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*Phylogenetic and functional diversity of aldehyde-alcohol dehydrogenases in microalgae*

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39 **Abstract**

40  
41 Aldehyde-alcohol dehydrogenases (ADHEs) are large metalloenzymes that typically perform the two-step  
42 reduction of acetyl-CoA into ethanol. These enzymes consist of an N-terminal acetylating aldehyde  
43 dehydrogenase domain (ALDH) and a C-terminal alcohol dehydrogenase (ADH) domain. ADHEs are present in  
44 various bacterial phyla as well as in some unicellular eukaryotes. Here we focus on ADHEs in microalgae, a  
45 diverse and polyphyletic group of plastid-bearing unicellular eukaryotes. Genome survey shows the uneven  
46 distribution of the *ADHE* gene among free-living algae, and the presence of two distinct genes in various species.  
47 We show that the non-photosynthetic Chlorophyte alga *Polytomella* sp. SAG 198.80 harbors two genes for  
48 ADHE-like enzymes with divergent C-terminal ADH domains. Immunoblots indicate that both ADHEs  
49 accumulate in *Polytomella* cells growing aerobically on acetate or ethanol. ADHE1 of ~105-kDa is found in  
50 particulate fractions, whereas ADHE2 of ~95-kDa is mostly soluble. The study of the recombinant enzymes  
51 revealed that ADHE1 has both the ALDH and ADH activities, while ADHE2 has only the ALDH activity.  
52 Phylogeny shows that the divergence occurred close to the root of the *Polytomella* genus within a clade formed  
53 by the majority of the Chlorophyte ADHE sequences, next to the cyanobacterial clade. The potential  
54 diversification of function in *Polytomella* spp. unveiled here likely took place after the loss of photosynthesis.  
55 Overall, our study provides a glimpse at the complex evolutionary history of the ADHE in microalgae which  
56 includes *i*) acquisition via different gene donors, *ii*) gene duplication and *iii*) independent evolution of one of the  
57 two enzymatic domains.

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59  
60 **Keywords:** Alcohol dehydrogenase (ADH), bifunctional enzyme, gene duplication, *Polytomella*, spirosome

61  
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67  
68 **Conflicts of interest**

69 The authors declare that they have no conflicts of interest with the contents of this article.

70  
71 **Ethics approval** (include appropriate approvals or waivers) N/A

72  
73 **Consent to participate** (include appropriate statements) N/A

74  
75 **Consent for publication** (include appropriate statements) N/A

76  
77 **Availability of data and material** (data transparency)

78 The ADHE1 cDNA sequence can be found in the DDBJ/EMBL/GenBank databases under the accession number  
79 PRJEB33479

80  
81 **Code availability** (software application or custom code) N/A

82  
83 **Authors' contributions**

84 AA and RvL conceived the study, designed and conducted the experiments. OV, NJT, BL and AA assembled and  
85 annotated genomes. YC and SB performed the proteomic studies. AA, RvL, WN and OV performed the  
86 bioinformatics analyses. AA, RvL and OV wrote the manuscript and generated the figures.

87 All authors participated and approved the final version of the manuscript.

88 **Introduction**

89

90 Microalgae represent a diverse and polyphyletic group of plastid-bearing unicellular eukaryotes. At the base of  
 91 aquatic ecosystems, microalgae also contribute significantly to carbon cycling and storage. Deciphering algal  
 92 genome sequences has shed light on unforeseen metabolic functions and identified targets for genetic  
 93 manipulations, both for laboratory reference algae such as *Thalassiosira pseudonana* (Armbrust et al. 2004) and  
 94 *Chlamydomonas reinhardtii* (Merchant et al. 2007) and for algae with a biotechnological potential such as  
 95 *Chlorella* spp. (Guarnieri et al. 2018; Wu et al. 2019) and *Nannochloropsis* spp. (Radakovits et al. 2012; Wang et  
 96 al. 2014). The availability of genomes of less studied algal species (Kim et al. 2014; Nelson et al. 2019; Blaby-  
 97 Haas and Merchant 2019) have greatly helped our understanding of *i*) the molecular mechanisms underlying  
 98 adaptation to adverse ecological niches and *ii*) the evolutionary history of the different lineages. Among the  
 99 surprises revealed by algal genome sequencing is the extended repertoire of fermentative enzymes in *C.*  
 100 *reinhardtii* (Atteia et al. 2006; Mus et al. 2007). Such a repertoire was long thought to be restricted to prokaryotes  
 101 and anaerobic protists (Rosenthal et al. 1997; Müller et al. 2012). Multidisciplinary studies have lifted the veil on  
 102 the significance of these enzymes and their integration in the global metabolism of the photosynthetic alga (Mus  
 103 et al. 2007; Hemschemeier et al. 2008; Catalanotti et al. 2012; Magneschi et al. 2012; van Lis et al. 2013; van Lis  
 104 et al. 2017; van Lis et al. 2020).

105 The aldehyde-alcohol dehydrogenase (ADHE) is one of the enzymes discovered in the green alga *C.*  
 106 *reinhardtii* that was thought to be lacking in aerobic eukaryotes. In bacteria and amitochondriate eukaryotes,  
 107 ADHE is a crucial fermentative enzyme that interconverts acetyl-CoA to ethanol with acetaldehyde as an  
 108 intermediate (Kessler et al. 1991; Bruchhaus and Tannich 1994; Sánchez 1998). ADHE is a fascinating example  
 109 of a non-cytoskeleton enzyme that can self-assemble into long filaments. The ability of the ADHE to auto-  
 110 assemble *in vivo* into large right-handed helices, also named spiroosomes, was first described in *Lactobacillus*  
 111 (Kawata et al. 1975), and later in *Escherichia coli* (Kessler et al. 1992) and in the anaerobic protist *Entamoeba*  
 112 *histolytica* (Bruchhaus and Tannich 1994). Recent cryo-electron microscopy studies of *E. coli* ADHE revealed  
 113 the mechanism of spiroosome formation and highlighted residues critical for oligomerization (Kim et al. 2019;  
 114 Pony et al. 2020).

