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On the dual nature of lichen-induced rock surface weathering in contrasting micro-environments

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Abstract

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Contradictory evidence from biogeomorphological studies has increased the debate on the extent of lichen contribution to differential rock surface weathering in both natural and cultural settings. This study, undertaken in Côa Valley Archaeological Park, aimed at evaluating the effect of rock surface orientation on the weathering ability of dominant lichens. Hyphal penetration and oxalate formation at the lichen-rock interface were evaluated as proxies of physical and chemical weathering, respectively. A new protocol of pixel-based supervised image classification for the analysis of periodic acid-Schiff stained cross-sections of colonized schist revealed that hyphal spread of individual species was not influenced by surface orientation. However, hyphal spread was significantly higher in species dominant on north-west facing surfaces. An apparently opposite effect was noticed in terms of calcium oxalate accumulation at the lichen-rock interface, detected by Raman spectroscopy and complementary X-ray microdiffraction on south-east facing surfaces only. These results suggest that lichen-induced physical weathering may be most severe on north-west facing surfaces by means of an indirect effect of surface orientation on species abundance, and thus dependent on the species, whereas lichen-induced chemical weathering is apparently higher on south-east facing surfaces and dependent on micro-environmental conditions, giving only weak support to the hypothesis that lichens are responsible for the currently observed pattern of rock-art distribution in Côa Valley. Assumptions about the drivers of open-air rock-art distribution patterns elsewhere should also consider the micro-environmental controls of licheninduced weathering, to avoid biased measures of lichen contribution to rock-art deterioration.

Keywords

Biodeterioration, Biogeochemistry, Biomineralization, Raman spectroscopy, XRMD, Image
 analysis, Schist

Introduction

The last three decades have been extremely rich in contributions to the knowledge of the various aspects of lichen-induced rock weathering, as seen by the number of reviews available (Adamo & Violante 2000, Chen et al. 2000, Seaward 2015, St. Clair & Seaward 2004). Alternative approaches to this subject have been focusing on: i) identifying individual species and species assemblages colonizing rock surfaces and making assumptions on their impact based on previous knowledge about their ecological requirements (e.g. Carballal et al. 2001); ii) determining the climatic constraints and habitat preferences of colonizing species based on field observations (e.g. Steinbauer et al. 2013, Viles & Cutler 2012) or controlled experiments (Adamson et al. 2013, Carter & Viles 2003, Kidron & Termina 2010) thus contributing to the knowledge of environmental factors that are also important for rock conservation; iii) detecting geophysical and geochemical changes at the lichen-rock interface associated with the growth of individual species including the occurrence of organic and mineral by-products of lichen activity (e.g. Arocena et al. 2007, Favero-Longo et al. 2005); iv) addressing the influence of human activities on such changes (e.g. Cámara et al. 2015) and v) developing methods to quantify the weathering rates induced by individual species or by a limited set of the most representative ones on the

- 69 surface of interest (e.g. Aghamiri & Schwartzman 2002, Bartoli et al. 2014, Gazzano et al.
- 70 2009a, b, Mcllroy de la Rosa et al. 2014).
- 71 The majority of work has been applied at characterizing the biodeterioration of a range of
- stonework in Europe and only sporadically in other regions of the globe. Few papers have dealt
- 73 with the relationship between lichen growth and the weathering of schist (Aghamiri &
- 74 Schwartzman 2002, Cann et al. 2012, Galvan et al. 1981, Fry 1924, 1927, Sanders et al. 1994)
- despite the increasing demand of schist as a building stone.
- 76 The processes of schist weathering, including those induced by lichen activity, are a primary
- concern in the Côa Valley Archaeological Park (Vila Nova de Foz Côa, north-east Portugal)
- 78 where one of the world's most important sets of Prehistoric rock-art is located, almost invariably
- on schistose surfaces. Efforts are being made there to understand the weathering dynamics acting
- on the schist outcrops that support the rock-art, integrating biological, geophysico-chemical and
- 81 environmental data in order to prevent major damages to the engraved surfaces (Aubry et al.
- 82 2012). There is general consensus about the combination of physical (mechanical) and chemical
- changes brought about by lichens to rock surfaces but the extent and relative contribution of their
- 84 weathering action is a fundamental, yet still unanswered question, both in the Côa Valley and in
- 85 the field of rock-art conservation in general. Adding to the debate over the threats of lichen-
- 86 induced processes is the contradictory evidence for lichen protection of rock surfaces against
- other deteriogenic agents (Carter & Viles 2005, McIlroy de la Rosa et al. 2013).
- 88 Recently, Aubry et al. (2012) suggested that aspect-related differences in the extent of lichen
- 89 (and bryophyte) colonization of vertical schist surfaces in the Côa Valley could be partly
- 90 responsible for the differential weathering of those surfaces and resulting pattern of rock-art
- 91 distribution in the Côa Valley, which is currently more concentrated at south-east facing than at

north-west facing slopes (Fernandes 2012). Provided that the interaction between lichens and the rock surface is species-dependent (e.g. Favero-Longo et al. 2005), a key uncertainty in the assumptions on the relationship between lichen colonization and rock weathering is precisely in the way that lichen species act under different weathering environments. Studies aimed at evaluating the response of individual species to changes in environmental conditions have typically demonstrated a shift in oxalate production (thus in the biodeteriorative action) in response to environmental variation (Caneva 1993, Edwards et al. 1995, Prieto et al. 2000). The extreme environments of cold and hot deserts have been particularly interesting for research (e.g. Wierzchos et al. 2013) since, besides acting differently in distinct weathering contexts, species are also expected to change their performance under the influence of environmental change. However, although useful first approximations, the existing studies commonly assume species neutrality, with substrate and climate as the primary controls. As a result, knowledge about lichen-induced rock weathering is mostly based on static views of the influence of environmental, geological or climatic parameters such as rock porosity, permeability and mechanical properties, or temperature, solar radiation and humidity. The variation of the diversity and composition of lichen assemblages with rock surface orientation is a well-known phenomenon in both natural and cultural contexts (Adamson et al. 2013) as orientation acts as a proxy of those environmental variables, mainly temperature and humidity, which affect the structure and dynamics of saxicolous lichen communities. Some components of that variation in the Côa Valley have been previously addressed from an ecological perspective (Marques et al. 2014). The present study directly assessed the effect of rock surface orientation on lichen-induced physical and chemical weathering in order to determine the contribution of lichen action to

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differential rock weathering in the Côa Valley Archaeological Park. Hyphal penetration and calcium oxalate formation at the lichen-rock interface were confidently assumed as proxies for physical and chemical weathering, respectively. Although how much hyphae penetrate through existing pores and fissures or actively produce discontinuities in the rocks is not definitely clear, microscopical observations have shown that penetrating hyphae induce mineral breakage and rock surface disaggregation by mechanical action (Ascaso & Wierzchos 1995, Favero-Longo et al. 2005). The close relationship between the chemical composition of colonized rocks and the type of oxalates accumulating immediately beneath or within the thalli of some lichen species have generally indicated an involvement of the mycobiont-secreted oxalic acid in lichen-induced chemical weathering of rock substrata (Adamo & Violante 2000 with refs. therein).

