UK Screening programmes

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Denver 2025





Disclosures

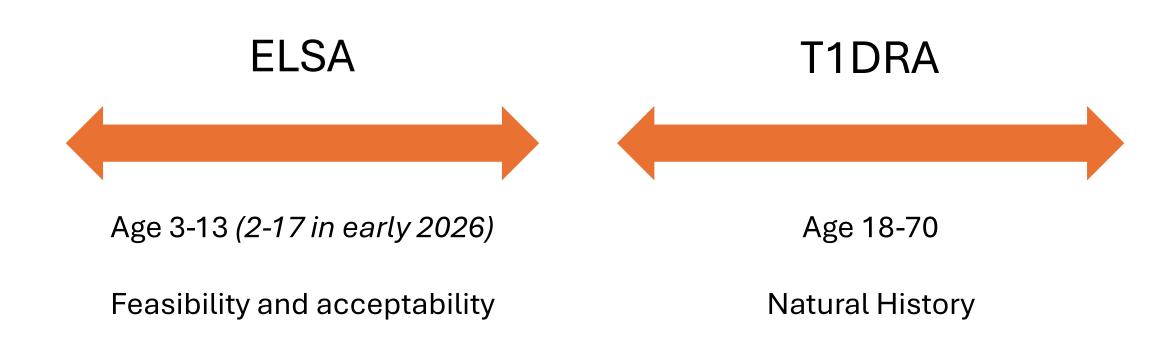
Fees paid to my institution:

