Screening for colorectal cancer risk biomarkers related to diet

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Background: Red and processed meat are associated with high risks of colorectal cancer due to the endogenous formation of O\(^6\)-carboxymethyl guanine (O\(^6\)CMG), a potent carcinogen. The aim of our research is to develop liquid chromatography tandem mass spectrometry (LC-MS/MS) analytical methods for the measurement of the DNA adducts, such as O\(^6\)CMG and its nucleoside O\(^6\)-carboxymethyl deoxyguanosine (O\(^6\)CMdG), in urine samples and correlate it to different diets.

Methods: Urine samples were collected from volunteers on three different diets (vegetarian as the control, with red and processed meat as the experimental group) over a period of 15 days at the Medical Research Council, Cambridge. Samples were analysed by LC-MS/MS either by direct injection or using a column-switching system with an on-line solid phase extraction (SPE) column.

Results: An LC-MS/MS method was developed and used initially to monitor and quantify O\(^6\)CMdG and O\(^6\)CMG using standards in synthetic urine. O\(^6\)CMG elutes at 4.7 min and have a limit of detection (LOD) of 0.3ng/mL, and O\(^6\)CMdG elutes at 14.1 min and have a LOD of 0.03ng/mL. The LC-MS/MS direct injection analysis of the clinical samples showed low sensitivity and the need for sample clean-up.

Conclusions: An efficient method for the separation and quantification of O\(^6\)CMdG and O\(^6\)CMG was developed. An on-line SPE column system is under development to allow an efficient and rapid processing of a large number of clinical urine samples.

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