Effect of AlKylGlycerols on angiogenic and inflammatory factors in Obese individuals

Submission date	Recruitment status	Prospectively registered		
20/02/2013	No longer recruiting	☐ Protocol		
Registration date	Overall study status	Statistical analysis plan		
04/04/2013	Completed	[X] Results		
Last Edited	Condition category	[] Individual participant data		
17/01/2019	Nutritional, Metabolic, Endocrine			

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s)

Scientific

Contact name

Dr Maria Isabel Covas

Contact details

IMIM Institut Hospital del Mar d'Investigacions Mèdiques Cardiovascular Risk and Nutrition Group Dr. Aiguader, 88 Barcelona Spain 08003

Additional identifiers

Protocol serial number 2006/2601/I/AGL2006-02031

Study information

Scientific Title

Effect of AlKylGlycerols on angiogenic and inflammatory factors in Obese individuals: a randomized double-blind cross-over clinical supplementation trial

Acronym

Study objectives

A rich alkylglycerol (AKG) preparation will be more effective on inflammatory markers than a low AKG preparation.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Clinical Research Ethical Committe of the "Institut de Recerca Hospital del Mar (IMIM)", 25 February 2008, ref: 2006/2601/I

Study design

Randomized double-blind cross-over single centre clinical supplementation trial

Primary study design

Interventional

Study type(s)

Treatment

Health condition(s) or problem(s) studied

Community-dwelling obese participants with a body mass index (BMI) \geq 30 kg/m2 to \leq 40 kg/m2

Interventions

Randomized, double-blind, cross-over, single centre clinical supplementation trial, in which 24 subjects (12 men and 15 women) will be randomised to one of two orders of administration of:

- 1. High alcoxyglycerol content (HAC: 20 mg AKG/ 6 ml olive oil)
- 2. Low alcoxyglycerol content (LAC: 10 mg AKG/ 6 ml olive oil)

There will be in three intervention periods of 3 weeks separated by 2-week wash-out periods avoiding shark oil or AKG supplements.

Intervention Type

Drug

Phase

Not Applicable

Drug/device/biological/vaccine name(s)

Alkylglycerol from olive oil

Primary outcome(s)

- 1. Inflammatory parameters measured at baseline and before and after each intervention:
- 1.1. Serum inmunoglobulines A, G, and M
- 1.2. Complement 3(C3) and 4 (C4)
- 1.3. Plasma interleukine-8 (IL8), monocyte chemotactic protein-1(MCP-1), and vascular endothelial growth factor (VEGF) .

Key secondary outcome(s))

Oxidative damage: Plasma lipids and lípid peroxides will be determined at baseline and 3 weeks after each intervention period.

Completion date

20/12/2012

Eligibility

Key inclusion criteria

- 1. Obese patients
- 2. Body mass index (BMI) \geq 30 kg/m2 to \leq 40 kg/m2
- 3. Aged 20 60 years, either sex

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Adult

Sex

All

Key exclusion criteria

- 1. Intake of antioxidant supplement or acetylsalicylic acid or any other drug with established antioxidative properties
- 2. Athletes with physical activity (PA >3000 kcal/day in leisure-time
- 3. BMI < 30 kg/m2 or > 40 kg/m2)
- 4. Diabetes, multiple allergies, intestinal diseases
- 5. Any condition that limits the mobility of the subject making study visits impossible
- 6. Life threatening illness such as cancer or severe disease with a lowered expected 3 year survival
- 7. Any other disease or condition that would worsen the adherence to the measurements or treatment

Date of first enrolment

02/05/2012

Date of final enrolment

20/12/2012

Locations

Countries of recruitment

Spain

Study participating centre
IMIM Institut Hospital del Mar d'Investigacions Mèdiques
Barcelona
Spain
08003

Sponsor information

Organisation

Spanish Ministry of Economy and Competitiveness (Spain)

Funder(s)

Funder type

Government

Funder Name

Spanish Ministry of Science and Technology (Spain)

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary

Not provided at time of registration

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article	results	01/06/2016	17/01/2019	Yes	No
Participant information sheet	Participant information sheet	11/11/2025	11/11/2025	No	Yes