** A schematized tree of life showing the high representation of photosynthetic organisms in biodiversity.

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