

NATIONAL CENTRE FOR DISEASE INFORMATICS AND RESEARCH

NATIONAL CANCER REGISTRY PROGRAMME

Indian Council of Medical Research

PATTERNS OF CARE AND SURVIVAL STUDIES

Patient Information Form - Cancer Cervix

A. IDENTIFYING, DEMOGRAPHIC AND DIAGNOSTIC INFORMATION

1.	Name of Participating Centre	Centre Code	<input type="text"/>	<input type="text"/>	<input type="text"/>
2.	Registration Number (as in HBCR) (First 2 digits are for year of registration and the next 5 digits for actual registration number)	<input type="text"/>	<input type="text"/>	<input type="text"/>	<input type="text"/>
		Year	Reg. No.		
3.1	(a) Name of Source of Registration (Reporting Institution (RI) / Hospital)	Code	<input type="text"/>	<input type="text"/>	<input type="text"/>
	(b) Name of Department / Unit etc.	Code	<input type="text"/>	<input type="text"/>	<input type="text"/>
	(c) Name of Physician	Mobile No.		
3.2	Hospital Registration Number				
3.3	Date of Registration at Source of Registration / Date of Reporting at this Hospital	<input type="text"/>	<input type="text"/>	<input type="text"/>	
		dd	mm	yy	
3.4	Case Registered As				
	(1) Out-patient (OP) <input type="checkbox"/>	(2) In-patient (IP) <input type="checkbox"/>			
	(3) OP and IP <input type="checkbox"/>	(4) Not Registered - Clinical Consultation / Opinion <input type="checkbox"/>			
	(5) Not Registered - Pathology Consultation / Opinion <input type="checkbox"/>	(8) Others (specify)..... <input type="checkbox"/>			
4.	Date of First Diagnosis (Date of first attendance to any hospital for this disease)	<input type="text"/>	<input type="text"/>	<input type="text"/>	
		dd	mm	yy	
5.	Full Name of Patient				
	(At least one name is compulsory)	First	Second	Last	
6.	Name of Spouse / Father / Mother / Caretaker (give any two names)				
	Name	Mobile No.	Name	Mobile No.
7.	Place of Residence: Permanent place of residence (where the person has been residing for the past one year (at least))				
	<u>Urban Areas</u> (Town / city / any other)	<u>Non-urban / Rural Areas</u>			
	House No.....	House No. and Ward			
	Road / Street Name.....	Name of Gram Panchayat / Village, etc:			
	Area / Locality.....	Name of Sub-Unit of District (Taluk / Tehsil / Other):			
	Ward / Corporation / Division <input type="text"/>			
	Name of City / Town	Name of PHC / Sub Centre			
	Name of District (in capitals)	Postal Pin Code <input type="text"/>			
	Telephone No(s): Off.	Res.			
	Mobile No.	Email ID			
	Aadhaar (Unique Identification) No.				

8. Duration of Stay (at the permanent place of residence (in years))

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9.1 Local Address

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Name of City/Town/District

Pin Code

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9.2 Name & Address of Referring / Family Doctor

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Name of City/Town/District

Pin Code

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10. Age (in years)

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Date of Birth

dd	mm	yy

11. Sex - Female

12. Method of Diagnosis

(1) Clinical Only ☐(2) Microscopic ☐(3) X-Ray / Imaging Techniques ☐(8) Others ☐(9) Unknown ☐

Microscopic (if 2 above)

(1) Histology of Primary ☐(2) Histology of Metastasis ☐(3) Cytology of Primary ☐(4) Cytology of Metastasis ☐

X-Ray / Imaging Techniques (if 3 above)

(1) X-Ray ☐(2) Isotopes ☐(3) Angiography ☐(4) Ultrasonogram ☐(8) All Others (specify)..... ☐

Others (if 8 above)

(1) Surgery or Autopsy without Histology ☐(2) Specific Biochemical and /
or Immunological Tests ☐(8) Others (specify)..... ☐

13. Anatomical Site of Specimen / Biopsy / Smear

14. Complete Pathological Diagnosis: (With complete description of Primary Site of Tumour and Morphological Diagnosis)

14.1 Primary Site of Tumour - Topography

14.2 Morphology

14.3 Pathology Slide No.

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Date

dd	mm	yy

15. Coding According to ICD-O-3:

15.1 Primary Site of Tumour - Topography C

(Include sub-site if any)

		.	
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15.2 Primary Histology - Morphology M

If morphology is that of metastasis mention Primary Site above and

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15.3 Secondary Site of Tumour C

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15.4 Morphology of Metastasis M

If the morphology diagnosis is only that of metastatic site, mention the Primary Site as decided by the treating clinician either through discussion or from case record.

