

RESEARCH PAPER

The capability to synthesize phytochelatins and the presence of constitutive and functional phytochelatin synthases are ancestral (plesiomorphic) characters for basal land plants

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Abstract

Bryophytes, a paraphyletic group which includes liverworts, mosses, and hornworts, have been stated as land plants that under metal stress (particularly cadmium) do not synthesize metal-binding peptides such as phytochelatins. Moreover, very little information is available to date regarding phytochelatin synthesis in charophytes, postulated to be the direct ancestors of land plants, or in lycophytes, namely very basal tracheophytes. In this study, it was hypothesized that basal land plants and charophytes have the capability to produce phytochelatins and possess constitutive and functional phytochelatin synthases. To verify this hypothesis, twelve bryophyte species (six liverworts, four mosses, and two hornworts), three charophytes, and two lycophyte species were exposed to 0–36 μM cadmium for 72 h, and then assayed for: (i) glutathione and phytochelatin quali-quantitative content by HPLC and mass spectrometry; (ii) the presence of putative phytochelatin synthases by western blotting; and (iii) *in vitro* activity of phytochelatin synthases. Of all the species tested, ten produced phytochelatins *in vivo*, while the other seven did not. The presence of a constitutively expressed and functional phytochelatin synthase was demonstrated in all the bryophyte lineages and in the lycophyte *Selaginella denticulata*, but not in the charophytes. Hence, current knowledge according to phytochelatins have been stated as being absent in bryophytes was therefore confuted by this work. It is argued that the capability to synthesize phytochelatins, as well as the presence of active phytochelatin synthases, are ancestral (plesiomorphic) characters for basal land plants.

Key words: Bryophytes, cadmium, charophytes, glutathione, hornworts, liverworts, lycophytes, metals, mosses, phytochelatins, phytochelatin synthase.

Introduction

Phytochelatins (PCs) are thiol-peptide compounds whose general structure is $(\gamma\text{-glutamate-cysteine})_n\text{-glycine}$, with n usually ranging from 2 to 5 (Grill *et al.*, 1985). Due to the thiol group of cysteine residues, PCs can bind cadmium (Cd) and other thiophilic metals, and prevent them from circulating in the cytosol (Grill *et al.*, 1985). PCs are synthesized from reduced glutathione (GSH), by means of the constitutively

expressed cytosolic enzyme phytochelatin synthase (PCS), a $\gamma\text{-glutamylcysteine dipeptidyl transpeptidase}$ (EC 2.3.2.15) (Grill *et al.*, 1989; Vatamaniuk *et al.*, 2004), belonging to clan CA of the papain-like cysteine proteases (Vivares *et al.*, 2005; Romanyuk *et al.*, 2006; Rea, 2012).

It is now well known that higher plants (Sanità di Toppi and Gabbrielli, 1999; Rea, 2012), a number of marine and

Abbreviations: ESI-MS, electrospray ionization mass-spectrometry; GSH, reduced glutathione; MS/MS, tandem mass spectrometry; Mya, million years ago; PC, phytochelatin; PCS, phytochelatin synthase.

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freshwater algae (*Chlorophyta*, *Chrysophyta*, *Phaeophyta*, *Rhodophyta*, etc.) (Gekeler et al., 1988; Pawlik-Skowrońska et al., 2007), some fungi (Gekeler et al., 1989; Kneer et al., 1992; Bolchi et al., 2011), lichens (Pawlik-Skowrońska et al., 2002), and even some animal species (Clemens et al., 2001; Vatamaniuk et al., 2001) do actually produce PCs in response to metal stress, particularly Cd.

In contrast, it has been recognized until now that the earliest diverged extant land plants, collectively referred to as bryophytes [liverworts (*Hepatophyta*, sin. *Marchantiophyta*), mosses (*Bryophyta*), and hornworts (*Anthocerotophyta*) (Vanderpoorten and Goffinet, 2009)] do not synthesize PCs under metal stress (Bruns et al., 1999, 2001; Rother et al., 2006; Leinenweber et al., 2009; Bleuel et al., 2011). As a matter of fact, the ‘model’ moss *Physcomitrella patens*, at present the only bryophyte with an entirely sequenced genome (Rensing et al., 2008), lacks the PCS gene (Kopriva et al., 2007). However, based on the limited number of species investigated (mainly mosses, very few liverworts, and no hornworts), the conclusion concerning a generalized absence of PCs in bryophytes appears to be quite premature and not aprioristically transferable *tout court* to all these plants. Indeed, with regards to liverworts, the very preliminary screening carried out by Gekeler et al. (1989) proved the thalloid species *Marchantia polymorpha* to be a weak producer of PCs under Cd stress. Such an interesting experimental result, however, was subsequently contradicted by Bruns et al. (2001), who exposed *M. polymorpha* and *Pellia epiphylla* to Cd for up to 10 d, and did not detect any PC synthesis. Likewise, although the presence of PCs in the metal-exposed freshwater moss *Rhynchostegium riparioides* and in the liverwort *Lunularia cruciata* was considered likely (Jackson et al., 1991; Carginale et al., 2004), no unequivocal proof of PC synthesis was ever provided in these species.

Recent studies have acquired a large body of evidence to support the hypotheses that: (i) land plants, and particularly bryophytes, have a charophytic ancestry [given in particular by the *Charophyta* orders *Charales*, *Coleochaetales*, and *Zygnematales* (McCourt et al., 2004; Qiu et al., 2006; Qiu, 2008; Becker and Marin, 2009; Wodniok et al., 2011)]; and (ii) hornworts are the sister group of tracheophytes (Qiu et al., 2006; Qiu, 2008; Villarreal et al., 2010; Ligrone et al., 2012). Concerning these points, with regard to charophytic algae, the synthesis of PCs has until now been definitely detected just in one species, namely *Micrasterias denticulata* (Volland et al., 2013), but no further investigation on PC synthesis occurrence in other charophytes, more phylogenetically close to (early) land plants, has been carried out. Secondly, no unequivocal data on PC production in early tracheophytes, namely lycophytes (*Lycophyta*), have been published, and the experiments performed by Gekeler et al. (1989) on Cd-induced PCs in *Lycopodium clavatum* and *Selaginella viticulosa* have not received solid confirmation in other lycophytes.

Interestingly, it has been well documented that the Early Devonian Rhynie Cherts of NE Scotland (~396 ± 12 Mya) contains the earliest and best preserved bryophytic and vascular flora (Rice et al., 1995; Kenrick and Crane, 1997; Pires and Dolan, 2012). This ancient site featured high levels of arsenic

(As), mercury (Hg), gold (Au), zinc (Zn), etc., all PC inducers to varying extents (Zenk, 1996, and references therein). In view of the above, the present work assumed the main hypothesis that the capability to produce PCs and the constitutive presence of functional PCS(s) enzyme(s) might represent useful conserved traits for all bryophyte lineages. *A fortiori*, the general reliance of bryophytes on aerial fallout for mineral nutrition, and hence their potential vulnerability to heavy metal pollution (Tyler, 1990; Harmens et al., 2012, and references therein), demand further caution in generalizing conclusions about an overall lack of PCs (and PCSs) in these plants.