115 The ADHE in the green alga *C. reinhardtii* resembles its bacterial and eukaryotic counterparts in several  
 116 aspects: *i*) it has an *N*-terminal acetylating aldehyde dehydrogenase domain and a *C*-terminal alcohol  
 117 dehydrogenase domain; *ii*) it catalyzes the sequential NADH-dependent reductions of acetyl-CoA to acetaldehyde  
 118 and then to ethanol and *iii*) has the ability to form spiroosomes (van Lis et al. 2017). However, the algal enzyme  
 119 differs from previously described ADHEs in its intracellular localization, *i.e.* it is a chloroplast protein, and in its  
 120 regulation, in particular the fact that it is expressed at ambient oxygen levels and the correlation to starch  
 121 accumulation (van Lis et al. 2017). In the unicellular alga *Polytomella* sp. SAG 198.80 (*Polytomella* sp.), a close  
 122 relative of *C. reinhardtii* that lacks photosynthetic pigments and a plastid genome (Pringsheim 1955; Smith and  
 123 Lee 2014), an ADHE-like protein was also identified (Atteia et al. 2003). However, the enzymatic activities of  
 124 the protein, which resides in the mitochondrion (Atteia et al. 2003), have never been investigated. Since the  
 125 discovery of *ADHE* genes in the Chlorophytes *Polytomella* and *C. reinhardtii*, homologous genes have been found  
 126 in other lineages, including in algae harboring secondary plastids such as the Rhizarian *Bigeloviella natans* and  
 127 the Cryptophyte *Guillardia theta* (Atteia et al. 2013). As in *C. reinhardtii* and *Polytomella* sp., the *ADHE* gene in  
 128 *B. natans* and *G. theta* is expressed at ambient oxygen levels (Gould et al. 2019), unlike in bacteria.

129

130 In the course of sequencing the genome of *Polytomella* sp., we found as expected the *ADHE* gene whose  
 131 product has been described earlier as a major mitochondrial protein (Atteia et al. 2003), but also a second gene  
 132 for a putative *ADHE*. In this study, we validate the presence of two distinct *ADHE*-like proteins in the non-  
 133 photosynthetic alga, and we provide a comparison of their biochemical and enzymatic characteristics. A genomic  
 134 and phylogenetic study of *ADHE*s in unicellular algae reveals novel aspects of the origin and evolution of *ADHE*s  
 135 in mitochondriate eukaryotes.

136

137

138 **Results**

139

140 **Structural features of *Polytomella* sp. ADHEs inferred by their primary sequences**

141

142 The genome sequence of *Polytomella* sp. revealed, in addition to the previously described *ADHE* gene (Atteia et al. 2003), another gene for a putative ADHE. For reasons detailed below, we call the newly uncovered gene  
 143 *ADHE1*, while the *ADHE* gene firstly described is renamed *ADHE2*. The overall primary sequence comparison  
 144 indicates that *PsADHE1* is 104-aa larger than *PsADHE2*, but also significantly less related to *PsADHE2* than to  
 145 its counterpart in the photosynthetic alga *C. reinhardtii* (*CrADHE1*), sharing respectively 60% and 72% amino  
 146 acid identity. To bring insights into the differences between the ADHEs in the colorless alga, their amino acid  
 147 sequences were aligned with that of *E. coli* (*EcADHE*), to date the best studied bacterial ADHE, and that of *C.*  
 148 *reinhardtii*, the only algal ADHE characterized so far. The most relevant diverging features are depicted in Fig.  
 149 1A. As compared to *PsADHE2*, *PsADHE1* possesses extended *N*- and *C*-termini. The *C*-terminal extension of  
 150 *PsADHE1* (55-aa) is characterized by a high content in proline and hydroxylated residues (Supplementary Fig.  
 151 S1). Compared to their homologues in *E. coli* (and other bacteria), the algal ADHEs have an *N*-terminal extension,  
 152 indicative of intracellular targeting. As shown earlier, the 27-aa *N*-terminus in *PsADHE2* serves as a targeting  
 153 sequence to the mitochondrion (Atteia et al. 2003). This localization is predicted with high confidence by the  
 154 public software programs MitoFate, TargetP and Predotar (Table 1). In contrast, a plastid localization is predicted  
 155 by the software Predalgo, designed to predict the intracellular targeting of precursor proteins of *C. reinhardtii* and  
 156 other Chlorophytes (Tardif et al. 2012). For *PsADHE1*, the programs agreed to various extents on a plastid  
 157 targeting prediction (Table 1). As no adequate protocol to obtain pure fractions of the nonphotosynthetic plastids  
 158 and mitochondria of the colorless alga is available at present, the plastid localization of *ADHE1* remains  
 159 hypothetical.

160  
 161 The multiple sequence alignment indicates that the ADHEs in *Polytomella* diverge most in the alcohol  
 162 dehydrogenase (ADH) domain (Supplementary Fig. S1). While *PsADHE1* harbors the two Fe-ADH signature  
 163 patterns (Prosite ADH\_iron\_1, PS00913; ADH\_iron\_2, PS00060), *PsADHE2* has a poorly conserved  
 164 ADH\_iron\_2 signature, lacking in particular the two histidine residues required for iron coordination (Montella  
 165 et al. 2005) (Conserved Domains database cd08178). Furthermore, in contrast to *PsADHE1*, *PsADHE2* lacks a  
 166 few residues identified by cryo-EM as critical for *E. coli* ADHE auto-assembly (Kim et al. 2019). In particular,  
 167 the aromatic residue between the two Fe-ADHE signatures (F670 in *EcADHE*, Y735 in *PsADHE1*), suggested to  
 168 be critical for spiroosome formation, is replaced by an arginine in *PsADHE2* (R699) (Supplementary Fig. S1). The  
 169 distinct features revealed by sequence analysis warranted a detailed biochemical and enzymatic comparative study  
 170 of the two *Polytomella* ADHEs.

171

172 **Kinetic analysis of recombinant *PsADHE1* and *PsADHE2***

173

174 ADHE typically catalyzes the reversible conversion of acetyl-CoA into ethanol with an acetaldehyde intermediate  
 175 (Fig. 1C). Four distinct reactions can be assessed *in vitro* by spectrophotometry measuring the oxidation of NADH  
 176 or the reduction of NAD<sup>+</sup>, as shown earlier with the *C. reinhardtii* ADHE (van Lis et al. 2017). To get insights  
 177 into the respective enzymatic properties of the two *Polytomella* ADHEs the proteins were produced  
 178 heterologously in *E. coli*. Coding sequences lacking the putative targeting sequences were cloned into the pET24a  
 179 expression vector (primers sequences are given in Supplementary Table S1). A truncated version of *PsADHE1*  
 180 lacking the last 41 residues (Supplementary Fig. S1) was also produced to investigate the significance of the *C*-  
 181 terminal extension. The recombinant ADHEs were purified via affinity chromatography (Fig. 1B), taking  
 182 advantage of the hexahistidine (6xHis)-tag introduced at their *C*-terminus by the vector. The ability of each of the  
 183 three enzymes to catalyze the different reactions ( $V_{max}$ ) was assessed. The acetaldehyde formed in reaction #1 is  
 184 in direct proximity to the ADH active site and reacts away to ethanol immediately (reaction #2) since both  
 185 reactions use NADH, producing NAD in a 1:1 stoichiometry. Therefore, it can be considered that the  $V_{max}$   
 186 measured in presence of acetyl-CoA and NAD represents the combined activities of reaction #1+2, with half of  
 187 this value corresponding to the actual  $V_{max}$  of reaction #1. Measurement of reaction #1 alone may be approached