				/	/
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16. Sequence

(0) One Primary Only ☐(3) Third of three or more primaries ☐(6) Sixth of six or more primaries ☐(9) Unspecified sequence number (Unknown) ☐(1) First of two or more primaries ☐(4) Fourth of four or more primaries ☐(7) Seventh of seven or more primaries ☐(2) Second of two or more primaries ☐(5) Fifth of five or more primaries ☐(8) Eighth or later primary ☐

B. DETAILS OF SOCIOECONOMIC STATUS, FAMILY INCOME, OCCUPATION, ETC.

Co-Morbid Conditions	Yes	No	Unknown	Co-Morbid Conditions	Yes	No	Unknown
(1) Tuberculosis	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	(2) Hypertension	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
(3) Diabetes	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	(4) Ischaemic Heart Disease	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
(5) Bronchial Asthma	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	(6) Allergic Conditions (specify).....	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
(7) Hepatitis / HBsAg +ve	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	(8) Others (specify).....	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
(10) AIDS/HIV +ve	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>				

C. DETAILS OF STAGE (Tick (✓) as appropriate)**1. Staging System Followed**

(1) FIGO staging ☐ (2) TNM staging ☐ (8) Others (specify)..... ☐ (9) Unknown ☐

2. Staging done at

(1) Reporting Institution ☐ (2) Previous Institution ☐ (8) Others (specify)..... ☐ (9) Unknown ☐

3. Stage

I <input type="checkbox"/>	IA <input type="checkbox"/>	IA1 <input type="checkbox"/>	IA2 <input type="checkbox"/>	
	IB <input type="checkbox"/>	IB1 <input type="checkbox"/>	IB2 <input type="checkbox"/>	
II <input type="checkbox"/>	IIA <input type="checkbox"/>	IIA1 <input type="checkbox"/>	IIA2 <input type="checkbox"/>	IIB <input type="checkbox"/>
III <input type="checkbox"/>	IIIA <input type="checkbox"/>	IIIB <input type="checkbox"/>		
IV <input type="checkbox"/>	IVA <input type="checkbox"/>	IVB <input type="checkbox"/>	Unknown <input type="checkbox"/>	

If TNM specify.....

T			

N	

M	

4. Investigations for Staging

	Yes	No	Unknown		Yes	No	Unknown
(1) Haemogram	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	(2) Biochemistry	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
(3) Chest X-ray	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	(4) Examination under Anaesthesia	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
(5) Cystoscopy	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	(6) Ultrasound of abdomen & pelvis	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
(7) Proctosigmoidoscopy	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	(8) Others (specify).....	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
(10) CT Scan	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	(11) MRI	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Specify any relevant abnormal findings _____

5. The actual assessment of staging was done by

(1) One Consultant Oncologist (CO) only	<input type="checkbox"/>	(2) Two COs from same department	<input type="checkbox"/>
(3) Two COs from different departments	<input type="checkbox"/>	(4) Tumour Board/Joint Clinic	<input type="checkbox"/>
(8) Others (specify).....	<input type="checkbox"/>	(9) Unknown	<input type="checkbox"/>

D. DETAILS OF CANCER DIRECTED TREATMENT (CDT) (Tick (✓) as appropriate)**6. Treatment given prior to Registration at Reporting Institution (RI)**(0) No ☐ (2) Yes ☐ (9) Unknown ☐

If yes,

6.1 Type of Prior Treatment Given:	Yes	No	Unknown	If yes, Date of completion of treatment			
(1) Surgery	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<table border="1" style="display: inline-table; border-collapse: collapse;"><tr><td style="width: 33px; height: 20px;"></td><td style="width: 33px; height: 20px;"></td><td style="width: 33px; height: 20px;"></td></tr></table>			
(2) Radiotherapy	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<table border="1" style="display: inline-table; border-collapse: collapse;"><tr><td style="width: 33px; height: 20px;"></td><td style="width: 33px; height: 20px;"></td><td style="width: 33px; height: 20px;"></td></tr></table>			
(3) Chemotherapy	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<table border="1" style="display: inline-table; border-collapse: collapse;"><tr><td style="width: 33px; height: 20px;"></td><td style="width: 33px; height: 20px;"></td><td style="width: 33px; height: 20px;"></td></tr></table>			
(8) Others (specify).....	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<table border="1" style="display: inline-table; border-collapse: collapse;"><tr><td style="width: 33px; height: 20px;"></td><td style="width: 33px; height: 20px;"></td><td style="width: 33px; height: 20px;"></td></tr></table> <div style="display: flex; justify-content: space-around; font-size: small;">dd mm yy</div>			