To verify this hypothesis, investigation was focused not only on mosses, the most studied group within bryophytes, but also on liverworts, placed in the basalmost position amongst land plants (Qiu et al., 2006; Qiu, 2008; Crandall-Stotler et al., 2009), and on hornworts, sisters of tracheophytes (Qiu et al., 2006; Qiu, 2008; Villarreal et al., 2010; Ligrone et al., 2012). Additionally, in order to provide a more comprehensive picture from a phylogenetic point of view, it was verified whether PCs (as well as PCSs) might also be present in charophytes, now thought to be the direct algal ancestors of land plants (McCourt et al., 2004; Qiu et al., 2006; Qiu, 2008; Becker and Marin, 2009; Wodniok et al., 2011), and in lycophytes, a very ancient lineage that diverged shortly after evolution of vascular tissues in land plants, and also present in the Early Devonian Rhynie Cherts (Rice et al., 1995; Kenrick and Crane, 1997; Pires and Dolan, 2012).

To provide a clear understanding of the taxa investigated in this work, a simplified cladogram is shown in Fig. 1.

Materials and methods

Experimental set-up and species employed

The material employed in the experiments [(charophytes; bryophyte gametophytes; lycophyte sporophytes (roots))] consisted of non-axenic cultures (see the species list below), collected in Italian locations (Apuan Alps, Massaciuccoli Lake, surroundings of Parma, and Salerno) in February–October of 2012 and 2013. In addition, specific monocultures of *Coleochaete scutata* were obtained from the Carolina Biological Supply Company, Burlington, NC, USA, and set up as the other cultures. All material was abundantly rinsed with double-distilled water, carefully checked for the absence of all potential soil/biological contaminants and endosymbiotic organisms (i.e. *Cyanobacteria*, *Glomeromycota*, etc.) under a stereomicroscope (WILD, Heerbrugg, Switzerland) and an Olympus BH2 microscope ×40 (Olympus Italia, Segrate, Italy), and then independently cultivated as specific monocultures in a growth chamber under the conditions detailed below.

All together, 17 species were investigated (Fig. 1): three charophytes (*Charophyta*), namely *Spirogyra* sp. Link, *Chara vulgaris* L., and *Coleochaete scutata* Bréb.; six liverworts (*Hepatophyta*), namely *Conocephalum conicum* (L.) Dumort., *Marchantia polymorpha* L., *Pellia epiphylla* (L.) Corda, *Radula complanata* (L.) Dumort., *Aneura pinguis* (L.) Dumort., and *Scapania undulata* (L.) Dumort.; four mosses (*Bryophyta*), namely *Sphagnum palustre* L., *Polytrichastrum formosum* (Hedw.) G.L.Sm., *Hypnum cupressiforme* Hedw., and *Fontinalis antipyretica* Hedw.; two hornworts (*Anthocerotophyta*), namely *Anthoceros punctatus* L. and *Phaeoceros laevis* (L.) Prosk; and two lycophytes, namely *Huperzia selago* (L.) Bernh. ex Schrank and Mart. and *Selaginella denticulata* (L.) Spring. As reference organisms for testing PC production and PCS presence/size/activity, monocultures of *Nostoc* sp. (*Cyanobacteria*), *Physcomitrella patens* (*Bryophyta*), and *Arabidopsis thaliana* (*Magnoliophyta*) *in vitro*

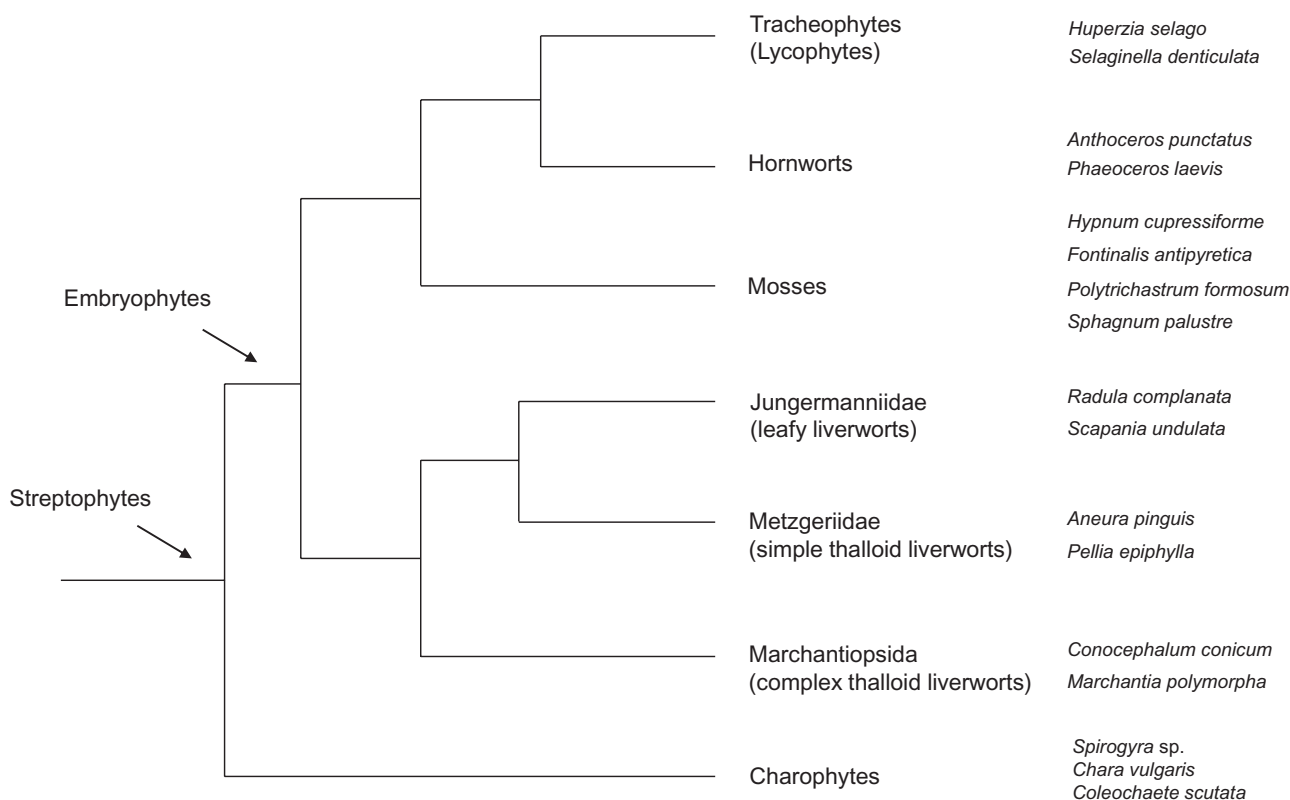


Fig. 1. Simplified phylogeny of charophytic algae and land plants, related to the species investigated in this work.

plants were grown in the same controlled conditions as the above material.

