Winter Meeting, 15 December 2009, Food supply and quality in a climate-changed world

Anthocyanin-rich berry-extract treatment decreases expression of dietary glucose transporter genes in human intestinal Caco-2 cells

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Berries are a rich source of polyphenolic compounds such as flavonoids, including anthocyanins, with wide-ranging potential therapeutic effects⁽¹⁾. It has been demonstrated previously that intestinal uptake of dietary glucose is impeded by the presence of polyphenolic compounds; however, little is known regarding the genes involved in this mechanism⁽²⁾. The aim of this study was to investigate the effects of an anthocyanin-rich berry-extract on dietary glucose transporting genes in Caco-2 cells.

Human intestinal Caco-2 cells were cultured for 19 d and were then treated for 16 h with an anthocyanin-rich berry-extract (OptiBerry; InterHealth Nutraceuticals, Benicia, CA, USA) at a final concentration of 0.5% (w/v). Subsequently mRNA was isolated and used for the quantitative real-time polymerase chain reaction (qRT-PCR), using 18S and GAPDH as housekeeping genes. Gene expression data are expressed as means (\pm sem) relative expression ratios of control; n = 5/6.

The expression of GLUT2 (apical/basolateral monosaccharide transporter), GLUT5 (apical fructose transporter) and SGLT1 (apical sodium/glucose co-transporter) was decreased by the berry-extract treatment. GLUT2, GLUT5 and SGLT1 expression as a ratio of the control was as follows: GLUT2 (0.20 \pm 0.02; $P = 1.9 \times 10^{-4}$), GLUT5 (0.57 \pm 0.08; $P = 2.5 \times 10^{-3}$) and SGLT1 (0.45 \pm 0.03; $P = 3.1 \times 10^{-2}$).

GLUT2, GLUT5 and SGLT1 have previously been identified as therapeutic targets for dysregulated glucose metabolism. In this study, the expression of these genes in Caco-2 cells was decreased by treatment with an anthocyanin-rich berry-extract. Studies are in progress to investigate the biological relevance of the observed effects in relation to berry consumption and the absorption of dietary sugar.

- 1. Zafra-Stone S, Yasmin T, Bagchi M et al. (2007) Mol Nutr Food Res 51, 675-683.
- 2. Kwon O, Eck P, Chen S et al. (2007) FASEB J 21, 366–377.