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and cytoplasmic catalases by lindane may cause ecies in *Saccharomyces cerevisiae*

T. Pita¹, I. Alves-Pereira^{1,2} and **R.** ¹Departmento de Química, Escola de C Évora, Portugal

Decline in glutathione per

an increase of reactive oxy

²Instituto de Ciências Agrárias e Ambie Évora, Portugal

Lindane or gamma $1\alpha, 2\alpha, 3\beta, 4\alpha, 5\alpha$, and aquifers, lipophilic, chemically commonly used on a wide variety of (with fungicides) as a seed treatme control of lice and mites (scabies) and mammals have been in the new clarify the toxicological mechanism was to evaluate the effects of line musts, Portugal, a unicellular eukar

Cells at mid-exponential phase wer 72 h in a water bath with orbital sha from each treatment were used to used for determination of reactive (glutathione peroxidase (GPx) [3], catalase (CAT T) [4] activities as w by spectrophotometry.

The results show that lindane inhib produced along 72 h, as well as cel the ROS content of post-12,000 g cells subjected to any exposure co which has become the detoxificati without significant changes in the a cerevisiae exposed to lindane, as w or cells signaling pathways that ass

Keywords: organochlorine; gluthation

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nlorocyclohexane is an organochlorine insecticide, persistent in soils ihemically inert that accumulates along the human food chain. It is n warehouses, in public health to control insect-borne diseases, and ne is also presently used in lotions, creams, and shampoos for the s. Several chemicals as lindane, toxic for aquatic organisms, birds y, since the European Union intend to ban it. Therefore it is urgent to compound in eukaryotic cells. Thus the main purpose of this work te wine wild-type *Saccharomyces cerevisiae* UE-ME₃ of Alentejo nism, described as resistant to the presence of pesticides or metals.

ted in YEPD medium with 2 % (w/v) glucose and incubated during 8 °C, in the absence or in presence of 5 and 50 μ M lindane. Samples owth curves, wet weight and to prepare post-12000 g supernatant, ecies (ROS) [1] by fluorimetry and alkaline phosphatase (ALP) [2], -dependent glutathione peroxidase (Se-GPx) [3] and cytoplasmic let for determination of peroxisomal catalase (CAT A) [4] activities

growth of *S. cerevisiae* UE-ME₃, causing a decrease in the biomass from 24 h of assay. On the other hand, was detected an increase in of cells exposed to 5 μ M lindane and post-12000 g supernatant of eventually conditioned by a decline in GPx and CAT T activities, lrogen peroxide less effective. The increase in the CAT A activity 8e-GPx activities justified, in part, the increase in ROS levels of S. loss of cell viability due to inadequate response of glutathione cycle iosynthesis.

se; yeast

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