Tritium influence on morphology, reactive oxygen species production and catalase gene expression in *Pseudendoclonium basilense* and *Stigeoclonium nanum* (Chlorophyta)

Petr HAŠLER, Vladan ONDŘEJ, Michaela ŠVÉCAROVÁ, Michaela SEDLÁŘOVÁ, Barbora VAIĐOVÁ & Aloisie POULÍČKOVÁ

Department of Botany, Faculty of Science, Palacký University, Šlechtitelů 27, CZ–783 71 Olomouc, Czech Republic; *Corresponding author e–mail: petr.hasler@upol.cz

**Abstract:** The response of two strains of *Pseudendoclonium basilense* ViscH & Stigeoclonium nanum (Dillwyn) Kötzing to ionizing radiation was studied under laboratory conditions. Both strains were isolated as epiphytic from stagnant water bodies under ambient levels of ionizing radiation and cultivated under laboratory conditions on a gradient of tritiated water diluted in liquid Bold’s basal medium. Exposure to low doses of ionizing radiation stimulated changes in cell dimensions and morphology of in both strains and catalase gene expression in *Pseudendoclonium basilense*. Moreover, the addition of tritiated water stimulated production of reactive oxygen species in *Pseudendoclonium basilense* which concentration decreased after increased gene expression of catalase.

**Key words:** oxidative stress, ionizing radiation, adaptation, tolerance, catalase, nuclear power plant, green algae

**Abbreviations:** (ROS) reactive oxygen species, (RONS) reactive oxygen and nitrogen species

**INTRODUCTION**

Ionizing radiation influences metabolism, ultrastructure, and the biology of organisms both directly and indirectly (KOVÁCS & KERESZTES 2002). The energy of ionizing radiation determines the extent of its impact on biological systems, leading to minor variations, severe damage or even to a cell death. Tritium, a soft β-emitter, is considered as one of the safest radioisotopes with low linear energy transfer causing a low degree of damage (Blaylock et al. 1986). Almost 99% of tritium produced by the action of cosmic rays in the upper layers of atmosphere is transformed into tritiated water which is spread in the surface water layer (CalmON & Garnier–Laplace 2010). Anthropogenic tritium is produced mainly in the form of tritiated water. Large amounts of tritium may be released into the environment from nuclear power plants (Fievet et al. 2013). In the Czech Republic, the Temelin nuclear power plant discharged nearly 30 Tbq of tritium in 2005 (Hanslík et al. 2009).

KIRCHMANN et al. (1979) proved that tritium concentration is equals in the environment, e.g. water, culture medium, and algal biomass. Similarly, the ratio of tritium fixed in algal organic matter (e.g. incorporated in membranes, organelles, structural macromolecules) and within the cytosol is equal. This is because algae persist in a stable state of uptake and release with respect to their short life and simple cell or thallus organization (Harrison 1971). Algae and higher plants can assimilate then incorporate tritium into macromolecules such as proteins and DNA through photosynthesis, especially water photolysis. The intensity of algal assimilation can be expressed as a function of incorporated tritium or specific activity inside cells. For example, *Acetabularia* sp. shows rapid growth after tritium incorporation followed by long and slow growth rate (Bonotto et al. 1982).

Tritium behavior in the environment and its incorporation into algal biomass has been studied both under laboratory conditions and in the field over last 50 years (KIRCHMANN et al. 1979; BELOT 1986; McCUBBIN et al. 2001; Kim et al. 2012; Fievet et al. 2013; Baglan et al. 2013; Jean–Baptiste & Fourre 2013). One can
perceive incorporated tritium, and its ionizing radiation respectively, as a form of abiotic stress. Tritium incorporation into algal DNA can cause chromosomal aberration, breakage, and mutation and subsequently can influence the physiological state of cells, particularly cell stress, biochemical pathways and overall metabolism. Abiotic stress (e.g. heat, salinity, heavy metals, desiccation or radioactive irradiation) can induce the formation of reactive oxygen species (e.g. \( \text{O}_2^·, \cdot \text{OH}, \text{H}_2\text{O}_2 \)) which can cleave biomolecules and also transduce signals. Cell stress responses including variation in metabolism, growth, and development are based on ROS interplay with plant hormones. Studies on higher plants confirmed stress–induced morphological responses to increased concentration of auxins in plant cells (e.g. Potters et al. 2007). Reactive oxygen species could enhance auxin–responsiveness in cell cycle reactivation, re–orientation of growth and polar cell expansion (Pasternak et al. 2005; Mangano et al. 2016). However, the effect of ionizing radiation on both algal cell morphology and ROS production during stress have not been previously studied.