After selection and cleaning, the youngest parts of the samples (n ranging from 6 to 24; Table 1) were transferred into sterilized plant culture vials (Phytacon™), filled with 100 ml of sterile modified Chiaudani–Vighi (Chiaudani and Vighi, 1977) culture medium (pH 6.0), with a final composition as follows: NaNO_3 2.55 mg l^{-1} , K_2HPO_4 0.10 mg l^{-1} , MgCl_2 0.57 mg l^{-1} , $\text{MgSO}_4 \cdot 7\text{H}_2\text{O}$ 2.94 mg l^{-1} , $\text{CaCl}_2 \cdot 2\text{H}_2\text{O}$ 0.44 mg l^{-1} , NaHCO_3 1.50 mg l^{-1} , FeSO_4 5.58 $\mu\text{g l}^{-1}$, H_3BO_3 18.55 $\mu\text{g l}^{-1}$, MnCl_2 26.43 $\mu\text{g l}^{-1}$, ZnCl_2 3.28 $\mu\text{g l}^{-1}$, CoCl_2 80 ng l^{-1} , CuCl_2 0.9 ng l^{-1} , $\text{Na}_2\text{MoO}_4 \cdot 2\text{H}_2\text{O}$ 730 ng l^{-1} , omitting the ethylenediaminetetraacetic acid (EDTA) in order to avoid undesirable Cd chelation. Then, all vials were placed in a growth chamber at 20 ± 1 °C, under a photoperiod of 16 h light [photosynthetic photon flux density (PPFD) 60 $\mu\text{mol m}^{-2} \text{s}^{-1}$]/8 h dark. With regard to the species to be analysed for their thiol-peptide content and PCS immunoreactivity, half of the samples were treated for 72 h with 36 μM Cd, in the form of $3\text{CdSO}_4 \cdot 8\text{H}_2\text{O}$ (*C. conicum* also with 72 μM Cd and for 144 h), whereas the other half (controls) were given deionized water in an identical volume as for the Cd solution. Moreover, the bryophyte cultures were daily sprayed with Cd at the same concentration as that in the medium (as well as deionized water in controls), because of their well-known capability to absorb elements also from the atmosphere (Tyler, 1990). At the end of the treatments, all plants were carefully rinsed with deionized water, gently blotted dry with filter paper, wrapped in aluminium foil, frozen in liquid nitrogen, and briefly stored at -80 °C for further analyses.

Thiol-peptide separation, detection, and quantification by HPLC

Cd-treated and control cultures [at least 250 mg fresh weight (FW) for each sample] were homogenized in a mortar in ice-cold 5% (w/v) 5-sulphosalicylic acid, containing 6.3 mM diethylenetriamine-pentaacetic acid, according to de Knecht *et al.* (1994). After centrifugation at 10 000 g for 10 min at 4 °C, the supernatant fraction was filtered through Minisart RC4 0.45 μm filters (Sartorius, Goettingen,

Germany) and immediately assayed by HPLC (Model 200, Perkin-Elmer, Wellesley, MA, USA). Thiol-containing peptides (GSH and PCs) were separated by a Supelco Ascentis Express reverse-phase C_{18} column (Sigma-Aldrich, Milan, Italy). The separation was achieved using a 0–26% acetonitrile gradient, with a flow rate set at 0.3 ml min^{-1} . The elution solutions contained 0.05% trifluoroacetic acid. Thiol-peptides were determined using post-column derivatization with 300 μM Ellman's reagent [5,5'-dithio(2-nitrobenzoic acid); DTNB] (Sigma-Aldrich), detected at 412 nm (PerkinElmer detector mod. 200), and quantified by calibration curves for standard SH groups. Identification of GSH and individual PCs was based on the comparison of their retention times with standard GSH (Merck, Darmstadt, Germany) and PC samples from *Silene vulgaris*.

Thiol-peptide characterization by mass spectrometry

The identity of putative GSH and PCs was verified by electrospray ionization mass spectrometry (ESI-MS) and tandem mass spectrometry (MS/MS). Liquid chromatographic elution was carried out on a Supelco Ascentis Express reverse-phase C_{18} column (Sigma-Aldrich), using a gradient solvent system [solvent A, aqueous 0.05% (v/v) trifluoroacetic acid; solvent B, 0.05% (v/v) trifluoroacetic acid in acetonitrile] as follows: solvent B was set at 2% for 12 min, raised with a linear gradient to 11% in 3 min, raised with a linear gradient to 26% in 16 min, and then raised with a linear gradient to 98% in 7 min. Solvent B was maintained at 98% for 6 min before column re-equilibration (10 min). The flow rate was 0.3 ml min^{-1} . The mobile phase was delivered by a Ultimate3000 system (ThermoElectron Corporation, San José, CA, USA). The injection volume was 20 μl . An LTQ Orbitrap XL (ThermoElectron Corporation) with a pneumatically assisted ESI interface was used. The system was controlled by Xcalibur software. The sheath gas (nitrogen, 99.999% purity) was delivered at a flow rate of 50 arbitrary units; sweep and the auxiliary gas (nitrogen, 99.999% purity) were delivered at a flow rate of 20 arbitrary units. The optimized conditions of the interface were as follows: ESI voltage, 3.5 kV; capillary voltage, 30 V; capillary

Table 1. Content of glutathione (GSH) and phytochelatin (PC) in charophytes, bryophytes (gametophytes), and lycophytes (roots), non-exposed (control) or exposed to 36 μM Cd for 72 h (in *Conocephalum conicum* also ^a36 μM Cd for 144 h and ^b72 μM Cd for 72 h)GSH and PC concentrations (means \pm SE) are expressed in nmol g^{-1} FW.

	GSH (control)	GSH (Cd-exposed)	PC ₂ (Cd-exposed)	PC ₃ (Cd-exposed)	PC ₄ (Cd-exposed)	Total PCs (Cd-exposed)
Charophytes						
<i>Spirogyra</i> sp. (n=10+10)	105.5 \pm 9.7	144.7 \pm 16.2	2.7 \pm 0.9	1.9 \pm 0.7	ND	4.6 \pm 0.8
<i>Chara vulgaris</i> (n=10+10)	92.5 \pm 10.5	99.7 \pm 7.5	4.1 \pm 0.3	3.8 \pm 0.6	2.7 \pm 0.7	10.6 \pm 0.8
<i>Coleochaete scutata</i> (n=5+5)	222.3 \pm 12.1	212.2 \pm 7.7	9.5 \pm 0.4	16.8 \pm 1.2	14.1 \pm 0.7	40.4 \pm 0.8
Liverworts						
<i>Conocephalum conicum</i> (n=6+6)	1.4 \pm 0.2	6.8 \pm 2.4**	6.1 \pm 3.5	1.2 \pm 0.5	0.8 \pm 0.3	8.1 \pm 1.4
		16.6 \pm 4.3** ^a	2.9 \pm 0.3 ^a	4.0 \pm 2.3 ^a	2.9 \pm 0.7 ^a	9.8 \pm 1.1 ^a
		10.7 \pm 3.2** ^b	4.2 \pm 3.5 ^b	2.6 \pm 1.4 ^b	1.7 \pm 0.4 ^b	8.5 \pm 1.8 ^b
<i>Marchantia polymorpha</i> (n=4+4)	2.9 \pm 0.2	3.6 \pm 0.8	1.8 \pm 0.4	1.3 \pm 0.9	ND	3.1 \pm 0.6
<i>Pellia epiphylla</i> (n=3+3)	132.2 \pm 36.8	89.4 \pm 26.5	1.0 \pm 0.3	ND	ND	1.0 \pm 0.3
<i>Radula complanata</i> (n=3+3)	2.7 \pm 1.1	6.3 \pm 2.1	1.2 \pm 0.3	ND	ND	1.2 \pm 0.3
<i>Aneura pinguis</i> (n=4+4)	3.3 \pm 1.0	5.6 \pm 1.7	ND	ND	ND	ND
<i>Scapania undulata</i> (n=3+3)	20.5 \pm 3.5	17.8 \pm 2.2	ND	ND	ND	ND
Mosses						
<i>Sphagnum palustre</i> (n=6+6)	23.4 \pm 3.1	34.2 \pm 2.1*	4.5 \pm 0.3	5.8 \pm 0.2	3.0 \pm 0.7	13.3 \pm 0.8
<i>Polytrichastrum formosum</i> (n=4+4)	121.7 \pm 12.7	367.7 \pm 25.6**	ND	ND	ND	ND
<i>Hypnum cupressiforme</i> (n=4+4)	302.9 \pm 46.1	123.6 \pm 28.1	ND	ND	ND	ND
<i>Fontinalis antipyretica</i> (n=3+3)	25.7 \pm 7.0	148.0 \pm 19.1*	ND	ND	ND	ND
Hornworts						
<i>Anthoceros punctatus</i> (n=5+5)	2.8 \pm 0.7	1.9 \pm 0.8	ND	ND	ND	ND
<i>Phaeoceros laevis</i> (n=12+12)	4.5 \pm 0.9	1.8 \pm 0.2*	1.6 \pm 0.3	3.3 \pm 0.9	ND	4.9 \pm 0.6
Lycophytes						
<i>Huperzia selago</i> (n=5+5)	337.4 \pm 23.6	275.9 \pm 31.3	ND	ND	ND	ND
<i>Selaginella denticulata</i> (n=5+5)	67.7 \pm 10.6	126.4 \pm 15.9*	3.8 \pm 1.6	3.1 \pm 1.0	ND	6.9 \pm 1.3

