# AFLATOXIN IN FEED, FARM GATE MILK AND SELECTED **COMMERCIALLY AVAILABLE MILK**

#### Lishia Daya Khilosia, Suretha De Kock, Michael Francis Dutton, Mulunda Mwanza

University of Johannesburg | Department of Food Technology | Email: lishia@ingredientworx.com or surethad@uj.ac.za

#### INTRODUCTION

#### METHODS

Mycotoxins are low molecular weight, non-antigenic fungal secondary metabolites, which are formed at the end of the exponential growth phase of an organism, and make no contribution to the development or metabolism of the producing organism. (Frazier & Westhoff, 1988). Mycotoxins are able to develop at any stage from farm to fork as shown in Figure 1.

The following diagram outlines the sampling procedure and methods of analysis employed for this study.

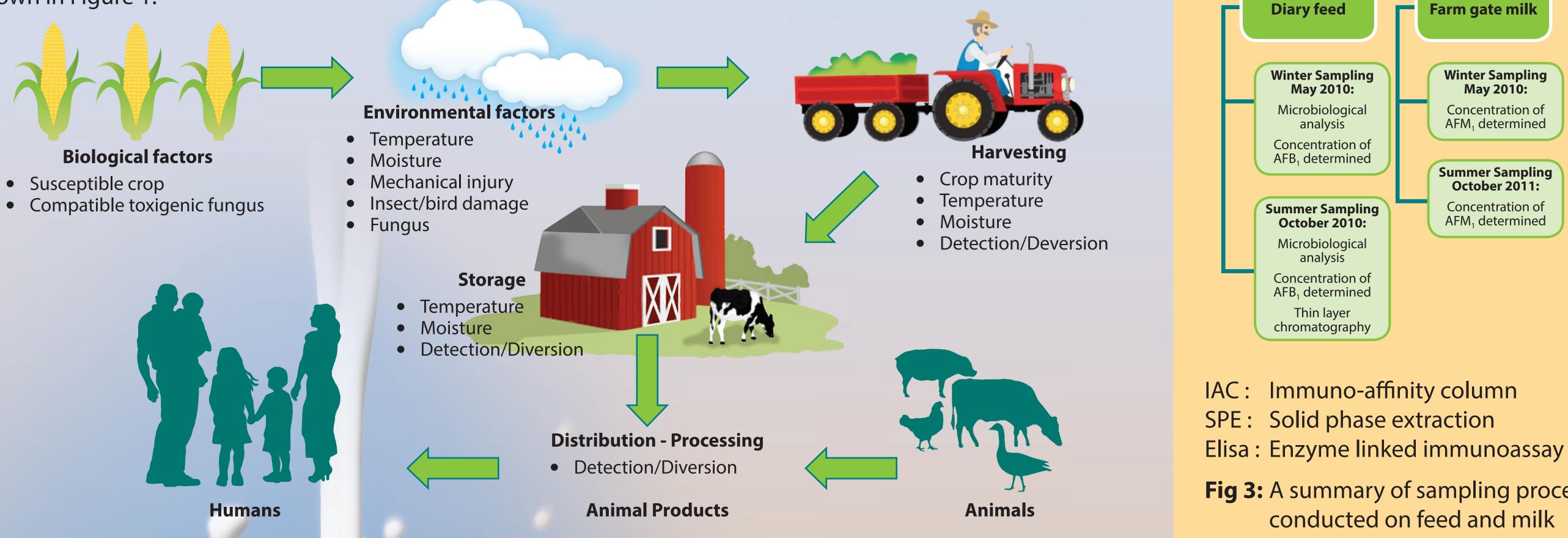


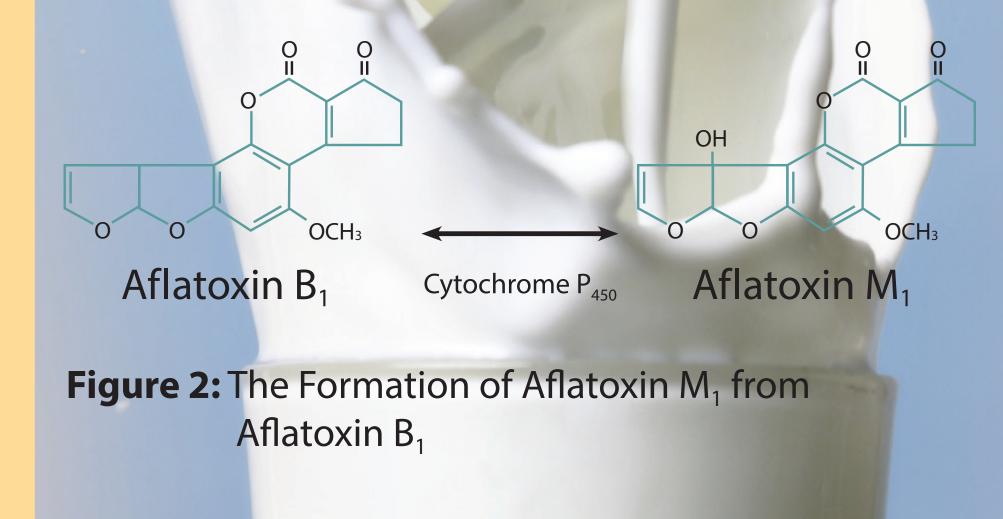
Figure 1: Factors which affect mycotoxin occurrence in the food chain (CAST, 2003).

Aflatoxin B<sub>1</sub> is a mycotoxin found in most feeds and foods and is highly carcinogenic (Eaton and Gallagher 1994) and once ingested by ruminants, it is converted to aflatoxin