

Efficacy of a combined acne treatment

Submission date 03/11/2021	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered <input checked="" type="checkbox"/> Protocol
Registration date 05/11/2021	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan <input checked="" type="checkbox"/> Results
Last Edited 24/08/2022	Condition category Skin and Connective Tissue Diseases	<input type="checkbox"/> Individual participant data

Plain English summary of protocol

Background and study aims

Acne is a multifactorial inflammatory skin disease affecting the quality of life of acne-prone subjects. Several therapeutic approaches are currently used to counteract this condition, mostly having side effects. As acne development has been recently linked to skin and gut dysbiosis, acting on both aspects could represent an alternative and a promising approach to ameliorate the acne clinical signs.

This study aims to assess the efficacy of a combined treatment (a cosmetic product + a food supplement containing probiotics) in improving skin appearance on adult subjects affected by acne through the positive modulation of the microbiota. Such a hypothesis is tested by combining the intake of the probiotics mixture or the placebo together with two different cosmetic products, a basic cream containing 1% Ectoin or a commercially available cosmetic product (reference cosmetic product) specifically intended for a cosmetic treatment of acne.

Who can participate?

Healthy adults of both sexes, aged 18 to 50 years, with acne severity from 1 to 3 according to IGA scale

What does the study involve?

Participants will be randomly allocated in 4 groups to receive for 56 days: i) the basic cosmetic cream containing Ectoin plus the placebo food supplement; ii) the basic cosmetic cream containing Ectoin plus the active food supplement (containing the probiotics mixture); iii) the reference cosmetic product plus the placebo food supplement; iv) the reference cosmetic product plus the active food supplement.

Assessment of parameters was evaluated at the beginning of the study and after 28 and 56 days.

What are the possible benefits and risks of participating?

The benefits associated with product use are amelioration of acne clinical signs.

Risks associated with the product's intake/application are considered from low to very low, in absence of allergy/intolerances to products ingredients; other ingredients in the formula of the product are commonly used in dietary supplements. All the carried out instrumental measurements are not invasive and no skin side effects are expected from the measurement process.

Where is the study run from?
Complife Italia SRL (Italy)

When is the study starting and how long is it expected to run for?
February 2019 to July 2019

Who is funding the study?
This study was realized in the frame of SCIDA project funded by Lombardy Region, Italy (2014IT16RFOP012-POR FESR 2014-2020-Project ID226149).

Who is the main contact?
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Contact information

Type(s)
Scientific

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Additional identifiers

EudraCT/CTIS number
Nil known

IRAS number

ClinicalTrials.gov number
Nil known

Secondary identifying numbers
E.HU.111-0080.17.003L_2019-684

Study information

Scientific Title

Clinical assessment of a combined treatment targeting subjects with acne-prone skin

Study objectives

A combined treatment based on a cosmetic product and food supplement containing selected probiotics could ameliorate clinical signs of acne?

Ethics approval required

Old ethics approval format

Ethics approval(s)

Approved 17/04/2019; Independent Ethical Committee for Non-Pharmacological Clinical Study Trials (Società Scientifica Italiana per le Indagini Cliniche Non Farmacologiche, Via XX Settembre 30/4, 16121 Genova, Italy; +39(0)10 5454842; a.scudieri@studiononfarmacologici.it); ref: 2019/04

Study design

Single-centre interventional double-blinded randomized placebo-controlled parallel-group clinical trial

Primary study design

Interventional

Secondary study design

Randomised parallel trial

Study setting(s)

Home

Study type(s)

Treatment

Participant information sheet

Not available in web format, please use the contact details below to request a patient information sheet

Health condition(s) or problem(s) studied

Acne

Interventions

A double-blind, randomized, placebo/reference product-controlled clinical study was carried out on eighty subjects of both sexes, enrolled by a dermatologist according to specific inclusion /exclusion criteria; subjects were equally divided in the following 4 groups according to a randomization list previously generated using an appropriate statistic algorithm ("Wey's urn"):

- Group 1 (G1): a basic cosmetic product containing 1% Ectoin + a placebo food supplement
- Group 2 (G2): a basic cosmetic product containing 1% Ectoin + an active food supplement (In&Out combined treatment)
- Group 3 (G3): a commercially available cosmetic product intended for cosmetic acne treatment available (cosmetic reference product)+ a placebo food supplement.
- Group 4 (G4): the cosmetic reference product + an active food supplement.

Subjects applied daily the cosmetics active/reference products and took one capsule of the active/placebo food supplement for 56 days.