Sanofi: Consultancy and speaker fees, support for meeting travel

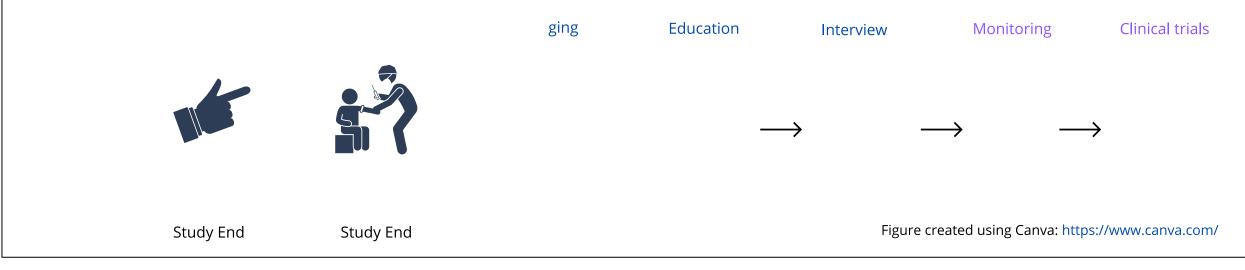
Lilly: Consultancy fees

Abbott: Consultancy fees

UK general population screening programmes







RSR-3 screen: GAD, IA2, ZnT8

GAD, IA2, ZnT8, + IAA GP informed SNOMED code

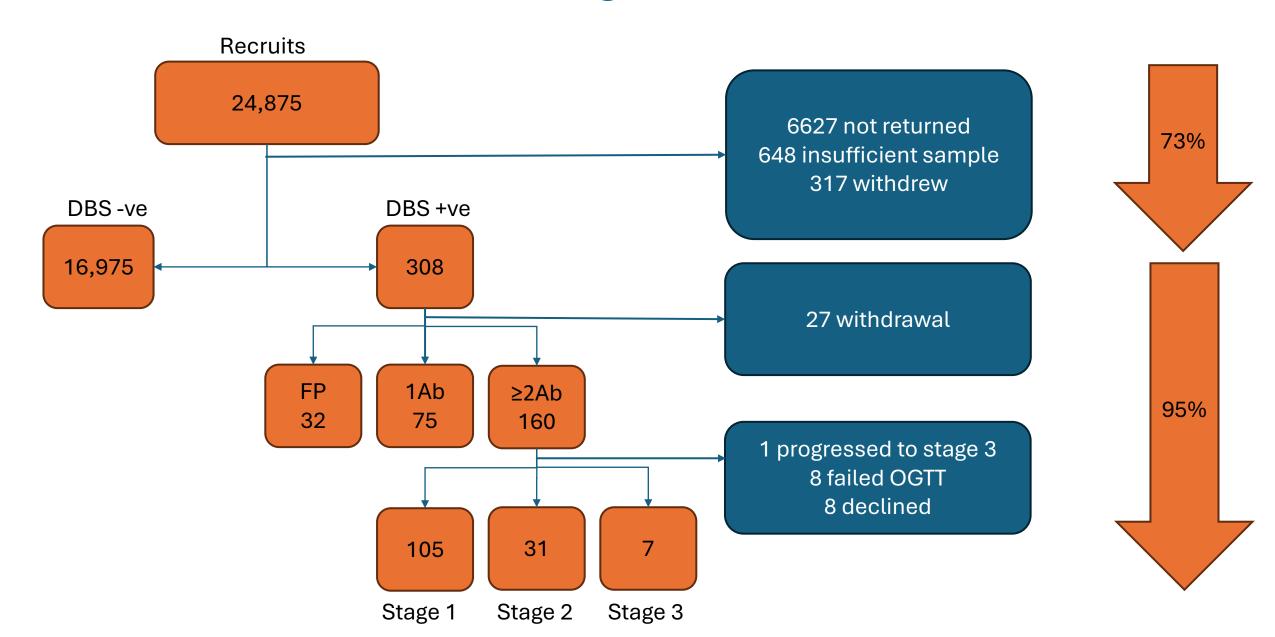








ELSA Consort diagram WDD 2022- WDD 2024



Dried blood spot testing followed by confirmatory testing at local sites is both feasible and acceptable

Feasibility

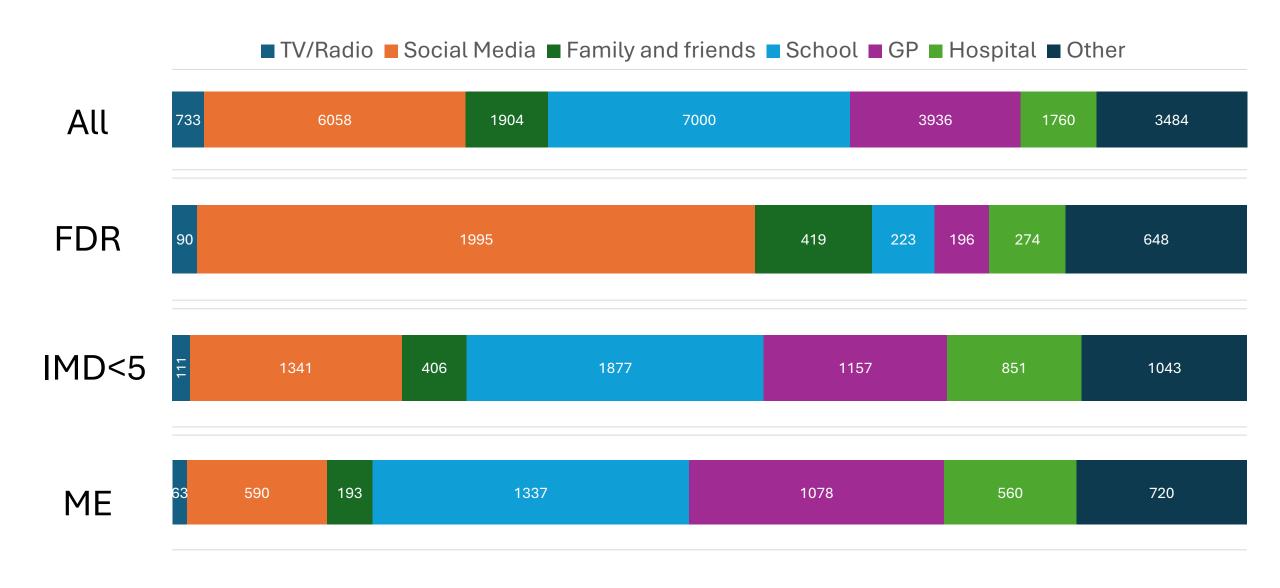
- 73% DBS return rate
- 95% of DBS positive proceeded to confirmatory testing and staging
- 85% completed education follow-up

Acceptability

98% found the screening acceptable, positive experience, would recommend

- High satisfaction
- Overall, easy, convenient process
- Trusted the study
- Benefit and process acceptable regardless of screening outcome (DBS positive or negative)
- Grateful for opportunity to take part and receive child's risk status
- Highly recommend

Where did you hear about the study?



