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# Effects of dietary saturated and n-6 polyunsaturated fatty acids on the incorporation of long-chain n-3 polyunsaturated fatty acids into blood lipids

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## Abstract

**Background/objectives:** Omega-3 polyunsaturated fatty acids (n-3PUFA) are better absorbed when they are combined with high-fat meals. However, the role of different dietary fats in modulating the incorporation of n-3PUFA in blood lipids in humans has not been previously explored. Omega-6 polyunsaturated fatty acids (n-6PUFA) are known to compete with n-3PUFA in the metabolic pathways and for the incorporation into phospholipids, whereas saturated fats (SFA) may enhance n-3PUFA incorporation into tissues.

**Subjects/methods:** In a randomized parallel-design trial, we aimed to investigate the long-term effects of n-3PUFA supplementation in subjects consuming a diet enriched with either SFA or n-6PUFA on fatty acid incorporation into plasma and erythrocytes and on blood lipid profiles (total cholesterol, low-density lipoprotein cholesterol (LDL-C), high-density lipoprotein cholesterol (HDL-C) and triglycerides).

**Results:** Dietary supplementation with n-3PUFA co-administered with SFA for 6 weeks resulted in a significant rise in total cholesterol ( $0.46 \pm 0.60$  mmol/L;  $P=0.020$ ) and LDL-C ( $0.48 \pm 0.48$  mmol/L;  $P=0.011$ ) in comparison with combination with n-6PUFA. The diet enriched with SFA also induced a greater increase in eicosapentaenoic acid ( $2.07 \pm 0.79$  vs  $1.15 \pm 0.53$ ;  $P=0.004$ ), a smaller decrease in docosapentaenoic acid ( $-0.12 \pm 0.23$  vs  $-0.30 \pm 0.20$ ;  $P=0.034$ ) and a similar increase in docosahexaenoic acid ( $3.85 \pm 1.14$  vs  $3.10 \pm 1.07$ ;  $P=0.128$ ) percentage in plasma compared with the diet enriched with n-6PUFA. A similar effect was seen in erythrocytes. N-3PUFA supplementation resulted in similar changes in HDL-C and triglyceride levels.

**Conclusions:** The results suggest that dietary substitution of SFA with n-6PUFA, despite maintaining low levels of circulating cholesterol, hinders n-3PUFA incorporation into plasma and tissue lipids.

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