

# Phase I Trial: Quotient Code QSC303217

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| <b>Submission date</b><br>24/06/2025   | <b>Recruitment status</b><br>No longer recruiting | <input checked="" type="checkbox"/> Prospectively registered<br><input type="checkbox"/> Protocol                       |
| <b>Registration date</b><br>01/07/2025 | <b>Overall study status</b><br>Deferred           | <input type="checkbox"/> Statistical analysis plan<br><input type="checkbox"/> Results                                  |
| <b>Last Edited</b><br>01/07/2025       | <b>Condition category</b><br>Other                | <input type="checkbox"/> Individual participant data<br><input checked="" type="checkbox"/> Record updated in last year |

## Plain English summary of protocol

The sponsor has confirmed that the trial meets the criteria for deferral of publication of the full details of the trial. Full details will be added to the study record within 30 months after the trial has ended.

## Contact information

### Type(s)

Public, Scientific

### Contact name

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### Type(s)

Principal investigator

### Contact name

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# Additional identifiers

## Clinical Trials Information System (CTIS)

Nil known

## Integrated Research Application System (IRAS)

1011885

## Protocol serial number

Sponsor Code: AVK-101-102, Quotient Code: QSC303217

# Study information

## Scientific Title

Phase I Trial: Quotient Code QSC303217

## Study objectives

The sponsor has confirmed that the trial meets the criteria for deferral of publication of the full details of the trial. Full details will be added to the study record within 30 months after the trial has ended.

## Ethics approval required

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## Ethics approval(s)

approved 19/06/2025, London – Hampstead Regulatory Ethics Committee (REC) (2 Redman Place, Stratford, London, E20 1JQ, United Kingdom; +44 (0)20 7104 8171; hampstead.rec@hra.nhs.uk), ref: 25/LO/0198

## Study design

Two-part single-centre mass balance recovery and food effect study in healthy volunteers

## Primary study design

Interventional

## Study type(s)

Other

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

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## Interventions

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## Intervention Type

Drug

**Phase**

Phase I

**Drug/device/biological/vaccine name(s)**

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**Primary outcome(s)**

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**Key secondary outcome(s)**

The sponsor has confirmed that the trial meets the criteria for deferral of publication of the full details of the trial. Full details will be added to the study record within 30 months after the trial has ended.

**Completion date**

22/08/2025

**Eligibility**

**Key inclusion criteria**

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**Participant type(s)**

Healthy volunteer

**Healthy volunteers allowed**

No

**Age group**

Adult

**Lower age limit**

18 years

**Upper age limit**

65 years

**Sex**

All

**Key exclusion criteria**

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**Date of first enrolment**

01/07/2025

**Date of final enrolment**

22/08/2025

## **Locations**

**Countries of recruitment**

United Kingdom

England

**Study participating centre**

**Quotient Sciences Limited**

Mere Way, Ruddington Fields, Ruddington

Nottingham

United Kingdom

NG11 6JS

## **Sponsor information**

**Organisation**

Aardvark Therapeutics, Inc.

## **Funder(s)**

**Funder type**

Industry

**Funder Name**

Aardvark Therapeutics, Inc.

## **Results and Publications**

### **Individual participant data (IPD) sharing plan**

The datasets generated and/or analysed during the current study are not expected to be made available because of their high commercial sensitivity and the negligible benefit to the public of publication of results of non-therapeutic clinical trials.

### **IPD sharing plan summary**

Not expected to be made available