Adverse events generally felt to be acceptable

DBS negative

No lasting impact
Worst parts were the DBS and the wait for results

DBS positive

Families still felt worthwhile and grateful for taking part
Anxiety requiring psychological support was rare
Most experienced screening anxiety which persisted after the
education session

(First Degree) Relatives are a good way to start

Greater kit return rates

More engaged

37% of ELSA participants

Enabler for study entry

Implications of negative test

Comprehension following education

More likely to screen positive

3.73% FDR 2.24% any relative 0.25% Gen Pop

Home DBS
testing
procedure
more
acceptable

FDR screening of all T1D children in England £7M over 5 years

(180M / year flu vaccine)

24,500 children with T1D

Screening FDRs at 3 age points: £3M

Monitoring those with early T1D for 5 years: £4M

ELSA 2025-2029

Increase screening age Age 2-17

Genetics

Health economics

Monitoring and support

Screening of relatives













TYPE 1 DIABETES RISK IN UK ADULTS



Study update 27 Oct 2025

bristol.ac.uk

Introduction

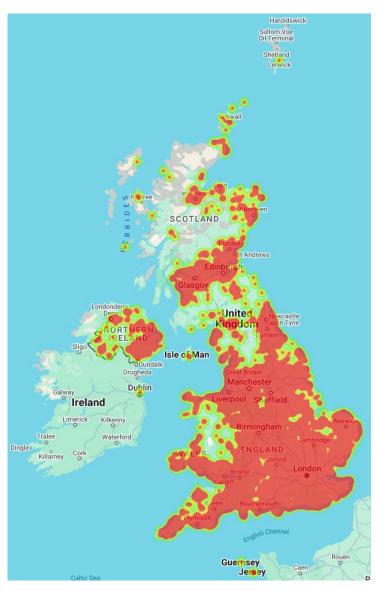
T1DRA was set up to understand the natural history of type 1 diabetes in adults by

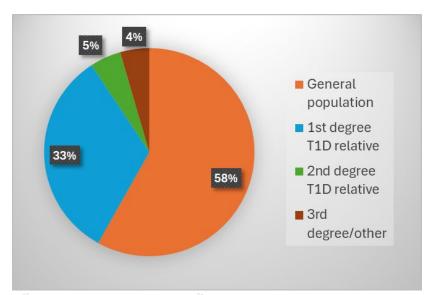
- screening individuals between the ages of 18 and 70 years for the presence of
- islet autoantibodies using high performance tests.
- Positive individuals enter an annual monitoring programme

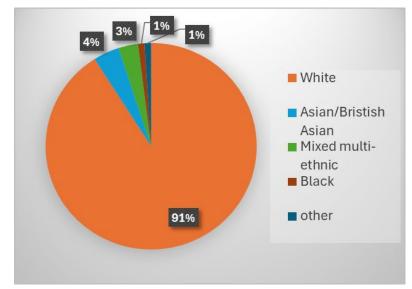


T1DRA Recruitment characteristics of 13535 adults

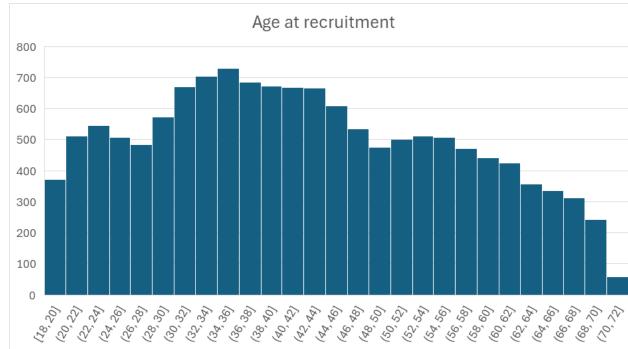
Data to 14 Oct 2025













Advertising







Recruitment

Consent online



Capillary sample kit posted (and returned)



