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# Effects of ultraviolet radiation on production of photoprotective compounds in microalgae of the genus *Pediastrum* from high Andean areas of Peru

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<b>ARTICLE INFO</b>	ABSTRACT		
Received on: 22/07/2021 Accepted on: 09/10/2021 Available Online: 05/03/2022	This study aims to collect microalgae from high Andean areas and evaluate their potential as natural ingredients in the cosmetics industry. The cell growth, pigment concentration, antioxidant activity, and polyphenols content of two microalgae of the genus <i>Pediastrum</i> (IMP-BG288 and IMP-BG255) collected from high Andean lagoons of more than 3,000 m.a.s.l. (Avacucho, Peru) were evaluated. The cultures were exposed to photosynthetically active radiation		
<i>Key words:</i> <i>Pediastrum</i> , antioxidant activity, pigments, polyphenols, lutein, cosmetic.	(PAR) (400–700 nm) and PAR with ultraviolet radiation A and B (PAB) (280–700 nm) in 2 l photobioreactors. Cultures of the genus <i>Pediastrum</i> sp. subjected to PAB radiation showed greater growth until the sixth day, when they reached the end of their exponential phase. The maximum concentrations of lutein and chlorophyll a and b were 6.96 $\pm$ 0.21, 1.64 $\pm$ 0.05, and 0.84 $\pm$ 0.06 mg/g, respectively; total antioxidant activity and total polyphenol content were 221.7 $\pm$ 11.76 µmol eq. Trolox/g and 7.36 $\pm$ 0.53 mg GAE/g, respectively, after being subjected to PAR. Conclusively, high Andean microalgae possess adaptation mechanisms that help them to cope with a high radiation flux. The high concentrations of antioxidants and pigments in the strain taken from 4,400 m.a.s.1 (IMP-BG288) suggest its greatest cosmetic potential.		

# **INTRODUCTION**

The study of the effect of ultraviolet radiation (UVR) on autotrophic organisms of Andean lakes has increased in the last few years. Studies were conducted to evaluate the adaptation mechanisms of these organisms against light stress (Bashevkin *et al.*, 2019; Cartuche *et al.*, 2019; de Oliveira *et al.*, 2016; Rojo *et al.*, 2017; Roleda *et al.*, 2012; Villafañe *et al.*, 2017; Villafañe *et al.*, 2015; Williamson *et al.*, 2019) and to search for novel industrially relevant compounds (Ariede *et al.*, 2017; Joshi *et al.*, 2018; Wang, 2015).

Generally, algae in the areas exposed to radiation possess mechanisms, including photorepair, antioxidant system activation, and biosynthesis of photoprotective compounds

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(Abu-Ghosh *et al.*, 2021; Häder *et al.*, 2007; Holzinger *et al.*, 2018; Ljubic *et al.*, 2021; Rastogi *et al.*, 2020; Singh *et al.*, 2019; Virtanen *et al.*, 2019). These mechanisms are employed to maximize the light capture, limit the damage caused by excessive amounts of solar radiation (Morgan-Kiss *et al.*, 2006), and optimize their photosynthetic performance against UVR (Franklin *et al.*, 2003).

UVR, known to produce reactive oxygen species (ROS), is classified according to wavelengths: UVA (320–400 nm), UVB (280–320 nm), and UVC (200–280 nm). Radiation increases by a combination of factors, such as latitudinal location, ozone, and altitude (Albarracín *et al.*, 2013; Feister and Häder, 2018; Helbling *et al.*, 2001; Michelutti *et al.*, 2015; Ruescas *et al.*, 2016; Villafañe *et al.*, 1999). On evaluating the altitude effect on Lake Titicaca (3,810 m.a.s.l.), one of the most important high Andean lakes in Peru, an increase of 4.8% in the quantity of UVR weighted for every 300 m altitude was recorded (Andrade *et al.*, 1998).



