

A clinical study to assess the safety and efficacy of poly-L-lactic acid dermal filler as a treatment for the rejuvenation of the upper arms

Submission date 12/03/2025	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered
Registration date 18/03/2025	Overall study status Completed	<input type="checkbox"/> Protocol
Last Edited 18/11/2025	Condition category Skin and Connective Tissue Diseases	<input type="checkbox"/> Statistical analysis plan
		<input checked="" type="checkbox"/> Results
		<input type="checkbox"/> Individual participant data

Plain English summary of protocol

Background and study aims

The aim of this study is to demonstrate the effectiveness and safety of Lanluma V in the treatment of age-related sagging of upper arm skin. Poly-L-lactic acid dermal fillers are designed to restore volume that has been lost through the aging process and may be used in the correction of the loss of elasticity and firmness in the skin of the upper arm.

Who can participate?

Men and women aged between 40 and 65 years old

What does the study involve?

Participants receive Lanluma V injections and consent to generate a clinical dataset to demonstrate the effectiveness and safety of Lanluma V as a treatment for the rejuvenation of the upper arms. Physicians with longstanding experience in the aesthetic field perform the injections.

The study consists of two visits on day 1 and in month 2 for treatment and one visit for an optional third treatment in month 4. There will be six visits to assess effectiveness and safety at months 2, 4, 6, 12, 18, and 25. A safety follow-up visit and check of participant diaries is performed in week 2.

What are the possible benefits and risks of participating?

There are risks related to the administration procedure per se (irrespective of the filler), including but not limited to bruising, swelling, redness, and infection. On rare occasions participants may experience hypersensitivity to Lanluma V.

Where is the study run from?

1. García Clínicas Dr. Jorge García (Spain)
2. Clínica de Medicina Estética Dra. Páez (Spain)

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

September 2022 to December 2025

Who is funding the study?
Sinclair Ltd (UK)

Who is the main contact?
Dr Stuart Boothman, sboothman@sinclair.com

Contact information

Type(s)

Public

Contact name

Dr Salvinia Mletzko

Contact details

RQM+ (Germany) GmbH
von-Geismar-Strasse 2
Ahlen
Germany
59229
+49 (0)1702062568
smletzko@rqmplus.com

Type(s)

Principal investigator

Contact name

Dr Jorge García

Contact details

C/ María de Molina, 13, 4º A. (Edif. Pza. Zorrilla)
Valladolid
Spain
47001
+34 (0)983 338 100
valladolid@esteticafacialjorgegarcia.com

Type(s)

Scientific

Contact name

Dr Stuart Boothman

Contact details

Eden House Lakeside
Chester Business Park
Chester
United Kingdom

CH4 9QT
+44 (0)1244 625127
sboothman@sinclair.com

Additional identifiers

Clinical Trials Information System (CTIS)

Nil known

ClinicalTrials.gov (NCT)

Nil known

Protocol serial number

CS-21-06

Study information

Scientific Title

A post-market follow-up study to assess the safety and efficacy of Lanluma V as a treatment for the rejuvenation of the upper arms

Study objectives

The primary hypothesis is that the efficacy of Lanluma V will be evident at 6 months post-treatment, while the secondary hypotheses are that Lanluma V will be efficacious and safe over the entire study period.

Ethics approval required

Ethics approval required

Ethics approval(s)

approved 15/09/2022, CEIm de las Áreas de Salud Valladolid (Hospital Clínico Universitario de Valladolid, Facultad de Medicina, Farmacología, Valladolid, 47005, Spain; +34 (0)983 423077; jalvarezgo@saludcastillayleon.es), ref: EOm 22-400

Study design

Multicentre prospective open-label (non-randomized) interventional post-market clinical follow-up (PMCF) study

Primary study design

Interventional

Study type(s)

Treatment

Health condition(s) or problem(s) studied

Ageing of the skin

Interventions

Lanluma V is a CE-marked medical device and is used within its intended use. Injection of Lanluma V at baseline is followed by retreatment at 2 months (following assessment of efficacy and safety) and an optional retreatment at 4 months.

During seven visits on site after the initial treatment, skin depression defects and the effect of treatment are assessed and documented using live GAIS evaluations and standardised photography. There is one phone call visit after the initial treatment.

Participants and physicians fill in satisfaction questionnaires. Safety data will be collected for up to 25 months.

