

Takakia possesses a key marker of embryophyte sporopollenin

Dae-Yeon Suh^{1§}, Damanpreet K Sraan¹, Neil W Ashton²

 1 Chemistry and Biochemistry, University of Regina, Regina, Saskatchewan, Canada

Abstract

The enigmatic moss, *Takakia lepidozioides*, possesses a particular type III polyketide synthase, ASCL (Anther-Specific Chalcone synthase-Like), that is an identifying marker for genuine sporopollenin in the walls of embryophyte spores and pollen grains. By contrast, a survey of all algae with sequenced genomes confirms that they do not possess ASCL and, therefore, their spore walls are not composed of sporopollenin.

²Biology, University of Regina, Regina, Saskatchewan, Canada

[§]To whom correspondence should be addressed: dae-yeon.suh@uregina.ca

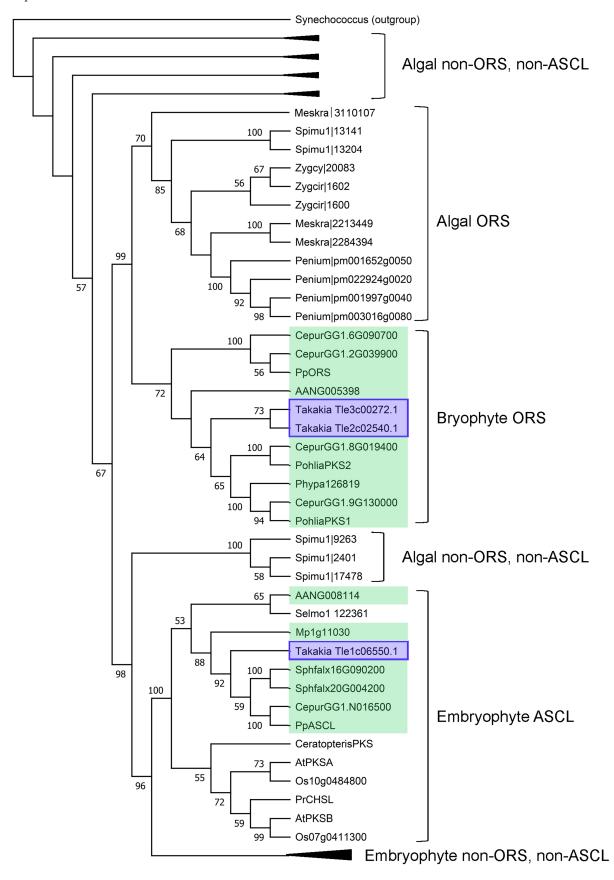


Figure 1. Maximum Likelihood tree of plant type III polyketide synthases:



Included in the tree are sequences of all known algal type III polyketide synthases, all known bryophyte enzymes that belong to either ORS (2'-OxoalkylResorcinol Synthase) or ASCL clades, representative ASCL enzymes from other major embryophyte groups, and some non-ORS and non-ASCL type III PKS enzymes from diverse embryophyte taxa. *Takakia* enzymes and other bryophyte enzymes are highlighted in blue and green, respectively. Bootstrap values (>50%) are displayed at the nodes. A cyanobacterial type III PKS was used as outgroup to root the tree. For brevity, four of the five algal non-ORS and non-ASCL clades and the embryophyte non-ORS and non-ASCL clade, which includes >20 diverse enzymes, e.g. chalcone synthase, stilbene synthase (Shimizu et al. 2017), have been collapsed. Sequences used for tree construction are provided in the Extended Data section (Fig. S1).

Description

Sporopollenin (SP) is the chemically resistant wall material of plant spores and pollen, which provides protection from subaerial stresses. For most of the time since the term was coined by Zetzsche in 1931 (Zetzsche and Kälin, 1931; Zetzsche and Vicari, 1931) resistance to acetolysis has been the sole criterion for the identification of SP. This chemically imprecise definition has resulted in numerous claims for the existence of SP or SP-like material in algae and various microorganisms (Suh and Ashton 2022 and references therein). Based on more recent chemical analyses (reviewed by Grienenberger and Quilichini 2021), the current view of embryophyte SP is that it is a polymer composed of polyhydroxylated polyketides, hydroxylated aromatics and fatty acid derivatives, crosslinked via ester and ether bonds and oxidative C–C coupling. A particular type III polyketide synthase (PKS), ASCL, plays a central role in the polyketide biosynthetic pathway that provides hydroxylated polyketides as SP precursors (Kim et al. 2010; Colpitts et al. 2011; Suh and Ashton 2022). Following an extensive BLASTp survey of plant genomes, we discovered that with an exception, the marine monocot, *Zostera marina*, which has exineless pollen (Olsen et al. 2016), representatives of all major embryophyte clades possess ASCL in stark contrast to algae, including Charophytes, which do not. This led us to propose a new definition for SP as follows to distinguish genuine SP in embryophytes from SP-like compounds such as the acetolysis-resistant algaenans possessed by some Chlorophytes and to use ASCL as an identifying marker for the presence of genuine SP in spore and pollen grain walls (Suh and Ashton 2022).