Asterisks indicate significant differences in GSH concentrations between controls and Cd-exposed samples at * $P < 0.05$ or ** $P < 0.01$, according to the Kruskal–Wallis non-parametric test; n for each experimental set (controls+Cd-exposed) is specified in parenthesis.

ND, no PCs detected by HPLC and ESI-MS analyses.

temperature, 300 °C; and tube lens, 110 V. MS experiments were carried out in the 200–1500 mass-to-charge ratio (m/z) range. MS/MS experiments were performed in the ion-trap collision cell with a normalized collision energy of 35 arbitrary units and an isolation width of 1 m/z ; the product ions were analysed with an Orbitrap analyzer, with the m/z set as a function of GSH and PC molecular mass.

Phytochelatin synthase activity assays

Bryophyte and lycophyte PCS activities were assayed in 250 mg of fresh Cd-untreated material, from gametophytes of *C. conicum*, *P. epiphylla*, *R. complanata*, *A. pinguis*, *S. undulata*, *S. palustre*, *P. formosum*, *F. antipyretica*, and *P. laevis*; and sporophytes (roots) of *H. selago* and *S. denticulata* (and *A. thaliana* plants as a positive control), grown as described above. The PCS activity was not tested in the charophytes, in the liverwort *M. polymorpha*, in the moss *H. cupressiforme*, and in the hornwort *A. punctatus*, essentially due to the lack of fresh material at the time of the revision of the present work. All the material was assayed *in vitro*, mainly following the protocol of [Wojas et al. \(2008\)](#), with some slight modifications. Each sample was put in a 1.5 ml Eppendorf tube (filled with 350 mg of glass microbeads with a diameter of 0.2 mm), placed in liquid nitrogen, and ground to a powder with a TAC 200/S amalgamator (Linea TAC s.r.l., Asti, Italy; oscillation frequency: 4200 strokes min^{-1} for 10 s); the use of the amalgamator gave a better yield compared with the mortar extraction. The powder was then added to 700 μl of extraction buffer ([Wojas et al., 2008](#)) and homogenized for 10 s with another cycle of amalgamator shaking. The homogenized samples were then centrifuged twice at 4 °C/13 000 g for 10 min, and

400 μl of the supernatants added to 100 μl of the reaction buffer, as in [Wojas et al. \(2008\)](#), containing 100 μM Cd and the protease inhibitor cocktail ‘complete mini EDTA-free’ (Roche Italia, Milan, Italy). After an incubation time of 90 min at 35 °C, and termination of the reaction with 125 μl of 20% trichloroacetic acid, the measurement of the PCS activity was immediately performed by HPLC–ESI-MS, set up as described in the previous paragraph. The PCS activity was determined from the integrated total PC peak areas, and expressed in pmol PCs min^{-1} g^{-1} FW, according to calibration curves set up with known concentrations of standard GSH (Merck, Darmstadt, Germany), and to [Ogawa et al. \(2010\)](#).

Western blots of phytochelatin synthases

Proteins were extracted from at least 150 mg of fresh tissues of *C. vulgaris* and *C. scutata*; gametophytes of *C. conicum*, *P. epiphylla*, *R. complanata*, *A. pinguis*, *S. undulata*, *S. palustre*, *P. formosum*, *F. antipyretica*, *P. laevis* (and *P. patens* as a negative control); and sporophytes of *P. laevis*, *H. selago* (roots), and *S. denticulata* (roots), by using 300 μl of lysis buffer, containing 50 mM TRIS-HCl pH 7.5, 2 M thiourea, 7 M urea, 2% (v/v) Triton X-100, 1% dithiothreitol (DTT), 2% (w/v) polyvinylpyrrolidone (PVPP), 1 mM phenyl-methylsulphonyl fluoride (PMSF), and 0.2% β -mercaptoethanol. Protein concentration was determined according to [Bradford \(1976\)](#), using bovine serum albumin as a standard. Aliquots of protein extracts (\sim 20 μg of total protein per well) were separated using a 12% SDS–polyacrylamide gel ([Laemmli, 1970](#)), and electrotransferred at 100 V for 60 min to a nitrocellulose membrane (GE Healthcare Bio-Sciences AB, Uppsala, Sweden), using the Mini Trans-Blot cell

apparatus (Bio-Rad Laboratories, CA, USA). Protein loading and transfer efficiency of extracts were verified by Ponceau-S staining. Western blot analyses were performed with a polyclonal antibody (diluted 1:5000 in blocking buffer and probed for 2 h) raised against *A. thaliana* PCS1. As references for the immunoblotting specificity/detection, recombinant *A. thaliana* PCS1 (AtPCS1, 56 kDa) (Ruotolo *et al.*, 2004), root samples of *in vitro* grown *A. thaliana* seedlings, and monocultures of the cyanobacterium *Nostoc* sp., were employed. No western blots of apo-*Glomeromycota* as accidental contaminants were performed, since these fungi do not possess any PCS (as shown by a search on the GenBank database for annotated PCS orthologues on 15 November 2013) and do not produce PCs (Bolchi *et al.*, 2011). Immunoreactivity was visualized using an anti-rabbit IgG antibody conjugated to horseradish peroxidase (GE Healthcare Bio-Sciences AB) and a chemoluminescence western blotting detection system (Pierce ECL Plus Western blot Substrate; Pierce, Rockford, IL, USA), according to the manufacturer's instructions.

Statistics

Statistical analyses were performed by SPSS Statistics, version 19. Comparisons amongst independent samples were done by the Kruskal–Wallis non-parametric test ($P < 0.05$ and $P < 0.01$).

Results

Bryophytes, charophytes, and lycophytes synthesize cadmium-induced phytochelatin *in vivo*

All charophytes investigated here were shown to possess relatively high levels of GSH, whose concentrations were not significantly affected by Cd treatment (Table 1). Two species, namely *C. vulgaris* and *C. scutata*, produced PCs under Cd stress up to PC₄, the latter species being the best PC producer amongst all the species tested in this work (Table 1). In contrast, *Spirogyra* sp. synthesized only traces of PCs, up to PC₃ (Table 1). In order to confirm the identity of the HPLC-detected thiol-peptides in charophytes, samples were subjected to ESI-MS and MS/MS analyses. Representative *C. vulgaris* and *C. scutata* ESI-MS chromatograms for GSH in controls, and for GSH and PCs in Cd-exposed material are shown in Supplementary Fig. S1 available at JXB online.