6.2 Details of Prior Treatment (including treatment interruption and complications) _____

7. Treatment at Reporting Institution

7.1 Intention to Treat

 (1) Curative/Radical ☐ (2) Palliative ☐
 (3) No treatment ☐ (9) Unknown ☐

7.2 If Palliative yes,

 (1) Palliative RT only ☐ (2) Palliative RT + CT ☐ (3) Palliative CT only ☐
 (4) Pain & Symptom Relief Drugs (specify)..... ☐ (5) Palliative Surgery ☐
 (8) Others (specify)..... ☐ (9) Unknown ☐

7.3 Type of Cancer Directed Treatment Planned at Reporting Institution:

	Yes	No	Unknown
(1) Surgery	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
(2) Radiotherapy	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
(3) Chemotherapy	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
(8) Others (specify).....	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

8. Performance Status (WHO) before treatment
 (0) Able to carry out all normal activity without restriction ☐
 (1) Restricted in physically strenuous activity but ambulatory and able to carry out light work ☐
 (2) Ambulatory and capable of all self-care but unable to carry out any work; up and about more than 50% of waking hours ☐
 (3) Capable of only limited self-care; confined to bed or chair more than 50% of waking hours ☐
 (4) Completely disabled; cannot carry on any self-care; totally confined to bed or chair ☐
 (9) Unknown ☐

9. Surgery

- 9.1 (0) Surgery not planned ☐ (1) Yes, done as planned/advised ☐
 (2) Surgery planned but not taken ☐ (8) Others (*specify*)..... ☐
 (9) Unknown ☐

9.2 If yes, Type of Surgical Procedure

- (1) Total (extra-fascial) Abdominal Hysterectomy (Type I) ☐ (5) Pelvic Excenteration ☐
 (2) Modified Radical Hysterectomy (Type II) ☐ (8) Others (*specify*)..... ☐
 (3) Radical Abdominal Hysterectomy (Type III) ☐ (9) Unknown ☐
 (4) Extended Radical Hysterectomy (Type IV) ☐

9.3 If yes, Lymphadenectomy (1) Not Done ☐ (2) Done ☐

9.4 Date of Surgical Procedure

<i>dd</i>	<i>mm</i>	<i>yy</i>

10. Surgical Histopathological Findings

10.1 Size of Tumour in Cervix

- (0) NA* ☐ (1) No Viable tumor ☐ (2) Tumor present (Size NK) ☐
 (3) Tumor < 2 cm ☐ (4) Tumor 2 - 4 cm ☐ (5) Tumor 4 - 6 cm ☐
 (6) Tumor > 6 cm ☐ (9) NK** ☐

10.2 Thickness of Cervical Invasion

- (0) NA ☐ (1) < 1/3 / in-situ ☐ (2) < 1/2 ☐
 (3) = 2/3 ☐ (4) Full thickness ☐ (9) NK ☐

10.3 Involvement of Uterus

- (0) NA ☐ (1) Not involved ☐
 (2) Involved ☐ (3) Endocervix extrn. ☐ (9) NK ☐

10.4 Thickness of Uterine Invasion

- (0) NA ☐ (1) Endometrium ☐ (2) Myometrium ☐
 (3) Serosa + ve ☐ (9) NK ☐

* Not Applicable, ** Not Known

10.5 Involvement of Vagina (*Cut edges*)

(0) NA ☐ (1) Negative ☐ (2) Positive ☐
 (3) Edge Close / +ve ☐ (9) NK ☐

10.6 Involvement of Parametrium

(0) NA ☐ (1) Negative ☐ (2) Positive ☐ (9) NK ☐

10.7 Tumor Emboli

(0) NA ☐ (1) No ☐ (2) Yes ☐ (9) NK ☐

10.8 Involvement of Ovaries

(0) NA ☐ (1) Not involved ☐ (2) Involved ☐ (9) NK ☐

10.9 Involvement of Regional Nodes

(0) NA ☐ (1) Not involved ☐ (2) Involved (Metastatic) ☐ (9) NK ☐

10.10 Site of Involved Regional Nodes

(0) NA ☐ (1) Not involved ☐ (2) Ilio- obtur. ☐
 (3) Iliac node ☐ (4) Obturator ☐ (5) Pre- Sacral ☐
 (6) Paracervical ☐ (7) Parametrial ☐ (9) NK ☐