188 using EDTA to chelate the Fe of the ADH-moiety and inhibit reaction #2. However, this approach was not  
 189 successful in previous studies on the CrADHE and was not attempted here (van Lis et al. 2017).  
 190 As shown in Table 2, purified recombinant ADHE1 (rADHE1) catalyzed the NADH-dependent ALDH and ADH  
 191 reactions (#1, 2) as well as the reverse reactions (#3, 4). Thus, *Ps*ADHE1 appears as a genuine bifunctional ADHE.  
 192 The study of the truncated ADHE1 (rADHE1\_t) did not reveal significant differences with respect to rADHE1  
 193 for reactions #1, 2 and 3 (Table 2). However, the activity of rADHE1\_t for reaction#4 was found to be 2.5 to 3  
 194 times higher than that of rADHE1, suggesting a regulatory role in the conversion of acetaldehyde to acetyl-CoA  
 195 for the C-terminal extension. Recombinant ADHE2 (rADHE2) catalyzed the acetaldehyde formation from acetyl-  
 196 CoA and NADH (reaction#1) but failed to form ethanol from acetaldehyde and NADH (reaction#2). The rate of  
 197 conversion of acetyl-CoA to acetaldehyde was slightly lower for rADHE2 than rADHE1, with 1.52 and 1.88  
 198 U/mg, respectively. In contrast, the CoA-dependent acetaldehyde oxidation activity (reaction#4) for rADHE2, of  
 199 23 U/mg, was significantly higher than that for rADHE1 of 0.27 U/mg. The high activity measured for the  
 200 conversion of acetaldehyde into acetyl-CoA by rADHE2 might be related to a specific physiological role (see  
 201 below). The fact that rADHE2 has no ADH activity was not so surprising in view of the absence of key residues  
 202 for metal coordination (Fig. 1A).

203

#### 204 **Blue native (BN) PAGE analysis of recombinant ADHE2**

205

206 BN-PAGE is a powerful method to analyze the oligomeric states and associated activities of soluble as well as  
 207 membrane-bound enzymes (Schägger and von Jagow 1991; Schägger et al. 1994). By applying soluble fractions  
 208 of *Polytomella* crude mitochondria on BN-PAGE, the native ADHE2 was previously identified as a dimer of 180  
 209 kDa (Atteia et al. 2003; van Lis et al. 2005). We used this method here to analyze the recombinant ADHE2 of  
 210 *Polytomella* with respect to that of *C. reinhardtii* (*Cr*-rADHE1), which was shown to separate in two predominant  
 211 oligomeric forms, *i.e.* a dimeric and a tetrameric form, with a NAD-dependent ADH activity (van Lis et al. 2017).  
 212 As shown in Fig. 2, the migration of rADHE2 is fuzzy with the prevalence of a monomeric and a dimeric form.  
 213 Unlike with *Cr*-rADHE1, none of these rADHE2 forms has the ability to oxidize ethanol (Fig. 2, “ADH activity”),  
 214 which corroborates the *in vitro* activity measurements (Table 2). In the light of the recently reported cryo-EM  
 215 structures (Kim et al. 2019; Pony et al. 2020), it may be possible that the oligomeric ADHE forms identified on  
 216 BN-PAGE for *C. reinhardtii* correspond to the basic unit (dimer) and the next immediate unit (tetramer) in the  
 217 spiroosome formation. While the dimer formation occurs in a head-to-head arm-crossing fashion independent of  
 218 the ADH domain, the tetramer results from the hydrophobic interactions between the ADH domains from a dimer  
 219 in a tail-to-tail manner (Kim et al. 2019). The seeming lack of a tetramer for rADHE2 might be related to the  
 220 replacement of several residues in the ADH domain that are key for the interactions, in particular the aromatic  
 221 residue (F670, *E. coli* numbering) and the residues which form the hydrophobic pocket (Fig. 1A; Supplemental  
 222 Fig. S1).

223

#### 224 **Accumulation of ADHEs in *Polytomella* sp. cells**

225

226 To gain insights into the growth conditions leading to the accumulation of ADHE1 and ADHE2 in the non-  
 227 photosynthetic alga, we carried out immunoblot experiments using a polyclonal antibody against bacterial and  
 228 algal ADHEs (anti-ALDH/ADH). As shown in Fig 5A, this serum detects with comparable signals the two  
 229 recombinant ADHEs, thereby allowing a relative quantification of the proteins in total cell extracts. We first  
 230 probed cells grown on ethanol at pH 6.0, in which ADHE2 was shown earlier to be abundant (Atteia et al. 2003).  
 231 In these cells, immunoblotting experiments identified two distinct proteins (Fig. 3A). Based on the relative  
 232 electrophoretic migration of the overexpressed proteins, we assigned the larger protein of ~105-kDa to ADHE1,  
 233 and the smaller protein of ~95-kDa to ADHE2. Looking at the relative intensities of their signals, it can be  
 234 concluded that ADHE1 is significantly less abundant than ADHE2. In *Polytomella* sp. cells grown on acetate at  
 235 pH 6.0, the relative abundance between ADHE1 and ADHE2 was found to be comparable to that in ethanol-grown  
 236 cells (Fig. 3B, lane 2), suggesting that the carbon source has no direct effect on the accumulation of the ADHEs.  
 237 In extracts from acetate-grown cells, mass spectrometry-based proteomics identified both ADHEs with a large

238 number of peptides (Supplementary Table S2). The estimation of the ADHE relative abundances using intensity-  
 239 based absolute quantification (iBAQ) (Schwanhäusser et al. 2011) is in agreement with ADHE2 being  
 240 significantly more abundant than ADHE1, with an extracted iBAQ value for ADHE2 more than five times higher  
 241 than for ADHE1 (Supplementary Table S3). As shown in Fig. 3B, *Cr*ADHE1 migrates slightly faster than  
 242 *Ps*ADHE1. This could indicate that the C-terminal extension of *Ps*ADHE1 (Fig. 1A) is present, at least partially,  
 243 in the mature protein. The effect of anoxia on the respective accumulation levels of both ADHE was assessed  
 244 following a protocol described earlier for *C. reinhardtii* (van Lis et al. 2017). Data indicate that anoxia has no  
 245 significant effect on the levels of ADHEs (Supplementary Figure S2). Similar findings were reported for the  
 246 ADHE of *C. reinhardtii* (van Lis et al. 2017).

247 It was reported earlier that *Ps*ADHE2 is a soluble protein (Atteia et al. 2003). Fractionation of  
 248 *Polytomella* cells after sonication and ultracentrifugation confirmed that ADHE2 is predominantly a soluble  
 249 protein (Fig. 3B, lane 4). In contrast, ADHE1 is found in the particulate fraction (Fig. 3B, lane 3), where it could  
 250 be associated with membranes or assembled in spiroosomes.

