



The effect of α - and γ -tocopherol on lipid oxidation and lipid stability of meat in broiler chickens

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INTRODUCTION

α -tocopherol

- antioxidant activity, used as a supplement
- NRC recommendations: 10 IU (10 mg α -tocopherol acetate = 6.7 mg RRR- α -tocopherol)

γ -tocopherol

- antiinflammatory activity, important in immune processes
- abundant in feed
- activity \rightarrow 10 % as α -tocopherol

\rightarrow Synergy??? \leftarrow

MATERIALS AND METHODS

46 one day old broilers \rightarrow 5 experimental groups

Table 1: Fat and vitamin E supplementation of the experimental diets

GROUP		SOURCE OF FAT	VITAMIN E SUPPLEMENTATION
Cont-	(N=10)	5% PALM FAT	10 mg α -tocopherol acetate/kg
Cont+	(N=10)		10 mg α -tocopherol acetate/kg
α	(N=10)	5% LINSEED OIL	67 mg RRR- α -tocopherol/kg
γ	(N=8)		67 mg RRR- γ -tocopherol/kg
α + γ	(N=8)		33.5 mg RRR- α -tocopherol + 33.5 mg RRR- γ -tocopherol/kg

ANALYSES:

- Oxidative stress *in vivo* \rightarrow lymphocyte DNA damage, plasma malondialdehyde (MDA), ferric reducing capacity (FRAP), antioxidant capacity of lipid soluble compounds (ACL)
- Plasma, breast and thigh muscle vitamin E concentrations
- Lipid stability of meat \rightarrow MDA in breast and thigh muscles stored under different conditions:
 - fresh
 - 2 months at -20°C
 - 6 days at 4°C
 - 4 months at -20°C

RESULTS

Table 2: Effect of dietary α - and γ -tocopherol on lymphocyte DNA damage (OTM), plasma malondialdehyde concentration (MDA), ferric reducing capacity (FRAP) and antioxidant capacity of lipid soluble compounds (ACL)

	Cont-	Cont+	α	γ	α + γ	Significance	SEM
OTM	3.16 ^{ab}	4.41 ^b	2.92 ^a	2.83 ^a	2.88 ^a	*	0.35
MDA (nmol/ml)	0.28 ^a	0.73 ^c	0.50 ^b	0.73 ^c	0.48 ^b	***	0.06
FRAP (nmol/ml)	685.3 ^a	679.9 ^a	908.0 ^b	650.9 ^a	779.7 ^{ab}	**	55.0
ACL (nmol/ml)	118.9 ^a	125.2 ^a	163.8 ^b	125.6 ^a	151.0 ^{ab}	***	11.1

^{abc} Least square means within a line with unlike superscripts differ significantly ($P < 0.05$).

¹OTM Olive Tail Moment (the product of the amount of DNA in the tail and the mean distance of migration in the tail)

Table 3: Effect of dietary α - and γ -tocopherol on concentrations of α - and γ -tocopherol in plasma, breast and thigh muscles

	Cont-	Cont+	α	γ	α + γ	Significance	SEM	
Plasma ($\mu\text{g/ml}$)	α -tocopherol	8.19 ^a	5.79 ^a	39.85 ^c	3.62 ^a	19.91 ^b	***	1.80
	γ -tocopherol	0.50 ^a	0.29 ^a	0.31 ^a	1.05 ^b	1.07 ^b	***	0.08
Breast ($\mu\text{g/g}$)	α -tocopherol	3.06 ^a	3.50 ^a	14.92 ^c	1.95 ^a	7.38 ^b	***	0.67
	γ -tocopherol	0.24 ^a	0.21 ^a	0.26 ^a	1.00 ^c	0.85 ^b	***	0.04
Thigh ($\mu\text{g/g}$)	α -tocopherol	9.94 ^a	13.48 ^a	48.89 ^c	6.79 ^a	26.29 ^b	***	2.33
	γ -tocopherol	0.84 ^a	0.93 ^a	0.88 ^a	3.46 ^b	3.06 ^b	***	0.13

^{abc} Least square means within a line with unlike superscripts differ significantly ($P < 0.05$).

