

Colposcopy manual form

Recorded by regional Te Whatu Ora colposcopy/oncology service or non-Te Whatu Ora colposcopy private specialist service.

Colposcopy clinic name:		HPI Facility ID:		<input type="checkbox"/> Te Whatu Ora Hospital site <input type="checkbox"/> Non-Te Whatu Ora site (Private)
Colposcopist		HPI CPN number		
Date referral received				
Date referral accepted				
Appointment date				

Participants details.

NHI:		Date of birth:		Ethnicity:	
Last name:			First name(s):		
Residential address:					

Referred by

Name:		Health practitioner HPI CPN:	<input type="checkbox"/> GP <input type="checkbox"/> Nurse <input type="checkbox"/> Other
HPI Facility ID and facility name:			
Method of referral:	Letter <input type="checkbox"/> Yes <input type="checkbox"/> No	Phone <input type="checkbox"/> Yes <input type="checkbox"/> No	Other (e-referral) <input type="checkbox"/> Yes <input type="checkbox"/> No

Type of referral

First assessment (new case)	<input type="checkbox"/> Yes	<input type="checkbox"/> No		
Subsequent assessment (follow-ups)	<input type="checkbox"/> 1st	<input type="checkbox"/> 2nd	<input type="checkbox"/> 3rd	<input type="checkbox"/> 4th

Note: If a participant is referred from another District or specialist for follow-up or treatment, this should be noted as a subsequent assessment (follow-up).

Assessment of the reason for referral

1.	<input type="checkbox"/> <input type="checkbox"/>	A. Clinical suspicion of invasion B. Any cytological glandular abnormality
2.	<input type="checkbox"/> <input type="checkbox"/> <input type="checkbox"/> <input type="checkbox"/>	Positive/detected high risk HPV test results A. HPV16/18 with or without cytology B. HPV other with HSIL cytology C. HPV where the participant is immune compromised D. HPV other with LSIL on two occasions if over 50 and 3 occasions if under 50
3.	<input type="checkbox"/> <input type="checkbox"/> <input type="checkbox"/> <input type="checkbox"/>	Clinical reasons only (e.g. abnormal cervical appearance) A. A low-grade clinical assessment B. High grade clinical assessment C. Suspicious of invasive cancer clinical assessment D. Other clinical assessment
4.		Optional comments about referral:

Colposcopy visit details.

Date of visit	
Admission type	<input type="checkbox"/> Outpatient <input type="checkbox"/> Day patient <input type="checkbox"/> Inpatient
First assessment (new case)	<input type="checkbox"/> Yes <input type="checkbox"/> No
Subsequent assessment (follow-ups)	<input type="checkbox"/> 1st <input type="checkbox"/> 2nd <input type="checkbox"/> 3rd <input type="checkbox"/> 4th
Pregnant	<input type="checkbox"/> Yes <input type="checkbox"/> No
Colposcopy performed	<input type="checkbox"/> Yes <input type="checkbox"/> No
Colposcopy site	<input type="checkbox"/> Cervical <input type="checkbox"/> Vaginal <input type="checkbox"/> Both cervical and vaginal <input type="checkbox"/> Other
Review / results discussed	<input type="checkbox"/> Yes <input type="checkbox"/> No
Arranged treatment	<input type="checkbox"/> Yes <input type="checkbox"/> No

Colposcopy findings

Squamocolumnar junction visible	<input type="checkbox"/> Completely	<input type="checkbox"/> Partially	<input type="checkbox"/> Not visible	<input type="checkbox"/> N/A
Transformation Zone Type	<input type="checkbox"/> 1	<input type="checkbox"/> 2	<input type="checkbox"/> 3	
Lesion present	<input type="checkbox"/> Yes	<input type="checkbox"/> No	<input type="checkbox"/> Inconclusive	
Number of quadrants involved	<input type="checkbox"/> 1	<input type="checkbox"/> 2	<input type="checkbox"/> 3	<input type="checkbox"/> 4
Normal findings noted	<input type="checkbox"/> Yes	<input type="checkbox"/> No		
Abnormal visible lesion	<input type="checkbox"/> Yes	<input type="checkbox"/> No		
Limits of lesion visible	<input type="checkbox"/> Yes	<input type="checkbox"/> No		
Predicted grade(s) of abnormality				
Low-grade squamous	<input type="checkbox"/> Yes	<input type="checkbox"/> No		
High-grade squamous	<input type="checkbox"/> Yes	<input type="checkbox"/> No		
Glandular (AIS)	<input type="checkbox"/> Yes	<input type="checkbox"/> No		
Micro-invasive cancer	<input type="checkbox"/> Yes	<input type="checkbox"/> No		
Invasive cancer (squamous/glandular)	<input type="checkbox"/> Yes	<input type="checkbox"/> No		