Material and methods

The study area

The Côa Valley Archaeological Park is a UNESCO world heritage site located in the confluence of River Côa with River Douro, approximately 200 km upstream of the mouth of the Douro, in the city of Porto (Portugal). Lithology in this part of the Côa Valley is dominated by metasedimentary rocks of the schist-greywacke complex ranging in age from the Precambrian to the Ordovician. River Côa and its tributary streams have cut deeply through the schist and greywacke basement, taking advantage of pre-existing major faults roughly NE-SW oriented (Aubry et al. 2012) and forming numerous steep-walled valleys that play a major influence on regional landscape. A special feature resulting from the down-cutting of the Côa Valley is the occurrence of massive vertical schist surfaces arranged in layers along the valley's slopes, which

have been gradually exposed by rock toppling, i.e. a sequence of gravity-induced block displacement after splitting of vertically orientated joints, along the schistosity plane. Climate is predominantly dry meso-Mediterranean, sheltered from the Atlantic influence by mountains to the north and the west, but thermo-Mediterranean microclimates are usually produced in the bottom of the valleys, where temperature is frequently above 40°C in late spring and summer (June-August), daily thermal amplitudes may reach 10° to 15° C and mean annual precipitation is often below 300 mm (Fernandes 2012).

Geological and mineralogical characterization of the studied lithotype

The target lithotype is a relatively low-grade metamorphic (greenshist facies) phyllite consisting of thin alternating layers of whitish psammitic and dark pelitic components (Aires et al. 2011, Búrcio 2004, Sousa 1982). The psammitic component is sometimes more abundant and the rock is then classified as a metaquartzwacke instead of a phyllite. For a matter of simplicity, this phyllite-metaquartzwacke sequence will be addressed under the broad term schist. This schist is mainly composed of quartz, sericite and/or muscovite, chlorite and biotite minerals, as well as plagioclase feldspars (mostly albite) in variable amounts depending of the psammitic contribution. Calcite is usually present in the matrix of these rocks in sufficient amounts to produce effervescence when treated with dilute hydrochloric acid. Magnetite and, more sporadically, pyrite crystals are present in both the psammitic and pelitic levels as accessory constituents (Sousa 1982). Additional accessory minerals including illite, kaolinite, montmorillonite, graphite, turmaline, zircon, apatite, epidote, hematite, leucoxene and alkali feldspars, such as microcline and orthoclase, were detected by polarized light (petrographic)

microscopy and X-ray diffraction (Aires et al. 2011, Búrcio 2004, Gomes & Almeida 2003, Sousa 1982). The general strike of the target fracture/joint surfaces is NE-SW, which is subparallel to the NNE-SSW sinistral strike-slip fault system that crosses the study area and formed by the same tectonic stress (Aubry et al. 2012). The plane of schistosity is consistently vertical and oriented subperpendicular to the fracture/joint surfaces.

Microclimatic characterization of vertical schist surfaces in the Côa Valley

To characterize the thermal and hydric contrasts of opposite slopes in the Côa Valley, Hygrochron iButton data-loggers (Maxim Integrated Products Inc., Sunnyvale (CA), USA) were placed on 12 vertical schist surfaces of varying orientations. The results were then grouped in the two aspect classes of interest for this study: north-west (NW) and south-east (SE). Data-loggers were synchronized and set to record both temperature (°C) and relative humidity (%), at hourly or half-hourly intervals during a 3-year period from late September 2010 to late September 2013.

Target species

The study dealt with the physical and chemical weathering driven by four locally-common lichens: *Aspicilia contorta* subsp. *hoffmanniana* S. Ekman & Fröberg (*Aspicilia hoffmanniana* hereafter), *Caloplaca subsoluta* (Nyl.) Zahlbr., *Lecanora pseudistera* Nyl. and *Peltula euploca* (Ach.) Poelt. Taxa selection was based on higher frequency and abundance (cover) on the vertical schist surfaces of the Côa Valley Archaeological Park (Marques et al. 2014). *Aspicilia hoffmanniana* is a crustose lichen varying in colour from greenish-grey in shaded NW facing

surfaces to light brown in exposed SE facing surfaces, where it is more abundant. Caloplaca subsoluta is a deep orange coloured crustose species that is common on siliceous rocks throughout the Mediterranean. In the Côa Valley it was found equally abundant on the vertical schist surfaces of the two opposing slopes. Lecanora pseudistera is a white crustose species proliferating on NW facing surfaces, although it can also be found less abundantly on SE facing schist surfaces. Peltula euploca is a widespread squamulose epilith characteristic of the raintrack communities of vertical schist surfaces and exclusive of SE facing exposures (Marques et al. 2014). Each squamule is attached to the substrate by a central umbilicus and its lower surface is pale to dark brown. The algal partner is *Trebouxia* spp. in the three crustose lichens and unicellular cyanobacteria (*Chroococcidiopsis* spp.) in *Peltula euploca*. The three crustose lichens are able to reproduce sexually, although Aspicilia hoffmanniana is most frequently sterile. Peltula euploca is often fertile in the study area but its characteristic mode of dispersal is through vegetative propagules (soredia). Secondary compounds in Aspicilia hoffmanniana are either lacking or include only aspicilin. Caloplaca subsoluta produces anthraquinones. The major secondary metabolites produced by *Lecanora pseudistera* are atranorin and 2'-O-methylperlatolic acid. Peltula euploca lacks secondary metabolites.

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Sampling strategy and in-field sample collection

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Colonized rock samples, with lichen thallus kept intact, were taken at random from non-engraved NW and SE facing schist surfaces, located at representative rock-art sites, namely Canada do Amendoal, Foz do Côa, Quinta das Tulhas, Vale do Forno and Vale de José Esteves, and are therefore very similar to the surfaces bearing rock-art in terms of their macro- and micro-

environmental constraints. Since *Peltula euploca* is virtually exclusive of SE facing surfaces, appropriate samples of rock colonized by this particular species could only be found and collected at SE. Uncolonized rock samples were also taken from the source outcrops to be used as controls. The use of bare-rock controls is a necessary condition for isolating the licheninduced effects from the ones induced by other weathering agents (either biotic or abiotic). The collected samples were cut perpendicular to the colonized or previously exposed surface, up to a depth of 3 to 4 cm and width of 5 cm, with an Isomet 1000 Precision Saw (Buehler, Düsseldorf, Germany). The surfaces of the resulting cross-sections were polished with sandpaper attached to an Ecomet 3000 (Buehler, Düsseldorf, Germany) polisher machine. Nine replicates were prepared for each combination of species vs orientation, and respective control (uncolonized) in order to evaluate if weathering associated with lichen colonization differs from the weathering produced on identical but lichen-free surfaces. Three subsets of three replicates each were taken from the initial sample set, and processed accordingly for Periodic acid-Schiff (PAS) staining, X-ray microdiffraction and FT-Raman spectroscopy. Almost all samples had been included in polyester resin (*Recapoli* 2196 styrene and phthalic anhydride, Methyl Ethyl Ketone peroxide as catalyst) before cutting, to avoid excessive loss of material due to the fragile nature of the schist samples, except for those used in Raman analysis, since resin inclusion would preclude from taking measures directly on the lichen thallus.