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The microalgae of the chlorophyte group are better adapted to UVR since they possess highly efficient mechanisms, such as accessory pigments, which protect the cytoplasm and chloroplast and allow individual cells to resist extreme conditions (Chiu *et al.*, 2020; Holzinger and Lütz, 2006). The pigments associated with the algal photosynthetic activity include chlorophylls,  $\beta$ -carotene, astaxanthin, xanthophylls, and phycobiliproteins. Most carotenoids have certain antioxidant activities due to the presence of conjugated double bonds, which eliminate UVR-induced free radicals (Huang *et al.*, 2018).

Generally, algae have a variety of bioactive compounds, such as pigments, polysaccharides, polyphenols, lipids, and proteins, as well as antioxidant, anti-inflammatory, antipigmentation, antiaging, and anticancer properties. Therefore, they are used as skincare alternatives as they are considered less harmful than synthetic compounds (Morocho-Jácome *et al.*, 2020b; Thiyagarasaiyar *et al.*, 2020). In addition, other secondary metabolites, such as flavonoids, could prevent UVR sunburn in humans with light-sensitive skin and low melanin levels (Morocho-Jácome *et al.*, 2020a).

Currently, some notable cosmetic applications of the extracts of *Spirulina* sp. and *Chlorella* sp. include their use as ingredients for sunscreen (Jha *et al.*, 2017), dandruff treatment, and hair growth stimulation (Joshi *et al.*, 2018). *Chlorogloeopsis* sp. and *Nannochloropsis* sp. protect keratinous tissue and prevent UVA- and UVB-induced damage, photoaging, wrinkle formation, and sagging skin (Ariede *et al.*, 2017). *Isochrysis* sp. is applied in sunscreen formulations containing organic and inorganic filters with sun protection factor 15 in order to prevent UVR transmission. *Thalassiosira* sp., *Monodus* sp., *Chlorococcum* sp., and *Chaetoceros* sp. are applied to prevent hair loss (Zanella *et al.*, 2018). Formulations with oil of *Chlorella* sp. make the skin and hair softer and more flexible (Brooks and Franklin, 2013). The hydroxy acids in *Pediastrum duplex* are potential skin moisturizers (Yarkent *et al.*, 2020).

This study aims to collect microalgae from high Andean areas and evaluate their potential as natural ingredients for the cosmetic industry. In this regard, microalgae of the genus *Pediastrum* were collected from lagoons of altitudes between 3,110 and 4,400 m.a.s.l. Adaptation of the microalgae to

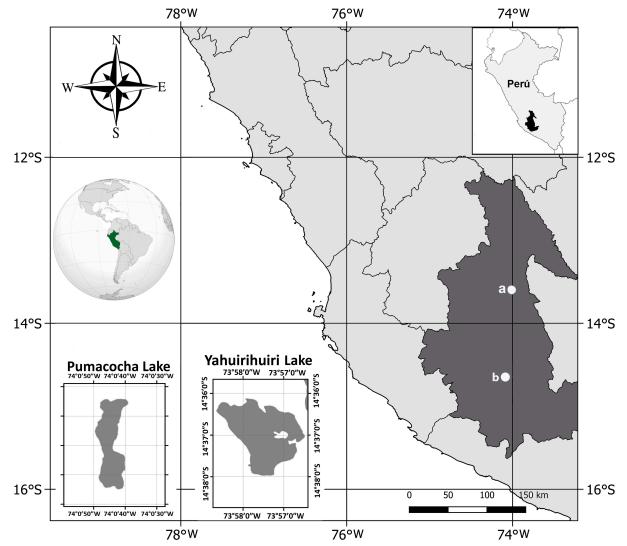


Figure 1. Location of the sampling areas in Ayacucho (Peru). a: Pumacocha Lake. b: Yahuirihuiri Lake.

photosynthetically active radiation (PAR) and PAR with ultraviolet A and B (PAB) was evaluated in the production of photoprotective compounds, such as pigments, antioxidant activity, and total polyphenols.