Actions taken during visit

Cervical/Vaginal Sample:	Cytology	<input type="checkbox"/> Yes	<input type="checkbox"/> No	HPV test	<input type="checkbox"/> Yes	<input type="checkbox"/> No
Biopsy:	<input type="checkbox"/> Yes <input type="checkbox"/> No					
Site of biopsy (biopsies) taken:						
If no biopsy taken, give reasons:						
Treatment this visit	<input type="checkbox"/> Yes	<input type="checkbox"/> No				
Type of treatment						
Wire loop excisional procedure	<input type="checkbox"/> Yes	<input type="checkbox"/> No				
Laser ablation	<input type="checkbox"/> Yes	<input type="checkbox"/> No				
Ablation by other means other than laser	<input type="checkbox"/> Yes	<input type="checkbox"/> No				
Cold knife cone	<input type="checkbox"/> Yes	<input type="checkbox"/> No				
Diathermy cone	<input type="checkbox"/> Yes	<input type="checkbox"/> No				
Laser cone	<input type="checkbox"/> Yes	<input type="checkbox"/> No				
Hysterectomy	<input type="checkbox"/> Yes	<input type="checkbox"/> No	<input type="checkbox"/> Total		<input type="checkbox"/> Subtotal	

Other (describe)	
Diagram/photo of lesion	<input type="checkbox"/> Yes <input type="checkbox"/> No
Local or general anaesthesia use	<input type="checkbox"/> Local <input type="checkbox"/> General <input type="checkbox"/> N/A
Reason for GA	
Follow-up management recommended	<input type="checkbox"/> Yes <input type="checkbox"/> No
Next visit recommended in	months

Data received from colposcopy visit.

Date histology specimen report received by colposcopy service	
Decision to treat date	
Date participant informed	
Histological specimen taken satisfactory for interpretation	<input type="checkbox"/> Yes <input type="checkbox"/> No
Biopsy result	<input type="checkbox"/> Negative <input type="checkbox"/> CIN1/HPV <input type="checkbox"/> CIN2 <input type="checkbox"/> CIN2/3 <input type="checkbox"/> CIN3 <input type="checkbox"/> AIS <input type="checkbox"/> Adenocarcinoma <input type="checkbox"/> Squamous carcinoma <input type="checkbox"/> Adenosquamous carcinoma <input type="checkbox"/> Other

Did not attend.

Scheduled visit date	
For	
1st assessment	<input type="checkbox"/> Yes <input type="checkbox"/> No
Treatment	<input type="checkbox"/> Yes <input type="checkbox"/> No
Follow-up after treatment/other	<input type="checkbox"/> Yes <input type="checkbox"/> No
Reason for DNA (if known)	
Rescheduled appointment date	

Support to screening provider referral	<input type="checkbox"/> Yes	<input type="checkbox"/> No
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Discharged from colposcopy.

To screen taker Name of health worker and HPI CPN Health facility HPI	<input type="checkbox"/> Yes <input type="checkbox"/> No	Date of discharge	<input type="checkbox"/> Recall 1 month <input type="checkbox"/> First TOC <input type="checkbox"/> Second TOC <input type="checkbox"/> HPV+/- Cyto 12 months <input type="checkbox"/> HPV 3 years <input type="checkbox"/> HPV 5 years <input type="checkbox"/> Cease screening
To oncology Name of health worker and HPI CPN Health facility HPI	<input type="checkbox"/> Yes <input type="checkbox"/> No	Date of discharge	
Other Specify service Name of health worker and HPI CPN Health facility HPI	<input type="checkbox"/> Yes <input type="checkbox"/> No	Date of discharge	

Please return this completed form to the NCSP-Register central team (Whakarongorau) by using one of the following:

1. **Email:** info@ncspregister.health.nz
2. **Postal (not for courier):**
 Whakarongorau Aotearoa
 NCSP-NCC CX team
 PO Box 5895
 Wellington 6140

3. **Physical:**

Whakarongorau Aotearoa
NSP-NCC CX team
Level 5, 36 Customhouse Quay
Wellington

NCSP Policies and Standards

Section 6: Providing a Colposcopy Service

Appendix 2 – Section 112M of Part 4A of the Health Act 1956

112M Duty of persons performing colposcopic procedures

- (1) Every person who performs a colposcopic procedure on a participant must—
- (a) explain the procedure to the participant; and
 - (b) provide information, to the extent that is reasonable in the circumstances, about the objectives of the NCSP and the NCSP register, the importance of having regular screening tests, who has access to information on the NCSP register, and the uses to which that information may be put; and
 - (c) if he or she believes that the participant is not enrolled in the NCSP, advise that they will be enrolled but that they may prevent or cancel that enrolment by notifying the NCSP manager under section 112G; and
 - (d) cause a report in relation to that colposcopic procedure to be forwarded to the NCSP manager.
- (2) A report under subsection (1)(d) must—
- (a) be provided free of charge; and contain the information requested.