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Periodic acid-Schiff staining

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This procedure aimed at highlightingthe Hyphal Penetration Component (HPC) of the target lichen species, as a proxy of lichen-induced physical weathering, following Favero-Longo *et al.*

(2005). Microphotographs of the stained cross-sections were then acquired at a ×10 magnification using a Nikon SMZ1000 stereomicroscope equipped with a Nikon DS Fi1 digital camera, at three random locations of each cross-section. Data on the depth of hyphal penetration was obtained by visual inspection of all hyphal bundles highlighted in the stained cross-sections under the same stereomicroscope. The mean and the maximum of measured data were calculated for each species at different aspects to provide the average and maximum depth of hyphal penetration sensu Favero-Longo et al. (2011). In order to quantify the extent of hyphal penetration, as well as the size of other important weathering-related features (e.g. weathering rind), the acquired images were submitted to a new protocol of pixel-based supervised classification using colour and texture features, which can be described briefly as follows: 1) Image pre-processing in ImageJ (http://imagej.nih.gov/ij/), including the resize of original images to 40% of the initial size using bilinear resampling (to increase computation speed and efficiency) and contrast and sharpening enhancement to allow a better discrimination of the HPC and weathering rind (when present), from the rock core; 2) Feature extraction, including a total of 162 texture features based on run-length, co-occurrence, image histogram and gradient matrices in MaZda (http://www.eletel.p.lodz.pl/programy/mazda/) and colour features based on several colour spaces namely RGB, HIS, YUV, YIQ and XYZ using 'adimpro' package (Polzehl & Tabelow 2007) in R; 3) Generation of training input data in ImageJ, by manually assigning points (i.e., XY coordinates of pixels) in the original images to the corresponding structures of interest, namely 'lichen thallus', 'hyphae', 'weathering rind' and 'rock core'; 4) Combination of the extracted features and training data, to calibrate a Random Forest (RF) classifier (Breiman 2001, Liaw & Wiener 2002) in R with 'ntree'= 200, 'mtry'= 6, 'nodesize' = 5 and the remaining parameters kept as default; the feature set included only a

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reduced subset containing the 20 top colour and texture features according to their relative predictive importance (Boulesteix et al. 2012, Oppel et al. 2009); 5) Evaluation of classifier's performance through Monte-Carlo cross-validation (Xu et al. 2004) with 70% for training and 30% for testing and a total of 100 replicates; 6) Prediction of the labels of the structures of interest for the whole image, using the RF classifier with highest overall test accuracy. The extent of the HPC and weathering rind (given in mm²) in each sample are among the descriptive statistics retrieved by the classifier.

FT-Raman spectroscopy

Spectroscopic analyses were performed in three locations of each colonized schist sample: 1) the upper layer (cortex) of the lichen thalli as in Prieto et al. (2000); 2) the surface of the rock in contact with the lichen thallus, named lichen-rock interface (Ascaso et al. 1976), where mineral neoformation, if taking place at all, is most likely to be due to lichen-rock interactions; and 3) the rock interior (at least 2 cm away from the surface) which is used as a specific control for each sample since lichen-induced oxalate formation is assumed not to reach such deeper areas beneath the lichen-rock interface (Adamo & Violante 2000). Control FT-Raman spectra were also recorded on non-colonized schist samples collected from the same schist outcrops, including 1) the exposed surface (taken from the top as in colonized samples); 2) a 5 mm deep virtual interface; and 3) the rock interior. A Bruker RFS 100/S FT-Raman spectrometer was used with a Nd:YAG laser operating at 1064 nm as the excitation light source and a resolution of 4 cm⁻¹. Spectral data were acquired after 1024 laser scans of 20 mW in lichen thallus, to minimize lichen damage, and 64 laser scans of 250 mW in rock interior and lichen-rock interface.

X-ray microdiffraction (XRMD)

XRMD measurements were performed on an Empyrean (PANalytical, Almelo, The Netherlands) diffractometer, equipped with a five-axis Chi-Phi-x-y-z stage goniometer, a copper sealed anticathode X-ray tube (*Empyrean Tube Cu Lff Hr*) and a PIXcel^{3D} (PANalytical, Almelo, The Netherlands) X-ray detector. Each sample was first submitted to a set of five random line scan readings along the polished cross-section in order to obtain a preliminary depth profile of the mineralogical composition of the studied schist. Yet no significant differences were detected between the surface of the sample and its interior so priority was given to the lichen-rock interface as in FT-Raman spectroscopy. The final measurements are based on ten random readings along the lichen-rock interface. Bragg angles were scanned between 4 and 47°, with steps of 0.02° (12 minutes time length) for exploratory measurements and between 3.5 and 60°, with steps of 0.02° (2 hours time length) for the final measurements, with a laser diameter of 0.6 mm.

Statistical analysis

The effects of species and orientation on the depth and extent of the Hyphal Penetration Component (HPC) was tested by means of two-way analysis of variance (ANOVA) and post-hoc Tukey HSD for pair-wise comparisons with 'agricolae' package (de Mendiburu 2013) in R. ANOVA's assumption of normality of residuals was checked graphically using qq-plots and that of homoscedasticity tested by means of the Levene test, also in R.

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Results and discussion

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Lichen-induced physical weathering

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The pattern of hyphal penetration into the rock substrate is highly influenced by the characteristics of the rock (Sanders et al. 1994) since hyphae tend to follow paths of least resistance. Depth of hyphal penetration of the four lichens analysed was on average 3.4 mm in samples coming from NW facing surfaces (n= 12), 4.6 mm in samples coming from SE facing surfaces (n= 9), and varied between 0.1 to 37 mm when considering all samples. Maximum depth of hyphal penetration is apparently higher at SE than at NW facing surfaces (Table 1). However, within sample variation, as depicted by the standard-deviation, was extremely high and there was no statistically significant effect of orientation or species in the depth of hyphal penetration. Additionally, although maximum depth of hyphal penetration detected among the analysed samples was 37 mm (in Aspicilia hoffmanniana from SE facing surfaces), hyphal bundles frequently reached the lower extremity of the samples, which were roughly between 30 and 40 mm long, and measure of real maximum depth of penetration could not be accurately estimated. As suggested by previous works (e.g. Favero-Longo et al. 2005, Wierzchos & Ascaso 1996, 1998) schistose rocks may be susceptible to greater lichen-induced weathering than igneous rocks due to their higher predisposition to break and the easier progression of hyphae along the planes of weakness, parallel to the schistosity (Fig. 1). Additional planes of weakness come from the intense tectonically-related fracturing that characterizes the studied rock. One feature observed in almost every sample, and also commonly referred in literature (e.g. Fry

1927), is the extremely penetrative type of hyphal bundles associated with the occurrence of apothecia, which are able to reach much deeper into the schist than adjacent hyphae (Fig. 1). Analysis of the effects of penetrating hyphae have long been relying on the use of scanning electron microscopy (SEM) and other high-resolution laboratory techniques (e.g. Jones et al. 1981, Strech & Viles 2002, Wierzchos & Ascaso 1994). Although extremely useful for examining the very specific changes occurring in rock minerals by direct contact with individual hyphae, the scale of these approaches is often too small to answer questions related with the performance of the entire lichen or lichen community at the scale of the whole surface (microscale) or site where these surfaces are located (meso-scale). Additionally, these techniques may only be considering worst (or best) case scenarios resulting from chance unless they are based on large sample sizes and use full random sampling as in Strech & Viles (2002) to achieve statistical robustness. Many studies have determined the maximum or average depth of hyphal penetration for a wide range of lithotypes, but usually ignore hyphal spread, with few exceptions (Bartoli et al. 2014, Casanova-Municchia et al. 2014, Gazzano et al. 2009a). Image analysis of colonized crosssections after PAS staining retrieved accurate values for the hyphal spread of each species inside the rock (Table 1; Fig. 1). These values could then be used to determine if surface orientation had any effect on species ability to spread into the rock interior and induce mineral breakdown.

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[Fig. 1 approximately here]

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The studied lichens differed significantly in the spread of the HPC irrespective of surface orientation (F-value= 10.974, p-value< 0.001). These differences were caused by a significantly

higher hyphal spread in *Lecanora pseudistera* than in any other of the lichens analysed, whereas *Aspicilia hoffmanniana*, *Caloplaca subsoluta* and *Peltula euploca* were very similar among each other (Table 1). The effect of interaction between species and surface orientation was not statistically significant (F-value= 0.867, p-value= 0.364) neither was the effect of orientation itself (F-value= 1.339, p-value= 0.262), which means that even if there were differences between species in terms of their ability to spread inside the rock, there are no statistically significant differences between NW and SE oriented surfaces in any of the studied species.