'Sporopollenin is a chemically resistant complex heteropolymer present in the outer walls of spores and pollen grains and is composed partly of hydroxylated polyketides derived from the conserved polyketide pathway, which involves ASCL.'

At the time we made this definition, the genome of the phylogenetically enigmatic plant, *Takakia*, of which there are only two species, *Takakia lepidozioides* and *Takakia ceratophylla*, was not available. Relatively recently, *Takakia* has been shown to be sister to all other extant mosses (Liu et al. 2019) and to have diverged from the Last Common Ancestor of embryophytes after hornworts and liverworts (Hu et al. 2023). Therefore, we were keen to discover whether the recently sequenced *Takakia lepidozioides* genome possesses an *ASCL* gene like the vast majority of other embryophyte genomes, indicating the probable presence of genuine SP in its spore wall.

Putative ASCL and other type III PKS sequences in *Takakia* were identified based on phylogeny and sequence analysis. Our Maximum Likelihood (ML) phylogenetic tree (Fig. 1) resolves type III PKS sequences into the following clades: (a) an embryophyte ASCL clade containing bryophyte ASCL sequences including one *Takakia* sequence plus ASCL sequences from other major embryophyte groups. In agreement with the other ASCL sequences, the *Takakia* ASCL possesses diagnostic sequence features in addition to those for type III PKSs, namely Gly225 and (Ala/Val)240 (numbering based on PpASCL (Colpitts et al. 2011)) (Fig. S1), (b) a bryophyte clade containing exclusively ORS sequences including two *Takakia* sequences. In agreement with the other ORS sequences, the *Takakia* ORSs possess diagnostic sequence features in addition to those for type III PKSs, namely Gln218, (Val/Ala)277 and Ala286 (numbering according to PpORS (Kim et al. 2013)) (Fig. S1), (c) an algal clade containing exclusively Charophyte ORS sequences, (d) an embryophyte clade comprised of non-ORS and non-ASCL type III PKS sequences including a representative *Takakia* type III PKS, (e) five separate algal clades, collectively comprising 35 non-ORS and non-ASCL type III PKS sequences.

Thus, *Takakia lepidozioides* has an *ASCL* gene in agreement with our contention that all embryophytes, with the possible exception of a few species, possess ASCL, which serves as a marker for genuine SP in their spore or pollen walls. The few species predicted to lack ASCL are likely to exist in habitats that do not require protection from subaerial stresses, e.g. *Zostera marina*, and are presumed to have lost ASCL and pollen wall SP secondarily by reductive evolution.

In this study, we have examined all the algal genomes in the PhycoCosm database and reinforced our discovery that, while algae possess type III PKS sequences, none of them falls within the ASCL clade and, therefore, algae don't possess genuine SP.

Another relevant discovery is that *Takakia* has two full-length *ORS* genes, whose distribution in the Plant Kingdom, unlike that of *ASCL* genes, is limited to Charophytes and bryophytes. Notably, however, *Marchantia* species lack *ORS* genes. As more data become available, it will be interesting to see whether ORS is missing in all liverworts or if the *Marchantia* genus is the



lone exception. We have shown previously that ORS is required for integrity of the leaf cuticle of *Physcomitrium patens* and for its resistance to dehydration (Li et al. 2018) and that (2'-oxo)alkylresorcinols restore dehydration tolerance in a PpORS knockout line (Aslam et al. 2022). We presume ORS has the same role in *Takakia* and at least some other bryophytes. Interestingly, although ORS sequences are present in Charophytes, they appear to be absent from other algae.