Instrumental evaluations of skin parameters and dermatological assessments of the subjects'

facial skin were carried out at baseline (T0), after 28 days, and after 56 days from the beginning of product use.

Intervention Type

Supplement

Primary outcome measure

1. Sebum levels, measured by using the Sebumeter® method (Sebumeter 815, Courage+Khazaka GmbH) at t=0, T28 and T56.
2. Skin pH measured by SKIN pH-METER 905® (Courage + Khazaka GmbH) at t=0, T28 and T56.
3. Skin moisturization, measured by the Corneometer® method (Corneometer® CM 825 (Courage+Khazaka, electronic GmbH) at t=0, T28 and T56.
4. Dermatological assessment of facial skin by counting acne lesions at t=0, T28 and T56
5. Evaluation of skin inflammatory status through face digital pictures at t=0, T28 and T56.

Secondary outcome measures

Products tolerability, efficacy, and acceptability were evaluated in a discussion with the dermatologist at the end of the treatment.

Overall study start date

04/02/2019

Completion date

19/07/2019

Eligibility

Key inclusion criteria

1. Good general health
2. Both sexes, caucasian ethnicity
3. Phototype I to IV
4. Age between 18 and 50 years old
5. Acne severity from 1 to 3 according to IGA, (Investigator's Global Assessment) severity scale,
6. Subjects who have not been recently involved in any other similar study
7. Willingness to use for face care only the creams that will be consigned at the beginning of the study
8. Willingness to submit before and after pictures
9. Willingness to use during all the study period only the products to be tested
10. Willingness not to use similar products that could interfere with the product to be tested
11. Willingness to not vary the normal daily routine (i.e. lifestyle, physical activity, etc.)
12. Subject is under effective contraception (oral/not oral); not expected to be changed during the trial
13. Subject aware of the study procedures and having signed an informed consent form
14. Subjects who accept not to expose in an intensive way to UV rays during the whole study duration

Participant type(s)

Healthy volunteer

Age group

Adult

Lower age limit

18 Years

Upper age limit

50 Years

Sex

Both

Target number of participants

80

Total final enrolment

80

Key exclusion criteria

1. Subjects who do not meet the inclusion criteria
2. Pregnant/breastfeeding female or who have planned pregnancy during the study period
3. Subjects under systemically pharmacological treatment
4. Subjects under locally pharmacological treatment on the skin area monitored during the test
5. Subjects with congenital or acquired immunodeficiency
6. Subjects under treatment with food supplements which could interfere with the functionality of the product under study
7. Subjects who show other skin alterations on the monitored area
8. Subjects considered as not adequate to participate in the study by the investigator
9. Subjects with known or suspected sensitization to one or more test formulation ingredients
10. Adult protected by law (under control or hospitalized in public or private institutions for reasons other than research, or incarcerated)
11. Subjects not able to communicate or cooperate with the investigator for problems related to language, mental retardation, or impaired brain function

Date of first enrolment

17/05/2019

Date of final enrolment

24/05/2019

Locations

Countries of recruitment

Italy

Study participating centre

COMPLIFE ITALIA SRL

Via Via Mons. Angelini 21

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Italy
27028

Sponsor information

Organisation

Complife Italia Srl

Sponsor details

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Sponsor type

Industry

Website

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Funder(s)

Funder type

Government

Funder Name

Regione Lombardia

Alternative Name(s)

Lombardy Region, Region of Lombardy

Funding Body Type

Government organisation

Funding Body Subtype

Local government

Location

Italy

Results and Publications

Publication and dissemination plan

Planned publication in a high-impact peer-reviewed journal

Intention to publish date

15/11/2021

Individual participant data (IPD) sharing plan

The datasets generated during and/or analysed during the current study will be stored in a non-publicly available repository. Raw data will be stored in Complife servers. A backup copy of the raw data will be also in a cloud-based backup server. Tables containing the raw data (output of the measurements) will be also included in the study report and shared with the study Sponsor by a pdf file electronically signed. The raw data will be stored for a minimum period of 10 years in Complife servers. In the raw data tables, subjects are identified by a means of a code generated by the Complife volunteer's management software. The code is composed of a letter, 4 digits, and a letter. The access to the study raw data is allowed only to the study director and the person designated by him to elaborate the raw data. Elaboration of the raw data includes descriptive statistics (mean and standard error) and the inferential analysis (data normality and statistical test).

IPD sharing plan summary

Stored in non-publicly available repository, Available on request, Published as a supplement to the results publication

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Protocol file	version 2	19/03/2019	04/11/2021	No	No
Results article		01/07/2022	18/07/2022	Yes	No