Weathering is quite obvious to the naked eye in most samples, where the black to more usually dark-grey rock core corresponding to unweathered parent rock gradually changes into the light brown to greyish-brown coloured weathering rind (Fig. 1b). All samples from NW facing surfaces showed such features, against only 13% of the samples from SE facing surfaces. The dimension of the weathering rind was unrelated with the extent of the HPC (Table 1: Spearman=-0.012).

[Table 1 approximately here]

Although the extent of the HPC is unrelated to surface orientation, as species differ in their ability to spread inside the rock, an indirect effect of orientation on lichen-induced physical weathering, as depicted by the proxy HPC, could be produced by means of the abundance of individual species. Since *Lecanora pseudistera* is the most effective in terms of hyphal spread among the four lichens analysed (Table 1), and having been the most frequent and abundant on NW facing slopes (Marques *et al.* 2014) one might assume a more intensive lichen-induced

physical weathering happening at NW than at SE facing surfaces in the Côa Valley. Among the four lichens analysed, *Peltula euploca* was probably the most surprising in terms of physical performance since the attachment of this peltate (shield-like) growth form is far from being limited to superficial layers, penetrating up to 3.9 mm into the rock.

Lichen-induced chemical weathering

FT-Raman and complementary X-ray microdiffraction analyses of bare-rock controls confirmed the presence of quartz, chlorite, muscovite and albite as the major minerals in all analysed locations of the presumably unweathered schist (Fig. 2 and Supporting Information Tables S1 and S2).

[Fig. 2 approximately here]

information about metabolic by-products of the lichen-induced weathering process (Edwards *et al.* 1995, Jorge-Villar *et al.* 2004).

Phyllosilicates and quartz are the strongest features in the internal layers of all lichen-colonized samples (Fig. 2), corresponding well with the presumably unweathered parent rock type.

Phyllosilicates have complex structures and highly variable compositions (Wang *et al.* 2002) reflected in complex FT-Raman spectra, but for the purpose of this paper, the occurrence of phyllosilicates can be usefully discussed in terms of the 100-600 cm⁻¹ spectral region, where, contrary to what happens in other regions of the spectra, peaks related to phyllosilicates are

The wavenumber region 100-1700 cm⁻¹ of FT-Raman spectra contains useful spectroscopic

easier to differentiate from those related to other substances (see below). According to Wang et al. (2002) di-octahedral phyllosilicates such as muscovite produce a strong FT-Raman peak at 260 cm⁻¹, which is depicted in the FT-Raman spectra of the rock interior and lichen-rock interface of almost all colonized samples. Mg-bearing phyllosilicates, such as chlorite, peak strongly at approximately 350 cm⁻¹. The peak at 356 cm⁻¹ visible in the FT-Raman spectra of the rock interior and lichen-rock interface of schist samples colonized by Aspicilia hoffmanniana (Fig. 2), can therefore be assigned to chlorite. Peaks at 200 cm⁻¹ are characteristic of trioctahedral phyllosilicates (Wang et al. 2002) and most probably indicate the presence of either chlorite or biotite. Quartz is very resistant to weathering and persists even at the surface of bare rock and at the lichen-rock interface of all samples (Fig. 2). It was detected in every sample by a sharp band at 464 cm⁻¹ in FT-Raman spectra and at 3.33 Å in X-ray microdiffraction (Supporting Information Tables S1 to S10). The presence of the same band in the FT-Raman spectra of Aspicilia hoffmanniana and Caloplaca subsoluta (Fig. 2a) indicates an incorporation of quartz particles by the thallus of these lichens. Evidence for the ability of Aspicilia hoffmanniana to incorporate phyllosilicate particles is also seen in its FT-Raman spectra, with characteristic bands at 200 and 260 cm⁻¹ (Wang et al. 2002). However, the occurrence of quartz and phyllosilicates in lichen thalli and lichen-rock interfaces can also have an exogenous origin from airborne dust, as suggested by Vingiani et al. (2013) after detecting the same kind of mineral incorporation in lichens growing on quartz- and phyllosilicate-free vulcanic rocks. Incorporation of quartz and phyllosilicate minerals by lichens is not exclusive of crustose lichens as can be inferred by the FT-Raman spectra of *Peltula euploca* showing a band at 432 cm⁻¹ (Fig 3), which is assignable to either of these silicate minerals. This would require further confirmation through higher resolution techniques such as scanning electron microscopy (SEM).

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Besides those minerals that are known to characterize the unweathered rock (see materials and methods section), X-ray microdiffraction and FT-Raman spectroscopy confirmed the occurrence of halloysite at both the virtual interface of bare rock samples and the lichen-rock interfaces, irrespective of the species and the orientation of the parent outcrop (Figs 2). Hallovsite is a common product of schist weathering, resulting from the transformation of chlorite, biotite, muscovite and feldspars (Banfield & Eggleton 1990, Kretzschmar et al. 1997, Parham 1969). Despite the differences stated above in terms of weathering rind formation, no differences were observed in terms of the occurrence of halloysite between NW and SE facing surfaces. Peak at 260 cm⁻¹ in FT-Raman spectra could also correspond to kaolinite, another product of schist weathering, but differentiation of kaolinite minerals from muscovite, and the latter from vermiculite, can be problematic because peaks shared by the three minerals are not fully differentiable by X-ray microdiffraction or FT-Raman spectroscopy (Wang et al. 2002). FT-Raman and complementary X-ray microdiffraction analyses also allowed detecting the occurrence of neoformation minerals commonly attributed to lichen activity, namely calcium oxalates, at the lichen-rock interface and thalli of some of the target species, and confirmed their absence at the rock interior of all samples. The pattern of occurrence of such minerals is, however, variable among the considered species, their origin and analysed location within samples. Key molecular signatures for oxalates occur in the 1400-1500 cm⁻¹ region of FT-Raman spectra (Edwards et al. 2003b) where 1476 cm⁻¹ is considered distinctive for weddellite (Fig. 2b). Other distinguishing bands for weddellite occur at 912 and 1634 cm⁻¹. The signature of weddellite was found either completely or partially in the thalli of Aspicilia hoffmanniana, Lecanora pseudistera

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and *Peltula euploca* from south-east facing surfaces (Fig 3a), but not in the thalli of *Caloplaca* subsoluta neither in any of the specimens taken from north-west facing surfaces (Fig 2a). In fact, previous experimental works had indicated that microclimatic factors could be important in determining the state of hydration of calcium oxalate in lichens (Edwards et al. 1995) and have associated the occurrence of the dyhidrate form with lichen's strategy for maintaining its water balance in dry exposed surfaces (Prieto et al. 2000, Prieto & Silva 2003). Calcium oxalate monohydrate, known as whewellite, has been detected in the thalli of Aspicilia hoffmanniana and Lecanora pseudistera on samples from NW facing surfaces, peaking in FT-Raman spectra at 1463 and 1631 cm⁻¹, respectively (Fig. 2). Variation in measured temperature and relative humidity between NW and SE facing surfaces is summarized in Table 2. There are similarities in the general pattern of annual rock surface temperature and relative humidity regimes. Both NW and SE facing surfaces exhibited a seasonal pattern of high temperature and low relative humidity values from around June to September followed by a much colder and moist period between November and February. Variation in temperature was huge at both orientations, but nevertheless higher at SE than at NW facing surfaces. The same happened with relative humidity, although relative humidity was always higher at NW facing surfaces than at SE facing surfaces. The occurrence of oxalates in the lichen thallus of Aspicilia hoffmanniana and Lecanora pseudistera is not completely unexpected since the fruiting bodies of the former are well known for being pruinose (i.e. covered by oxalate crystals) and the later belongs to a group of lichens that are characterized precisely by the presence of large amphithecial crystals. The occurrence of the monohydrate form in specimens that grow under less contrasting humidity and temperature regimes is consistent with the physiological role attributed to calcium oxalate.