Regulation of oxidative stress tolerance in plant cells, and likely in green algae as well, represents a complex regulatory system among plant hormones and enzymes. This strictly balanced system maintains cell homeostasis and protects against cell damage or death. Catalase (\( \text{H}_2\text{O}_2\text{,} \text{H}_2\text{O}_2 \) oxidoreductase; EC 1.11.1.6) is a tetrameric heme–containing enzyme that converts hydrogen peroxide into oxygen and water thus protecting the cell from the damaging effects of hydrogen peroxide accumulation. These enzymes can catalyze direct degradation of hydrogen peroxide or its depletion by different substrates (e.g., methanol, ethanol, formaldehyde, nitrite) oxidation (Dát et al. 2000; McClung 1997).

The aim of this study is to investigate morphological changes, ROS production, and catalase gene expression in two green algae, *Pseudendoclonium basilense* and *Stigeoclonium nanum*, under a gradient of soft \( \beta^– \) irradiation from tritiated water. Both *P. basilense* and *S. nanum* are periphytic species frequently inhabiting stones or submerged macrophytes in stagnant and flowing waters which can be polluted by human nuclear waste.

**MATERIAL AND METHODS**

**Experimental design.** Two strains of *Pseudendoclonium basilense* (strain Vaidová 2014, isolated as epiphytic, pond near Dolní Benešov, Czech Republic) and *Stigeoclonium nanum* (strain Vaidová 2014/1, isolated as epiphytic, pond near Velká Bystřice, Czech Republic) were maintained for three weeks under laboratory conditions as follows: continuous irradiation of 20 mmol.m\(^{-2}\).s\(^{-1}\), temperature of 15 °C, liquid Bold’s basal medium (BBM; Bold 1949). Harvested biomass (approx. 500 mg) was split into twelve parts which were exposed to tritiated water with activity 0, 100, 500 and 1000 Bq (three replicates) for 21 days. After 1 day of exposure sub-samples (100 mg) were taken and tested for ROS activity and expression of catalase gene. After 21 days following exposure the algal biomass was tested for morphological changes, expression of catalase gene and ROS production.

**Evaluation of morphological variability under a gradient of soft \( \beta^– \) irradiation.** Morphological changes of *P. basilense* and *S. nanum* such as cell shape, dimensions, cell cytology or cell division and reproduction were evaluated after 21 days of exposure as follows: 1) samples were photographed using with Zeiss AxioImager (obj. C–Apochromat 40×/NA. 0.75, HRc camera 13MPS); 2) photographed cultures were analyzed using with Zeiss AxioVision 4.9.1. software; 3) dimensions of one hundred cells (width, length) were measured for all replicates. Descriptive statistics and One Way ANOVA were used for statistical evaluation of taken data (NCSS statistical package, Hintze 2007).