Methods

We performed **BLASTp** searches the Takakia lepidozioides genome database against (https://www.takakia.com/blast/blast_cs.html (v3.1)) with PpASCL and PpORS (Physcomitrium patens 2'-OxoalkylResorcinol Synthase), a bryophyte/charophyte-specific type III PKS, as query sequences. Sixteen putative type III PKS models were identified based on the presence of the catalytic Cys-His-Asn triad and signature sequences (G/A)FGPG (Suh et al. 2000). Among these sixteen sequences, one ASCL and two ORS sequences were recognised based on phylogeny (Fig. 1) and possession of additional enzyme-specific residues as described in the text and in Fig. S1. Similarly, putative algal type III PKS retrieved by BLASTp searches against each algal genome sequences were (https://phycocosm.jgi.doe.gov/phycocosm/home (accessed on 01 September 2023)). In cases of fusion proteins, portions of sequences that matched type III PKS sequences from the same or related species were taken for further analysis. Representative embryophyte type III PKS sequences were retrieved from Phytozome 13 (https://phytozome-next.jgi.doe.gov/ (Physcomitrium patens v3.3)) as described previously (Aslam et al. 2022). The sequences used for tree reconstruction (Table 1) were aligned by the MUSCLE method in MEGA 11 (Tamura et al 2021), and a ML phylogenetic tree (Fig. 1) was reconstructed in MEGA 11 using the JTT substitution model. The initial tree was created using the default NJ/BioNJ method, and tree improvement was performed using the nearest-neighbor-interchange ML heuristic method. Support for the tree was measured using 1,000 bootstrap replicates.

Reagents

Table 1. Plant type III polyketide synthases used for tree reconstruction

Type III PKSs are listed in the same order of their appearance (from the top) in the ML tree (Fig. 1) before collapsing some of the clades. *Takakia* enzymes are shown in bold. Fusion proteins containing a type III PKS domain are indicated with asterisks.

Enzyme	Species	Classification	
Outgroup			
Synechococcus PKS	Synechococcus sp.	Cyanobacteria, Synechococcales	
Algal type III PKS (non-ORS, non-ASCL)			

Chrveli1 20057*	Chromera velia	SAR, Chromerida
Pico_ML_1 52161	Picocystis sp.	Chlorophyta, Picocystales
Semro1 36990*	Seminavis robusta	Ochrophyta, Naviculales
Ochro1393_1_4 754228*	Ochromonas sp.	Ochrophyta, Ochromonadales
Ochro2298_1 456847	Ochromonadaceae sp.	Ochrophyta, Ochromonadales
Ochro2298_1 419265*	Ochromonadaceae sp.	Ochrophyta, Ochromonadales
Ochro1393_1_4 932390	Ochromonas sp.	Ochrophyta, Ochromonadales
Ochro1393_1_4 179269	Ochromonas sp.	Ochrophyta, Ochromonadales
Mesen1 9713	Mesotaenium endlicherianum	Charophyta, Zygnematales
Ectsil1 17490	Ectocarpus siliculosus	SAR, Ectocarpales
Claok1 5931	Cladosiphon okamuranus	SAR, Ectocarpales
Sacja1 7224	Saccharina japonica	SAR, Laminariales
Macpyr2 5041534	Macrocystis pyrifera	SAR, Laminariales
Undpi1 10741	Undaria pinnatifida	SAR, Laminariales
Alaesc1 15363	Alaria esculenta	SAR, Laminariales
MonC141_1 1230	Monodopsis strain	Ochrophyta, Eustigmatales
VisC74_1 13231*	Vischeria strain	Ochrophyta, Eustigmatales
Ochro1393_1_4 905391	Ochromonas sp.	Ochrophyta, Ochromonadales
Ochro2298_1 408009	Ochromonadaceae sp.	Ochrophyta, Ochromonadales
Ochro2298_1 458546	Ochromonadaceae sp.	Ochrophyta, Ochromonadales
Pelago2097_1 478370	Pelagophyceae sp.	SAR, Pelagomonadales
Ectsil1 30595	Ectocarpus siliculosus	SAR, Ectocarpales
Claok1 7458*	Cladosiphon okamuranus	SAR, Ectocarpales
Macpyr2 9688433	Macrocystis pyrifera	SAR, Laminariales
Sacja1 14235	Saccharina japonica	SAR, Laminariales
Alaesc1 3438	Alaria esculenta	SAR, Laminariales
Undpi1 3722	Undaria pinnatifida	SAR, Laminariales
Undpi1 3721	Undaria pinnatifida	SAR, Laminariales
SymretSc1 46235	Symbiochloris reticulata	Chlorophyta, Trebouxiales
Coccomyxa PKS	Coccomyxa subellipsoidea	Chlorophyta, Trebouxiales
Sceob152z_1 1656	Scenedesmus obliquus	Chlorophyta, Sphaeropleales
Tetrob172_l 3940502	Tetradesmus obliquus	Chlorophyta, Sphaeropleales
Spimu1 9263*	Spirogloea muscicola	Charophyta, Spirogloeales
Spimu1 2401*	Spirogloea muscicola	Charophyta, Spirogloeales
Spimu1 17478*	Spirogloea muscicola	Charophyta, Spirogloeales

