

Comparative efficacy and tolerability of a novel botanical formulation LI12507F in the treatment and control of obesity

Submission date

09/04/2010

Recruitment status

No longer recruiting

☐ Prospectively registered

☐ Protocol

Registration date

30/07/2010

Overall study status

Completed

☐ Statistical analysis plan

☒ Results

Last Edited

20/01/2014

Condition category

Nutritional, Metabolic, Endocrine

☐ Individual participant data

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s)

Scientific

Contact name

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Contact details

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520007

Additional identifiers

Protocol serial number

10-002/Ob/Sp

Study information

Scientific Title

Comparative efficacy and tolerability of a novel botanical formulation LI12507F in the treatment and control of obesity: a randomised, double-blind placebo controlled clinical study

Acronym

Anti-obese Nutraceuticals

Study objectives

Supplementation of herbal formulations might be useful for management of body weight in obese human subjects.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Internal Review Board of ASR Academy of Medical Sciences (India) approved on the 10th March 2010 (ref: ASRIRB-10-002)

Primary study design

Interventional

Study design

Randomised double blind placebo controlled study

Study type(s)

Treatment

Health condition(s) or problem(s) studied

Obesity

Interventions

A total 60 human subjects is randomised and divided into two groups:

1. LI12507F
2. Placebo

The treatment dosage is 800 mg daily, consisting of two 400 mg doses daily for the active group. Each subject included in the second group receives matched placebo per day.

The study duration will be of 56 days. The visits and the evaluations are at the baseline, 14th day, 28th day and the 56th day.

Intervention Type

Drug

Phase

Not Specified

Drug/device/biological/vaccine name(s)

LI12507F

Primary outcome(s)

1. Physical/anthropometric parameters:
 - 1.1. Body weight
 - 1.2. Body mass index
 - 1.3. Waist hip ratio

2. Biochemical parameters (serum/plasma):

- 2.1. Fasting glucose
- 2.2. Fasting insulin
- 2.3. Triglycerides
- 2.4. Cholesterol
- 2.5. High density lipoprotein (HDL) cholesterol
- 2.6. Low density lipoprotein (LDL) cholesterol

The primary and secondary outcomes will be measured on baseline, 14th day, 28th day and 56th day.

Key secondary outcome(s)

Obesity biomarkers (serum/plasma):

1. Leptin
2. Adiponectin
3. Ghrelin

The secondary outcomes will be measured on the baseline, and on 56th day.

Completion date

10/08/2010

Eligibility

Key inclusion criteria

1. Participants must understand the risks and benefits of the study
2. Adults ages 21 - 50 years, either sex
3. Body mass index (BMI) greater than 30 kg/m²
4. Willingness to participate in an exercise-walking program, supervised by a trained exercise specialist
5. Willingness to consume the prescribed study diet of approximately 2,000 KCal per day as outlined in the protocol (meals will be provided at free of cost by the study sponsor)
6. Ability to provide written informed consent for participation in the trial
7. Willingness to complete standard health history questionnaire before induction into the study
8. Willingness to participate in 5 clinic visits (Screening, baseline, 2, 4 and 8 weeks)
9. Subject willing to participate in health exercise program (30 minutes walking) monitored by the study physical trainer
10. Subjects must be instructed to abstain from alcoholic products during the study
11. If female:
 - 11.1. Should be negative in pregnancy test
 - 11.2. Should not be nursing
 - 11.3. Of childbearing potential, should agree to follow an acceptable method of birth control for the duration of the study, such as condoms, foams, jellies, diaphragm, intrauterine device (IUD), etc., or postmenopausal for at least 1 year, or surgically sterile (bilateral tubal ligation, bilateral oophorectomy, or hysterectomy)

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Adult

Sex

All

Key exclusion criteria

1. History of thyroid disease or cardiovascular disease or diabetes (uncontrolled)
2. Subjects having chronic diarrhoeal disorders, cancer, neurological disorders, hepatic dysfunction
3. Respiratory tract infection and other serious medical respiratory illnesses (i.e., chronic obstructive pulmonary disease [COPD], sarcoidosis)
4. History of allergy to spices and herbal products
5. Intractable obesity or uncontrolled body weight, BMI greater than 40 kg/m²
6. Non obese (BMI less than 30 kg/m²) and morbidly obese (BMI greater than 40 kg/m²)
7. Presently using other weight loss medications, as well as stimulants, laxatives or diuretics taken solely for the purpose of weight loss
8. Pregnant or nursing or lactating females
9. Recent, unexplained weight loss or gain
10. Women with a positive pregnancy test
11. Human immunodeficiency virus (HIV) or other known immunodeficiency
12. Undergone surgery before 30 days of screening or planning to undergo surgery within the study days
13. History of hepatitis, pancreatitis, lactic acidosis or hepatomegaly with steatosis
14. History of motor weakness or peripheral sensory neuropathy
15. Any evidence of organ dysfunction or any clinically significant deviation from the normal, in physical or clinical determinations

Date of first enrolment

14/03/2010

Date of final enrolment

10/08/2010

Locations**Countries of recruitment**

India

Study participating centre

Department of General Medicine

Vijayawada

India

520007

Sponsor information

Organisation

Laila Nutraceuticals (India)

ROR

<https://ror.org/05q6g7072>

Funder(s)**Funder type**

Industry

Funder Name

Laila Nutraceuticals (India)

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	01/06/2013		Yes	No