# MATERIALS AND METHODS

# Sampling and isolation of microalgae strains

In the Andean lagoons of Yahuirihuiri and Pumacocha, located in the Department of Ayacucho, Peru (Fig. 1), microalgae samples were collected with a 20 µm phytoplankton network. The samples were placed in 50 ml Falcon® tubes, stored in thermal boxes with a gel pack, and then transferred to the Laboratory of the Germplasm Bank of Aquatic Organisms (BGOA) of the Marine Institute of Peru. Monoclonal strains of microalgae of the genus Pediastrum were obtained through the cell isolation methodology described by Andersen and Kawachi (2005) and using inverted phase-contrast microscopy (Nikon, Eclipse TS100, Japan). For the cultivation of *Pediastrum* sp, a Wright's cryptophyte (WC) culture medium (Andersen et al., 2005; Guillard and Lorenzen, 1972) was used in a growth chamber (Binder KBW400, Germany) under the following conditions:  $18^{\circ}C \pm 1^{\circ}C$ , 80 µmol. s<sup>-1</sup>. m<sup>-2</sup>, and a 2:12 hours light:dark photoperiod. These monoclonal strains were classified based on their morphology (Lenarczyk and Saługa, 2018; Zamaloa and Tell, 2005) and assigned codes (Table 1) to integrate them into the BGOA collection.

# **Experimental design**

Strains of *Pediastrum* were cultivated inside a culture chamber in 2 l rectangular parallelepiped photobioreactors ( $26 \times 15.1 \times 5.1$  cm) with the WC culture medium. The average initial cell density of the cultures was  $9.35 \times 10^5$  cells/ml.

For a period of 8 days, two radiation levels were applied to the cultures. All the following cultures were carried out in triplicate: a) PAR (80  $\mu$ mol. S<sup>-1</sup> m<sup>-2</sup>) with a 12:12 hours (light:dark) photoperiod and b) PAB with a 12:12 hours (light:dark) photoperiod; 3 hours after starting the PAR luminous phase (80  $\mu$ mol. S<sup>-1</sup>m<sup>-2</sup>), UV radiation (UVA + UVB, 4 mW/cm<sup>2</sup>) was activated for 3 hours.

PAR was recorded using a quantometer (Liquor Quantimeter), whereas UVR was recorded using a radiometer (UVP UVX, USA). All treatments were carried out at  $18^{\circ}C \pm 1^{\circ}C$  and at constant aeration for 8 days. Samples (1 ml) were taken from each experimental unit every 24 hours and were fixed with Lugol for cell quantification.

Finally, cultures were harvested on the eighth day and the biomass was concentrated by centrifugation (3,500 rpm for 10 minutes at 5°C) in a refrigerated centrifuge (Hettich, Germany). The wet biomass was dried in a lyophilizer (Labconco, EEUU) and stored at 8°C in a dark and cold environment.

# Specific growth rate

The cell density of the cultures was evaluated through the cell counting method with a Neubauer camera on a compound microscope (Nikon, Japan). The specific growth rate ( $\mu$ ) was determined using the following formula:

$$\mu = (\ln x_2 - \ln x_1) / (t_2 - t_1),$$

Where  $t_2$  is the final time,  $t_1$  is the initial time,  $x_1$  is the initial cell density, and  $x_2$  is the final cell density.

