- Interventional Longitudinal Study on The Effects of Long-Chain Omega-3 Docosahexaenoic Acid on Platelet Function and **Redox Status**

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Last update: 09/05/2017 | Version: 2 | ID: 4789

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Identification

Detailed name Interventional Longitudinal Study on The Effects of

Long-Chain Omega-3 Docosahexaenoic Acid on

Platelet Function and Redox Status

CNIL registration number, number and date of CPP agreement, AFSSAPS (French Health Products Safety Agency) authorisation

CCPPRB LYON A (03/11/2004).

General Aspects

Medical area Hematology

Health determinants Nutrition

Keywords omega-3 fatty acids, docosahexaenoic acid, platelet

function, redox status, healthy males, intake

Scientific investigator(s)

(Contact)

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Mixed
INSERM, grant from the Lipids and Nutrition Group (Neuilly-sur-Seine, France), grant from the French Ministry of Research and Education. Funding from the Lipids and Nutrition Group.
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Public
Study databases
Longitudinal study (except cohorts)
A selection of health institutions and services
No

sample selection.

Sample Sciedalom	
Database objective	
Main objective	To determine the effects of increased intake of docosahexaenoic acid on platelet function and oxidative stress in a group of healthy male volunteers.
Inclusion criteria	Healthy male volunteers.
Population type	
Age	Adulthood (45 to 64 years)
Population covered	General population
Gender	Male
Geography area	Local
French regions covered by the database	Auvergne Rhône-Alpes
Detail of the geography area	Lyon.
Data collection	
Dates	
Date of first collection (YYYY or MM/YYYY)	2005
Date of last collection (YYYY or MM/YYYY)	2007
Size of the database	
Size of the database (number of individuals)	< 500 individuals
Details of the number of individuals	12 (2014)
Data	
Database activity	Data collection completed
Type of data collected	Clinical data Biological data
Clinical data (detail)	Medical registration

Details of collected clinical data	
Biological data (detail)	Blood lipid assay, study of platelet function and redox status parameters.
Presence of a biobank	No
Health parameters studied	Others
Other (detail)	
Procedures	
Data collection method	Blood and urine samples collected before and after each supplementation dose and 8 weeks after arrest of supplementation. Supplementation: daily intake of successive doses: increments of 200, 400, 800, and 1600 mg DHA (Decola, Maldegem, Belgium) for 2 weeks for each dose without interruption.
Participant monitoring	Yes
Details on monitoring of participants	8 weeks after treatment.
Links to administrative sources	No
Links to administrative sources Promotion and access	No
	No
Promotion and access	No http://tinyurl.com/Hal-Effets-chaine-omega3
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