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The origin of Ca ions for calcium oxalate production in lichens, however, is still an unsolved matter. Calcite (CaCO₃) is usually present in the matrix of the studied rock type in sufficient amounts to produce effervescence when treated with dilute hydrochloric acid. Calcite is highly alterable and a potential source of Ca ions for calcium oxalate formation promoted by biological colonization. However, calcite, with characteristic features in FT-Raman spectra being a strong, sharp band at 1086 and weaker bands at 712 and 286 cm⁻¹ (Edwards et al. 1995, 2003b), is missing in all schist samples. Although it has been proved that lichens are able to uptake Ca ions from calcareous rocks such as limestones and marbles for calcium oxalate production (e.g. Seaward & Edwards 1995), the rock is definitely not the only source of this element as the occurrence of both forms of calcium oxalate has been reported in lichens colonizing non-calcareous substrates such as granites (e.g. Prieto & Silva 2003), serpentinites (Favero-Longo et al. 2005) or even tree-bark (Edwards et al. 2005) and tree leafs (de Oliveira et al. 2002). The presence of calcium oxalates inside the lichen thallus is therefore not necessarily indicative of its biodeteriogenic activity and, when the calcium source is exogenous, calcium oxalate patinas were even indicated as bioprotective (McIlroy de la Rosa et al. 2013). Occurrence of calcium oxalates in lichens growing on iron- and magnesium-rich siliceous rocks instead of the most expected ferrous oxalate dehydrate (humboldtine) and magnesium oxalate (glushinskite), respectively, has also been reported before (e.g. Prieto et al. 1997, 2000, Prieto & Silva 2003, Favero-Longo et al. 2005). This phenomenon can, according to Prieto & Silva (2003) and Favero-Longo et al. (2005), be explained by the higher water solubility of ferrous and magnesium oxalates as well as their higher susceptibility to oxidation. The detection of calcium oxalates at the lichen-rock interface by X-ray microdiffraction (Table 3 and Supporting Information Table S7, S9 and S10), however, is not as

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easily assignable to external sources of Ca. Only weddellite has been detected at the lichen-rock interface and that was by X-ray microdiffraction exclusively on samples taken from SE facing surfaces, colonized by Caloplaca subsoluta, Lecanora pseudistera and Peltula euploca (Table 3). Weddellite was apparently absent from the lichen-rock interface of Aspicilia hoffmanniana although present in the thallus (Table 4). The opposite was observed in the samples colonized by Caloplaca subsoluta, with weddellite detected at the lichen-rock interface by X-ray microdiffraction (Table 3) and no form of calcium oxalate detected inside the thallus by FT-Raman (Table 4). These results suggest that calcium oxalates at the lichen-rock interface and inside the lichen thallus may have different origins and/or functions. Except for the occurrence of weddellite, FT-Raman spectroscopy and X-ray microdiffraction of the lichen-rock interface retrieved quite similar results to that of the virtual interface in bare rock controls. The relevance of these results in the search for the causes of differential weathering of schist surfaces in the Côa Valley is opposite to those already mentioned for lichen-induced physical weathering. Assuming that calcium oxalate at the lichen-rock interface is being produced from Ca taken from the rock and considering its formation as a proxy of lichen-induced chemical weathering, such type of weathering might be more intense on SE facing surfaces. The possibility that due to differences in micro-environmental conditions, calcium oxalates on SE surfaces are subject to less dissolution and are consequently more stable than those on NW surfaces cannot be ruled out by present evidence. However it seems unlikely because the poorly soluble Ca oxalates were found to persist at the lichen-rock interface of thalli exposed to significantly higher precipitation regimes (Favero-Longo et al. 2005). Other bands present in FT-Raman spectra include 1158, 1552 and 1612 cm⁻¹, which are characteristic of parietin and therefore present in the FT-Raman spectra of Caloplaca subsoluta.

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The resemblance of the FT-Raman spectra of *Caloplaca subsoluta* and that of other members of the lichen family Teloschistales (Jorge-Villar *et al.* 2004) is quite obvious mainly due to the profile of parietin.

The FT-Raman spectra of the thallus of *Aspicilia hoffmanniana* and *Lecanora pseudistera* on

The FT-Raman spectra of the thallus of *Aspicilia hoffmanniana* and *Lecanora pseudistera* on samples taken from NW and SE facing surfaces, respectively, contain a series of bands that could not be assigned to any substance for the time being, but are probably related with other secondary metabolites produced by these lichens, which may further act as biodeteriogenic factors (Adamo & Violante 2000, with refs. therein). Also unknown were the peaks at 5.96 and 9.09 Å in X-ray microdiffraction of samples colonized by *Aspicilia hoffmanniana* and 2.32 Å in samples colonized by *Aspicilia hoffmanniana* and *Caloplaca subsoluta* (Supporting Information Table S7 and S9). The very broad bands at 1332 and 1595 cm⁻¹ of all FT-Raman spectra are due to amorphous carbon.

Conclusions

Variation in microclimatic factors related to surface orientation produces different effects depending on the nature of lichen-induced weathering. Analysis of stained polished cross-sections of schist samples colonized by the crustose *Aspicilia hoffmanniana*, *Caloplaca subsoluta*, *Lecanora pseudistera* and the squamulose *Peltula euploca* showed that hyphae originating from the medulla of these lichens penetrate more than 30 mm and follow a unidirectional pattern along the depth rock profile. The lamellar nature of schist minerals offering pathways of least resistance along intermineral voids, probably favours this deep penetration. Therefore the effects of hyphal penetration on schist should go far beyond the

surface and also involve the minerals in the deeper layers. The spread of the hyphal penetration component in the analysed species was similar on NW and SE facing surfaces, but may turn out to be more severe at NW facing surfaces, due to the higher frequency and abundance of species with higher penetrative ability, such as *Lecanora pseudistera*. Orientation is thus likely to have an indirect effect on lichen-induced physical weathering by means of the abundance patterns of individual species, highlighting the importance of accurate estimates of the relative abundances of colonizing species, stemming from community ecology approaches, for rock-art condition assessments. Other evidences of lichen-induced weathering produced in these rocks are related with the incorporation of quartz and phyllosilicate particles by the thalli of all the lichens studied, including the squamulose *Peltula euploca*. The external origin of these particles cannot be ruled out, however, as the mechanisms of airborne mineral incorporation by the lichen thallus are not fully understood. Also of interest for the purpose of this study is the presence of kaolinite and halloysite, two common products of schist weathering, at the lichen-rock interface and on the surface of bare rock controls, irrespective of surface orientation. Variations in the amount of these minerals depending on the colonizing species and microclimatic factors remain to be tested. Evidence for the occurrence of metabolic by-products of lichen activity in the analysed samples is limited to calcium oxalates. Specimens of Aspicilia hoffmanniana from dry SE facing surfaces produced weddellite exclusively, while those from moist NW facing surfaces produced a mixture of weddellite and whewellite. Specimens of Lecanora pseudistera produced weddellite on SE facing surfaces and whewellite on NW facing surfaces. Weddellite was also detected inside the thalli of squamulose *Peltula euploca*, occurring only on SE facing surfaces. None of these forms