#### **Determination of pigments**

Pigments were extracted from 50 mg of the lyophilized microalgae with 6 ml of 100% acetone as per the protocol of Pasquet et al. (2011). Samples were filtered using a 4.5 µm polytetrafluoroethylene syringe filter, and 20 µl of the filtrate was used to carry out high-performance liquid chromatography (Hitachi, Tokyo). The Merck LiChrospher® C18 250  $\times$  4.6 mm  $\times$  5  $\mu$ m analytical column was used. The mobile phases were A = 100%methanol, B = methanol:0.5 N ammonium acetate (80:20), and C =tetrahydrofuran. The ramp was 0.0 min (0% A, 100% B, and 0% C), 5.0 minutes (98% A, 0% B, and 2% C), 17.2 minutes (80% A, 0% B, and 20% C), 25.0 minutes (80% A, 0% B, and 20% C), 26.0 minutes (98% A, 0% B, and 2% C), and 34.0 minutes (0% A, 100% B, and 0% C). The column temperature was maintained at 25°C. Spectra at the 210 and 690 nm wavelengths were recorded. The analytical standards (Sigma-Aldrich) of lutein, chlorophyll a, chlorophyll b, and  $\beta$ -carotene were used as references to quantify the pigments.

#### **Determination of total polyphenols**

The polyphenols were extracted from 50 mg of the lyophilized biomass using 6 ml of 100% acetone and subsequently with 6 ml of ultrapure water. Total polyphenols in the extracts were analyzed following the methods described by Singleton and Rossi (1965). The calibration curve was prepared using gallic acid at a concentration range of 100–500  $\mu$ g/ml. The absorbance was measured at 760 nm using a spectrophotometer (Varian, Australia). The values obtained in triplicate were expressed in mg GAE/g. The total polyphenol content was considered as the sum of the aqueous and acetone fractions.

#### **Determination of antioxidant activity**

Overall, 50 mg of the lyophilized biomass was used to determine the antioxidant activity and exhaustive extraction was carried out using three solvents: hexane, acetone, and water. The hexane and acetone extracts were evaporated by injecting

 Table 1. Pediastrum strain from high Andean lakes from the Department of Ayacucho, geographic coordination, and BGOA code.

Strain	Code	Lake	Altitude (m.a.s.l.)	Latitude, longitude
Pediastrum sp.	IMP-BG288	Yahuirihuiri Lake	4,400	14° 37.85′ S, 73° 57.17′ W
Pediastrum sp.	IMP-BG255	Pumacocha Lake	3,311	13° 35.49′ S, 74° 00.76′ W

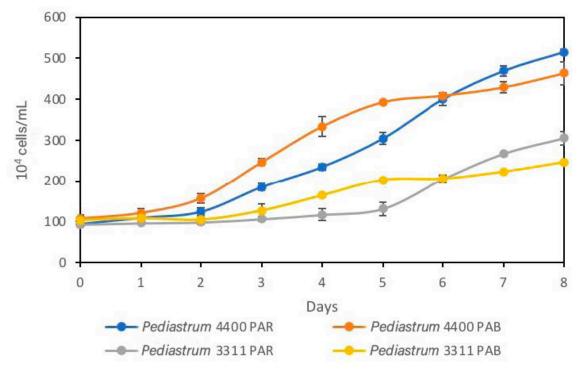


Figure 2. Effect of PAR and PAB radiations on growth of the strains IMP-BG288 (4,400 m.a.s.l) and IMP-BG255 (3,111 m.a.s.l).

nitrogen gas and resuspended in 2 ml of absolute ethanol. This was carried out in triplicate to quantify the antioxidant activity. The antioxidant activity of the extracts was determined using the (2,2'-azino-bis(3-ethylbenzothiazoline-6-sulfonic acid)) + (2,2'-azinobis-(3-ethylbenzothiazoline-6-sulfonic acid)) radical cation method, as described by Re *et al.* (1999). Absorbance was measured at 714 nm using a spectrophotometer (Varian, Australia). The values obtained were expressed in µmol eq. Trolox/g. The total antioxidant activity was considered as the sum of the hexane, acetone, and aqueous fractions.

#### Statistical analysis

Data obtained from the growth, antioxidant activity, pigments, and polyphenol assays were analyzed using the analysis of variance at a significance level of  $\alpha = 0.05$ . Means were compared using Tukey's post hoc test. All statistical analyses were carried out using the MINITAB 16 program.