10.11 Para-aortic Node(s)

(0) NA ☐ (1) Not involved ☐ (2) Involved (Metastatic) ☐ (9) NK ☐

10.12 Laterality of Positive Nodes

(0) NA ☐ (1) Not involved ☐
 (2) Unilateral (R / L) ☐ (3) Bilateral ☐ (9) NK ☐

10.13 No. of Positive Nodes

0 ☐ 1 ☐ 2 ☐ 3 ☐ 4 ☐ >4 ☐ NK ☐

10.14 Tumour Grade

(0) NA ☐ (1) Grade I ☐ (2) Grade II ☐
 (3) Grade III ☐ (4) Grade IV ☐ (9) NK ☐

10.15 Lymphovascular Space Involvement

(0) NA ☐ (1) Not involved ☐ (2) Involved ☐ (9) NK ☐

11. Radiotherapy

- 11.1 (0) Radiotherapy (RT) not planned ☐ (1) Yes, RT given as planned ☐
 (2) Yes, RT given, but incomplete ☐ (3) RT planned but not taken ☐
 (specify reason)..... (specify reason).....
 (8) Others (specify)..... ☐ (9) Unknown ☐

11.2 Type of RT

- (1) Teletherapy (External RT) ☐ (2) Brachytherapy ☐ (3) 3D Conformal ☐
 (4) IMRT ☐ (5) Electron Beam ☐
 (8) Others (specify)..... ☐ (9) Unknown ☐

11.3 Details of External RT

	I	II	III
Technique (specify)
Type of beam (Photon/Electron)
Energy
Field Size
Field/day
Total Tumour Dose (cGy)
Total No. of Fractions
Fractions / week
Region(s) of Irradiation
Interruption - YES (Y) / NO (N)

Date first started

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Date last ended

dd	mm	yy

dd	mm	yy

dd	mm	yy

11.4 Details of Brachytherapy

	I	II	III	IV	V	>V
Date	<div>dd mm yy</div>	<div>dd mm yy</div>	<div>dd mm yy</div>	<div>dd mm yy</div>	<div>dd mm yy</div>	<div>dd mm yy</div>
Type of applicators
Type of Dose Rate (LDR/MDR/HDR/PDR)
Dose in cGy. to point A/ ICRU Reference Points/ Any other specify....
Dose rate in cGy.
Dose to Rectum in cGy.
Dose to Bladder in cGy.

12. Chemotherapy

12.1 (0) Chemotherapy (CT) not planned	<input type="checkbox"/>	(1) Yes, CT given as planned	<input type="checkbox"/>
(2) Yes, CT given, but incomplete	<input type="checkbox"/>	(3) CT planned but not taken	<input type="checkbox"/>
(8) Others (specify).....	<input type="checkbox"/>	(9) Unknown	<input type="checkbox"/>

If Yes,

12.2 Type of CT

(1) Anterior/neo-adjuvant/induction	<input type="checkbox"/>	(2) Concurrent	<input type="checkbox"/>
(3) Adjuvant	<input type="checkbox"/>	(4) Combination of any of the above	<input type="checkbox"/>
(8) Others (specify).....	<input type="checkbox"/>	(9) Unknown	<input type="checkbox"/>

12.3 Drug(s) comprising CT Regimen

(1) Single (specify).....	<input type="checkbox"/>	(2) Two Drug (specify).....	<input type="checkbox"/>
(3) More than two drug (specify).....	<input type="checkbox"/>	(8) Others (specify).....	<input type="checkbox"/>
(9) Unknown	<input type="checkbox"/>		