251

252

### 253 **Overview of the distribution of the ADHE genes among microalgae**

254

255 The growing availability of sequenced genomes and transcriptomes allowed reassessment of the distribution and  
 256 evolution of aldehyde-alcohol dehydrogenase genes among microalgae. The unicellular algae surveyed here  
 257 belong to the supergroup of Archaeplastida, which harbor a primary plastid, as well as to supergroups or clades  
 258 bearing secondary plastids (SAR, Haptista and Cryptista clades). As shown in Supplementary Table 4, less than  
 259 half of the species analyzed (20/49) have a gene for a *bona fide* ADHE, defined here by the presence of two  
 260 consecutive domains with similarities to ALDH and ADH (Fig. 1A). This is clearly not due to incomplete genome  
 261 annotation, but reflects the gene loss in most algal clades. As observed in Supplementary Table S4, the majority  
 262 of the algae harboring an *ADHE* gene fall within the Chlorophyte division (*i.e.* green algae) of Archaeplastida, as  
 263 do *Polytomella* and *C. reinhardtii*. In Chlorophyceae and Trebouxiophyceae, the *ADHE* gene is common although  
 264 some species seem to lack it. Amid the Mamiellophyceae, the tiniest Chlorophytes known to date (Piganeau et al.  
 265 2011), no *ADHE* gene could be found. In the algae with secondary plastids, a scattered distribution of the gene is  
 266 observed (Supplementary Table S4). To date, *B. natans* is the only species with a secondary plastid deriving from  
 267 a green alga for which the genome sequence is publicly available. In this alga, a genuine *ADHE* gene is present,  
 268 as reported earlier (Atteia et al. 2013). As shown in Supplementary Table S4, several species with a plastid  
 269 deriving from a red alga contain an *ADHE* gene. This is unexpected as of today the gene for the bifunctional  
 270 enzyme has not been described in red algae (Rhodophytes) (Supplementary Table S4). Among the eight  
 271 Stramenopiles surveyed here, the marine diatom *Fistulifera solaris* is the only one with an *ADHE* gene (in fact it  
 272 has two highly related genes). In the genomes of the two free-living chromerids (Alveolates) sequenced to date,  
 273 *Chromera velia* and *Vitrella brassicaformis*, genuine *ADHE* genes were found. Finally, the survey indicates the  
 274 absence of an *ADHE* gene in the Haptista *Emiliana huxleyi* but its presence in the Cryptista *G. theta*. If we  
 275 consider the environment in which the microalgae thrive, it seems that the *ADHE* gene is more common in  
 276 freshwater species than in marine species (Supplementary Table S4).

277 Based on the multiple alignment of 105 eukaryotic and bacterial protein sequences (Supplementary Table  
 278 S5) and the reconstruction of an unrooted Maximum Likelihood phylogenetic tree (Fig. 4), we found that ADHEs  
 279 from the most evolved Chlorophyte clades (Trebouxiophyceae, Chlorophyceae) group together, close to the clade  
 280 of Cyanobacteria. This proximity suggests a vertical inheritance of the *ADHE* gene in this algal division, including  
 281 for *Polytomella* sp. SAG 198.80. The remaining green algal ADHEs sequences, which belong to the other clades  
 282 (Chlorodendrophyceae, Palmophyllophyceae, Prasinophyceae) are found within a composite eukaryotic group  
 283 made of sequences from amitochondriate eukaryotes and from the algae *B. natans* and *G. theta*, described earlier  
 284 (van Lis et al. 2017). Within this same group are also found the sequences of the Alveolates *C. velia* and *V.*  
 285 *brassicaformis*. The position of the two *ADHE* sequences of *F. solaris* in a clade formed by the proteobacterial  
 286 sequences might indicate gene acquisition by lateral transfer or a contaminant bacterial DNA.

287 Our gene survey revealed that half of the algal species (10/19) harbor two distinct ADHE genes  
 288 (Supplemental Table S4). In subsections of the tree containing algal ADHEs, the paralogs of a given species  
 289 usually group together, suggesting recent gene duplication or possibly allelic divergence as for *Picochlorum*  
 290 NBRC 102739 or *Pyramimonas amyliifera*. In *C. reinhardtii*, two pseudogenes were found next to one another on  
 291 chromosome 16, highly similar to *CrADHE1*, testifying of the tendency of ADHE to duplicate. Chlorellales were  
 292 different in that the divergence of the two paralogs occurred early, *i.e.* before the divergence of *Micractinium* and  
 293 *Chlorella*. An ancestral duplication appears very likely for the *Polytomella* genus. Indeed, the tree shown in Fig.  
 294 4, the ADHEs of the colorless algae are found close to each other surrounded by most green algal ADHEs. This  
 295 suggests that ADHE1 and ADHE2 proteins have evolved by duplication from a cyanobacterial-type ancestral  
 296 gene. The long branch length of *PsADHE2* (Fig. 4, sequence 1) strongly suggests an unusually rapid sequence  
 297 divergence in *Polytomella* sp. (which translated into unusual biochemical properties as discussed above) in clear  
 298 contrast with the situation in the photosynthetic algal species.

299 Illumina sequencing of genomic DNA is available for four known species of the *Polytomella* genus, *P.*  
 300 *capuana*, *P. magna*, *P. parva*, and *P. piriformis* (Smith and Lee 2014), which are distinguished by the structure  
 301 of their mitochondrial DNA (Smith et al. 2013). The genomic data of these species were searched for the presence  
 302 of *ADHE* genes. In all four species, two *ADHE* genes were found. While one gene codes for a genuine ADHE  
 303 (named ADHE1), the other codes for an atypical ADHE showing divergence in the C-terminal part of the protein  
 304 (named ADHE2). The predicted ADHE2 sequences were analyzed for the deriving features revealed in this study,  
 305 *i.e.* the lack of the Fe-coordinating residues and of the aromatic residue required for auto-assembly. As depicted  
 306 in Fig. 5, the ADHE2 sequences in *P. piriformis* and *P. parva* are comparable to *PsADHE2* (*Polsp\_ADHE2*). In  
 307 contrast, the sequences of the other two species show only one of the two distinctive features. In *P. magna*, the  
 308 ADHE2 lacks the aromatic residue (a valine instead) necessary for spiroosome formation but has residues required  
 309 for iron-binding. In *P. capuana*, the opposite is observed.

310 The comparison of the C-terminal ADH domain of ADHE1 paralogs in algal species indicates a high  
 311 degree of conservation, including between *CrADHE1* and one of the pseudogenes (- $\Psi$ 1, Fig. 5). It is however  
 312 interesting to note that all but one *Polytomella* species (*P. magna*) have a longer C-terminus.