# **RESULTS AND DISCUSSION**

#### Growth of Pediastrum sp. after PAR and PAB radiation

The *Pediastrum* cultures subjected to PAB radiation showed a short latency phase: 1 day for IMP-BG288 and 2 days for IMP-BG255. Concurrently, the same strains subjected to PAR radiation showed latency phases of 2 and 4 days, respectively (Fig. 2). This indicates that the strains from these high Andean areas are genetically conditioned to modulate UVR stress. In this case, growth stimulation was observed. Also, UVR photostress reduced the latency phase in *Chlorella vulgaris* (NIOT5 India) (Balan and Suraishkumar, 2014). Between the third and fourth days, the cultures of *Pediastrum* sp. taken from 4,400 and 3,311 m.a.s.l. and exposed to PAB radiation (Table 2) showed the highest specific growth rate. This is due to the adaptation ability of this strain to high radiation (Andrade *et al.*, 1998). The differences in the growth rates of *Pediastrum* sp. strains could be as a result of an adaptive photoprotection response that is associated with specific species (Janknegt *et al.*, 2009; Wong *et al.*, 2007). In *C. vulgaris*, the specific growth rate ( $\mu$ ) was 0.39 when it was exposed to UVA+B radiation. This was higher than the control, in which UVB radiation generated an increase in the growth rate during the early growth phase. This is as a result of changes in the photosystem stoichiometry and the size of the antenna pigments (Balan and Suraishkumar, 2014).

On the sixth day, the cultures of *Pediastrum* sp. exposed to PAR radiation were found in the exponential phase. However, the cultures exposed to PAB radiation showed a delay in their growth, entering a short period of stationary phase between days 5 and 6 (Fig. 2). It should be mentioned that similar densities were observed on day 6 of the cultivation in the strains exposed to PAR and PAB radiation. This short period of stationary phase was because UVR exposure induces oxidative stress in microalgae

**Table 2.** Effect of PAR and PAB radiations on specific growth rate  $(\mu)$  in the exponential phase (third and fourth days) of *Pediastrum* sp.

Treatments	μ	
DAD DC299 (4.400 1)	PAR	$0.19\pm0.10^{\rm a}$
IMP-BG288 (4,400 m.a.s.l.)	PAB	$0.30\pm0.04^{\rm b}$
D (D D C 255 (2 211 1)	PAR	$0.09\pm0.09^{\circ}$
IMP-BG255 (3,311 m.a.s.l.)	PAB	$0.26\pm0.15^{\rm b}$

Different letters mean statistically significant differences (p  $\leq 0.05$ , Tukey's test).

(Paliwal *et al.*, 2017; Tian and Yu, 2009). This shows that the repair mechanisms for the genes regulating photosynthesis and metabolic stress are rapid, which activate the antioxidant systems and DNA repair (Brosché and Strid, 2003; Buma *et al.*, 2003; Helbling *et al.*, 2001; Portwich and Garcia-Pichel, 2000; Sijil *et al.*, 2020). Nevertheless, between days 7 and 8, the cultures of both strains showed significant increases in their cell density. Similar specific growth rates were observed in the cultures of IMP-BG288 (0.08 and 0.09) and IMP-BG255 (0.10 and 0.13) strains exposed to PAB and PAR, respectively.

The adaptive response of *Pediastrum* sp. strains derived from lagoons at different heights above sea level to UVR was expressed in the growth stages. Xiong *et al.* (1996) evaluated the growth of microalgae from several regions and demonstrated that the microalgae from high altitude areas adapt to radiation. This was because this factor induces different genes that encode for proteins involved in the biosynthesis of photoprotective pigments, such as those involved in the regulation of photosynthesis and metabolic stress, as well as those involved in the stimulation of enzyme systems, antioxidants, and DNA repair (Barrado-Moreno *et al.*, 2017; Brosché and Strid, 2003; Buma *et al.*, 2003; Helbling *et al.*, 2001; Portwich and Garcia-Pichel, 2000).