M<sub>1</sub> (AFM<sub>1</sub>), a carcinogen under the influence of the cytochrome P<sub>450</sub> oxidase system found in rumen micro-flora (Yoshikawa et al. 1982) (Figure 2).

The graph below demonstrates that all the feed samples tested for Aflatoxin B<sub>1</sub> exceeded the permitted level as prescribed by South African legislation

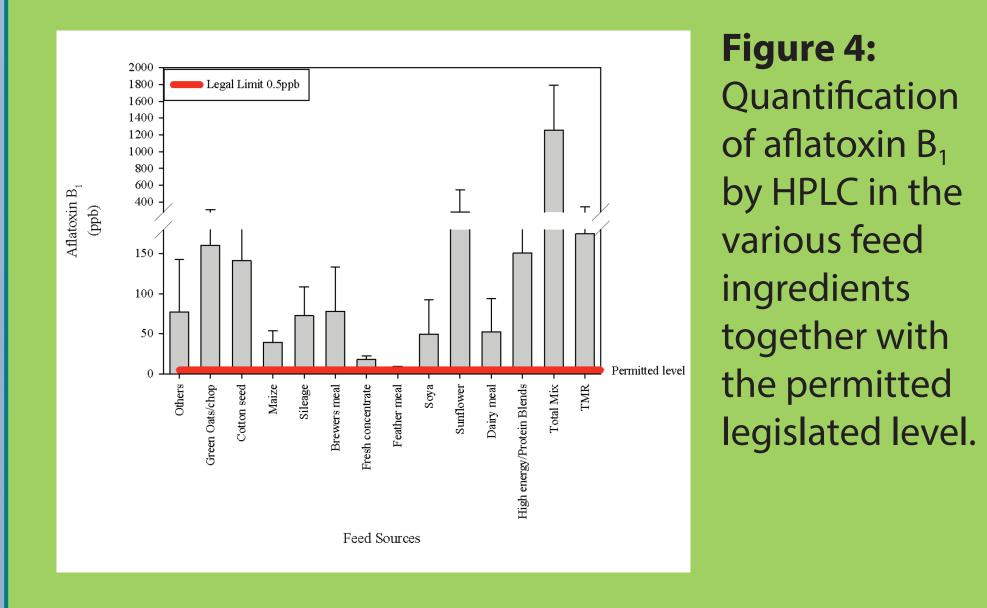
**Retail milk** surveillance **Diary feed** Farm gate milk study Winter Sampling Winter Sampling Winter Sampling May 2010: May 2010: **April 2010: Concentration of** Concentration of Microbiological analysis AFM<sub>1</sub> determined AFM<sub>1</sub> determined Concentration of AFB<sub>1</sub> determined **Summer Sampling Summer Sampling October 2011:** September 2010: Concentration of Concentration of **Summer Sampling** AFM<sub>1</sub> determined October 2010: AFM<sub>1</sub> determined Microbiological analysis **Summer Sampling** Concentration of September 2010: AFB<sub>1</sub> determined Various methods Thin layer investigated for chromatography analysis of AFM<sub>1</sub>: SPE IAC: Immuno-affinity column ELISA SPE: Solid phase extraction

