The I TRANSCEND Study: A Clinical Research Study in Adults Investigating a Potential Treatment Option for ANCA-Associated Vasculitis

Who We Are

At Alexion AstraZeneca Rare Disease, the work we do is guided by people living with rare and devastating diseases. We are driven to continuously innovate and create meaningful value in all we do to help patients and families fully live their best lives.

Our Values and Principles

We believe in improving access to clinical research studies for all eligible participants, at no cost to them.

Removing Barriers to Participate in Clinical Research Studies

If you meet the study's eligibility criteria but experience barriers to participate, please reach out to your study site to explore ways you can join.

Barriers to study participation can include:

- ❑ Any transportation or travel-related issues
- ❑ Other barriers, such as workplace or family obligations

Cost and Reimbursement Specifics

Costs associated with study participation will be billed to the study sponsor.

Eligible patient expenses will be reimbursed for the study participant and one caregiver, where permitted.
Reimbursement will be provided by an external vendor or the study site per the approved budget and informed consent form (ICF).

Approved reimbursements can be expected within 24-48 hours of submission and may include:

Per visit participant stipend, as allowed by local regulation	Travel companion or caregiver costs, eligible under the same rules
Study related travel costs, such as mileage, tolls, train/plane/car service, meals, and any other out-of-pocket expenses, as approved	Other expenses will be considered on a case-by-case basis. For example, overnight hotel stays or other relevant expenses for out-of-town participants

Costs not outlined in the reference list should be estimated and discussed with the study sponsor before a decision is made. For example:

Other considerations for out-of-town participants.

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AstraZeneca Rare Disease

Activities covered by the study sponsor in the I TRANSCEND Study:*

General assessments/procedures	• Informed consent • Eligibility Assessments • Demographics • Review of medical history
Study intervention	• Assigned study treatment as an add-on to the standard of care
Safety assessments	• Weight and height • Vital signs • Blood sample collections • Pregnancy tests • In-home procedures (if applicable)
Effectiveness and quality of life assessments	• Review of health and potential side effects • Questionnaires • Blood sample collections • Urine sample collections

*Refer to the Informed Consent Form (ICF) for specific information. This list may vary by site

Thank you for your interest in the I TRANSCEND Study.