Serum isolated/volume recorded



Samples tested for IAA (RBA)/tGADA/IA-2A/ZnT8A LIPS



Confirmation sample requested on positives



T1DRA PPIE Group

Education about T1D and annual follow up including HbA1c offered to confirmed positive participants



Confirmed Positive adults (no diagnosis of Diabetes)

n= 290 without T2D (3.7% of participants screened)

Multiple islet autoantibody positives

n= 41 (**0.5**% of screens); 25x2abs, 15x3abs 1x4abs

(3 participants Dx T1D since screening)

Relatives 54.7%; General population 45.3%

Age: Median 36 years, (Range 19-67 years); 70% female

Single islet autoantibody positives

n= 249 (**3.1%** of screens).159 tGADA; 40 ZnT8A, 39 IAA, 11 IA-2A

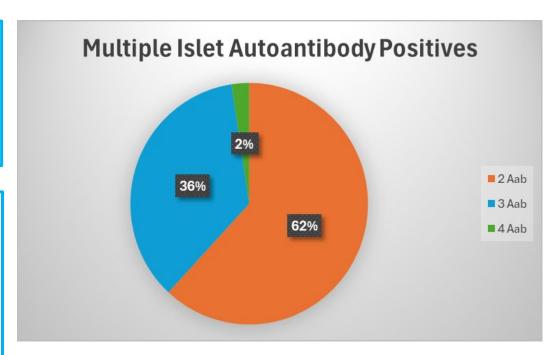
(2 participants Dx T1D since screening)

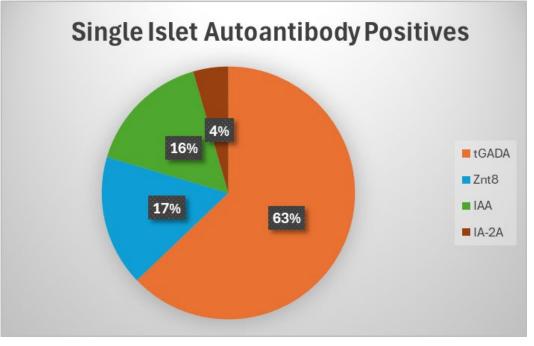
Relatives 59%; General population 41%

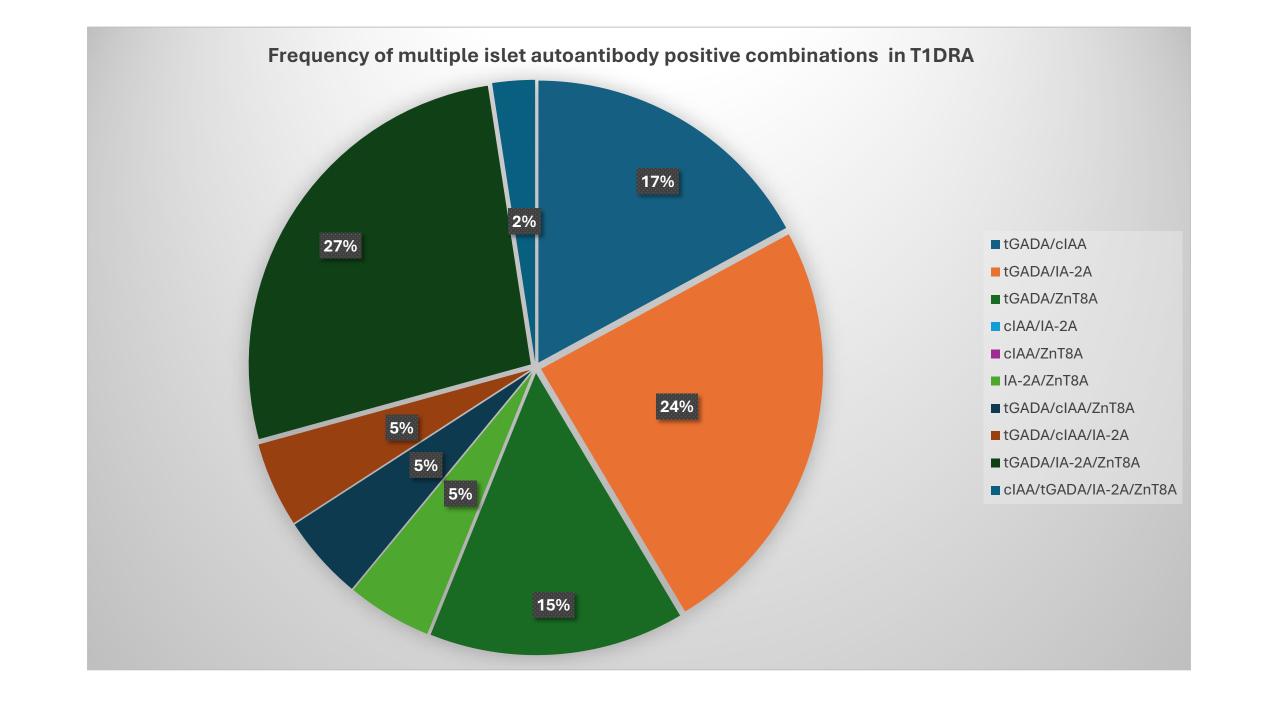
Age: Median 41.2 years, (Range 18-70 years)