As regards bryophytes, amongst the liverworts belonging to the class Marchantiopsida (Fig. 1), in *C. conicum* the GSH level under Cd exposure was much higher than in controls ($P = 0.004$, Table 1), and PCs (PC₂, PC₃, and PC₄) were detected, even with longer exposure time (144 h) or higher Cd concentration (72 μ M) (Table 1). In *M. polymorpha*, no differences in GSH levels between controls and Cd-exposed samples were found, and PC₂ and PC₃, but not PC₄, were produced (Table 1). With regard to the Jungermanniopsida (comprising Jungermanniidae and Metzgeriidae) (Fig. 1), the highest levels of GSH amongst all liverworts investigated here were found in *P. epiphylla*, with no differences between controls and Cd-exposed samples (although there was a non-significant, but manifest downward trend under Cd stress) (Table 1); as far as PCs are concerned, only PC₂ was produced *in vivo* by this liverwort (Table 1). In *R. complanata*, the levels of GSH were much lower than those of *P. epiphylla*, and no significant differences were observed in GSH levels between controls and Cd-exposed samples; with

regard to PCs, in *R. complanata* only PC₂ was induced *in vivo* by Cd (Table 1). In *A. pinguis* and *S. undulata*, no differences in GSH levels were found between controls and Cd-treated samples (Table 1), and no PC *in vivo* production was detected.

As far as mosses are concerned, in *S. palustre*, belonging to the very ancient Sphagnopsida moss clade (Fig. 1) (Newton *et al.*, 2009), the GSH level under Cd stress was higher than in controls ($P = 0.048$), and the production of PC₂, PC₃, and PC₄ was detected (Table 1). In contrast, in *P. formosum*, *H. cupressiforme*, and *F. antipyretica*, no PC production under Cd stress was evidenced, but marked differences as regards the GSH content between control and Cd-exposed samples were observed (Table 1). In particular, *P. formosum* and *F. antipyretica* showed large increases in GSH levels under Cd stress compared with controls ($P = 0.015$ and $P = 0.042$, respectively), whereas *H. cupressiforme* displayed an opposite trend (Table 1).

In the two hornworts studied (*A. punctatus* and *P. laevis*), the thiol-peptide production in response to Cd stress varied qualitatively between the two species. In *A. punctatus*, no significant differences in GSH levels between controls and Cd-exposed plants were measured, and no PC synthesis was detected (Table 1). In contrast, in *P. laevis* the GSH concentration under Cd stress significantly diminished ($P = 0.023$), whereas synthesis of PC₂ and PC₃ was induced (Table 1).

Finally, in the two lycophytes, namely *H. selago* and *S. denticulata*, the thiol-peptide levels varied to a large extent between them. In *H. selago*, high constitutive levels of GSH, without variation under Cd stress, were detected, and no PC synthesis was measured *in vivo* (Table 1). In contrast, in *S. denticulata* the GSH concentrations were lower than in *H. selago*, but they significantly increased ($P = 0.031$) under Cd stress, which also induced the synthesis of PC₂ and PC₃ (Table 1).

Representative *C. conicum*, *S. palustre*, and *P. laevis* HPLC and ESI-MS chromatograms for GSH in controls, and for GSH and PCs in Cd-exposed samples, are shown in Fig. 2. Moreover, representative *C. conicum* MS/MS product-ion spectra and the relative fragmentation patterns for GSH, PC₂, PC₃, and PC₄ are shown in Fig. 3. Finally, ESI-MS chromatograms of GSH and PCs produced by *P. epiphylla* and *S. denticulata* are shown in Supplementary Fig. S2 available at JXB online.

Basal land plants possess constitutive and functional phytochelatin synthases

The gametophytes of the liverworts *C. conicum* and *S. undulata*, and of the moss *S. palustre*, showed the constitutive presence of a putative PCS band of almost the same molecular mass, that is ~36 kDa (Fig. 4, lanes 5–7); in contrast, the gametophytes of the hornwort *P. laevis* and the sporophytes (roots) of the lycophyte *S. denticulata* were shown to possess, respectively, a lighter and a heavier (than those above) putative PCS band, with molecular masses of ~28 kDa and ~40 kDa (Fig. 4, lanes 3 and 9). A band of ~28 kDa was also immunodetected in the sporophytes of *P. laevis* (Fig. 4, lane 2). All the basal land plant PCSs were lighter than the PCSs from *A. thaliana* (roots and recombinant), whose bands were shown to possess a molecular mass of ~56–58 kDa (Fig. 4, lanes 4 and 10).