313

314

## 315 Discussion

316

### 317 The non-photosynthetic alga *Polytomella* sp. harbors two enzymes related to bacterial ADHEs

318

319 Aldehyde-alcohol dehydrogenases are bifunctional enzymes long thought to be restricted to Bacteria and  
 320 eukaryotes devoid of mitochondria, where they play a crucial role in fermentative metabolism (Kessler et al. 1991;  
 321 Bruchhaus and Tannich 1994; Boxma et al. 2004). The occurrence of ADHE-like proteins in microalgae, which  
 322 are mitochondriate eukaryotes, was interesting in several respects, especially for their integration in an aerobic  
 323 metabolic context and their evolutionary origin. We have shown recently that the photosynthetic alga *C.*  
 324 *reinhardtii* harbors an ADHE that resembles its bacterial counterparts in *i)* the enzymatic activities catalyzed and  
 325 *ii)* the ability to oligomerize into spiroosomes. The phylogeny pointed to the vertical inheritance of the algal enzyme  
 326 from a cyanobacterial ancestor (van Lis et al. 2017). With the study of the ADHE-related enzymes in *Polytomella*  
 327 sp. SAG 198.80 and other *Polytomella* species, we disclosed fascinating facets of the evolution of the so-called  
 328 bifunctional enzyme.

329 Significant molecular and biochemical data on the oxidative phosphorylation complexes of *Polytomella*  
 330 sp. have been obtained over the years that provided further evidence for the phylogenetic relationship of the  
 331 colorless alga with the green alga *C. reinhardtii* (Atteia et al. 1997; Funes et al. 2002; van Lis et al. 2005; van Lis  
 332 et al. 2007). The metabolic routes existing in *Polytomella* have in contrast been poorly investigated and  
 333 consequently, the knowledge on their mechanisms underlying highly versatile heterotrophic capacities is scarce  
 334 (Wise 1955; Wise 1959; Atteia et al. 2000). Through this study we provide insights into the aldehyde/alcohol  
 335 metabolism unexplored so far in *Polytomella* species. Based on its sequence similarity to *C. reinhardtii* ADHE1  
 336 (Fig. 1A, Supplementary Fig. S1) and enzymatic activities (Table 2), it seems plausible that *Polytomella* sp.



ADHE1 functions in the same way, that is in the ethanolic fermentation fueled by starch glycolysis (Magneschi et al. 2012; van Lis et al. 2017). Although *Polytomella* sp. ADHE2 shares a number of features with *CrADHE1* and bacterial ADHEs, our experimental data clearly show that it is not a typical aldehyde-alcohol dehydrogenase. First of all, *PsADHE2* is impaired in ethanol dehydrogenase activity (Table 2; Fig. 2). Furthermore, *PsADHE2* is a soluble protein occurring as a dimer (Atteia et al. 2003; van Lis et al. 2005) which likely follows from the lack of the key residues required for spiroosome formation (Fig. 1A; Supplemental Fig. S1). The absence of an ADHE2 in *C. reinhardtii* illustrates the fact that making predictions on the metabolic enzymes in *Polytomella* sp. from *C. reinhardtii* studies may have its limits, and emphasizes the importance de develop studies dedicated to the colorless alga. The ADHE gene duplication and divergence (of function and possibly of cellular localization) in the non-photosynthetic alga might be seen as beneficial for the adaptation to a new lifestyle (Zimorski and Martin 2014; Copley 2020).

To our knowledge, this work is the first report of a full-length ADHE that performs only one of the two typical enzymatic reactions. The presence of such an enzyme in the colorless alga is puzzling and raises questions about its metabolic integration. Indeed, in nature there exist acetylating aldehyde dehydrogenases (AcAldDH) (EC 1.2.1.10) which catalyze the same overall reaction as *PsADHE2*, that is the interconversion of acetaldehyde and acetyl-CoA using NADH as a co-factor (Burton and Stadtman 1953). These mono-enzymatic proteins of ~45 kDa can catalyze the formation of acyl-CoAs, such as the conversion of acetaldehyde to acetyl-CoA in the utilization of ethanolamine by *Salmonella typhimurium* (Roof and Roth 1992), or the formation of aldehydes in alcohol-producing bacteria such as *Clostridium beijerinckii* (Toth et al. 1999). The reason for keeping a full-length ADHE with a non-functional alcohol dehydrogenase domain is at present unclear, but it is tempting to propose that ADHE2 bears an as yet unknown additional function. It is worth mentioning here that the ADHE in *E. coli* was found to act as an ATP-dependent RNA helicase through the associate with the small subunit head domain of the ribosome (Shasmal et al. 2016; Kurylo et al. 2018).

In *Polytomella*, we found that the relative abundance of the ADHEs is comparable in cells growing on ethanol and on acetate (pH 6.0) (Fig. 3), which is not in favor of the involvement of ADHE2 (and ADHE1) in ethanol assimilation. Earlier results on ethanol-growing *Polytomella* sp. indicated that the pH of the medium had an impact on the expression and accumulation levels of ADHE2, with high levels at pH 6.0 but much lower at pH 3.7 (Atteia et al. 2003). The external pH as a regulatory factor may provide hints as to the connection between ADHE2 and mitochondrial respiration, since it was found that *i)* ADHE2 steady-state levels were inversely regulated to those of respiratory complexes I, III, IV and to the levels of cytochrome *c*, and *ii)* the respiratory rates were 20-25% lower in isolated mitochondria from cells grown at pH 6.0 ('high' ADHE2) than from cells grown at pH 3.7 ('low' ADHE2) (Atteia et al. 2000). It may be proposed that a downregulation of OXPHOS activity by the external pH alters TCA cycle activity necessitating a metabolic rerouting involving reductant and CoA pools, in which ADHE2 has a role to play.

Assuming that native ADHE2 only catalyzes the interconversion of acetyl-CoA and acetaldehyde as is the case for the recombinant enzyme (Table 2), the question arises as to the directionality of the reaction catalyzed *in vivo* by this mitochondrial protein. At pH 6.0 on acetate or on ethanol, *PsADHE2* is very abundant. When functioning in the forward direction (Fig. 1C, reaction #1), its activity would lead to the production of acetaldehyde from acetyl-CoA. Acetaldehyde is highly toxic for the mitochondrial function, and must be metabolized quickly to prevent damage of proteins and DNA (Cederbaum et al. 1973; Cederbaum et al. 1974). In the case of *E. coli* ADHE, it is proposed that the acetaldehyde remains sequestered inside the spiroosome structure and is converted into ethanol at the inner surface without being released into the cell's cytoplasm (Kim et al. 2019). In mitochondria, acetaldehyde detoxification typically occurs via CoA-independent aldehyde dehydrogenase, and results in the production of NADH and acetate. In the *Polytomella* sp. genome, seven distinct mon-enzyme ALDH genes were identified, and two of them code for enzymes predicted to reside in the mitochondrion. Setting up tools to follow the accumulation and activity of ALDH enzymes in isolated organelles is required to clarify their role in relation to ADHE2. In the proteomic-based analysis of the acetate-grown cells, no ALDH was identified at levels comparable to those of ADHE2 (Supplementary Table S3). When functioning in the reverse direction (Fig. 1C, reaction #4), ADHE2 would use acetaldehyde to produce acetyl-CoA, a central metabolite (Pietrocola et al. 2015). It has been shown earlier that *Polytomella caeca* can use acetaldehyde as