**Quantitative reverse transcriptase polymerase chain reaction (qRT–PCR).** Total RNA was isolated from 100 mg of fresh biomass per each sample of green algae after 1 and 21 days of tritium treatment (control, 100, 500, 1000 Bq) using the Spectrum Plant Total RNA kit (SigmA–Aldrich, St. Louis, MO). The RNA was quantified by spectrophotometric analysis. Prior to amplification all RNA samples were treated with RNase–free DNase (Promega, Madison, WI) to eliminate genomic DNA contamination. The isolated RNA was transcribed to cDNA by Transcriptor High Fidelity cDNA Synthesis Kit (Roche, Basel, Switzerland). Reverse transcription (RT) was carried out in total volume of 20 µl containing 1 µl anchored–oligo(dT)18 primers (30 µmol.l\(^{-1}\)), 9 µl total RNA, 1.4 µl RNase Free dH2O, 4 µl RT reaction buffer (5 times conc.), 0.5 µl protector RNase inhibitor (40 U.µl\(^{-1}\)), 2 µl deoxyribonucleotide mix (10 mmol.l\(^{-1}\)), 1.1 µl reverse transcriptase (20 U.µl\(^{-1}\)) and 1 µl dithiotreitol. The RT reaction was carried out at 45 °C one hour and enzyme was inactivated by heating to 85 °C for 5 minutes. The qRT–PCR was done to analyze catalase expression in the green algae using LightCycler® Fast Start DNA MasterPlus SYBR Green I kit (Roche, Basel, Switzerland) in total volume 20 µl containing 5 µl of complementary DNA (250 ng). The amplification was performed in LightCycler Nano Real– Time PCR System. The appropriate primer sequences are: (forward) 5´– CTC GAA CTA CCG CCA CAT GCG–3´ and (reverse) 5´– GCC GTC TGC AGC AGC TGT TGTC T–3´ for catalase; (forward) 5´– GAA GAC CTT CCG TCG AGG AGG–3´ and (reverse) 5´– CCC ACT CGT TGT CG TAC CA–3´ for GAPDH as a housekeeping gene. These primers were synthetized by Generi Biotech (Hradec Kralové, Czech Republic). The PCR reaction mixture contained forward and reverse primers in final concentration 0.5 µmol.l\(^{-1}\) (1 µl), Master Mix Stimes conc. (4 µl), water PCR grade (10 µl) and 5 µl cDNA. PCR conditions were as follows: pre-incubation at 95 °C for 10 min, followed by 40 cycles of denaturation at 95°C for 10 s, annealing at 49°C for 30 s, extension at 72 °C for 20 s and finally 5 min incubation at 72 °C. The primary data (Cq) were analyzed by LightCycler Nano Software 1.1. Gene expression was normalized per Glyceraldehyde–3–phosphate dehydrogenase as a housekeeping gene. Data were processed by the delta–delta method, compared to the expression levels in control samples. The experiment was repeated three times. Melting curve analysis was carried out to confirm specificity of each product. Melting curve conditions were 60 °C to 97 °C at 0.1 °C.s\(^{-1}\).
Localization of oxygen radicals by confocal microscopy. The free radical sensor 2′,7′-dichlorodihydrofluorescein diacetate (DCDHF DA; Molecular Probes, ThermoFisher Scientific, Inc.), was used for in–vivo imaging of RONS generated and 21 days after application of tritium. After hydrolysis of the diacetate groups by cytosolic esterases the non–fluorescent fluorescein derivative is transformed to fluorescent dichlorofluorescein (DCF) upon reaction with peroxynitrite, hydroxyl radical and peroxyl radical. Formation of DCF (maximum λex = 495 nm, λem = 523 nm) was monitored by confocal laser scanning microscopy.

Pseudendoclonium basilense and Stigeoclonium nanum cells from different treatments were incubated either in the absence or presence of 20 μM DCDHF DA in darkness for 10 minutes. Immediately after staining, the cells were transferred to fresh water and visualized by confocal laser scanning microscopy (Fluoview 1000 unit attached to IX80 microscope; Olympus Czech Group, Prague, Czech Republic). The excitation of fluorochrome was achieved by a 488 nm line of an argon laser and signal was detected by a 505–550 nm emission filter. Chlorophyll autofluorescence was visualized by chlorophyll excitation with 543 nm helium–neon laser, and emission recorded with a 655–755 nm band pass filter. Cell morphology was visualized by a transmitted light detection module with 405 nm excitation using a near ultraviolet diode laser and Nomarski DIC filters. The proper intensity of all lasers was set according to unstained samples at the beginning of each experiment. Microscopic analysis was repeated for two sets of algal samples from independent growth and tritium treatment. Integral distribution of signal intensity (0–4096) in 12–bit microphotographs was evaluated by image analysis software FV10–ASW Viewer 4.0 (Olympus).