12.4 Other Details of CT

Height in cms.	<input type="text"/>	<input type="text"/>	<input type="text"/>	Weight in Kg.	<input type="text"/>	<input type="text"/>	
Cycles	I	II	III	IV	V	VI	>VI
Regimen	<hr/>	<hr/>	<hr/>	<hr/>	<hr/>	<hr/>	<hr/>
Date(s)	<input type="text"/> <i>dd mm yy</i>	<input type="text"/> <i>dd mm yy</i>	<input type="text"/> <i>dd mm yy</i>	<input type="text"/> <i>dd mm yy</i>	<input type="text"/> <i>dd mm yy</i>	<input type="text"/> <i>dd mm yy</i>	<input type="text"/> <i>dd mm yy</i>
Day(s)	<hr/>	<hr/>	<hr/>	<hr/>	<hr/>	<hr/>	<hr/>
Drug (s)							
Name	Dose	Dose	Dose	Dose	Dose	Dose	Dose
.....
.....
.....
.....
.....
.....
.....
Date of start of First Cycle of CT	<input type="text"/> <i>dd mm yy</i>			Date of completion of Last Cycle of CT	<input type="text"/> <i>dd mm yy</i>		

13. Response of Disease (Adopted from WHO) to RT / RT + CT (6-12 weeks after completion of treatment)

- | | | | |
|--------------------------|--------------------------|---|--------------------------|
| (0) RT / CT not received | <input type="checkbox"/> | (1) Complete response -
No Evidence of Disease | <input type="checkbox"/> |
| (2) Partial response | <input type="checkbox"/> | (3) No change | <input type="checkbox"/> |
| (4) Progressive disease | <input type="checkbox"/> | (5) Post Surgical - Adjuvant | <input type="checkbox"/> |
| (9) Unknown | <input type="checkbox"/> | | |

13.1 Date(s) of assessment of
response to RT / RT + CT

dd mm yy

dd mm yy

14.1 Date of Completion of Initial Cancer Directed Treatment at RI

<i>dd</i>	<i>mm</i>	<i>yy</i>

14.2 Complications During Treatment

(0) No ☐ (2) Yes ☐ (9) Unknown ☐

If Yes,

Nature of Complication(s)	Maximum Grade	Date of Onset	Resolved		Date last seen (if resolved)												
			Yes	No													
_____	_____	<table border="1"><tr><td></td><td></td><td></td></tr><tr><td><i>dd</i></td><td><i>mm</i></td><td><i>yy</i></td></tr></table>				<i>dd</i>	<i>mm</i>	<i>yy</i>	<input type="checkbox"/>	<input type="checkbox"/>	<table border="1"><tr><td></td><td></td><td></td></tr><tr><td><i>dd</i></td><td><i>mm</i></td><td><i>yy</i></td></tr></table>				<i>dd</i>	<i>mm</i>	<i>yy</i>
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<i>dd</i>	<i>mm</i>	<i>yy</i>															
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<i>dd</i>	<i>mm</i>	<i>yy</i>															
<i>dd</i>	<i>mm</i>	<i>yy</i>															

15. Performance Status at 6-12 Weeks of Completion of all of CDT

- (0) Able to carry out all normal activity without restriction ☐
- (1) Restricted in physically strenuous activity but ambulatory and able to carry out light work ☐
- (2) Ambulatory and capable of all self-care but unable to carry out any work; up and about more than 50% of waking hours ☐
- (3) Capable of only limited self-care; confined to bed or chair more than 50% of waking hours ☐
- (4) Completely disabled; cannot carry on any self-care; totally confined to bed or chair ☐
- (9) Unknown ☐

15.1 Date of Assessment of Performance Status

<i>dd</i>	<i>mm</i>	<i>yy</i>

E. FOLLOW-UP INFORMATION (USE SEPARATE PAGE FOR EACH VISIT)**16. Due Date for Follow up**

<i>dd</i>	<i>mm</i>	<i>yy</i>

Date of Actual Follow-up

<i>dd</i>	<i>mm</i>	<i>yy</i>

Follow-up Visit No.