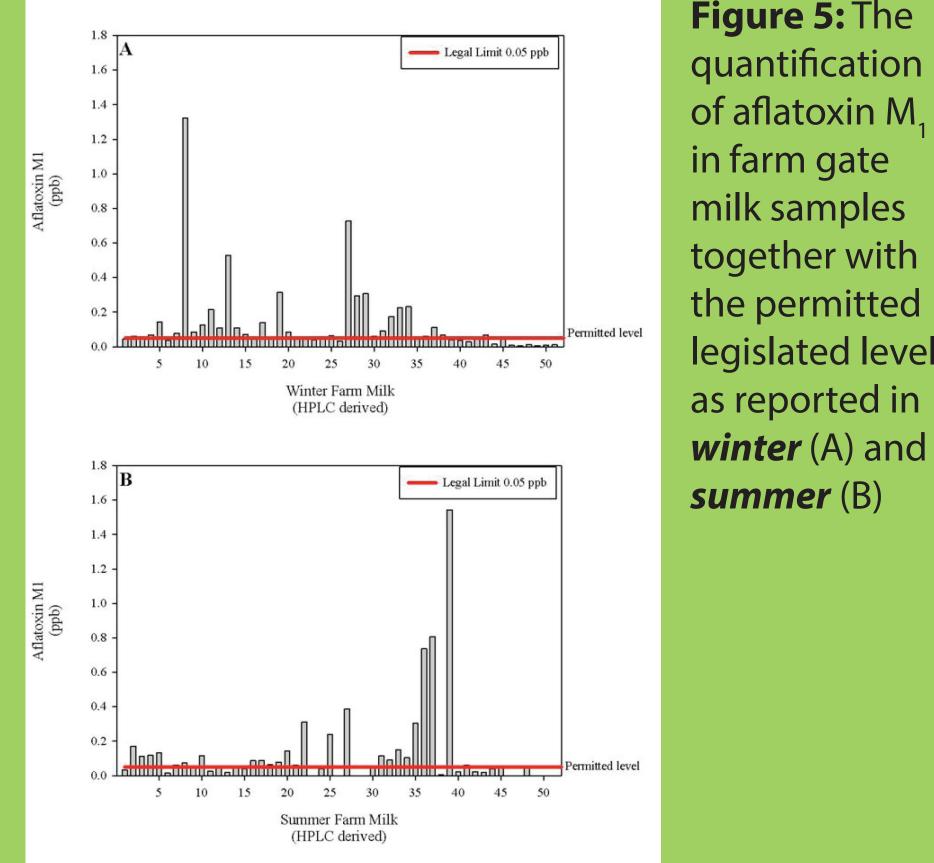
**Fig 3:** A summary of sampling procedure and analysis conducted on feed and milk