387 growth substrate (Wise 1968; Wise 1970), but the assimilation pathway was not described. In bacteria,  
 388 acetaldehyde is involved in metabolic pathways involving various carbohydrate, lipid, and aromatic compounds  
 389 (Lees and Jago 1978). In *Polytomella* sp. ADHE2 may be used to assimilate other C-sources, but a source of  
 390 acetaldehyde in the mitochondrial matrix must be envisioned. In *Polytomella* cells grown on acetate or on ethanol,  
 391 acetaldehyde could be produced in the matrix from ethanol, as in the yeast *Saccharomyces cerevisiae*, by one of  
 392 at least nine mono-enzymatic ADH predicted from the *Polytomella* sp. genome. Ethanol is indeed a good carbon  
 393 source for growth of *Polytomella* (Atteia et al. 2000). However, we showed here that ADHE2 is also highly  
 394 expressed in cells growing on acetate (Fig. 3). Acetate assimilation occurs as in any acetoflagellate via acetyl-  
 395 CoA synthase likely in the cytosol. Without a better knowledge of the metabolic enzymes and routes existing in  
 396 the colorless alga, it is difficult to propose a pathway from cytosolic acetyl-CoA to mitochondrial acetaldehyde.  
 397 In any case, the abundance of ADHE2 prompts further examination of acetaldehyde as important metabolite in  
 398 *Polytomella*. Elucidation of the metabolic integration of the ADHE-like enzyme in the colorless alga will have to  
 399 rely on the use of large-scale analyses including proteomics and metabolic fluxes but also on biochemical  
 400 approaches.

401

#### 402 **Diversity of ADHEs in unicellular plastid-bearing eukaryotes**

403

404 The genome survey carried out in this work shows that ADHE is a common enzyme among microalgae. Our  
 405 phylogenetic tree does not allow a full reconstitution of the deep phylogeny of ADHEs, but does reveal a complex  
 406 history within Chlorophytes (Archaeplastida supergroup). As illustrated here, there seems to be a tendency for  
 407 gene duplication and maintenance of *bona fide* ADHE duplicates among Chlorophytes, especially in the  
 408 Chlorophyceae and Trebouxiophyceae (Fig. 4, Supplementary Table S4,). Sometimes, diverging evolution of one  
 409 of the ADHE duplicates can occur as exemplified here with the nonphotosynthetic *Polytomella* species.  
 410 Interestingly in these species, most mutations seem to have occurred in the C-terminal with two distinct and non-  
 411 exclusive outcomes: *i*) the loss of ADH activity (mutations of the iron-coordinating residues) and *ii*) the loss of  
 412 spiroosome formation ability (mutation of the aromatic residue). While the ADHE2 in *P. magna*, the most ancestral  
 413 of the *Polytomella* species (Smith et al. 2013), retains the iron-binding and active site residues of the ancestor, it  
 414 has lost the key aromatic residue for spiroosome formation. We therefore predict that ADHE2 in *P. magna*, while  
 415 retaining its capacity for ethanol production/oxidation, will not be found in the particulate fraction but rather as a  
 416 soluble protein. This raises the question of the potential toxicity of the acetaldehyde produced in the absence of  
 417 spiroosome sequestration. Conversely, *P. capuana* ADHE2, which conserves the aromatic residue but has lost the  
 418 residues necessary for ADH activity, is likely to be able to self-assemble. Without the residues for iron-  
 419 coordination and the aromatic residue for the spiroosome formation, the ADHE2 in *Polytomella* sp. SAG 198.80  
 420 and in the species *P. piriformis* and *P. parva* represent the most divergent duplicates.

421 Since *Polytomella* sp. is the only alga known to date to harbor a full-length ADHE with a catalytically-  
 422 defective ADH domain, it is tempting to correlate this evolution to the loss of photosynthetic activity in some  
 423 way. However, it is clearly not a prerequisite for loss of photosynthesis, as it has not occurred in *P. magna*. The  
 424 different *Polytomella* species represent different stages of functional adaptation of one of the duplicates, but the  
 425 link between acquisition of mitochondrial targeting sequence and the loss of ADH activity or ability to auto-  
 426 assemble, and the generalized metabolic consequences of the ADHE2 evolution, remains to be studied.

427

#### 428 **Concluding remarks**

429

430 With the study of ADHEs in the algae *C. reinhardtii* and *Polytomella* we just begin to understand the significance  
 431 of this fascinating multifunctional enzyme amid eukaryotes that thrive in oxic conditions, and to decipher the  
 432 evolutionary steps since gene acquisition in the Archaeplastida supergroup. The original properties of *Polytomella*  
 433 ADHE2 unveiled by this work are likely to serve as an inspiration for enzymatic engineering, since ADHEs are  
 434 promising protein targets in biotechnological applications in ethanol/butanol production, especially in *Clostridium*  
 435 spp.

436

437  
438439 **Materials and Methods**440  
441 **Algal strain and growth conditions**  
442

443 *Polytomella* strain SAG 198.80 (*Polytomella* sp.) was obtained from the SAG algal culture collection of the  
 444 University of Göttingen (<http://sagdb.uni-goettingen.de/>) and axenized using ampicilline. *Polytomella* sp. was  
 445 cultivated either on acetate (MAP medium) or ethanol (MEP medium) in low light at room temperature (24°C)  
 446 without shaking. Both MAP and MEP culture media contain 35 mM MES (Sigma Aldrich), 0.8 mM K<sub>2</sub>HPO<sub>4</sub>, 0.2  
 447 mM KH<sub>2</sub>PO<sub>4</sub>, 7.5 mM NH<sub>4</sub>Cl, 0.3 mM CaCl<sub>2</sub>, 0.5 mM MgSO<sub>4</sub>. Oligoelements are present at 1.39 µM ZnSO<sub>4</sub>, 0.8  
 448 µM H<sub>3</sub>BO<sub>3</sub>, 2.65 µM MnSO<sub>4</sub>, 0.74 µM FeCl<sub>3</sub>, 0.83 µM NaMoO<sub>4</sub>, 0.16 µM CuSO<sub>4</sub>, 0.6 µM KI. The pH was  
 449 adjusted to 5.8 with KOH. Thiamine and cyanocobalamin are added to a final concentration of 20 µg/L and 1  
 450 µg/L, respectively. In both media, carbon source is added to a final concentration of 30 mM. One-liter Erlenmeyer  
 451 flasks containing 0.2 L of culture medium were inoculated with 4 mL of preculture in exponential growth phase.  
 452 Cells were harvested in their late log phase, after ~3 days (Atteia et al. 2000).

