Letter to the Editor Obesity and related medical conditions: a role for functional foods

Sir.

Dietary fat intake, especially saturated fat, has long been associated with both coronary heart disease risk factors and obesity^{1,2}. In addition, consumption of high levels of sugar and other refined carbohydrates has been reported to cause an increase in blood triglycerides³. High triglyceride levels in the blood have also been associated with coronary heart disease and hypertension^{4,5}. Hypertension is the most common cardiovascular risk factor in the USA^{6,7}, and a number of studies have shown unequivocally a powerful association between high blood pressure and risk of cardiovascular disease^{8,9}. Dietary salt reduction has been recommended to reduce hypertension and cardiovascular disease mortality and morbidity^{10,11}.

In an attempt to reduce the impact of a negative diet on heath status, we have developed a functional food designed to reduce selected risk factors associated with obesity and coronary heart disease. A functional food is one that has demonstrated a significant effect on the body beyond adequate nutrition, in a way that improves health and well-being and reduces disease. The developed product is based on one of the UK's best-selling biscuits, which traditionally is high in both fat and sugar (hence high in energy) but contains little fibre. The first part of the study was to modify the ingredient profile of the standard product. This included increasing fibre while reducing both fat and sugar levels and removing salt to the lowest level. This was undertaken in conjunction with maintaining commercial viability.

Biscuits are traditionally high in both fat (including saturated fat) and sugar, and the biscuit market is dominated by biscuits having fat levels in excess of 20%.

The base biscuit (jam not included) contains no sugar (100% reduction) and the fat level has been reduced by 66%, based on the standard product. Salt has been completely removed from the modified biscuit formulation.

A number of studies have reported an association between low dietary fibre intake and risk of coronary heart disease 12,13 . The level of fibre in the modified biscuit is double that found in the standard product (3.4 vs. 1.7 g). Vitamins were also added to the biscuit in an attempt to reduce plasma levels of homocysteine. Homocysteine is one of the more contemporary risk factors that have been identified as an independent risk factor for coronary heart disease $^{14-16}$. Vitamin B_6 , B_{12} and folic acid are factors that

can modify plasma homocysteine levels¹⁷ and were added to the biscuit to help reduce homocysteine concentrations.

The second part of the study was designed to ascertain and acquire consumer acceptability, which was achieved by relevant organoleptic assessment. We used consumers who regularly ate the standard product and asked them to eat the modified and standard product in a randomised double-blind trial. No preference (P > 0.05) was shown for either of the biscuits, and it was concluded that the modified product was acceptable to the consumer.

The final part of the biscuit design was confirmation of the functionality of the product by biomedical assessment. This part of the study was a randomly designed, 12 h fasting study to measure differences in blood levels of a number of risk factors associated with heart disease. A two-way repeated analysis of variance was used for normally distributed data and the Wilcoxon matched pairs signed rank test was applied to data that were not normally distributed. We found a significant decrease in homocysteine, glucose and, surprisingly, lipoprotein(a) (P < 0.05) following consumption of the modified biscuit (see Table 1). The lower level of serum homocysteine is consistent with the findings of earlier reports relating to the effects of dietary folate, vitamins B_6 and $B_{12}^{16,17}$.

We are currently investigating further the relationship between the biscuit and lipoprotein(a). We can think of no content in the biscuit that would have caused a decrease in this individual cardiovascular risk factor other than the probiotic. This is of particular interest to us because there has been little experimentation investigating the influence of diet on this lipid–protein complex. Since its discovery by Berg in 1963, high levels of blood lipoprotein(a) have been shown to be associated with heart disease ^{18–20} and, to date, niacin appears to be the only effective treatment in the control of lipoprotein(a) concentrations ^{21–23}.

Biscuits are typically high in both fat and sugar and have been identified as a food contributing to negative health. They are consumed by a large part of the UK population and are popular with both children and adults. Biscuits are therefore a good choice, and one example of many popular food products that can be healthily redesigned and consumed favourably as opposed to discouraged in society. We are aware of the food industry's attempts to modify the food content of certain high-risk foods to make them more acceptable to the ever-increasing health demands of society. To our knowledge, no food has been as vigorously changed as the biscuit we describe

Letter to the Editor 1329

Table 1 Homocysteine and lipoprotein(a) concentrations before (pre) and after (post) consumption of modified (mod) and non-modified (non-mod) biscuits

Variable	Pre (mod)	Post (mod)	Pre (non-mod)	Post (non-mod)
Homocysteine (μ I ⁻¹)	15.3 ± 6.9*	11.2 ± 4.3*	12.4 ± 5.9	11.9 ± 1.5
Lipoprotein (a) (mg l ⁻¹)	193.5 ± 76.8*	149.6 ± 27.5*	251.5 ± 97.9	259 ± 103

Values are means ± standard deviation.

whilst still maintaining its taste and attractiveness to the consumer.

Table 1 presents the results for homocysteine and lipoprotein(a) concentrations (n=10) following a 12 h, randomly assigned, fasting study when the modified biscuits were compared with the non-modified product. Significant reductions (P < 0.05) in concentrations were observed after consumption of the modified biscuit only. Pre indicates blood values prior to biscuit administration. Post indicates blood values following the intervention.