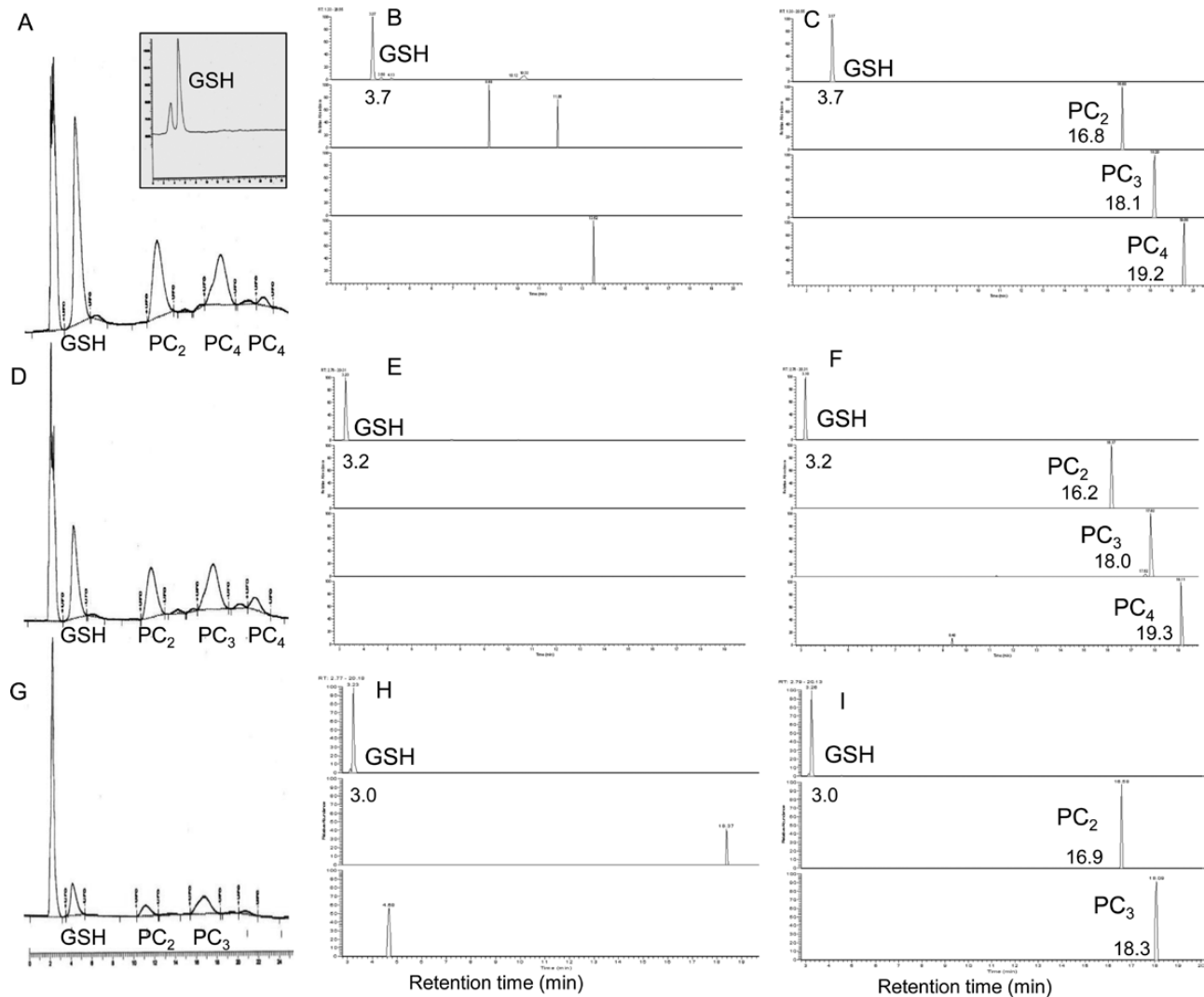


Fig. 2. Representative Ellman-derivatized HPLC chromatograms of thiol-peptides [GSH, PC₂, PC₃ (and PC₄)] from Cd-exposed (A) *Conocephalum conicum* (control in the insert), (D) *Sphagnum palustre* and (G) *Phaeoceros laevis*. Relative ESI-MS chromatograms of: *C. conicum* (B) GSH in controls and (C) GSH, PC₂, PC₃, and PC₄ in Cd-exposed plants; *S. palustre* (E) GSH in controls and (F) GSH, PC₂, PC₃, and PC₄ in Cd-exposed plants; *P. laevis* (H) GSH in controls and (I) GSH, PC₂, and PC₃ in Cd-exposed plants. ESI-MS retention times are indicated below each thiol-peptide peak. Unlabelled peaks are non-specific and do not represent signals for thiol-peptide characterization.

As expected, no PCS band was detected in *P. patens* (Fig. 4, lane 8). Western analysis of the other species, including the charophytes, did not reveal the presence of any immunoreactive band. Also, in all the above blots, no presence of aspecific bands was detected, thus allowing it to be excluded that accidental contamination with other species had occurred. In particular, no putative PCS-like bands were immunodetected in extracts from monocultures of *Nostoc* sp. (Fig. 4, lane 1).

Furthermore, the assayed bryophyte and lycophyte species had *in vitro* PCS activity varying to different extents (Table 2), but in general much lower than that measured in *A. thaliana* plants *in vitro*. In fact, if the latter activity was fixed equal to 100%, the relative PCS activities of bryophytes and lycophytes (extracted and incubated in identical conditions to those of *A. thaliana*) ranged from a minimum of 1.0% in *S. palustre*, to a maximum of 13.1% in *H. selago* (Table 2). In general, the highest PCS activities were assayed in the two lycophytes, followed in decreasing order by the liverworts, the hornwort *P. laevis*, and the moss *S. palustre*. No PCS activity

was detected in the mosses *P. formosum* and *F. antipyretica*. Omission of Cd supply in the extraction and reaction buffers resulted in no detectable or very low synthesis of PCs.

Discussion

The results of this study differ from the currently held view that basically affirms: 'all bryophytes do not produce PCs and do not possess a PCS enzyme' (Bruns *et al.*, 1999, 2001; Rother *et al.*, 2006; Kopriva *et al.*, 2007; Leinenweber *et al.*, 2009; Bleuel *et al.*, 2011). In fact, PC *in vivo* production and western-immunoreactive PCS bands were found in: (i) a number of liverworts; (ii) the moss *S. palustre*; and (iii) gametophytes and sporophytes of the hornwort *P. laevis*. Cd-induced PCs were also detected *in vivo* in three charophytes, and in one lycophyte, namely *S. denticulata*, in which a PCS band was also immunodetected.

Other than PCs and PCSs, a number of the basal land plants investigated here (particularly lycophytes and liverworts) showed a detectable, and sometimes not negligible,

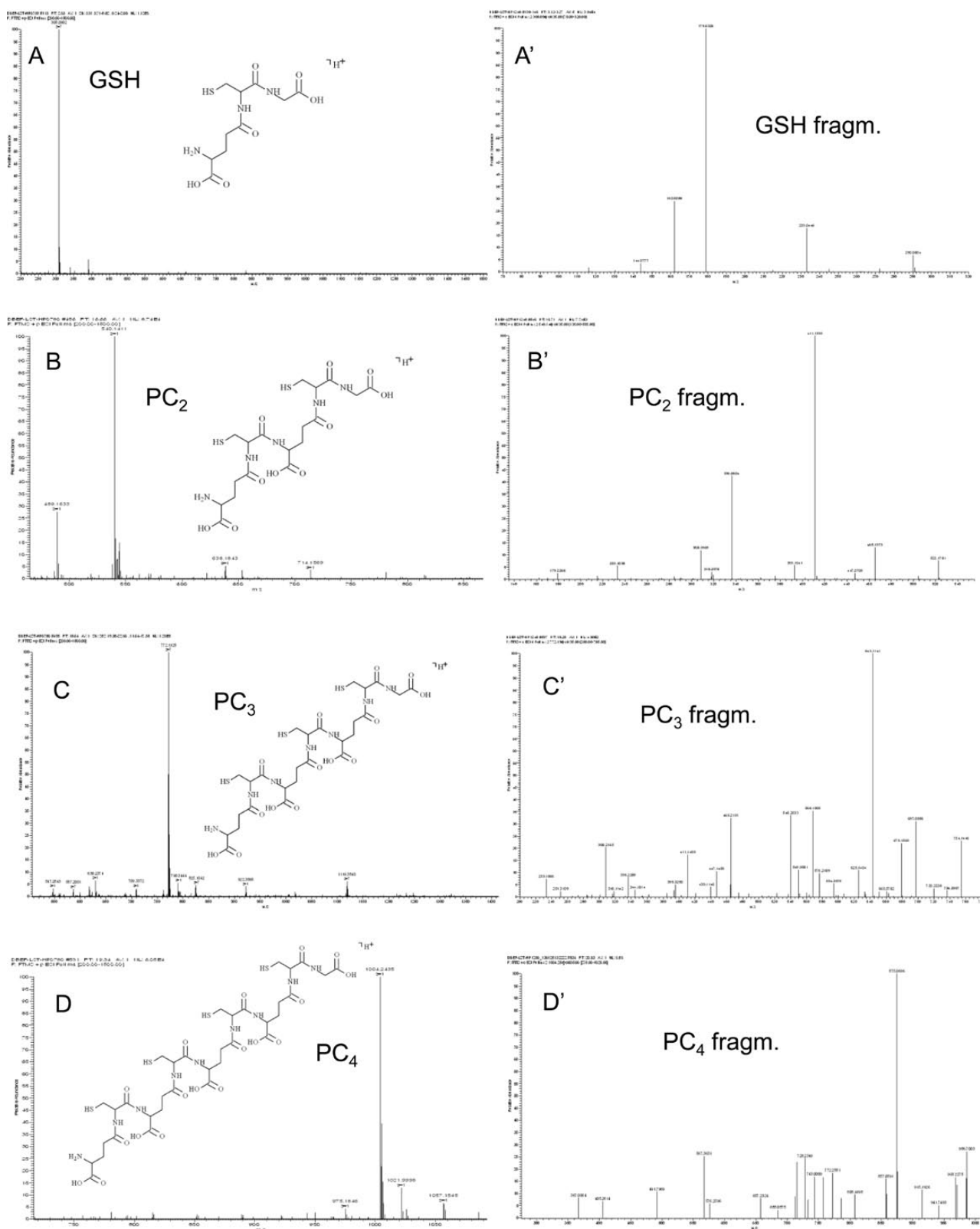


Fig. 3. Representative *Conocephalum conicum* MS/MS product-ion spectra of thiol-peptides (A, m/z 308=GSH; B, m/z 540=PC₂; C, m/z 772=PC₃; D, m/z 1004=PC₄), deduced formulae, and relative fragmentation patterns at a normalized collision energy of 35 arbitrary units (A', GSH; B', PC₂; C', PC₃; D', PC₄). The MS/MS thiol product-ion spectra from all the other species were almost identical to that shown here.