453

454 **Cell fractionation and preparation of cell extracts for mass spectrometry**

455

456 Acetate-grown cells were harvested (3,000 x g for 10 min) at 4°C and resuspended in HEPES 50 mM (pH 7.5)  
 457 supplemented with the protein inhibitors 1mM amino caproic acid, 0.5 mM benzamidine). Cells were broken by  
 458 sonication and briefly further centrifuged at 1,500 rpm to remove insoluble material (starch) and unbroken cells.  
 459 The resulting supernatant was either analyzed directly by mass-spectrometry to identify and evaluate the relative  
 460 abundance of the algal ADHEs. Alternatively, the supernatant was ultracentrifuged at 4°C (130,000 x g for one  
 461 hour) to separate the soluble and particulate fractions. The resulting pellet was washed once in HEPES medium  
 462 (with protease inhibitors).

463

464 **PAGE analysis, immunodetection of ADHEs, and in-gel activity**

465

466 Proteins in whole cells and cell extracts were precipitated by chloroform/methanol (Wessel and Flügge 1984), and  
 467 then resuspended in 2% SDS (w/v). Protein concentrations were determined using the bicinchoninic acid Pierce™  
 468 Protein assay kit (ThermoFisher) using bovine serum albumin as standard. For SDS-PAGE analysis, protein  
 469 samples (40 µg) were resuspended in 2% SDS (w/v) and 2% beta-mercaptoethanol, and boiled for 90 sec.  
 470 Insoluble material was removed via a 2 min-centrifugation at 13,000 x g. Apparent molecular masses were  
 471 estimated using Prestained protein ladder Plus (Euromedex). After electrophoresis, proteins were stained with  
 472 Coomassie Brilliant Blue R250 or transferred onto nitrocellulose membranes (GE Healthcare,) and stained with  
 473 Ponceau S. Immunoblotting was carried out as described (van Lis et al. 2017). The anti-ADH/ALDH (Agrisera)  
 474 was used at 1/1,000.

475

476 Blue Native-PAGE was carried out using a 3-12% linear polyacrylamide gradient as described (van Lis et al.  
 477 2017). The molecular mass of the rADHE oligomers were estimated using NativeMark™Unstained Protein  
 478 Standard (Thermofisher Scientific). Recombinant CrADHE, used as control protein, was prepared as described  
 479 previously (van Lis et al. 2017). For ADH activity, BN-gel was washed twice in 100 mM Tris (pH 8.8) and then  
 480 incubated in 25 ml of 100 mM Tris-Cl (pH 8.8) containing 20 mg of NAD, 5 mg of nitroblue tetrazolium, 1 mg  
 481 of phenazine methosulfate, and 25 µl of absolute ethanol.

482

483 **Mass spectrometry-based proteomics**

484

485 Protein extract from acetate-grown cells was stacked in the top of a 10% SDS-PAGE gel and stained with  
 486 Coomassie Brilliant blue. The band was excised and submitted to in-gel digestion using modified trypsin

487 (sequencing grade) (Promega) as previously described (Salveti et al. 2016). Resulting peptides were analyzed by  
 488 online nanoliquid chromatography coupled to tandem MS (UltiMate 3000 and LTQ-Orbitrap Velos Pro, Thermo  
 489 Scientific). Peptides were sampled on a 300  $\mu\text{m}$  x 5 mm PepMap C18 precolumn and separated on a 75  $\mu\text{m}$  x 250  
 490 mm PepMap C18 column (Thermo Scientific) using a 25-min gradient for individual gel bands and a 80-min  
 491 gradient for mitochondrial soluble fraction. MS and MS/MS data were acquired using Xcalibur (Thermo  
 492 Scientific). Peptides and proteins were identified using Mascot (version 2.6) through concomitant searches against  
 493 homemade *Polytomella* protein sequence database, classical contaminants sequence database and the  
 494 corresponding reversed databases. The Proline software (<http://proline.profi-proteomics.fr>) was used to filter the  
 495 results: conservation of rank 1 peptides, peptide length  $\geq 7$  amino acids, false discovery rate of peptide-spectrum  
 496 matches  $< 1\%$  as calculated on scores by employing the reverse database strategy, minimum peptide-spectrum  
 497 match score of 25, and minimum of 1 specific peptide per identified protein group. Proline was then used to  
 498 perform a MS1-based label-free quantification of the protein groups identified in the sample.

499

### 500 ***Polytomella* sp. ADHE gene cloning, expression and purification of recombinant enzymes**

501 Via rapid cDNA amplification-based techniques, both full-length coding sequences were cloned. In brief, total  
 502 RNA was isolated from acetate-grown cells using the RNeasy Plant Mini Kit (Qiagen, Hilden, Germany).  
 503 Complementary DNA (cDNA) amplifications were done using the FirstChoice®RLM-RACE Kit (Thermo Fisher  
 504 Scientific) and specific primers designed based on the *Polytomella* draft genome sequence. Using the 3' RACE  
 505 DNAs as templates, the full-length sequences coding *Polytomella* ADHEs were amplified by PCR using *Pfu*  
 506 polymerase (Promega). The primers used to amplify ADHE1 cDNA (2973 bp) and ADHE2 cDNA (2717 bp) are  
 507 listed in Supplementary Table S1. Sequencing of *PsADHE2* cDNAs revealed a few differences with the nucleotide  
 508 sequence published earlier (Atteia et al. 2003), which are believed to result from an inappropriate assembly of  
 509 partial cDNA sequences.

510

511 The full-length cDNAs were used as templates to amplify the coding sequence for predicted mature proteins using  
 512 the primers listed in Supplementary Table S1. Amplified fragments were cloned into the pET-24a expression  
 513 vector (Merck Novagen) using the NdeI and XhoI restriction site within the multiple cloning site to incorporate a  
 514 6xHis tag at the C-terminus of the proteins. Constructs with confirmed nucleotide sequences (at GATC Biotech)  
 515 were used to transform *Escherichia coli* BL21 (DE3). Transformed cells from plates were inoculated in the auto-  
 516 induction medium ZYM5052 (Studier 2005) supplemented with kanamycin (30  $\mu\text{g}/\text{ml}$ ), and grown overnight  
 517 either at 25°C for rADHE1, or at 37°C for rADHE2. The cells were harvested by centrifugation, washed once in  
 518 Buffer 2 (300 mM NaCl, 50 mM sodium phosphate, pH 8.0) and flash-frozen and stored at -20°C until use. Protein  
 519 purification was carried out at 4°C as follows. Cell pellets were resuspended in Buffer 1 supplemented with 0.5  
 520 mg/ml lysozyme, 0.5 mg/ml DNase and 2 mM  $\text{MgCl}_2$ , and protease inhibitors (PMSF, amino caproic acid,  
 521 benzamidine) and subjected to three cycles of freezing (liquid nitrogen) and thawing. The lysates were clarified  
 522 via centrifugation (10,000 x  $g$  for 15 min, at 4°C) and the supernatant was gently mixed with HIS-Select Nickel  
 523 Affinity gel (Sigma-Aldrich) equilibrated with Buffer 2. The mixture was poured into an empty column, and the  
 524 gel was extensively washed with 20 mM imidazole in Buffer 2. His-tagged ADHEs were eluted with 100 mM  
 525 imidazole in Buffer 2 rapidly desalted on PD10 Desalting columns (GE Healthcare) pre-equilibrated with 50 mM  
 526 Tricine, pH 8.0, and finally concentrated on a Vivaspin 50 (Sartorius) to reach a final concentration of 5-7 mg  
 527 protein/ml.