Intervention Type

Device

Phase

Not Applicable

Drug/device/biological/vaccine name(s)

Lanluma V

Primary outcome(s)

The proportion of participants (%) with an improvement (i.e., improved, much improved or very much improved) over baseline at 6 months in Global Aesthetic Improvement Scale (GAIS) assessments in the treated region(s) of the upper arm, as assessed by an on-site independent evaluator

Key secondary outcome(s)

1. The proportion of participants (%) with an improvement (i.e., improved, much improved or very much improved) over baseline in Global Aesthetic Improvement Scale (GAIS) assessments of the treated upper arm at 2, 4, 12, 18 and 25 months, as assessed by an on-site independent evaluator and participant
2. The proportion of participants (%) exhibiting an improvement of ≥ 1 point from baseline on a photographic Arm Visual Analog scale at 2, 4, 6, 12, 18 and 25 months as assessed by a blinded remote independent evaluator
3. Treatment satisfaction based on questionnaires filled by the participants (at 2, 4, 6, 12, 18 and 25 months) and the investigators (straight after treatment)

Completion date

12/12/2025

Eligibility

Key inclusion criteria

1. Subjects with type II or type III upper arms on the Arm Visual Analog Scale (Arm VAS) seeking treatment for rejuvenation of the upper arms
2. Female and male subjects between 40-65 years of age
3. Subjects who are willing to provide written informed consent including approval of the use of their data and photographs in this study and any subsequent publications or presentations
4. Subjects willing to commit to having no further aesthetic treatments, such as body contouring procedures, for the duration of the study period
5. Subjects willing to commit to having no further tattoos on the upper arms for the duration of

the study period

6. Women of childbearing potential should use a medically accepted contraceptive regimen since at least 12 weeks prior to study entry and during the entire study

Participant type(s)

Healthy volunteer

Healthy volunteers allowed

No

Age group

Mixed

Lower age limit

40 years

Upper age limit

65 years

Sex

All

Total final enrolment

72

Key exclusion criteria

1. Subjects who, in the 12 months prior to their enrolment assessment have undergone in the region of the upper arms:
 - 1.1. Tissue grafting (e.g. fat injections)
 - 1.2. Lipolysis
 - 1.3. Tissue lifting implants (e.g. threads, barbs) or other implants
 - 1.4. Augmentation with any permanent or semipermanent filler (e.g. silicone, PMMA, PLLA)
 - 1.5. Temporary dermal filler injections (e.g., HA, CaHA, PCL)
 - 1.6. Neuromodulator injections
 - 1.7. Mesotherapy
 - 1.8. Resurfacing (laser, radio frequency, dermabrasion, or chemical peel)
2. Subjects who have received treatment with a permanent filler (e.g. silicone, PMMA, PLLA) in the region of the upper arms
3. Subjects who have received any other aesthetic procedures in the region of the upper arms at any time during the study period
4. Subject who had been deprived of their freedom by administrative or legal decision or who is under guardianship
5. Subject is an employee of the aesthetic surgery department on the investigational site, the Clinical Research Organisation (CRO) or the study sponsor
6. Subjects who in the opinion of the investigator are unsuitable to take part in the study for scientific or medical reasons
7. Subjects currently enrolled in other clinical trials
8. Excessive skin laxity in the region of the upper arms
9. Subjects who, if female, are pregnant or plan to become pregnant during the study period
10. Subjects with known allergies to the product ingredients (i.e. poly-L-lactic acid (PLLA), sodium carboxymethyl cellulose (CMC), Mannitol)

11. Subjects must avoid receiving a Coronavirus Disease (COVID)-19 vaccination for the 14 days before and following the injection
12. Subject with a cutaneous disorder, inflammation, infection, significant scarring, open wounds, lesions or tattoos in the region of the upper arms
13. Subjects taking thrombolytics or anticoagulants
14. Subjects with bleeding disorders
15. Subjects with a history of severe allergy or anaphylactic shock
16. Subjects with active (or a history of) autoimmune disease
17. Subjects with porphyria
18. Subjects with a tendency to form keloids, hypertrophic scars or any other healing disorders
19. Subjects with a history of lymphedema in the arms

Date of first enrolment

13/01/2023

Date of final enrolment

08/05/2023

Locations

Countries of recruitment

Spain

Study participating centre

García Clínicas Dr. Jorge García

C/ María de Molina, 13, 4º A. (Edif. Pza. Zorrilla)

Valladolid

Spain

47001

Study participating centre

Clínica de Medicina Estética Dra. Páez

Avenida Condes de San Isidro, Pasaje La Camboria Ed. Fuengirola 1, 1º izda

Fuengirola, Málaga

Spain

29640

Sponsor information

Organisation

Sinclair Pharmaceuticals Ltd

Funder(s)

Funder type
Industry

Funder Name
Sinclair Ltd

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary
Not expected to be made available

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Basic results		18/11/2025	18/11/2025	No	No