PCS *in vitro* activity. Hence, a functional PCS enzyme is present in all the lineages of bryophytes, as well as in early tracheophytes.

Overall, the total amount of PCs produced *in vivo* under Cd stress by the above basal land plants appears to be much lower (sometimes by one or two orders of magnitudes or

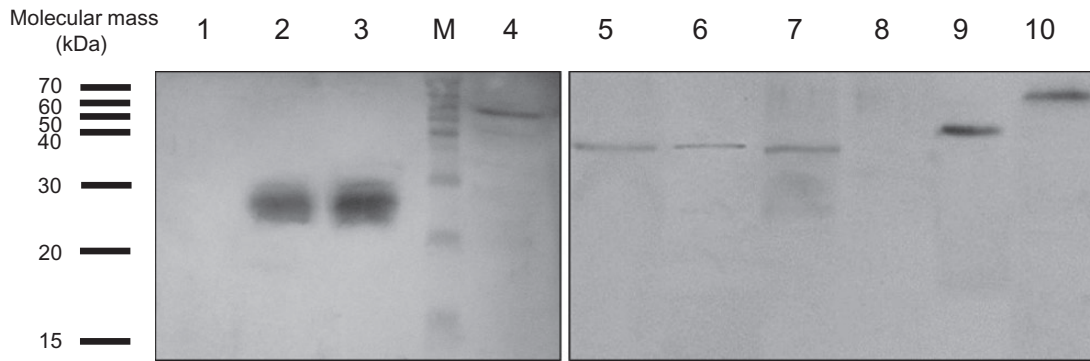


Fig. 4. Western blot of phytochelatin synthases (PCS) from bryophytes and one lycophyte species. A polyclonal antibody raised against *Arabidopsis thaliana* PCS1 was employed. From left to right: (1) *Nostoc* sp. (Cyanobacteria); (2) ~28kDa PCS, sporophytes of *Phaeoceros laevis* (hornwort); (3) ~28kDa PCS, gametophytes of *Phaeoceros laevis* (hornwort); M, molecular mass marker (Pink Prestained Protein Ladder, range 15–75kDa, Nippon Genetics Europe GmbH, Düren, Germany), merged with the autoradiographic membrane; (4) recombinant ~56kDa PCS1, *A. thaliana* (AtPCS1, 50 ng, as an internal reference); (5) ~36kDa PCS, *Conocephalum conicum* (liverwort); (6) ~36kDa PCS, *Scapania undulata* (liverwort); (7) ~36kDa PCS, *Sphagnum palustre* (moss); (8) *Physcomitrella patens* (moss, negative control); (9) ~40kDa PCS, *Selaginella denticulata* roots (lycophyte); (10) ~58kDa PCS, *A. thaliana* roots (positive control).

Table 2. In vitro activity of phytochelatin synthase (PCS) in extracts from bryophytes (gametophytes) and lycophytes (roots), treated with 100 μ M Cd for 90 min at 35 °C in the proper reaction mixture (see the Materials and methods).

PCS activity (mean \pm SE) is expressed in pmol PCs $\text{min}^{-1} \text{g}^{-1}$ FW. For each species, the percentage of PCS activity relative to that of *Arabidopsis thaliana* (=1018.7 \pm 318.5 pmol PCs $\text{min}^{-1} \text{g}^{-1}$ FW, fixed equal to 100%) is also given, assayed in identical conditions. $n=3$.

	PCS activity (pmol PCs $\text{min}^{-1} \text{g}^{-1}$ FW)	Relative PCS activity (% of <i>A. thaliana</i> PCS activity)
Liverworts		
<i>Conocephalum conicum</i>	52.2 \pm 10.6	5.1
<i>Pellia epiphylla</i>	77.7 \pm 10.0	7.6
<i>Radula complanata</i>	28.8 \pm 4.5	2.8
<i>Aneura pinguis</i>	96.1 \pm 12.7	9.4
<i>Scapania undulata</i>	59.6 \pm 7.6	5.8
Mosses		
<i>Sphagnum palustre</i>	10.6 \pm 0.7	1.0
<i>Polytrichastrum formosum</i>	0	0
<i>Fontinalis antipyretica</i>	0	0
Hornworts		
<i>Phaeoceros laevis</i>	15.9 \pm 0.6	1.6
Lycophytes		
<i>Huperzia selago</i>	133.7 \pm 19.7	13.1
<i>Selaginella denticulata</i>	102.3 \pm 48.9	10.0

Abbreviations: ESI-MS, electrospray ionization mass spectrometry; GSH, reduced glutathione; MS/MS, tandem mass spectrometry; Mya, million years ago; PC, phytochelatin; PCS, phytochelatin synthase.

even more) than the amount quantified in several angiosperms (*Magnoliophyta*), grown and tested in very similar experimental conditions by our group and colleagues (Paradiso et al., 2008; Brunetti et al., 2011; Vurro et al., 2011; Sanità di Toppi et al., 2012). Accordingly, it seems reasonable to postulate that the lower amount of PCs may be related, at least in general, to the lower basal PCS activity of land plants, compared for instance with that of

A. thaliana (Table 2). A little more speculatively, it is also proposed that, from the lighter and less active (than that of *A. thaliana*) ‘early’ PCS of basal land plants, an evolutionarily driven trend has led to dimensional increases of the enzyme length and an improvement of its functionality, found in angiosperms. To this end, it is hypothesized that such dimensional variations might be mainly ascribed to increases in the length of the C-terminal domain (as detailed by Rea, 2012). It has in fact been proven that the *A. thaliana* truncated PCS1 (shorter than the full-length enzyme, since it was obtained after partial proteolysis of the C-terminal domain) possesses both a substantially decreased thermal stability and a significant decline in supporting a broad response to metal ions, particularly to Zn and Hg (Ruotolo et al., 2004); it has also recently been postulated that the C-terminal domain plays a role in protecting against the metal-induced oxidative damage possibly occurring in the N-terminal domain (Matsumoto et al., 2009). Land plants might have encountered, amongst other factors: (i) the need to counteract rapid changes in temperature, particularly high temperature excesses (Rensing et al., 2008; Banks et al., 2011) compared with the much more ‘buffered’ environment provided by water; (ii) the exigency of regulating the fluxes of metals in palaeo-soils; and (iii) the need to better protect themselves against the light-induced oxidative stress. Consequently, a longer PCS (with a better thermal stability and an improved response to metal ions and to oxidative agents) might potentially have been promoted in the course of evolution, up to the achievement of the full-length dimension of the angiosperm PCSs.

