

# TRANS-FOODS: Reducing the risk of developing peanut allergy through the skin by improved understanding, modifying peanut snack production, and adjusting skincare practices.

<b>Submission date</b> 11/09/2023	<b>Recruitment status</b> No longer recruiting	<input checked="" type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
<b>Registration date</b> 25/09/2023	<b>Overall study status</b> Ongoing	<input type="checkbox"/> Statistical analysis plan <input type="checkbox"/> Results
<b>Last Edited</b> 03/02/2025	<b>Condition category</b> Skin and Connective Tissue Diseases	<input type="checkbox"/> Individual participant data <input type="checkbox"/> Record updated in last year

## Plain English summary of protocol

### Background and study aims

Allergic diseases, including atopic eczema, and food allergies affect over a quarter of all children across Europe. The way the immune system responds to ingested foods are well-established, and regular eating of allergy-causing foods can prevent food allergies to a degree. However, recent research also shows that food allergies can develop through the skin, especially in the presence of dry skin and eczema. We do not fully understand how this happens at the moment.

This project aims to study the immune responses to peanut allergen in those with a skin barrier defect with and without skin massage.

### Who can participate?

Anyone without a history of peanut allergy.

Anyone either with a history of dry skin/eczema or healthy skin.

Anyone aged 18 years or over.

### What does the study involve?

60 healthy adult volunteers and 60 adult volunteers with atopic dermatitis and dry skin will be asked to either massage, or not massage the skin after applying peanut extract twice daily for 4 weeks at home. Half of the participants will also apply a barrier enhancing cream 30 minutes before applying the peanut extract.

Samples of interstitial fluid (ISF) will be collected from the same location that the cream and peanut extracts are being applied and at another site that is remote to this site during a visit to King's College London. The amount of water that evaporates from the skin will be measured with a small hand-held machine. A chamber will then be placed on your forearm and a small volume of water will be applied. A hand pump will be used to apply suction on your forearm for about 10 minutes at a pressure we know will not damage the skin. The suction will pull molecules called cytokines out of the skin and we will remove the liquid from the skin containing

these cytokines using a plastic pipette. The extracted samples will be collected for analysis of the extracted chemicals.

Participation in this study will require presence in two extraction sessions with an interval of 4 weeks, each session will last for approximately 1 h.

What are the possible benefits and risks of participating?

The participants will be tested to peanut and environmental allergens. Otherwise, there is no direct benefit, however, the scientific information that is obtained from this research will be used to help scientists and doctors learn more about the causes of peanut allergy and could benefit people in the future.

There is a very low risk of an allergic reaction to the skin prick test or the peanut preparation used in this study. The use of the interstitial fluid device may result in the formation of a transient indentation mark on the skin caused by the suction chamber. Bruising may occur with the use of this device, which should go away after a couple of days.

Where is the study run from?

The study is run from King's College London

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

July 2022 to May 2026

Who is funding the study?

1. Biotechnology and Biological Sciences Research Council (UK)
2. Food Standards Agency (UK)
3. Joint Programming Initiative (UK)

Who is the main contact?

If you have questions about this research study, contact the research team via email  
TransFoods@kcl.ac.uk

## Contact information

### Type(s)

Public

### Contact name

Miss Preeti Khurana

### Contact details

Unit for Population-Based Dermatology Research  
St John's Institute of Dermatology  
1st Floor  
C Staircase  
South Wing  
St. Thomas' Hospital  
Westminster Bridge  
London  
United Kingdom  
SE1 7EH

+44 2071887188  
Transfoods@kcl.ac.uk

**Type(s)**  
Scientific

**Contact name**  
Prof Carsten Flohr

**ORCID ID**  
<https://orcid.org/0000-0003-4884-6286>

**Contact details**  
Unit for Population-Based Dermatology Research  
St John's Institute of Dermatology  
1st Floor  
C Staircase  
South Wing  
St. Thomas' Hospital  
Westminster Bridge  
London  
United Kingdom  
SE1 7EH  
+44 2071887188  
Transfoods@kcl.ac.uk

**Type(s)**  
Principal investigator

**Contact name**  
Prof Carsten Flohr

**Contact details**  
Unit for Population-Based Dermatology Research  
St John's Institute of Dermatology  
1st Floor  
C Staircase  
South Wing  
St. Thomas' Hospital  
Westminster Bridge  
London  
United Kingdom  
SE1 7EH  
+44 2071887188  
Transfoods@kcl.ac.uk

## **Additional identifiers**

**Clinical Trials Information System (CTIS)**  
Nil known

**ClinicalTrials.gov (NCT)**

NCT05407012

**Protocol serial number**

Nil known

## Study information

### Scientific Title

TRANS-FOODS: Preventing Peanut Allergy Through Improved Understanding of the Transcutaneous Sensitisation Route, Novel Food Processing and Skin Care Adaptations (TRANS-FOODS)

### Acronym

TRANS-FOODS

### Study objectives

1. When a peanut protein-containing solution is regularly massaged into the skin, peanut protein components can be detected in ISF.
2. Higher concentrations of peanut protein are detected in ISF in those with skin barrier impairment (dry skin and AD).
3. Peanut proteins present in ISF retain their allergenicity and can interact with immune cells and induce activation of blood basophils and mast cells from peanut-allergic donors.
4. Regular skin massage with peanut protein extract induces detectable skin inflammation (raised Th2 cytokines, TSLP, and IL-33), more so in those who have dry skin and AD/raised TEWL at baseline (compared to no massage).
5. The cutaneous uptake of peanut protein is attenuated by the prior application of a barrier-enhancing moisturising cream.