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of calcium oxalate were detected inside the thalli of Caloplaca subsoluta. These results indicate, as others beforehand, that there is some preference among the studied lichens for the production of the dehydrate form of calcium oxalate under the highly variable microclimate conditions provided by SE facing surfaces, while on the slightly less variable NW facing surfaces, a mixture of both monohydrate and dehydrate forms can occur. Given the possibility of calcium uptake from airborne particles, it is impossible to state unequivocally whether calcium ions used in the formation of calcium oxalates were acquired from the substrate, but the probability that this might have happened is higher in those cases where calcium oxalates were also detected at the lichen-rock interface. Weddellite was detected at the interface of all species except Aspicilia hoffmanniana from SE facing surfaces. This study therefore suggests that lichen-induced physical weathering in the Côa Valley is species-specific and may be stronger on north-east facing surfaces, whereas lichen-induced chemical action is microclimatically controlled and may be more severe on SE facing surfaces. There is probably some variation in the relative abundance of alteration minerals and calcium oxalates at different portions of the samples but according to present evidence, the lichens currently dominant on the vertical schist surfaces in the Côa Valley are unlikely to be responsible for the differential weathering (and presumably consequent distribution pattern) of engraved schist surfaces. Furthermore, calcium oxalate production by lichens not attributable to any kind of lichen activity, as it happens with Aspicilia hoffmanniana, adds to the doubts concerning its importance in lichen-induced weathering, especially since this seems to be limited to a few species, and, as demonstrated by this study, changes with microclimatic conditions. Assumptions about the drivers of open-air rock-art distribution elsewhere should therefore also consider the micro-

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environmental controls of lichen-induced weathering and associated deteriogenic activity in order to avoid biased measures of the influence of lichen activity in the deterioration process, or superficial decisions in terms of rock-art conservation practices.

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Literature cited

Adamo P., Violante P. (2000) Weathering of rocks and neogenesis of minerals associated with lichen activity. *Applied Clay Science* 16, 229-256.

596	Adamson C., McCabe S., Warke P.A., McAllister D., Smith B.J. (2013) The influence of aspect
597	on the biological colonization of stone in Northern Ireland. International Biodeterioration
598	and Biodegradation 84, 357-366.
599	Aghamiri R., Schwartzman D.W. (2002) Weathering rates of bedrock by lichens, a mini
600	watershed study. Chemical Geology 188, 249-259.
601	Aires S., Carvalho C., Noronha F., Ramos J.F., Moura C., Sant'Ovaia H., Sousa M. (2011) Os
602	xistos do complexo xisto-grauváquico – grupo do Douro. Potencial como recurso
603	geológico. In Actas do VI Seminário Recursos Geológicos, Ambiente e Ordenamento do
604	Território, Vila Real, 6-8 October 2011, pp. 159-165. UTAD, Vila Real.
605	Arocena J.M., Siddique T., Thring R.W., Kapur S. (2007) Investigation of lichens using
606	molecular techniques and associated mineral accumulations on a basaltic flow in a
607	Mediterranean environment. Catena 70, 356-365.
608	Ascaso C., Galvan J., Ortega C. (1976) The pedogenic action of <i>Parmelia conspersa</i> ,
609	Rhizocarpon geographicum and Umbilicaria pustulata. The Lichenologist 8, 151-171.
610	Aubry T., Luís L., Dimuccio L.A. (2012) Nature vs. Culture, present-day spatial distribution and
611	preservation of open-air rock-art in the Côa and Douro River Valleys (Portugal). Journal of
612	Archaeological Science 39 (4), 848-866.
613	Banfield J.F., Eggleton R.A. (1990) Analytical transmission electron microscope studies of
614	plagioclase, muscovite, and K-feldspar weathering. Clays and Clay Minerals 38 (1), 77-89.
615	Bartoli F., Municchia A.C., Futagami Y., Kashiwadani H., Moon K.H., Caneva G. (2014)
616	Biological colonization patterns on the ruins of Angkor temples (Cambodia) in the
617	biodeterioration vs bioprotection debate. International Biodeterioration and
618	Biodegradation 96, 157-165.

619	Boulesteix AL., Janitza S., Kruppa J., König I.R. (2012) Overview of Random Forest
620	methodology and practical guidance with emphasis on computational biology and
621	bioinformatics. Wiley Interdisciplinary Reviews, Data Mining and Knowledge Discovery 2
622	493-507.
623	Breiman L. (2001) Random Forests. Machine Learning 45, 5-32.
624	Búrcio M. (2004) Controle estrutural da localização de pedreiras de esteios de xisto para vinha
625	em Vila Nova de Foz Côa. MSc Thesis. University of Évora.
626	Caneva G. (1993) Ecological approach to the genesis of calcium oxalate patinas on stone
627	monuments. Aerobiologia 9, 149-156.
628	Cann J.H. (2012) Physical weathering of slate gravestones in a Mediterranean climate.
629	Australian Journal of Earth Sciences 59 (7), 1021-1032.
630	Carballal R., Paz-Bermúdez G., Sánchez-Biezma M.J., Prieto B. (2001) Lichen colonization of
631	coastal churches in Galicia, biodeterioration implications. International Biodeteriorarion
632	and Biodegradation 47, 157-163.
633	Carter N.E.A., Viles H.A. (2003) Experimental investigations into the interactions between
634	moisture, rock surface temperatures and an epilithic lichen cover in the bioprotection of
635	limestone. Building and Environment 38 (9-10), 1225-1234.
636	Carter N.E.A., Viles H.A. (2005) Bioprotection explored: the story of a little known earth surface
637	process. Geomorphology 67 (3), 273-281.
638	Casanova Municchia A., Percario Z., Caneva G. (2014) Detection of endolithic spatial
639	distribution in marble stone. Journal of Microscopy 256 (1), 37-45.
640	Chen J., Blume H.P., Beyer L. (2000) Weathering of rocks induced by lichen colonization - a
641	review. Catena 39 (2), 121-146.

642 de Mendiburu F. (2013) agricolae, statistical procedures for agricultural research. R package 643 version 1.1-5. de Oliveira L.F.C., Edwards H.G.M., Feo-Manga J.C., Seaward M.R.D., Lücking R. (2002) FT-644 645 Raman spectroscopy of three foliicolous lichens from Costa Rican rainforests. The 646 Lichenologist 34 (3), 259-266. Edwards H.G.M., Garcia-Pichel F., Newton E.M., Wynn-Williams D.D. (2000) Vibrational 647 Raman spectroscopic study of scytonemin, the UV-protective cyanobacterial pigment. 648 649 Spectrochimica Acta Part A 56 (1), 193-200. Edwards H.G.M., Newton E.M., Wynn-Williams D.D., Lewis-Smith R.I. (2003a) Non-650 651 destructive analysis of pigments and other organic compounds in lichens using Fourier-652 transform Raman spectroscopy: a study of Antarctic epilithic lichens. Spectrochimica Acta Part A 59, 2301-2309. 653 Edwards H.G.M., Russell N.C., Seaward M.R.D. (1997) Calcium oxalate in lichen 654 655 biodeterioration studied using FT-Raman spectroscopy. Spectrochimica Acta Part A 53, 99-105. 656 Edwards H.G.M., de Oliveira L.F.C., Seaward M.R.D. (2005) FT-Raman spectroscopy of the 657 658 Christmas wreath lichen, Cryptothecia rubrocincta (Ehrenb., Fr.) Thor. The Lichenologist 659 37 (2), 181-189. 660 Edwards H.G.M., Russell N.C., Seaward M.R.D., Slarke D. (1995) Lichen biodeterioration 661 under different microclimates, an FT Raman spectroscopic study. Spectrochimica Acta 662 *Part A* 51, 2091-2100.