81% female (64.5% tGADA); 19% male (54.3% tGADA)

Results as of 14th October







Follow up for T1DRA participants

- Annual islet autoantibodies
- HbA1c
- UCPCR/remote OGTT
- Make participants aware of upcoming clinical trials

Follow up

Nov 2023 Nov 2024 Oct 2025

Recruitment/Screening/Confirmation = 291 with islet abs

Annual follow up samples n=103

Review > Lancet Diabetes Endocrinol. 2025 Nov;13(11):980-986. doi: 10.1016/S2213-8587(25)00260-8.

Managing adults with screen-detected islet autoantibody positivity: a pragmatic framework

Nicholas Thomas ¹, Danijela Tatovic ², Angus Jones ³, Parth Narendran ⁴
Affiliations + expand

PMID: 41101877 DOI: 10.1016/S2213-8587(25)00260-8

FULL TEXT LINKS

ELSEVIER

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ACTIONS

66 Cite



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Same result – n=83

Single ab lost – n=13 – usually GADA

Single gained two abs - n=3

Single gained one – n=2

One three became 2

One two became 1

- -

HbA1c in Islet autoantibody positives close to confirmation

Stage 1 n=36
Stage 2 n= 4 – 2 offered Teplizumab (two tipped over to Stage 3)
Abnormal glucose tolerance - n=4 (2 GADA/IA-2A/ZnT8A; 1 IA-2A/ZnT8; 1 GADA/IA-2A)
Stage 3 n= 1

Stage 0 (Single positive) HbA1c data available on n=38

Type 1 diabetes diagnosis - n=2 (high single GADAs)
One participant did not send confirmation sample (3 abs) has been diagnosed
Year 0 follow up – High single GADA – raised Hba1c
Year 1 follow up – High Single IA-2A – raised HbA1c

Summary

- Almost 14,000 adults have been recruited to the T1DRA study
- Three quarters of participants are female
- Approximately one third of recruits are FDRs
- In the dataset thus far,
 - 0.5% of adults are multiple islet autoantibody positive
 - 3.1% of adults are single islet autoantibody positive
 - tGADA are the most prevalent
 - IAA are unexpectedly common
- Annual monitoring is ongoing in all positive participants
- Increasing focus on recruiting individuals from diverse backgrounds in the general population

Acknowledgements

Many thanks to the T1DRA Study Participants



Study Coordinator: Rachel Aitken

Study nurse: Clare Megson

Database Manager: Stu Toms

Antibody testing: The Alistair Williams Antibody Facility

Study support: Diabetes and Metabolism Research Group

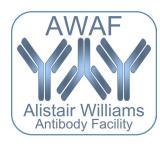
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