CONCLUSIONS

- High levels of polyunsaturated fatty acids \rightarrow \uparrow lipid oxidation
- The retention of vitamin E \rightarrow α -tocopherol $>$ γ -tocopherol
- Lipid oxidation \rightarrow \downarrow in α -tocopherol, but not in γ -tocopherol supplemented group
- DNA damage was prevented in all supplemented groups.
- The combination of both antioxidants showed some synergistic effects.

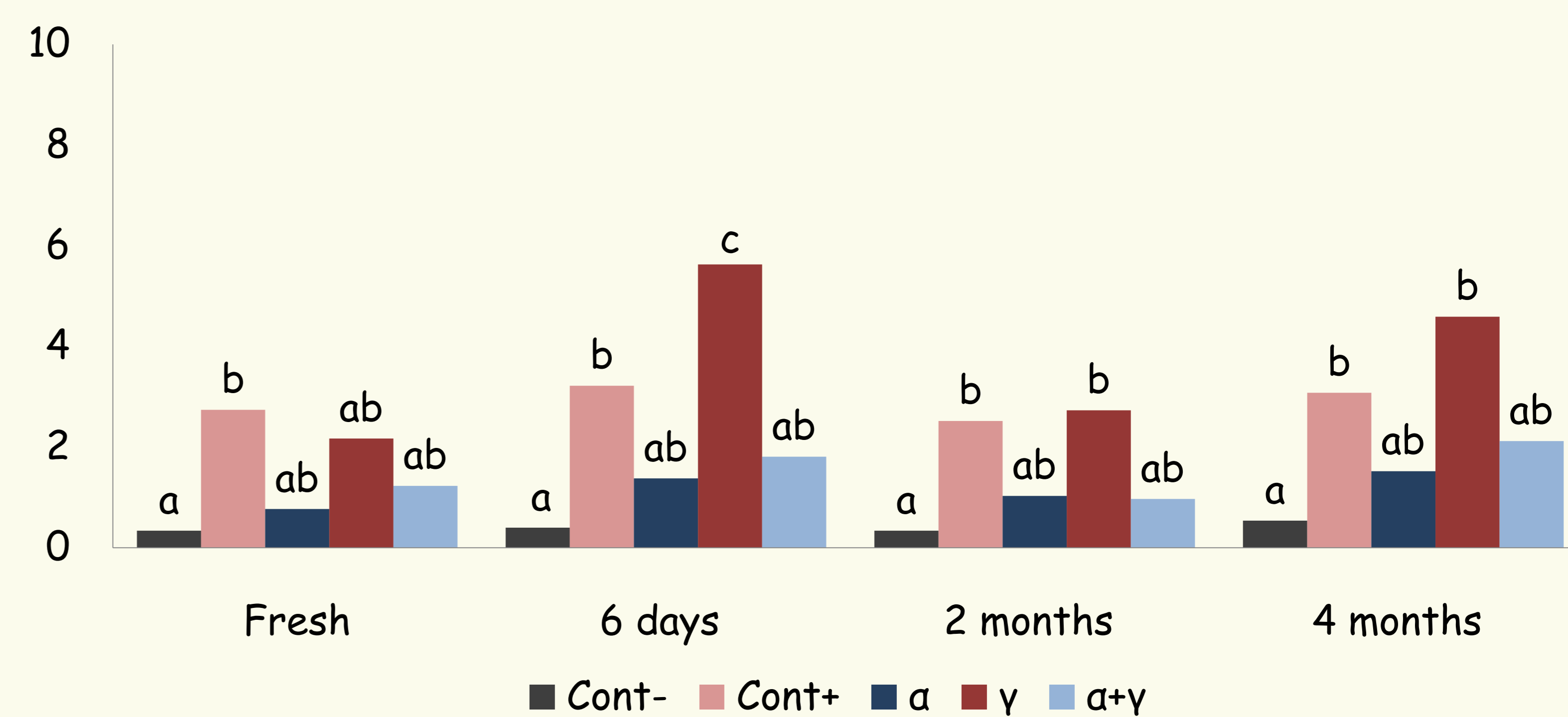


Figure 1: Effect of dietary α - and γ -tocopherol on malondialdehyde concentration (nmol/g) in breast muscle under different storage conditions