663	Edwards H.G.M., Seaward M.R.D., Attwood S.J., Little S.J., de Oliveira L.F.C., Tretiach M.
664	(2003b) FT-Raman spectroscopy of lichens on dolomitic rocks, an assessment of metal
665	oxalate formation. The Analyst 128, 1218-1221.
666	Favero-Longo S.E., Castelli D., Salvadori O., Belluso E., Piervittori R. (2005) Pedogenetic
667	action of the lichens Lecidea atrobrunnea, Rhizocarpon geographicum gr. and Sporastatia
668	testudinea on serpentinized ultramafic rocks in an alpine environment. International
669	Biodeterioration and Biodegradation 56 (1), 17-27.
670	Favero-Longo S.E., Gazzano C., Girlanda M., Castelli D., Tretiach M., Baiocchi C., Piervittori
671	R. (2011) Physical and Chemical Deterioration of Silicate and Carbonate Rocks by
672	Meristematic Microcolonial Fungi and Endolithic Lichens (Chaetothyriomycetidae).
673	Geomicrobiology Journal 28 (8), 732-744.
674	Fernandes A.P.B. (2012) Natural processes in the degradation of open-air rock-art sites: an
675	urgency intervention scale to inform conservation. PhD Dissertation. Bournemouth
676	University.
677	Freeman J.J., Wang A., Kuebler K.E., Jolliff B.L., Haskin L.A. (2008) Characterization of
678	natural feldspars by Raman spectroscopy for future planetary exploration. The Canadian
679	Mineralogist 46, 1477-1500.
680	Frost (1997) The structure of the kaolinite minerals – a FT-Raman study. <i>Clay Minerals</i> 32, 65-
681	77.
682	Fry E.J. (1924) A suggested explanation of the mechanical action of lithophytic lichens on rock
683	(shale). Annals of Botany 38, 175-196.
684	Fry E.J. (1927) The mechanical action of crustaceous lichens on substrate of shale, schist, gneiss,
685	limestone and obsidian. Annals of Botany 41, 437-460.

686	Galvan J., Rodriguez C., Ascaso C. (1981) The pedogenic action of lichens in metamorphic
687	rocks. Pedobiologia 21, 60-73.
688	Gazzano C., Favero-Longo S.E., Matteucci E., Piervittori R. (2009a) Image analysis for
689	measuring lichen colonization on and within stonework. The Lichenologist 41 (3), 299-
690	313.
691	Gazzano C., Favero-Longo S.E., Matteucci E., Roccardi A., Piervittori R. (2009b) Index of
692	lichen potential biodeteriogenic activity (LPBA), a tentative tool to evaluate the lichen
693	impact on stonework. International Biodeterioration and Biodegradation 63 (7), 836-843.
694	Giordani P., Modenesi P., Tretiach M. (2003) Determinant factors for the formation of the
695	calcium oxalate minerals, weddellite and whewellite, on the surface of foliose lichens. The
696	Lichenologist 35 (3), 255-270.
697	Gomes L. M. F., Almeida P. G. (2003) As pedreiras do Poio (Foz Côa) - a região e o turismo. In
698	A Geologia de Engenharia e os Recursos Geológicos. Vol. 1. Geologia de Engenharia (ed
699	M.P.V. Ferreira), pp. 299-316. Imprensa da Universidade de Coimbra, Coimbra.
700	Jones D., Wilson M.J., McHardy W.J. (1981) Lichen weathering of rock-forming minerals,
701	application of scanning electron microscopy and microprobe analysis. Journal of
702	Microscopy 124, 95-104.
703	Jorge-Villar S.E., Edwards H.G.M., Seaward M.R.D. (2004) Lichen biodeterioration of
704	ecclesiastical monuments in northern Spain. Spectrochimica Acta Part A 60, 1229-1237.
705	Kidron G.J., Temina M. (2010) Lichen colonization on cobbles in the Negev Desert following 15
706	years in the field. Geomicrobiology Journal 27 (5), 455-463.

- 707 Kretzschmar R, Robarge W.P., Amoozegar A., Vepraskas M.J. (1997) Biotite alteration to 708 halloysite and kaolinite in soil-saprolite profiles developed from mica schist and granite 709 gneiss. Geoderma 75 (3-4), 155-170. 710 Liaw A., Wiener M. (2002) Classification and regression by Random Forest. R News 2, 18-22. 711 Marques J., Hespanhol H., Paz-Bermúdez G., Almeida R. (2014) Choosing between sides in the 712 battle for pioneer colonization of schist in the Côa Valley Archaeological Park, a 713 community ecology perspective. Journal of Archaeological Science 45, 196-206. 714 McIlroy de la Rosa J.P., Warke P.A., Smith B.J. (2013) Lichen-induced biomodification of 715 calcareous surfaces: bioprotection versus biodeterioration. *Progress in Physical Geography* 716 37 (3), 325-351. 717 McIlroy de la Rosa J.P., Warke P.A., Smith B.J. (2014) The effects of lichen cover upon the rate 718 of solutional weathering of limestone. Geomorphology 220, 81-92. 719 Oppel S., Strobl C., Huettmann F. (2009) Alternative methods to quantify variable importance in 720 ecology. Technical Report Number 65. University of Munich, Munich. 721 Parham W.E. (1969) Formation of halloysite from feldspar, low temperature, artificial 722 weathering versus natural weathering. Clays and Clay Minerals 17, 13-22. 723 Polzehl J., Tabelow K. (2007) Adaptive smoothing of digital images, the R package adimpro. 724 Journal of Statistical Software 19, 1-17. 725 Prieto B., Edwards H.G.M., Seaward M.R.D. (2000) A Fourier transform-Raman spectroscopic
 - Acta Científica Compostelana (Bioloxía) 13, 35-45.

study of lichen strategies on granite monuments. Geomicrobiology Journal 17, 55-60.

Prieto V.B., Silva B. (2003) Neoformed calcium minerals in granite colonised by lichens. *Nova*

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729 Prieto B., Silva B., Rivas T., Wierzchos J., Ascaso C. (1997) Mineralogical transformation and 730 neoformation in granite caused by the lichens *Tephromela atra* and *Ochrolechia parella*. 731 *International Biodeterioration and Biodegradation* 40 (2-4), 191-199. 732 Sanders W.B., Ascaso C., Wierzchos J. (1994) Physical interactions of two rhizomorph-forming 733 lichens with their rock substrate. Botanica Acta 107 (6), 432-439. 734 Seaward M.R.D. (1997) Major impacts made by lichens in biodeterioration processes. 735 *International Biodeterioration and Biodegradation* 40 (2-4), 269-273. 736 Seaward M.R.D., Edwards H.G.M. (1995) Lichen-substratum interface studies, with particular reference to Raman microscopic analysis. 1. Deterioration of works of art by *Dirina* 737 738 massiliensis forma sorediata. Cryptogamic Botany 5 (3), 282-287. 739 Seaward M.R.D. (2015) Lichens as agents of Biodeterioration. In Recent advances in 740 Lichenology. Modern methods and approaches in biomonitoring and bioprospection (eds 741 D.K. Upreti, P.K. Divakar, V. Shukla & R. Bajpai), pp. 189-211. Springer India, New 742 Delhi. Sousa M.B. (1982) Litostratigrafia e estrutura do complex xisto-grauváquico ante-Ordovícico – 743 grupo do Douro (nordeste de Portugal). PhD thesis. University of Coimbra. 744 745 St. Clair L.L., Seaward M.R.D. (2004) Biodeterioration of stone surfaces. Kluwer Academic 746 Press, Dordrecht. 747 Steinbauer M., Gohlke A., Mahler C., Schmiedinger A., Beierkuhnlein C. (2013) Quantification 748 of wall surface heterogeneity and its influence on species diversity at medieval castles -749 implications for the environmentally friendly preservation of cultural heritage. *Journal of* Cultural Heritage 14 (3), 219-228. 750

