



### FACULTY OF BIOSCIENCE ENGINEERING

# Stable carbon isotope fractionation may be diet dependent

S. De Smet<sup>1</sup>, K. Raes<sup>1</sup>, E. Claeys<sup>1</sup> and P. Boeckx<sup>2</sup>

<sup>1</sup>Laboratory for Animal Nutrition and Animal Product Quality, Ghent University, Proefhoevestraat 10, 9090 Melle, Belgium <sup>2</sup>Laboratory of Applied Physical Chemistry-ISOFYS, Coupure Links 653, 9000 Gent, Belgium

# Conclusions

Stable carbon isotope fractionation may depend on the diet. Across trials, there was a significant linear decrease in fractionation with increasing (less negative) dietary  $\delta^{13}$ C values.

# Background

□ Stable isotope analysis of animal tissues has potential for discriminating between diets, e.g. the  $\delta^{13}$ C value is known to reflect the proportion of C<sub>3</sub> and C<sub>4</sub> plants in the diet.

 $\Box$  During metabolism, depletion of  $^{13}C$  occurs (fractionation), resulting in  $\delta^{13}C$  values in animal samples that differ from the corresponding dietary values (trophic shift).

□ This trophic shift is often assumed to be constant for a given tissue.

#### Objective

 $\square$  Verify whether the fractionation of  $^{13}C$  in plasma is constant or depends on the dietary  $\delta^{13}C$  value

#### Results

 $\Box$  Across trials and treatments, mean plasma  $\delta^{13}C$  fractionation was +1.4‰, ranging between -1.3‰ and +3.8‰.

 $\Box$  There was a significant linear decrease in fractionation with increasing (less negative) dietary  $\delta^{13}C$  values and increasing proportions of C<sub>4</sub> plant material in the diet.

 $\Box$  However, this dietary effect on plasma  $\delta^{13}C$  fractionation appeared in two trials, wheras it was not obvious in two other trials.

□ A non-constant trophic shift may have implications for backcalculation of diets, as was shown by Focken (2004).



<u>Figure 1.</u> Plasma  $\delta^{13}C$  values in relation to dietary  $\delta^{13}C$  values

#### References

Balcaen et al. (2003). Proc. Annual Meeting BSAS, p. 159. De Smet et al. (2004). Rapid Communications in Mass Spectrometry 18: 2087-2092. Focken (2004). Rapid Communications in Mass Spectrometry 18: 1227-1232.

# Material and methods

□ Data from four trials with different animal types and diets differing in the proportion of C<sub>4</sub> plant material (maize). Values are treatment means. □ Determination of  $\delta^{13}$ C values of plasma and dietary samples by ANCA-SL elemental analyser, connected to an isotope ratio mass spectrometer (PDZ-Europa, UK). Ratios  $^{13}C/^{12}$ C are expressed as  $\delta^{13}$ C values in per mill (‰) relative to VPDB standard. Values for fractionation = plasma – dietary values. □ Plasma samples had been taken after at least two months following a dietary change, so that equilibrium could be assumed.

Trial	Animal type	Group	# Animals	%C <sub>4</sub> in diet	Diet	
•	Adult wethers	1	4	0	Ryegrass hay only (1) or with	
(Balcaen et al., 2003)		2	4	46	maize silage (2)	
	Growing bulls	1	16	0	High concentrate diets differing	
(De Smet et al., 2004)		2	7	13,5	in content of grass/maize silage	
		3	8	35	concentrates	
	Growing lambs	1	4	0	Hay/concentrate (50/50) without	
(unpublished)		2	4	17,8	(1) or with maize (2)	
•	Growing pigs	1	6	0	Fattening diet without (1) or with	
(unpublished)		2	6	15	maize (2)	



<u>Figure 2.</u> Plasma  $\delta^{13}$ C fractionation in relation to % C<sub>4</sub> plant material in diet (upper graph) and dietary  $\delta^{13}$ C value (lower graph)

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Laboratory for Animal Nutrition and Animal Product Quality http://www.lanupro.UGent.be – Stefaan.DeSmet@UGent.be

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