

Long term dietary supplementation with microalgae increases plasma docosahexaenoic acid in milk and plasma but does not affect plasma 13, 14-dihydro-15-keto PGF_{2α} concentration in dairy cows

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SUPPLEMENTARY FILE

Supplementary Material and Methods

The FA content of the TMR was determined according to Jenkins (2010), whilst milk FA analysis followed the method described by Feng et al. (2004) for lipid extraction, with the methylation of the lipids conducted as described by Lock et al. (2006). Individual FAME were determined by GLC (Hewlett Packard 7820, Wokingham, UK) fitted with a CP-Sil 88 column (100 m×0.25 mm i.d.×0.2 µm film, Agilent Technologies, Santa Clara, California, USA) as described previously (Lock et al. 2006). Total lipid was extracted from plasma samples and methylated using the method of Burdge et al. (2005). Resulting FA methyl esters were resolved on a CP-Sil 88 column using a gas chromatograph (GC; Bruker 350, Bruker, Germany) equipped with a flame ionisation detector. The GC conditions were as previously described (Kliem et al., 2013), and plasma FAME were identified based on retention time comparisons with an authentic standard (GLC463, Nu-Chek Prep Inc., Elysian, MN), and cross-referencing with previously published chromatograms (Kliem et al., 2013). Carbon deficiency in the detector response was accounted for using a combined correction factor (Ulberth et al., 1999), and results were expressed as g/100 g total FA.

References

Burdge GC, Tricon,S, Morgan,R. Kliem,KE, Childs C, Jones E, Russell JJ, Grimble RF, Williams CM, Yaqoob P & Calder PC 2005 Incorporation of *cis*-9, *trans*-11 conjugated linoleic acid and vaccenic acid (*trans*-11 18:1) into plasma and leucocyte lipids in healthy men consuming dairy products naturally enriched in these fatty acids. *British Journal of Nutrition* **94** 237-243

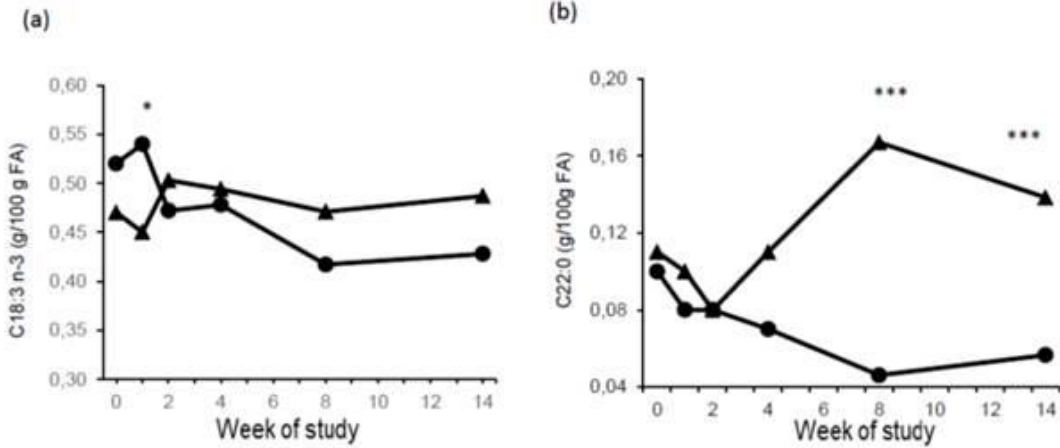
Feng S, Lock AL & Garnsworthy PC 2004 Technical note: A rapid lipid separation method for determining fatty acid composition in milk. *Journal of Dairy Science* **87** 3785-3788

Jenkins TC 2010 *Technical note*: Common analytical errors yielding inaccurate results during analysis of fatty acids in feed and digesta samples. *Journal of Dairy Science* **93** 1170-1174

Kliem, KE, Shingfield, KJ, Livingstone, KM & Givens, DI 2013 Seasonal variation in the fatty acid composition of milk available at retail in the United Kingdom and implications for dietary intake. *Food Chemistry* **141** 274-281

Lock AL, Teles BN, Perfield II, Bauman DE & Sinclair LA 2006 A conjugated linoleic acid supplement containing *trans*-10, *cis*-12 reduces milk fat synthesis in lactating sheep. *Journal of Dairy Science* **89** 1525- 1532

Supplementary Fig. 1. Milk fat concentration of (a) C18:3 n-3 and (b) C20:0. SED = 0.034, 0.022 respectively. Within time points, treatments that differ at $P < 0.05$ or $P < 0.001$ are denoted by *, or *** respectively.



Supplementary Fig. 2. Blood plasma fat concentration of (a) C18:0 (b) C18:1 *trans*-10 (c) C18:3 n-3, (d) C20:4 n-6 (e) C20:0 (f) total saturated fatty FA and (g) total n-3 PUFA in dairy cows fed no SCIM (Control ●) or 100 g per cow per day of microalgae (SCIM ▲). SED = 0.44, 0.066, 0.19, 0.09, 0.05, 0.68 and 0.34 respectively. Within time points, treatments that differ at $P < 0.05$, $P < 0.01$ or $P < 0.001$ are denoted by *, **, or *** respectively.