751 Stretch R., Viles H.A. (2002) Lichen weathering on Lanzarote lava flows. Geomorphology 47, 752 87-94. 753 Viles H.A., Cutler N.A. (2012) Global environmental change and the biology of heritage 754 structures. Global Change Biology 18(8), 2406-2418. Vingiani S., Terribile F., Adamo P. (2013) Weathering and particle entrapment at the rock-lichen 755 interface in Italian volcanic environments. Geoderma 207-208, 244-255. 756 Wang A., Freeman J., Kuebler K.E. (2002) Raman spectroscopic characterization of 757 phyllosilicates. 33rd Annual Lunar and Planetary Science Conference, March 11-15, 758 Houston (TX), USA. 759 Wierzchos J., Ascaso C. (1994) Application of back-scattered electron imaging to the study of 760 761 the lichen-rock interface. Journal of Microscopy 175 (1), 54-59. Wierzchos J., Ascaso C. (1996) Morphological and chemical features of bioweathered granitic 762 biotite induced by lichen activity. Clays and Clay Minerals 44 (5), 652-657. 763 764 Wierzchos J., Ascaso C. (1998) Mineralogical transformation of bioweathered granitic biotite, studied by HRTEM, evidence for a new pathway in lichen activity. Clays and Clay 765 Minerals 46 (4), 446-452. 766 Wierzchos J., de los Ríos A., Ascaso C. (2013) Microorganisms in desert rocks: the edge of life 767 768 on Earth. *International Microbiology* 15 (4), 172-182. 769 Xu Q.-S., Liang Y.-Z., Du Y.-P. (2004) Monte Carlo cross-validation for selecting a model and 770 estimating the prediction error in multivariate calibration. Journal of Chemometrics 771 18,112-120.

773 Tables

Table 1. Hyphal penetration: measures taken out of colonized cross-sections after PAS staining. Depth: estimated depth of hyphal penetration in mm; HPC: area occupied by the hyphal penetration component in mm²; WR: area occupied by the weathering rind in mm². The area analysed in each cross-section was 57.5 mm². Different letters in brackets indicate statistically significant pair-wise comparisons (p-value< 0.05).

Species	Aspect	Depth	Depth	HPC (mean ±	WR
		(mean ± sd)	(max)	sd)	
Aspicilia	South-east	5.7 ± 7.6	37.0	2.96 ± 2.03 (a)	0.00
hoffmanniana					
	North-west	3.5 ± 3.5	12.0	2.37 ± 0.81 (a)	0.00
Caloplaca subsoluta	South-east	4.1 ± 5.3	30.0	2.09 ± 1.71 (a)	4.07
	North-west	1.8 ± 1.9	6.8	1.74 ± 0.71 (a)	7.56
Lecanora pseudistera	South-east	5.1 ± 6.6	31.0	8.01 ± 2.02 (b)	0.00
	North-west	4.1 ± 3.8	13.0	5.79 ± 2.97 (b)	30.36
Peltula euploca	South-east	2.4 ± 1.0	3.9	0.69 ± 0.60 (a)	6.72
				Spearman: – 0.0	12

Table 2. Data on rock surface microclimate at opposite orientations in the Côa Valley.

		North-west	South-east
Temperature	Average	17 °C	20 °C
	Minimum	13 °C	15 °C
	Maximum	23 °C	30 °C
Relative humidity	Average	70 %	63 %
	Minimum	55 %	45 %
	Maximum	81 %	77 %

Table 3. Summary of the neoformed calcium minerals detected in the lichen-rock interface (FT-

788 Raman spectroscopy and X-ray microdiffraction).

	Aspicilia		Caloplaca subsoluta		Lecanora		Peltula
	hoffmanniana				pseudistera		euploca
	North-	South-	North-	South-	North-	South-	South-east
	west	east	west	east	west	east	
Whewellite							
Weddellite				X		X	X

Table 4. Summary of the minerals detected in lichen thalli (FT-Raman spectroscopy)

	Aspicilia		Caloplaca		Lecanora		Peltula
	hoffmann	iana	subsoluta		pseudistera		euploca
	North-	South-	North-	South-	North-	South-	South-east
	west	east	west	east	west	east	
Whewellite	X				X		
Weddellite	X	X				X	X
Quartz		X	X				X
Phyllosilicates		X				X	

Figure legends

Figure 1. On the top, cross-sections of schist samples colonized by *Lecanora pseudistera* from a north-west facing surface (a) and by *Aspicilia hoffmanniana* from a south-east facing surface (b); and on the bottom, cross-section of a schist sample colonized by *Aspicilia hoffmanniana* from a south-east facing surface after PAS staining (c) and respective image classification (d) with medium grey areas corresponding well with the area occupied by the hyphal penetration component (in purple in the original image).

Figure 2. FT-Raman spectra of bare rock control samples (first row) and colonized samples (remaining rows). On the first row: external surface (thick line), internal surface (thin line) and core (dotted line) of bare rock control samples from north-west (a) and south-east (b) facing vertical schist surfaces. On the remaining rows: thallus surface (thick lines), rock-lichen interface (thin lines) and rock core (dotted lines) of, from top to bottom, *Aspicilia hoffmanniana*, *Caloplaca subsoluta*, *Lecanora pseudistera* and *Peltula euploca* from north-west (a) and south-east (b) facing vertical schist surfaces. Conditions as explained in material and methods.

Wavenumber region: 100-1700 cm⁻¹. Wavenumber assignments are based on Freeman et al. (2008), Frost (1997), Wang et al. (2002), Jorge-Villar et al. 2004, Edwards et al. (1997, 2000, 2003a). Additional information on the wavenumbers for the Raman spectra of all species and respective wavenumber assignment is given in Supporting Information Tables S1-S5.

817 Figure 1

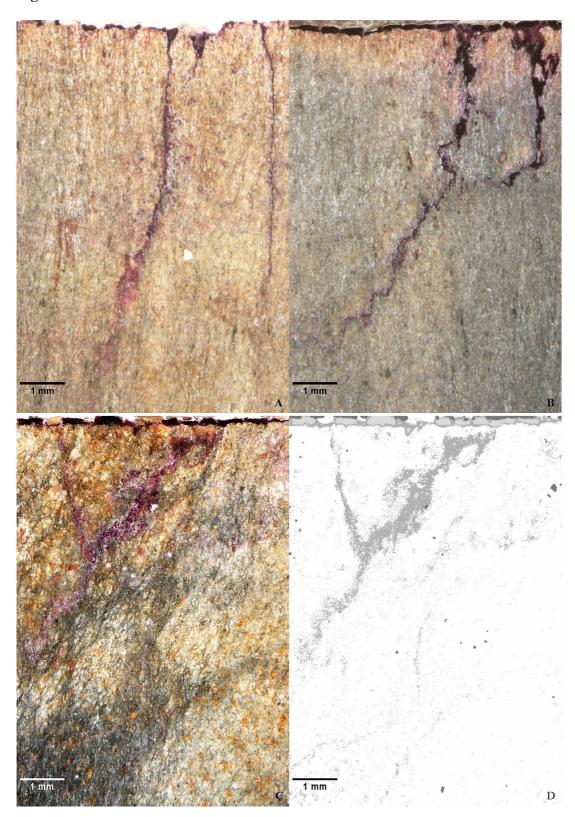


Figure 2