As schematically depicted in Fig. 5, the present data demonstrate that the capability for PC production under metal stress and the presence of a constitutive and functional PCS enzyme actually represent ancestral (plesiomorphic) characters. They occur not only in the dominant gametophyte generation of bryophytes (and in hornwort sporophytes), but also in the dominant sporophyte generation of basal tracheophytes. The present experiments also suggest that the capability for PC production under metal stress was, in some cases,

probably independently lost during evolution, as supported by the absence of a putative functional PCS and a detectable PC synthesis in the investigated mosses, with the exception of *S. palustre*. In this context, it would therefore appear coherent that the ‘model’ moss *P. patens* (Bryopsida) has been proven not to possess the gene for the PCS (Kopriva *et al.*, 2007), and this might also be the same for some other species investigated here. At the same time, the apparent discrepancy between the lack of *in vivo* detectable PC production in some species (i.e., *S. undulata*, *A. pinguis*, and *H. selago*) and the demonstrated activity/presence in them of the PCS enzyme might possibly be due to morphological/ultrastructural characteristics (i.e. cell wall thickness, mucilage canals, etc.) that could reduce or prevent Cd from reaching the cytosolic PCS in amounts sufficient for inducing an *in vivo* detectable synthesis of PCs.

Regarding specifically the hornworts, in *P. laevis* (game-topophytes and sporophytes) a PCS enzyme with an apparent molecular mass of ~28 kDa was immunodetected. Considering its molecular mass, the *P. laevis* PCS would theoretically be compatible in size with a cyanobacterial PCS-like enzyme (Harada *et al.*, 2004; Tsuji *et al.*, 2004; Vivares *et al.*, 2005). As a consequence, from an evolutionary standpoint, an ancestral horizontal transfer of the PCS from an archaic (*Nostoc*-like?) cyanobacterium to an early hornwort might not be excluded in principle, with the subsequent further full acquisition, only in the hornwort, of a PC synthesis capability, since cyanobacteria do not produce PCs (Harada *et al.*, 2004; Vivares *et al.*, 2005), or at the most they synthesize traces of PC₂ alone (Tsuji *et al.*, 2004). In contrast, a direct immunodetection in *P. laevis* of a ‘contaminant’ PCS-like *Nostoc* sp. band should be ruled out completely, as: (i) the hornwort was thoroughly cleaned of all biological (endo)contaminants under two types of microscopes; (ii) *P. laevis* produced PCs up to PC₃; and (iii) a PCS band of the same size was also detected in the hornwort sporophytes, not colonized by *Nostoc* sp. (Ligrone *et al.*, 2012). In any case, in the *Nostoc* sp. monocultures set

up specifically for this purpose, no PCS-like band was ever detected by the *A. thaliana* PCS1 polyclonal antibody used here (Fig. 4).

The present results also demonstrated that PCs were synthesized in charophytes (sisters of land plants), but, unexpectedly, no putative PCS bands were detected by western blots in these algae. This might be attributed to the nature of the antibody used which probably fails to recognize cyanobacterial/algal PCS epitope(s), instead recognizing only those of (early land) plant. In any case, the present study extended the investigation by Volland *et al.* (2013) carried out on the desmid *M. denticulata*, demonstrating that an unequivocal PC production capability is also a defining character of three further charophyte taxa, namely *Spirogyra* sp., *C. vulgaris*, and *C. scutata*, evolutionarily closer to land plants than *M. denticulata* (McCourt *et al.*, 2004; Qiu *et al.*, 2006; Qiu, 2008; Becker and Marin, 2009; Wodniok *et al.*, 2011).

In conclusion, assuming that early bryophytes and lycophytes spread in palaeo-environments rich in PC-inducing metal(loid)s (Meharg and Hartley-Whitaker, 2002; Rice *et al.*, 2005; Ernst *et al.*, 2008), then the synthesis of PCs and the presence of constitutive and functional PCS enzymes, particularly in the lineages originated during the Devonian period (Fig. 5), might represent a remnant of that time. Due to their ‘benefits’, these traits have been retained up till now, perhaps with the main aim of controlling Zn [and perhaps copper (Cu)] homeostasis (Thumann *et al.*, 1991; Tennstedt *et al.*, 2009; Vurro *et al.*, 2011), other than performing Cd and As detoxification.

Supplementary data

Supplementary data are available at *JXB* online.

Figure S1. ESI-MS chromatograms of: *Chara vulgaris* (A) GSH in controls and (B) GSH, PC₂, PC₃, and PC₄ in Cd-exposed plants; *Coleochaete scutata* (C) GSH in controls

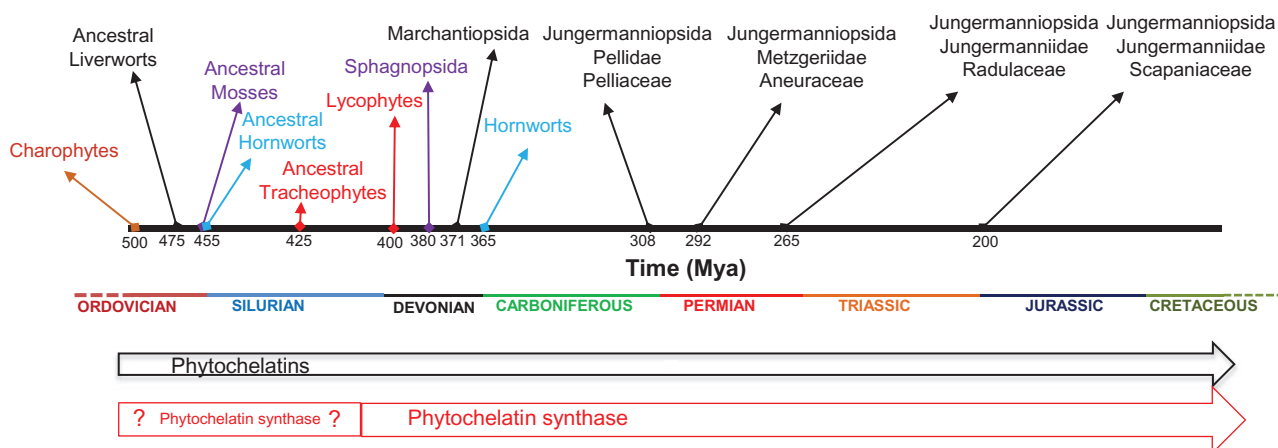


Fig. 5. Key timing events regarding charophyte and basal land plant origin and diffusion, related in particular to the lineages investigated here (chronological data are from McCourt *et al.*, 2004; Qiu *et al.*, 2006; Qiu, 2008; Becker and Marin, 2009; Crandall-Stotler *et al.*, 2009; Wikström *et al.*, 2009; Villarreal *et al.*, 2010; Banks *et al.*, 2011; Ligrone *et al.*, 2012). The postulated ancestral presence of constitutive and functional phytochelatin synthases and the occurrence of a phytochelatin synthesis capability are also indicated by the arrows. (This figure is available in colour at *JXB* online.)

and (D) GSH, PC₂, PC₃, and PC₄ in Cd-exposed plants. Unlabelled peaks are non-specific and do not represent signals for thiol-peptide characterization.

Figure S2. ESI-MS chromatograms of: *Pellia epiphylla* (A) GSH in controls and (B) GSH and PC₂ in Cd-exposed plants; *Selaginella denticulata* (C) GSH in controls and (D) GSH, PC₂, and PC₃ in Cd-exposed plants. Unlabelled peaks are non-specific and do not represent signals for thiol-peptide characterization.

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