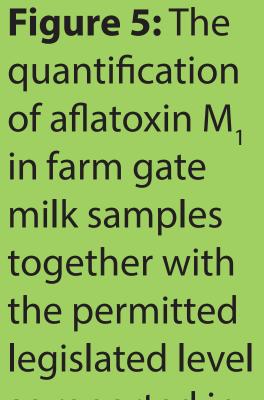


## OBJECTIVES

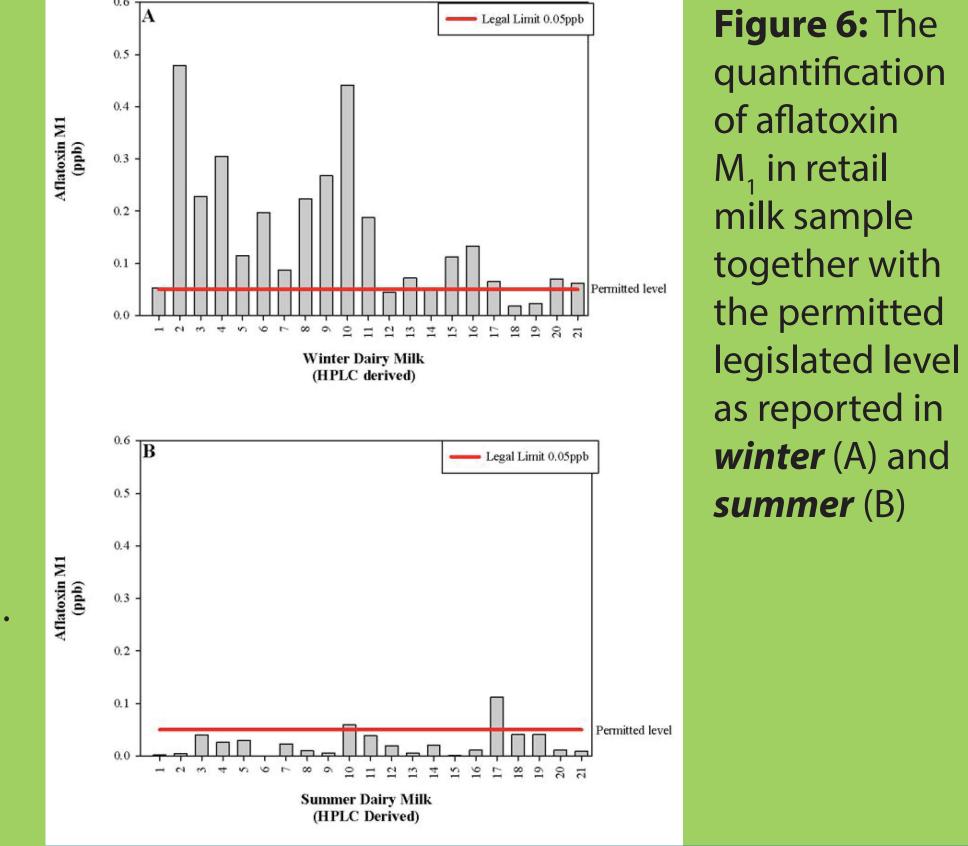
- To conduct a surveillance study of selected retail milk samples
- To determine levels of AFB<sub>1</sub> in feed and AFM<sub>1</sub> in milk from selected farms in Kwa-Zulu Natal







RESULTS



### CONCLUSION

There are serious concerns regarding the levels of AFB<sub>1</sub> contamination in the dairy feed supply chain which contributes to the presence of AFM<sub>1</sub> in commercially availablemilk.Farmersneedtobeeducatedremycotoxins their prevention, control and effects. Intervention from

#### REFERENCES

CAST. (2003). Mycotoxins: Risk in plant, animal and human systems. Ames IA Council for Agricultural Science and Technology.

Eaton, D.L. & Gallagher, E.P. (1994). *Mechanisms of aflatoxins carcinogenesis*. Annual Review of Pharmacology and Toxicology **34** pp 135-172.

Frazier W.C., Westhoff D.C. (1988). Micro-organisms important in Food Microbiology. In: Food Microbiology. Fourth edition. McGraw Hill International, pp 17-32.

Yoshikawa, H., Uchimaru, R., Kamataki, T. Kato, R. & Ueno, Y. (1982). Metabolism and activation of aflatoxin  $B_1$  by reconstituted cytochrome  $P_{450}$  system of rat liver. Cancer Research **42** pp 1120-1124.

#### government and import officials is needed to ensure stringent quality control over imported materials. UNIVERSITY **JOHANNESBURG**