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Nanomaterials include all substathis size, the characteristics of differ substantially from macroto the sidelines. Although, the gearth crust can lead to suppose a unique charateristics of nanopar. The industrial development couraise their environmental level nanoparticles in certain regions years, since its reactivity with biphysicochemical factors such as

Thus, the main objective of the antioxidant response of *S. ce* nanoparticles (TiO2-NPs). Cells (w/v) glucose at 28 °C are experiment at 40 °C. Samples from each used for protein content, DPPH, determinations.

The results show that the prese evidenced by a decrease of pr 3.1.3.1) activity, loss of redox GSH+GSSG contents and GSH loss of ability to scavenge free I lipoxygenase (LOX, EC 1.13.11 a loss of antioxidant power mec slowdown of β -oxidation. Final (CAT T, EC 1.11.1.6) in cells exthis enzyme activity in cells expe a loss of proliferative capacity b for 1 mg/mL TiO₂-NPs level, ap

Keywords: yeast; alkaline phosphatase

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fluence of titanium dioxide nanoparticles in Saccharomyces cerevisiae BY4741?

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contain nanoscale structures sized between 1 and 100 nm. At s change: their strength, conductivity, and reactivity, which m-sized materials, shifting the rules of physics and chemistry 1 origin and the ubiquitous occurrence of nanoparticles in the nylogenetic adaptation of living beings to such substances, the IPs) bring a new dimension to environmental effects testing. I vast new applications of nanomaterials, have contributed to 1 because, concern over the environmental pressure of the rld as well as its effects on the biosphere has grown in recent es mainly depends on the surface area/molecular size ratio and emperature.

was to evaluate how heat-shock affects cell survival and BY4741, a Eurocast strain, exposed to titanium dioxide nential phase were inoculated in liquid YEPD medium 2 % .1 or 1.0 μ g/mL NP-TiO₂ prepared by sonication, during 200 nt were used to obtain the post-12000 g fractions, which were ne antioxidant capacity and, ALP, catalase and LOX activities

iO₂-NPs in the culture medium induced cell death, response e capacity detected by the alkaline phosphatase (ALP, EC pacity mediated by glutathione, evidenced by a decrease of atio. On the other hand, cell death also appears depend on the estimated by DPPH method. We also observed an increase of /ity, a marker of lipid peroxidation, which may be related with peroxisomal catalase (CAT A, EC 1.11.1.6), probably due a observed an increase of the antioxidant cytoplasmic catalase concentrations of 0.1 mg/mL, but a significantly decrease of mg/mL TiO₂-NPs. This apparently bimodal response indicates we process when the level exposure was 0.1 mg/mL. However, occur a transition for necrosis.

ne; lipoxygenase; catalase

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