RESULTS

Changes in cell dimensions and morphology under gradient of soft β irradiation

Pseudendoclonium basilense showed significant morphological differences along the gradient of soft β–irradiation, both in cell length and width but not in length/width ratio (Figs 1–3, Table 1). However, differences in cell dimensions did not exhibit a dose–dependent pattern to the tritium gradient. The longest and widest cells were found in untreated medium and under activity 1000 Bq, whilst cells under 100 and 500 Bq were the smallest both in length and width. Length/width ratios did not show any significant variation. In contrast to characteristic shape of natural thallus, cultured algae usually formed spherical to oval cells and pseudoparenchymatous aggregates, with rare lateral branches. Control samples most closely resembled the natural appearance of environmental thalli (Fig. 4).

Cells tended to form similar aggregation of spherical or hemispherical cells into irregular colonies with lateral branches. On the other hand, along gradient of β–irradiation cells became more elliptical and the thallus lost its characteristic arrangement (Figs 5–7). No zoospores were produced under tritium treatments.

Stigeoclonium nanum showed more obvious morphological changes under tritium treatment than Pseudendoclonium basilense (Figs 1–14). Untreated samples of both taxa resembled samples collected from natural populations (e.g., field collected)? Increasing activity of tritiated water stimulated cell elongation of S. nanum from 11.99±8.82 (activity 0 Bq) to 23.01±3.85 μm (activity 1000 Bq; Fig. 8, Table 1), along with changes in cell structure (especially development of chloroplasts, which are smaller under 1000 Bq in comparison to 0 Bq of triitated medium) and shape of apical cells (from pointed to rounded or almost spherical; see arrows at Figs 11–14). Cell width was not affected oppositely to length/width ratio which showed increasing trend similarly to cell length (Fig 9–10). Asexual reproduction by zoospores was stimulated only at 500 Bq of soft β–irradiation whilst production of zoospores at 100 and 1000 Bq was rarely observed. Filament branching underwent a conspicuous change from regularly branched filaments at 0 Bq to irregularly branched filaments at 100 and 500 Bq. The highest value of tritiated water activity (1000 Bq) caused stub formation of branches usually consisting of one elongated and rounded cell (Fig. 14). Cells of P. basilense remained spherical or elliptical shape under gradient of tritiated medium. Chloroplasts did not show any marked changes and infilled almost the whole cell volume. In contrast to S. nanum, we did not observed any zoospores or zoosporangia formation.

Effect of tritium on the catalase expression in the green algae

We compared the effect of tritium exposure on the expression of catalase in Pseudendoclonium basilense and Stigeoclonium nanum. We did not find any significant changes in catalase expression in case of 1 day exposure. Conversely, the activity of tritium (100, 500, 1000 Bq) induced synthesis of catalase RNA. After 21 days of incubation, catalase activity in P. basilense was 2.2×, 1.4×, and 5.1× higher than in untritiated medium (see Fig. 15). The highest level (5.1-fold) of catalase RNA was observed under 1000 Bq. No evident changes in catalase expression were found in S. nanum after 21 days of incubation.

Imaging of reactive oxygen species

To elucidate sites of oxidative stress, the fluorescence signal of DCF was visualized within cells of both algal species and complemented by analysis of corresponding integral distribution of DCF signal intensities.