#### **Pigments**

Chlorophyll a and b in both *Pediastrum* strains decreased after PAB exposure (p < 0.05) (Fig. 3). *Chlorella vulgaris, Chlorococcum infusionum* (Schrank) Meneghini, and *Chlorogonium elongatum* (Dang.) showed similar results in terms of chlorophyll concentrations when exposure to UVB was increased (Agrawal, 1992; Ganapathy *et al.*, 2017).

In addition, lutein was the most abundant pigment (>5 mg/g) in the two *Pediastrum* strains that were subjected to PAR and PAB radiation. The pigment content decreased in cultures exposed to PAB radiation (p < 0.05). The concentration of lutein in

the cultures of the algae *Aglaothamnion uruguayense* exposed to PAB and PAR was observed to be higher in the cultures after PAB radiation (Ouriques *et al.*, 2017).

The  $\beta$ -carotene concentrations of the two *Pediastrum* strains exposed to PAR and PAB showed no significant differences (p > 0.05). However, there were no apparent changes (p > 0.05) in *Dunaliella salina* when exposed to UVR. This demonstrated that carotenoids were more associated with antioxidants than photosynthetic pigments (Tian and Yu, 2009). Generally, the accumulation of pigments at high radiation and light intensity occurs in a short period of time in some algae regulated by photomorphogenic photoreceptors (Wang *et al.*, 2007).

In addition, the percentages of lutein and chlorophylls when exposed to PAB radiation decreased by 20%–24% and 45%–67%, respectively, for both strains. The concentrations of chlorophyll a and carotenoids significantly decreased in the cultures containing *Phormidium tenuous* and *Nostoc muscorum* by over 70% and 80%, respectively (Agrawal and Chaudhary, 1996; Bhandari and Sharma, 2011).

In summary, our results indicate that when microalgae *Pediastrum* cultures were subjected to PAB radiation, the concentrations of lutein, chlorophyll a, and chlorophyll b were significantly (p < 0.05) low on the eighth day compared with those exposed to PAR.

In addition, algae pigments have antiaging, antioxidant, and neuroprotective properties, which make them suitable for use in cosmetics industries (Berthon *et al.*, 2017; Yarkent *et al.*, 2020). Also, pigments, such as chlorophyll a, chlorophyll b,  $\beta$ -carotene, and lutein, can be used in the cosmetic industry as antioxidants, anti-inflammatories, antiwrinkles products, and moisturizers, as well as in collagen synthesis, for skin elasticity, and to prevent premature aging (Joshi *et al.*, 2018; Wang, 2015).

According to our results, the strain IMP-BG288 exposed to PAR radiation can possibly have great cosmetic potential. This is

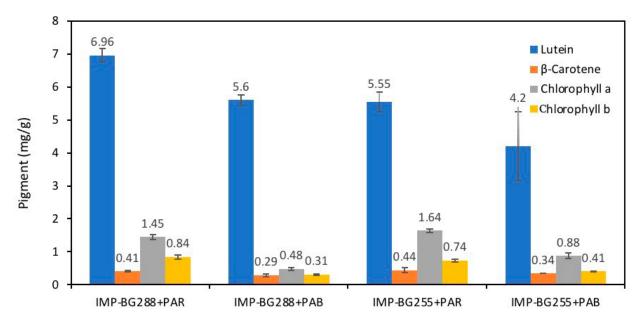


Figure 3. Concentration of lutein,  $\beta$ -carotene, chlorophyll a, and chlorophyll b (mg/g) in cultures of IMP-BG288 and IMP-BG255 strains of *Pediastrum* sp. exposed to PAR and PAB radiation.

