

Effect of lipid-rich plant extract on the fatty acids composition and meat quality of Belgian-Blue cross bred steers

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Introduction We have previously demonstrated the ability of a lipid-rich plant extract to enhance levels of polyunsaturated fatty acid (PUFA) in beef (Kim *et al.*, 2008). The plant extract (PX) is developed from the liquid fraction extracted from fresh lucerne (*Medicago sativa* L.), and then heat-treated and dried. The PX has a high degree of rumen protection. This study investigated the effect of incremental inclusion of PX in steers fed on grass silage relative to concentrate on the fatty acid composition of beef and meat quality. Effects of additional vitamin E within the concentrate were also investigated.

Materials and methods Following summer grazing on perennial ryegrass/white clover swards, forty Belgian-Blue × Holstein steers (~ 400 kg liveweight) were allocated to one of five dietary treatments: 1) grass silage *ad libitum* (GS), 2) grass silage *ad libitum* plus 75 g PX/dry matter intake (DMI; GS-LPX), 3) grass silage *ad libitum* plus 150 g PX/DMI (GS-HPX), 4) restricted barley straw and control concentrate (40:60 on a DM basis; S-CC), and 5) restricted barley straw and concentrate with PX (25% in concentrate on a DM basis) (40:60 on a DM basis; S-PXC) and additional vitamin E (~ 300 mg/kg). The PX was manufactured by Désialis–France Luzerne, France. Liveweight was monitored every 28 days and the liveweight gain was used to modify feed intake of the S-CC and S-PXC animals to achieve a similar growth rate to those fed on *ad libitum* forage. Animals were slaughtered when they achieved fat class 3 and samples of *longissimus thoracis et lumborum* were taken at 48 h post-mortem for fatty acid analysis, vitamin E analysis, 10-d aged samples for shelf life studies in modified atmosphere packs. An analysis of variance was conducted with diet as the main factor using GenStat (11th edition) statistical software.

Results Liveweight gain, conformation score and fatness score were similar across diets averaging 1.1 kg/d, 76.7 and 56.5, respectively. Small differences in carcass weight were noted (Table 1). Total fatty acids and amounts of the major saturated fatty acids were not different (Table 1). Feeding grass silage relative to concentrate increased deposition of n-3 relative to n-6 PUFA. Incremental PX on grass silage resulted in additional deposition of 18:3n-3 (and 18:2n-6) and longer chain derivatives EPA and DHA resulting in improvements in P:S and n-6:n-3 ratio. Additional vitamin E in the diet of S-PXC increased its content in muscle impacting on lipid stability (TBARS).

Table 1 Animal performance, fatty acid composition (mg/100 g muscle) of *longissimus thoracis et lumborum*, and TBARS and vitamin E content of muscle in Belgian-Blue cross bred steers given experimental diets

	GS	GS-LPX	GS-HPX	S-CC	S-PXC	s.e.d.	P
Right-side cold carcass (kg)	169 ^{ab}	174 ^b	170 ^{ab}	177 ^b	165 ^a	3.8	0.032
Total fatty acids	2551	2510	2433	2532	1999	379.3	NS
16:0	665	623	596	654	488	107.6	NS
18:0	325	332	339	344	274	57.2	NS
18:1n-9	880	850	794	825	581	134.8	NS
CLA (<i>cis</i> -9, <i>trans</i> -11)	12.9	14.4	14.1	14.7	13.4	2.46	NS
18:2n-6	56.7 ^a	70.2 ^{ab}	72.7 ^b	121.3 ^c	134.5 ^c	7.37	<0.001
18:3n-3	26.9 ^a	38.3 ^b	41.3 ^b	13.8 ^a	35.7 ^b	3.05	<0.001
20:5n-3 (EPA)	14.3 ^{bc}	16.1 ^{cd}	18.3 ^d	9.8 ^a	12.3 ^b	0.79	<0.001
22:6n-3 (DHA)	2.42 ^{bc}	2.28 ^b	2.77 ^c	1.58 ^a	1.63 ^a	0.176	<0.001
P:S ratio	0.08 ^a	0.11 ^{ab}	0.13 ^b	0.13 ^b	0.23 ^c	0.018	<0.001
n-6:n-3 ratio	1.83 ^b	1.69 ^{ab}	1.62 ^a	6.28 ^c	3.36 ^c	0.049*	<0.001
Vitamin E (mg/kg muscle)	3.86 ^{ab}	4.41 ^b	4.16 ^b	3.17 ^a	7.75 ^c	0.373	<0.001
TBARS, d10 (mg/kg muscle)	0.59 ^{ab}	0.79 ^b	1.41 ^c	1.39 ^c	0.28 ^a	0.236	<0.001

TBARS=Thiobarbituric acid reactive substances; *on log scale; Means within a row with different superscripts differ (P<0.05).

Conclusions Feeding incremental PX to grass silage-fed animals resulted in enhancement of 18:3n-3 and increased the longer chain derivatives EPA and DHA, resulting in improved P:S and n-6:n-3 ratios. Within the concentrate treatments PX also increased 18:3n-3 and EPA. The highest P:S ratios were noted on S-PXC treatment reflecting the high levels of 18:2n-6 and 18:3n-3 deposited. Under these circumstances the additional vitamin E fed helped to control oxidative stability as reflected in the lower TBARS.

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References

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