Radicals formed during Pseudendoclonium basilense metabolism were detected in control cells, especially in those dying or undergoing division (Figs 16, 17). Activity of 1000 Bq caused increase of DCF signal when applied for 1 day. Conversely, the signal was decreased after application of 100 Bq and 500 Bq and almost completely abolished in 1000 Bq cells.
Figs 1–7. Evaluation of morphological variability of *Pseudendoclonium basilense* under gradient of tritium activity: (1) variability of cell length, (cont.) untritiated medium, 13.27±2.98 μm, (100) 100 Bq, 12.13±2.10 μm, (500) 500 Bq, 12.35±2.35 μm, (1000) 1000 Bq, 13.59±2.13 μm, F = 5.14, p = 0.0018; (2) variability of cell width, (cont.) untritiated medium, 8.56±1.75 μm, (100) 100 Bq, 7.91±1.28 μm, (500) 500 Bq, 7.64±1.46 μm, (1000) 1000 Bq, 8.10±2.50 μm, F = 2.73, p = 0.045; (3) variability of length/width ratio, (cont.) untritiated medium, 1.61±0.46 μm, (100) 100 Bq, 1.58±0.40 μm, (500) 500 Bq, 1.67±0.43 μm, (1000) 1000 Bq, 2.20±3.11 μm, F = 2.05, p = 0.108; (4) thallus cultivated at activity A = 0 Bq; (5) thallus cultivated at activity A = 100 Bq; (6) thallus cultivated at activity A = 500 Bq; (7) thallus cultivated at activity A = 1000 Bq. Scale bar 10 μm.

Stressed for 21 days (Figs 18, 19). The combination of Nomarski DIC, DCF fluorescence, and chlorophyll fluorescence channels revealed localization of oxidative stress both in chloroplasts and also in cytoplasm. On the other hand, strains of *Stigeoclonium nanum* produced only low amounts of RONS which gave very low DCF signal. No significant changes in the DCF fluorescence signal intensity were detected among different treatments (Figs 20–23).

**DISCUSSION**

Stress, both abiotic and biotic, represents one the most effective tools of selection and evolutionary pressure. Depending on the extent and intensity, a stress factor can affect organisms on local or global level. Ionizing radiation as a global abiotic factor influences organisms during their entire lifespan. Plant reactions to ionizing radiation include death, growth inhibition, or
morphogenetic abnormalities and vary as a function of dose and time (Gunckel et al. 1953; Gunckel & Sparrow 1954). Low energy soft β-irradiation has not been considered as harmful irradiation causing death or significant irreversible changes of organisms. Activities of the tritiated water used in our experiments are similar to surface waters in close proximity to nuclear power plants (NPP) in the Czech Republic (~10 – 500 Bq.l⁻¹, data from website Monitoring of Radiation Situation, Czech Office for Nuclear Safety) and in freshwater in rivers downstream of NPP (HanSlík et al. 2009). These levels of radiation have not been believed harmful for organisms living in such environment. Thus, our results evoke a question as to whether these low levels of activity really are safe or whether they can cause stress, which would inevitably lead to adaptation or death. In our study, we found changes in algae exposed to tritiated water on both morphological and
molecular levels. The most noticeable were changes in the cell shape and cell dimensions. Previous, similar investigations on vascular plants (i.e., apples, bananas) showed structural changes of cells, such as in cell walls and cytoplasm with organelles. Irradiated cell walls usually lose firmness because of pectin degradation (Kovács & Kereczs 2002). Thus, short stump–like branches and spherical to globose apical cells in algae can be related to irregular growth because of changes of cell wall structure. Similarly, irradiated chloroplasts of vascular plants exhibit changes their ultrastructure, dilatations of intra thylakoid space, and loss of granal stacking (Kovács & Kereczs 2002). The strain of Stigeoclonium nanum exhibited the most visible changes of chloroplasts especially at 1000 Bq. On the other hand, P. basilense did not show any obvious changes in chloroplast shape and arrangement. It seems that morphological answer to ionizing radiation at low level of activity is specific depending on particular species and its adaptability.