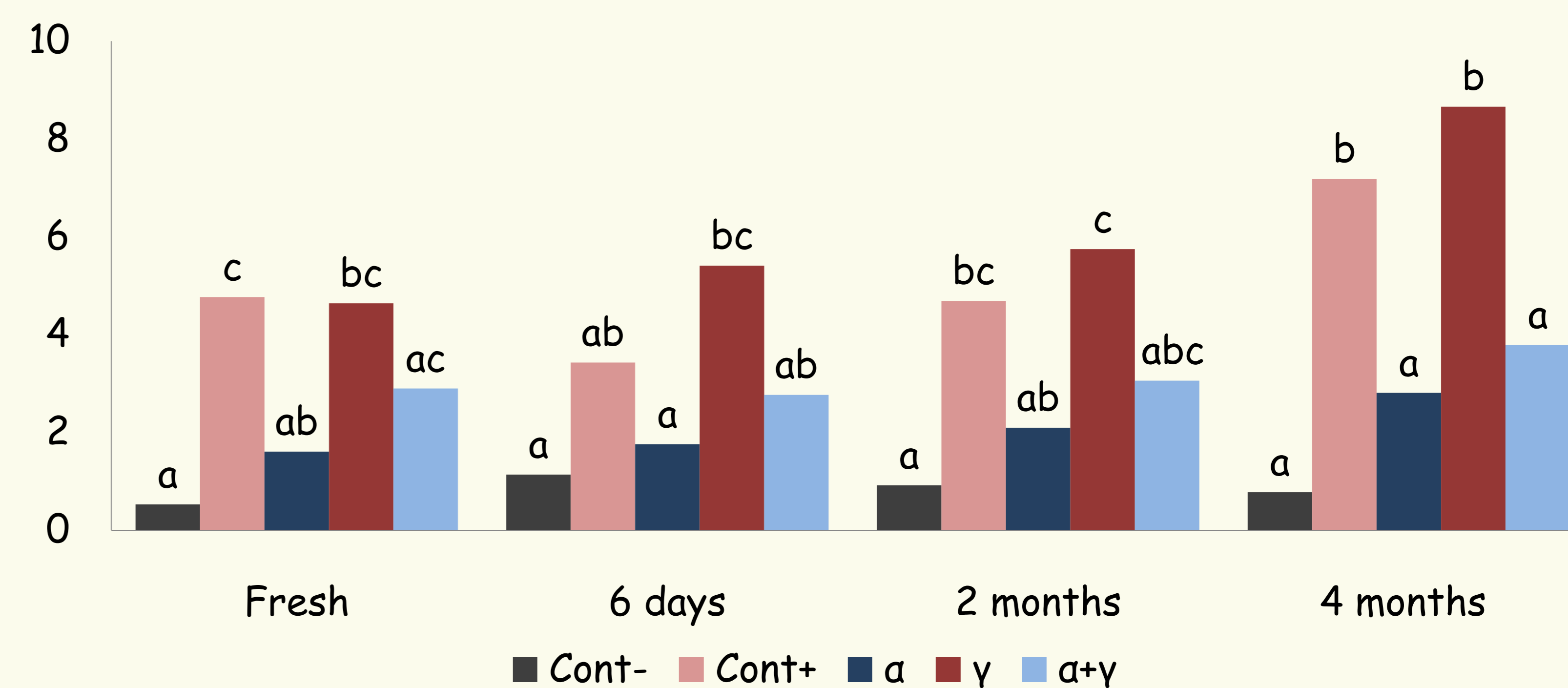


Figure 2: Effect of dietary α - and γ -tocopherol on malondialdehyde concentration (nmol/g) in thigh muscle under different storage conditions