

[Date]

[Our GP ref]

[GP address]

Dear Dr [GP name],

ASCEND

Clinical Trial Service Unit (CTSU)
Richard Doll Building
University of Oxford
Old Road Campus
Headington
Oxford
OX3 7LF

Office telephone: 01865 743888

Office fax: 01865 743981

Freefone: 0800 585323

e-mail: ascend@ctsu.ox.ac.uk

Website: www.ctsu.ox.ac.uk/ascend

**[Patient name], [DOB], [address]
ASCEND Study Ref: [100-1234]
[Blood] [and] [urine] results from the ASCEND trial**

As you know [Mr Patient] [is participating in the ASCEND study. He recently provided [a non-fasting blood sample] [and] [a urine specimen] which was mailed to our central laboratory. Routine analysis of [this sample] [these samples] revealed the following results:

Total cholesterol:	mmol/L
HbA _{1c} DCCT:	%
HbA _{1c} IFCC :	mmol/mol
HDL-cholesterol:	mmol/L

Urinary microalbumin/creatinine ratio*: mg/mmol creatinine

[†] NICE guidelines: microalbuminuria defined as ratio ≥ 2.5 mg/mmol [men] or 3.5 mg/mmol [women]]

Please feel free to telephone Freefone 0800 585323 if you wish to discuss any aspect of the study. Many thanks for your help.

Yours sincerely



Dr Jane Armitage



Dr Louise Bowman

Study Coordinators