Ionizing radiation plays an important role in cell stress, and may be moderated through the network of metabolic pathways. Reactive oxygen species such as H$_2$O$_2$, O$_2^*$ or ·OH play important roles in signaling pathways (e.g. Azzam et al. 2012; Tognetti et al. 2012; Krishnamurthy & Rathinasabapathi 2013). At low levels, ROS can act as secondary messengers that transfer stress signals and facilitate responses to various environmental conditions or stress. The concentration of ROS depends on both production and scavenging efficacy, influenced by internal and external stimuli. ROS production is rapid and localized, initiated immediately after stimulus action. The degree of reaction depends on individual factors, such as intensity and timing. For example, Arabidopsis thaliana can produce ROS in waves within as little as 10 minutes following wounding (Mittler et al. 2011). Réty et al. (2012) observed peak of ROS production between 14 and 15 min after tritiated water exposure in the alga Chlamydomonas reinhardtii, followed by a rapid decrease and leveling off within 45 minutes. Chlamydomonas likely reduced ROS production due to activity of protective enzymatic system. Neither cell growth or size of C. reinhardtii was affected by the addition of tritiated water. Herein, both tested strains P. basilense and S. nanum showed stress responses within 1 day after exposure. However, only P. basilense was stimulated by the addition of tritiated water for ROS production. On the other hand, S. nanum was unaffected by tritiated water. In both of our strains, ROS concentrations later equilibrated to a constant level within 21 days after inoculation of both strains, similar to that seen in C. reinhardtii (Réty et al. 2012). Changes of ROS concentrations indicate a capability to respond to environmental stress. In our study, P. basilense exhibited a low but permanent response to tritiated water via upregulation of catalase gene expression. This adaptation led to decreasing of ROS concentration in cells making new balanced homeostasis as found in Chlamydomonas reinhardtii. On the other hand, S. nanum exhibited changes in cell morphology (e.g., cell...
elongation and the presence of spherical apical cells), possibly due to the role of auxins. Direct water radiolysis via ionizing radiation is known to induce the production of both ROS species and H$_2$O$_2$ (Bensasson et al. 1993). ROS are known to be involved in both cell wall loosening and reinforcement. Plant peroxidases utilize ROS and H$_2$O$_2$ to produce phenolic radicals in cell walls and consequently create cross-links via these radicals. Cross-linking of the cell wall components leads to cell wall tightening and could result in a growth restriction (Kärkönen & Kuchitsu 2015). We speculate that similar process might contribute to H$_2$O$_2$ removal from $S$. nanum cells. If expression of the catalase gene was not upregulated then the altered thickness of the cell wall could lead to changes in cell shape.

Moreover, ROS, together with Ca$^+$ and pH, were reported to sustain polar growth in plant cells over the time (Mangano et al. 2016) and thus affect development of Stigeoclonium nanum fibers through changes in cell wall dynamics.

Studies (Pasternak et al. 2002, 2005) have demonstrated that mild oxidative stress could mimic the effect of auxin on plant development. For example, inhibition of catalase activity, as well as direct H$_2$O$_2$ treatment, of Lycium barbarum calli stimulated embryogenesis (Cui et al. 1999) in a manner similar to auxin application. Auxins also inhibit shoot branching and promote lateral root formation (Pasternak et al. 2002) and polar auxin transport is thought to play a key role in such processes. Auxin transport has recently
been described in the green alga Chara (ZHANG & VAN DUIN 2014). We hypothesize that oxidative stress induced by tritiated water treatment also mimics auxins which could disturb polar auxin transport in S. nanum leading to changes in branching.

The level of tritiated water in our study was low without perceivable harmful or damaging effects. The two different algal species reacted differently to low dose exposure to tritiated water. They exhibited different morphological responses and gene expression as adaptive stress response. Thus, soft β radiation should be considered as low energy radiation capable of causing cell stress which influences cell morphology, metabolism, and gene expression.

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