Wyndham J Boobier*, Julien S Baker and Bruce Davies

Health and Exercise Science Research Unit School of Applied Sciences University of Glamorgan, Pontypridd, CF37 1DL, UK

*Email: wboobier@glam.ac.uk DOI: 10.1079/PHN2005832

References

- 1 Keys A. Coronary heart disease in seven countries. *Circulation* 1970; **41**: 11–211.
- 2 Gotto AM Jr. High density lipoprotein cholesterol and triglycerides as therapeutic targets for preventing and treating coronary artery disease. *American Heart Journal* 2002; 144: S33–S42.
- 3 Yudkin J, Kang SS, Bruckdorfer KR. Effects of high dietary sugar. British Medical Journal 1980; 281: 1396.
- 4 Jeppesen J, Hein HO, Saudicani P, Gyntelberg F. High triglycerides/low high-density lipoprotein cholesterol, ischemic electrocardiogram changes, and risk of ischemic heart disease. *American Heart Journal* 2003; **145**: 103–8.
- 5 Zieden B, Kaminskas A, Kristenson M, Olsson AG, Kucinskiene Z. Long chain polyunsaturated fatty acids may account for higher low-density lipoprotein oxidation susceptibility in Lithuanian compared to Swedish men. Scandinavian Journal of Clinical and Laboratory Investigation 2002; 62: 307–14.
- 6 Nisbett EF, Rossiter AH, Miller AR, Thacker D. Fat is short dough biscuits. Flour Milling and Baking Research Association 1986; (2): 63–77.

7 Davis MM, Jones DW. The role of lifestyle management in the overall treatment plan for prevention of management of hypertension. *Seminars in Nephrology* 2002; **22**: 35–43.

- 8 Franklin SS, Khan SA, Wong ND, Larson MG, Levy D. Is pulse pressure useful in predicting risk factors for coronary heart disease? The Framingham Heart Study. *Circulation* 1999; **100**: 354–60.
- 9 Van den Hoogen PC, Feskens EJ, Nagelkerke NJ, Menotti A, Nissinen A, Kromhout D. The relation between blood pressure and mortality due to coronary heart disease among men in different parts of the world. Seven Countries Study Research Group. New England Journal of Medicine 2000; 342: 1–8
- 10 He J, Ogden LG, Vupputuri S, Bazzano L, Loria C, Whelton PK. Dietary sodium intake and myocardial infarction. *Journal of the American Medical Association* 1999; 282: 2027–34.
- 11 Krauss RM, Deckelbaum RJ, Ernst N, Fisher E, Howard BV, Knopp RH, et al. Dietary guidelines for healthy American adults: a statement for health professionals from the nutrition committee, American Heart Association. Circulation 1996; 94: 1795–800.
- 12 Pietinen P, Rimm EB, Korhonen P, Hartman AM, Willett WC, Albanes D, et al. Intake of dietary fibre and risk of coronary heart disease in a cohort of Finnish men. The alpha-Tocopherol Beta-Carotene Prevention Study. Circulation 1996; 94: 2720-7.
- 13 Rimm EB, Ascherio A, Giovannucci E, Spiegelman D, Stampfer MJ, Willett WC. Vegetable, fruit and cereal fibre intake and risk of coronary heart disease among men. *Journal of the American Medical Association* 1996; 275: 447–51
- 14 McCully KS. Vascular pathology of homocysteinemia: implications for the pathogenesis of arteriosclerosis. *American Journal of Pathology* 1969; **56**: 111–28.
- 15 Klerk M, Verhoef P, Verbruggen B, Schouten EG, Blom HJ, Bos GM, et al. Effect of homocysteine reduction by B vitamin supplementation on marker of clotting activation. *Throm*bosis and Haemostasis 2002; 88: 230–5.
- Billion S, Tribout B, Cadet E, Queinnec C, Rochette J, Wheatley P, et al. Hyperhomocysteinemia, folate and vitamin B₁₂ in unsupplemented haemodialysis patients: effect of oral therapy with folic acid and vitamin B₁₂. Nephrology, Dialysis, Transplantation 2002; 17: 455–61.
- 17 Genest F Jr. Homocysteine: to screen and treat or wait and see? *Canadian Medical Association Journal* 2000; **163**: 37–8
- 18 Hoefler G, Harnoncourt F, Pashcke E, Mirtl W, Pfeiffer KH, Kostner GM. Lipoprotein(a) Lp(a). A risk factor for myocardial infarction. Arteriosclerosis 1988; 8: 394–401.
- 19 Sandkamp M, Funke H, Schulte EK, Assman G. Lipoprotein(a) is an independent risk factor for myocardial infarction at a young age. *Clinical Chemistry* 1990; **36**: 20–3.
- 20 Rosengren A, Wilhelmsen L, Eriksson E, Risberg B, Wedel H. Lipoprotein (a) and coronary heart disease: a prospective case control study in a general population sample of middleaged men. *British Medical Journal* 1990; 301: 1248–51.
- 21 Batiste MC, Schaefer EJ. Diagnosis and management of lipoprotein abnormalities. *Nutrition in Clinical Care* 2002; **5**: 115–23.
- 22 Pieper JA. Understanding niacin formulations. American Journal of Managed Care 2002; 8: \$308-14.
- 23 Pan J, Van JT, Chan RL, Lin M, Charles MA. Extended-release niacin treatment of the atherogenic lipid profile and lipoprotein (a) in diabetes. *Metabolism* 2002; **52**: 1120–7.

^{*} Significant at P < 0.05.