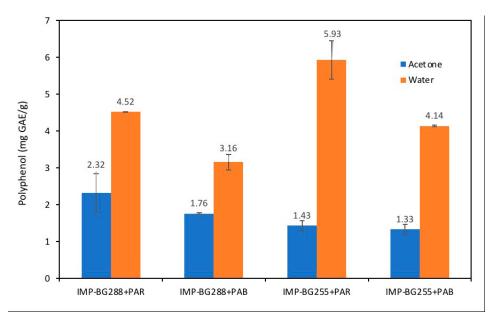


Figure 4. Phenolic contents (mg GAE/g) of different fractions of *Pediastrum* sp.

because lutein is an intracellular algae product that has a protective effect on the skin when combined with other antioxidants and immunoprotective substances (Wang, 2015). It can also be applied as a colorant in the cosmetic industry (Sathasivam *et al.*, 2019).

## Polyphenols

Polyphenols show antioxidant properties and antiradical, antiallergic, anti-inflammatory, and UV protective properties which can react directly with ROS, forming less reactive products (Abu-Ghosh *et al.*, 2021; Berthon *et al.*, 2017; Moskaug *et al.*, 2005).

The aqueous fraction of both strains (IMP-BG288 and IMP-BG255) of *Pediastrum* recorded values between 3.0 and 6.0 mg GAE/g, respectively. These values were higher than those reported by Li *et al.* (2007) for the aqueous fraction of the *Chlorella* group (0.97 to 2.67 mg GAE/g) and lower than that of *Nostoc ellipsosporum* CCAP 1453/17 (10.68 mg GAE/g). It should be noted that the polyphenol concentrations of cultures of both strains exposed to PAB radiation significantly decreased (p < 0.05) when compared with those exposed to PAR (Fig. 4).

However, the polyphenols extracted using acetone did not show significant differences (p > 0.05) in the two *Pediastrum* strains subjected to PAR and PAB radiation.

Duval *et al.* (2000) observed similar results when UVA radiation was emitted in a species of *Chlamydomonas nivalis* collected from 3,100 m.a.s.l. These polyphenol extracts (ethanol) did not show significant differences with UVA exposure for up to 15 days.

In relation to total polyphenols, both strains (IMP-BG288 and IMP-BG255) of *Pediastrum* had values between 4.90 and 7.40 mg GAE/g. Li *et al.* (2007) evaluated the total polyphenols in four different strains of *N. ellipsosporum*, reporting values between 7.2 and 60.4 mg GAE/g. In three strains of *Chlorella pyrenoidosa*, values between 10.46 and 17.24 mg GAE/g were reported. Also, the three strains of *C. vulgaris* had values between 8.6 and 15.0 mg GAE/g.

Polyphenols are necessary for the cosmetic industry due to the essential biological properties they possess. Hence, in this

study, both *Pediastrum* strains have similar cosmetic potential based on their total polyphenols content.

# Antioxidant activity

In this study, for *Pediastrum* sp., the UVR dose of 4 mW.cm<sup>-2</sup> acts as an inducer of the biosynthesis of photoprotective and antioxidant compounds. This allows the species to achieve a homeostatic balance against stress generated by exposure to UVR (Rastogi *et al.*, 2020). However, the response will be different, depending on the place of origin and species. Wong *et al.* (2007) observed that the sensitivity of microalgae to UVB stress varied according to the species and geographical location. The strain IMP-BG288 from the lagoon at 4,400 m.a.s.l. showed significantly higher antioxidant activity (p < 0.05) than that of the strain IMP-BG255 from 3,100 m.a.s.l. This result confirms that microalgae species that inhabit lagoons exposed to high UVR have better protection mechanisms, which is reflected in the outstanding antioxidant activity of the strain IMP-BG288 (Fig. 5).