528

### 529 **Enzyme assays**

530

531 Freshly purified recombinant ADHEs were assessed for the different enzymatic activities at 25°C. Standard assay  
 532 medium for NADH-dependent reactions consisted of 50 mM potassium phosphate (pH 7.0) and 0.42 mM NADH.  
 533 Acetyl-CoA reductase activity was determined in the presence of 0.21 mM acetyl-CoA, acetaldehyde reductase  
 534 activity with 1 mM acetaldehyde and 0.21 mM CoASH. The rates of both reductase reactions were monitored  
 535 spectrophotometrically following the disappearance of NADH at 340 nm. Standard assay medium for  $\text{NAD}^+$ -  
 536 dependent reactions contained 1.5 mM  $\text{NAD}^+$  in 50 mM Glycine/NaOH buffer pH 9.0. Ethanol dehydrogenase

537 activity was assayed in 170 mM ethanol (1% v/v); acetaldehyde dehydrogenase activity was assayed in 1 mM  
538 acetaldehyde and 0.2 mM CoASH. The rates of both dehydrogenases were monitored by the formation of NADH  
539 at 340 nm. All reactions were initiated by the addition of purified recombinant enzyme (25-40 µg). In all the  
540 assays, one unit of enzyme activity (U) was defined as 1 µmol of NADH (or NAD) converted per min.

541

#### 542 **Bioinformatic analysis**

543

544 The genome of *Polytomella* sp. SAG 198.80 was assembled from paired-end Illumina and PacBio sequences,  
545 using a combination of the programs ABYSS (1.9.0), CERULEAN (0.1), PBJELLY (15.2.20) and BESST (2.1).  
546 Details will be published elsewhere. *Ab initio* annotation was performed with AUGUSTUS, using available  
547 transcriptome assemblies to generate the species models. Protein sequences were aligned using MAFFT (7.130)  
548 or Clustal Omega 1.2.4. (<https://www.ebi.ac.uk/Tools/msa/clustalo/>). Enzyme domain searches were done at  
549 <https://www.ncbi.nlm.nih.gov/Structure/cdd/cdd.shtml>. The iron-containing alcohol dehydrogenases signatures  
550 (ADH\_iron\_1, PS00913; ADH\_iron\_2, PS00060) were obtained at Prosite (<https://prosite.expasy.org/>) (Bairoch  
551 1992). Intracellular targeting predictions were made using TargetP (Plant network v2.0) (Emanuelsson et al.  
552 2000), Mitofates (plant) v1 (Fukasawa et al. 2015), Predotar (Small et al. 2004) and Predalgo (Tardif et al. 2012).

553

#### 554 **Phylogenetic dataset construction and analysis**

555

556 ADHE proteins sequences were searched among microalgae via BLASTP using the *C. reinhardtii* ADHE  
557 sequence (UniprotKB entry Q1RS84\_CHLRE) as a query against the proteomes available at the US National  
558 Center for Biotechnology and Information (NCBI) (<https://blast.ncbi.nlm.nih.gov/Blast.cgi>), at Algoblast  
559 ([https://giavap-genomes.ibpc.fr/cgi-bin/AlgoBLAST/algoBlast\\_mainpage.php](https://giavap-genomes.ibpc.fr/cgi-bin/AlgoBLAST/algoBlast_mainpage.php)),  
560 at CryptoDB (<https://cryptodb.org/cryptodb/>), at the Joint Genome Institute (JGI) genome Portal  
561 (<https://genome.jgi.doe.gov/portal/>), and at Phytozome from the JGI (<https://phytozome.jgi.doe.gov/>). All the  
562 retrieved sequences were checked manually for the length and the presence of the key residues and domains of  
563 ADHE. A total of 49 algal genome sequences were surveyed. Phylogenetic analyses were conducted using  
564 MEGAX and MEGA7 software (<http://www.megasoftware.net>), and phylogenetic relationships were inferred  
565 using the maximum likelihood (ML) method applied with the JTT amino-acid substitution matrix-based model  
566 (Jones et al. 1992).

567

#### 568 **Nucleotide sequence accession numbers**

569 The ADHE1 cDNA sequence can be found in the DDBJ/EMBL/GenBank databases under the accession number  
570 PRJEB33479. The sequence of ADHE2 cDNA (UniProtKB entry- Q70YJ9) has been updated.

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**Table 1. Targeting predictions for *Polytomella* and *C. reinhardtii* ADHEs.**

Predictions were made using four on-line localization predictors.

	MitoFate		TargetP			Predotar v1.04			PredAlgo		
	M probability	Cleavage site	M score	C score	Loc - RC	M score	C score	prediction	M score	C score	Target
PsADHE1	0.047	nd	0.047	0.882	C - 2	0.07	0.34	possibly P	0.00861	0.890	C
PsADHE2	0.991	25-26	0.784	0.134	M - 2	0.7	0.06	M	0.0911	1.824	C
CrADHE1	0.017	nd	0.184	0.597	C - 3	0.49	0.49	possiby M	0.0299	3.946	C

**Table 2. Enzymatic assays monitoring different activities of *Polytomella* sp. recombinant ADHEs.**

The different reactions assessed are depicted in Figure 2. Enzymatic activities were determined spectrophotometrically following the production or consumption of NADH. Results are presented as the mean +/- standard deviation of triplicate samples measured from three independent protein preparations. Note that the assays for reaction 1 were performed in absence of an iron chelator. Since acetaldehyde formed in reaction #1 reacts away to ethanol immediately, and reactions #1 and #2 both produce NAD in a 1:1 stoichiometry, the real Vmax for reaction 1 can be extrapolated, corresponding to 50% to the determined Vmax.

Reaction	Substrates	Products	Vmax ( $\mu\text{mol}\cdot\text{min}^{-1}\cdot\text{mg}^{-1}$ )		
			rADHE1	rADHE1_t	rADHE2
#1	acetyl-CoA	acetaldehyde, CoASH (ethanol)	1.88 $\pm$ 0.0085	2.221 $\pm$ 0.096	1.52 $\pm$ 0.087
#2	acetaldehyde	ethanol	2.095 $\pm$ 0.171	2.246 $\pm$ 0.080	0.0034 $\pm$ 0.002
#3	ethanol	acetaldehyde	2.887 $\pm$ 0.257	2.552 $\pm$ 0.075	0.0087 $\pm$ 0.001
#4	acetaldehyde, CoASH	acetyl-CoA	0.271 $\pm$ 0.0103	0.619 $\pm$ 0.073	23.45 $\pm$ 1.104