

# THE IN VITRO EFFICACY OF AN ANTI-MYCOTOXINS AGENT AGAINST ERGOT ALKALOIDS AND OTHER MYCOTOXINS

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## INTRODUCTION

The contamination of crops with mycotoxins is a major worldwide challenge in livestock production. **Ergot alkaloids** are mycotoxins mainly produced by fungal *Claviceps* species that usually affect cereals and can cause adverse health effects in animals, standing out their neurotoxic effects.

**Fumonisins and aflatoxins** are generated by other mycotoxin-producing fungi, contaminating cereals. **Sterigmatocystin** (STC) is a precursor for the synthesis of aflatoxin B1 and has been associated with several toxic effects, such as hepatotoxicity, nephrotoxicity and pulmonary injuries.

# OBJECTIVE

The aim of this study was to evaluate the *in vitro* efficacy and the selectivity (no adsorption of vitamins or amino acids) of an anti-mycotoxins agent based on selected binding material combined with natural extracts and yeasts.

### MATERIALS AND METHODS





## CONCLUSIONS

The anti-mycotoxins agent tested in this study is selective and has a high efficacy to mitigate the ergot alkaloids, fumonisins, aflatoxins and STC, being a promising strategy to reduce the negative impact of mycotoxins.





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