The total antioxidant activity in the cultures of the strains of *Pediastrum* sp. when exposed to PAB radiation did not show any significant differences (p > 0.05) compared with when exposed to PAR. The IMP-BG288 *Pediastrum* strain recorded high values in the acetone fraction (>149.0 µmol eq. Trolox/g) when exposed to both PAR and PAB radiations compared to those of the water and hexane fraction. In addition, the strain IMP-BG255 showed higher values in the aqueous fraction (> 49.0 µmol eq. Trolox/g) when exposed to both PAR and PAB radiations (Fig. 5). This is because the antioxidant compounds of microalgae can vary significantly in polarity, with marked differences even within the same families and species (Li *et al.*, 2007).

The total antioxidant activity of the strain IMP-BG288 for both treatments had values greater than 210.0 umol eq. Trolox/g with regard to the records of microalgae species, such as *Botryococcus braunii* (53.90 umol eq. Trolox/g), *Chlorella* sp. (59.57 umol eq. Trolox/g), *Tetraselmis suecica* (56.46 umol eq. Trolox/g), *Neochloris oleoabundans* (64.30 umol eq. Trolox/g),

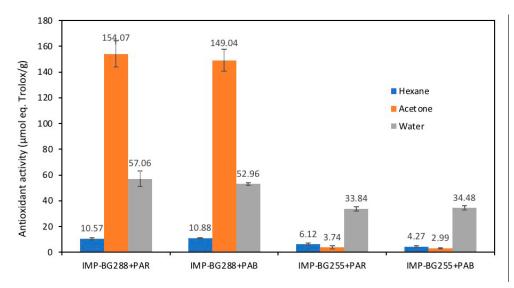


Figure 5. Antioxidant activity (µmol eq. Trolox/g) of different fractions in *Pediastrum* sp.

*Anabaena cylindrica* (54.81 umol eq. Trolox/g), *C. vulgaris* (58.24 umol eq. Trolox/g), *Fischerella ambigua* (74.34 umol eq. Trolox/g), and *Nostoc piscinale* (65.84 umol eq. Trolox/g) (Goiris *et al.*, 2012).

These results imply that habitat is vital for the synthesis of antioxidants as the antioxidant content was higher in the strains from 4,400 m.a.s.l. than those from 3,100 m.a.s.l. As a result of the increase in the photoprotection and antioxidant capacity of the skin, these antioxidants could be extracted from the algal biomass and used to develop cosmetic products (Ariede *et al.*, 2017; Mourelle *et al.*, 2017; Puchkova *et al.*, 2020; Thiyagarasaiyar *et al.*, 2020; Wang, 2015). In addition, investigations on *Pediastrum boryanum* have shown high potential in natural antioxidants, which has other areas of application (Corrêa da Silva *et al.*, 2020; Lee *et al.*, 2009). It is important to note that plants from environments exposed to UVR and only PAR do not activate their metabolic routes (Holzinger and Lütz, 2006). However, this induces possible side effects, such as low concentrations of the antioxidants, which enable them to have defensive mechanisms against pathogens.

#### CONCLUSION

Microalgae from high Andean lagoons have adaptation mechanisms, which enable them to cope with the high radiation flux. The *Pediastrum* sp. strain from the lagoon at 4,400 m.a.s.l. showed greater growth after PAB radiation treatment compared with PAR. Also, the strain from the lagoon at 3,310 m.a.s.l. showed prolonged growth in the adaptation phase.

*Pediastrum* strains from the high Andean lagoons are potential natural ingredients for the cosmetic industry. The strain from 4,400 m.a.s.l. (IMP-BG288) demonstrated the greatest potential due to its high concentrations of antioxidants and pigments. However, the use of UV radiation to achieve an increase in the concentrations of these metabolites is not recommended. Continuous investigation of the strain from 4,400 m.a.s.l. through cosmetic efficacy tests is suggested as this will help in evaluating its application as a cosmetic ingredient.

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# **CONFLICT OF INTERESTS**

The authors report no financial or any other conflicts of interest in this work.

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#### ETHICAL APPROVALS

This study does not involve experiments on animals or human subjects.

# DATA AVAILABILITY

All data generated and analyzed are included within this research article.

# **PUBLISHER'S NOTE**

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