Algal ORS

Meskra 3110107	Mesotaenium kramstae	Charophyta, Zygnematales
Spimu1 13141	Spirogloea muscicola	Charophyta, Spirogloeales
Spimu1 13204	Spirogloea muscicola	Charophyta, Spirogloeales
Zygcyl6981a_1 20083	Zygnema cf. cylindricum	Charophyta, Zygnematales
Zygcir1559_1 1602	Zygnema circumcarinatum	Charophyta, Zygnematales
Zygcir1559_1 1600	Zygnema circumcarinatum	Charophyta, Zygnematales
Meskra 2213449	Mesotaenium kramstae	Charophyta, Zygnematales
Meskra 2284394	Mesotaenium kramstae	Charophyta, Zygnematales
Penium pm001652g0050	Penium margaritaceum	Charophyta, Desmidiales
Penium pm022924g0020	Penium margaritaceum	Charophyta, Desmidiales
Penium pm001997g0040	Penium margaritaceum	Charophyta, Desmidiales
Penium pm003016g0080	Penium margaritaceum	Charophyta, Desmidiales
Bryophyte ORS		·
CepurGG1.6G090700	Ceratodon purpureus	Bryophyta, Dicranales
CepurGG1.2G039900	Ceratodon purpureus	Bryophyta, Dicranales
PpORS	Physcomitrium patens	Bryophyta, Funariales
AANG005398	Anthoceros angustus	Anthocerotophyta, Anthocerotales
Takakia Tle2c02540.1	Takakia lepidozioides	Bryophyta, Takakiales
Takakia Tle3c00272.1	Takakia lepidozioides	Bryophyta, Takakiales
CepurGG1.8G019400	Ceratodon purpureus	Bryophyta, Dicranales
Pohlia PKS2	Pohlia nutans	Bryophyta, Bryales
Phypa 126819	Physcomitrium patens	Bryophyta, Funariales
	Countedon money	Bryophyta, Dicranales
CepurGG1.9G130000	Ceratodon purpureus	Dryophyta, Dictanaics

AANG008114	Anthoceros angustus	Anthocerotophyta, Anthocerotales
Mp1g11030	Marchantia polymorpha	Marchantiophyta, Marchantiales
Takakia Tle1c06550.1	Takakia lepidozioides	Bryophyta, Takakiales
Sphfalx16G090200	Sphagnum fallax	Bryophyta, Sphagnales
Sphfalx20G004200	Sphagnum fallax	Bryophyta, Sphagnales
CepurGG1.N016500	Ceratodon purpureus	Bryophyta, Dicranales
PpASCL	Physcomitrium patens	Bryophyta, Funariales
Selmo1:122361	Selaginella moellendorffii	Lycopodiophyta, Selaginellales
Ceratopteris PKS	Ceratopteris richardii	Polypodiopsida, Polypodiales
PrCHSL	Pinus radiata	Gymnosperms, Pinales
AtPKSA	Arabidopsis thaliana	eudicots, rosids, Brassicales
AtPKSB	Arabidopsis thaliana	eudicots, rosids, Brassicales
Os10g0484800 (YY2)	Oryza sativa	monocots, Poales
Os07g0411300	Oryza sativa	monocots, Poales
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Embryophyte type III PKS (non-ORS, non-ASCL)

Takakia Tle2c05338.1	Takakia lepidozioides	Bryophyta, Takakiales
PpCHS	Physcomitrium patens	Bryophyta, Funariales
Mp4g23190	Marchantia polymorpha	Marchantiophyta, Marchantiales
AANG010604	Anthoceros angustus	Anthocerotophyta, Anthocerotales
MsCHS2	Medicago sativa	eudicots, rosids, Fabales
AhSTS	Arachis hypogaea	eudicots, rosids, Fabales
Gh2PS	Gerbera hybrid cultivar	eudicots, asterids, Asterales

Extended Data

Description: Fig. S1 Amino acid sequences of plant type III polyketide synthases included in the phylogenetic tree (Figure 1). Resource Type: Dataset. File: Revised Extended Data (22 June 2024).docx. DOI: 10.22002/1tdp3-92698

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