



CervicalCheck
AN CLÁR NAISÍONTA SCAGTHÁSTÁLA CEIM
THE NATIONAL CERVICAL SCREENING PROGRAM

This section is not for
GP practices

CervicalCheck

Detach the vial number
label from the vial and
place it here

Incomplete forms may be returned

Please verify with the client that the form are correct.
Once verified please attach the label from the sample vial and

Please use
every effort
to provide
the PPSN

Complete MCRN of
CervicalCheck
CONTRACT HOLDER

A. Client's Details

Personal Public
Service Number

CSP ID

Hospital Number
(if applicable)

Date of Birth

Surname Use BLOCK CAPITALS

First Name

Middle Name

Surname at Birth

Mother's Maiden Name

Full Postal Address (The result letter will be sent to this address)

Eircode:

Contact
Telephone No.

To ensure accurate
identification, please
confirm details with
the woman and
complete this section
in its entirety

Ensure that consent is
recorded here
(signature, witnessed
mark, verbal with note
of doctor /nurse)

B. Consent

I have checked that all of the information on this form is correct.
I have read and understood the information and I consent to take part in CervicalCheck.

Client's Signature:

CervicalCheck does not accept third party consent for a client unless a family member or carer have specific legal authority to do so.

C. Details of Contracted Clinic

Medical Council Registration
Number of contracted doctor:

OR

Clinic code:
(CLIN COLP GYN PPCC STI or ONC)

Contracted Doctor
or Clinic's Name:

Address:

Telephone No.

Complete name,
address &
phone number
of CervicalCheck
CONTRACT
HOLDER

D. Sampletaker's details

MCRN or NMBI

Sampletaker's name:

Complete Section D with the details
of the HEALTH PROFESSIONAL WHO
TOOK THE TEST

E. Cervical Screening Test Information

Date of Test

Sample site

☐ Cervix

☐ Vault (post total hysterectomy)

Identify the sample site

Where the cervix is present, the sampletaker must visualise the entire cervix and sample it correctly with 5 x 360° rotations of the broom/brush. Submission of the sample is confirmation that this has been done.

F. Relevant clinical details

LMP

☐ OCP/Hormones/HRT

☐ Pre/Post Transplant

☐ Post-coital bleeding

☐ IUCD

☐ Dialysis

☐ Post-menopausal bleeding

☐ Post-menopausal

☐ HIV Positive

☐ Sub-total Hysterectomy

☐ Total Hysterectomy

Tick ONLY clinically
appropriate boxes

G. Screening and Treatment History

LABORATORY USE ONLY

Date Received in
Laboratory

Accession number

1°

2°

TZ Cells

Yes ☐

No ☐

Date Reported

Path

Management recommendation

Signature