### Ethics approval required

Ethics approval required

### Ethics approval(s)

approved 23/06/2023, KCL REC (3rd Floor, 5-11 Lavington Street, London, SE1 0NZ, United Kingdom; -; rec@kcl.ac.uk), ref: HR/DP-22/23-37419

### Study design

Pilot investigational interventional randomized parallel open label

### Primary study design

Interventional

### Study type(s)

Prevention

### Health condition(s) or problem(s) studied

Peanut allergy and atopic dermatitis

### Interventions

Using block randomisation, the two groups of healthy volunteers and participants with skin barrier impairment (AD/dry skin) will be evenly split into the following groups:

- application of the barrier enhancing preparation around 30mins before application of the peanut protein extract +/- massage after extract application
- no application of the barrier enhancing preparation, just application of the peanut protein extract +/- massage after extract application

This will generate 8 intervention groups of 15 participants each. Participants will apply the test interventions to the central area of the left outer forearm twice daily with 10 mins massage throughout the 4-week study period. The control semi-solid formulation (containing no extracts) will be applied without massage to the central right outer forearm twice daily as control. We will provide detailed instructions for all participants on the following:

- hand hygiene prior to the application of the study intervention (left outer forearm) and the sunflower oil control (right outer forearm),
- clear instructions on how the massage is to be performed (if participants are randomised to this intervention) to standardise this across participants. Participants will follow the massaging technique at home using video guidance, which we will send to the participant at the start of every week as a reminder, and
- that no other topical preparation must be applied for at least 4 hours after the application to allow for take up into the skin. (This is based on our extensive experience in studying topical preparations and to avoid cross-contamination and ensure adequate cutaneous uptake of the peanut extract).

### **Intervention Type**

Other

### **Primary outcome(s)**

1. Detection of peanut protein components ( $\mu\text{g}$  of the proteins per  $\text{cm}^2$  of skin) in retrieved interstitial fluid. [ Time Frame: 8 weeks ]
2. Activation of blood basophil/mast cell from peanut allergic donors (measured as %CD63-positive basophils/mast cells) by peanut proteins present in interstitial fluid. [ Time Frame: 8 weeks ]

### **Key secondary outcome(s)**

1. Detection of inflammatory cytokine markers (IL-4, IL-13, IL-33 and TSLP) in interstitial fluid. [ Time Frame: 8 weeks ]
2. Raised transepidermal water loss. [ Time Frame: 8 weeks ]

### **Completion date**

31/05/2026

## **Eligibility**

### **Key inclusion criteria**

1. Adult healthy volunteers (50% of the cohort), and adults with dry skin and AD (fulfilling the refined Hanifin and Rajka criteria, 50% of the cohort).
2. Written informed consent for study participation.
3. Competent use of English language.
4. Willingness to comply with all study requirements.

### **Participant type(s)**

Healthy volunteer, Patient

**Healthy volunteers allowed**

No

**Age group**

Adult

**Lower age limit**

18 years

**Upper age limit**

65 years

**Sex**

All

**Key exclusion criteria**

1. Patients unable to give informed consent.
2. History of peanut allergy.
3. Positive skin prick test to peanut (>0 mm).
4. No regular consumption of peanut products.
5. Widespread AD, in particular if this involves the test sites of the forearms.

**Date of first enrolment**

05/04/2024

**Date of final enrolment**

31/07/2025

## **Locations**

**Countries of recruitment**

United Kingdom

England

**Study participating centre**

**King's College London, Franklin-Wilkins Building.**

150 Stamford Street

London

United Kingdom

SE19NH

## **Sponsor information**

**Organisation**

King's College London

**ROR**

<https://ror.org/0220mzb33>

**Funder(s)****Funder type**

Research council

**Funder Name**

Biotechnology and Biological Sciences Research Council

**Alternative Name(s)**

UKRI - Biotechnology And Biological Sciences Research Council, BBSRC UK, Biotechnology and Biological Sciences Research Council (BBSRC), BBSRC

**Funding Body Type**

Government organisation

**Funding Body Subtype**

National government

**Location**

United Kingdom

**Funder Name**

Food Standards Agency

**Alternative Name(s)**

The Food Standards Agency, FSA

**Funding Body Type**

Private sector organisation

**Funding Body Subtype**

Other non-profit organizations

**Location**

United Kingdom

**Funder Name**

## Results and Publications

### Individual participant data (IPD) sharing plan

The datasets generated during and/or analysed during the current study will be available upon request from TransFoods@kcl.ac.uk.

### IPD sharing plan summary

Available on request

### Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
<a href="#">Participant information sheet</a>	Participant information sheet	11/11/2025	11/11/2025	No	Yes