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16.1 Method of Follow-up

- | | | |
|--|---|---|
| (0) No follow-up <input type="checkbox"/> | (1) Hospital visit <input type="checkbox"/> | (2) By post <input type="checkbox"/> |
| (3) Through telephone <input type="checkbox"/> | (4) Home visit <input type="checkbox"/> | (8) Others (<i>specify</i>)..... <input type="checkbox"/> |
| | | (9) Unknown <input type="checkbox"/> |

16.2 Vital Status

- (1) Alive ☐ (2) Dead ☐ (9) Unknown ☐

16.3 Disease Status (at Follow-up)

- | | |
|---|--|
| (1) No Evidence of Disease <input type="checkbox"/> | (2) Residual Disease only <input type="checkbox"/> |
| (3) Local recurrence <input type="checkbox"/> | (4) Regional/Nodal recurrence <input type="checkbox"/> |
| (5) Distant metastasis : <i>specify site</i> <input type="checkbox"/> | (9) Unknown <input type="checkbox"/> |

16.4 If Disease is present, indicate Basis of Diagnosis

- | | | |
|---|---|---|
| (1) Histopathology <input type="checkbox"/> | (2) Cytopathology <input type="checkbox"/> | (3) FNAC <input type="checkbox"/> |
| | (Other than FNAC) | |
| (4) Bone Marrow <input type="checkbox"/> | (5) Peripheral Smear <input type="checkbox"/> | (6) Radiological <input type="checkbox"/> |
| (7) Clinical <input type="checkbox"/> | (8) Others (<i>specify</i>)..... <input type="checkbox"/> | (9) Unknown <input type="checkbox"/> |

16.5 Treatment if 16.3 above indicates presence of disease

- (0) No Treatment ☐ (2) Yes, Treatment given ☐ (9) Unknown ☐

16.6 If yes, Details of Treatment and Outcome (*Use separate sheet if necessary*)

16.7 Late Complications of CDT

- (0) Nil ☐ (2) Yes ☐ (9) Unknown ☐

If Yes,

Nature of Complication(s)	Maximum Grade	Date of Onset	Resolved		Date last seen (if resolved)												
			Yes	No													
		<table border="1"><tr><td></td><td></td><td></td></tr><tr><td><i>dd</i></td><td><i>mm</i></td><td><i>yy</i></td></tr></table>				<i>dd</i>	<i>mm</i>	<i>yy</i>	<input type="checkbox"/>	<input type="checkbox"/>	<table border="1"><tr><td></td><td></td><td></td></tr><tr><td><i>dd</i></td><td><i>mm</i></td><td><i>yy</i></td></tr></table>				<i>dd</i>	<i>mm</i>	<i>yy</i>
<i>dd</i>	<i>mm</i>	<i>yy</i>															
<i>dd</i>	<i>mm</i>	<i>yy</i>															
			<input type="checkbox"/>	<input type="checkbox"/>													
			<input type="checkbox"/>	<input type="checkbox"/>													
			<input type="checkbox"/>	<input type="checkbox"/>													
			<input type="checkbox"/>	<input type="checkbox"/>													
			<input type="checkbox"/>	<input type="checkbox"/>													
			<input type="checkbox"/>	<input type="checkbox"/>													
			<input type="checkbox"/>	<input type="checkbox"/>													

17. Second Primary(0) No evidence of second primary ☐ (2) Yes, evidence of second primary ☐ (9) Unknown ☐

If Yes,

17.1 Primary Site of Tumour (ICD-O-3) (Topography)

C

		.	
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17.2 Primary Histology (ICD-O-3) (Morphology)

M

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17.3 Secondary (Metastatic) Site of Tumour (ICD-O-3)

C

		.	
--	--	---	--

17.4 Histology of Metastasis

M

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17.5 Method of Diagnosis

(1) Clinical Only ☐ (2) Microscopic ☐ (3) X-Ray / Imaging Techniques ☐ (8) Others ☐ (9) Unknown ☐

Microscopic (If 2 above)

- (1) Histology of Primary ☐
- (2) Histology of Metastasis ☐
- (3) Autopsy with Histology ☐
- (4) Bone Marrow ☐
- (5) Blood Film ☐
- (6) Cytology of Primary ☐
- (7) Cytology of Metastasis ☐

X-Ray / Imaging Techniques (If 3 above)

- (1) X-Ray ☐
- (2) Isotopes ☐
- (3) Angiography ☐
- (4) Ultrasonogram ☐
- (8) All Others (specify)..... ☐

Others (If 8 above)

- (1) Endoscopy ☐
- (2) Surgery or Autopsy without Histology ☐
- (3) Specific Biochemical and /
or Immunological Tests ☐
- (8) Others (specify)..... ☐

17.6 Date of Diagnosis

dd	mm	yy

17.7 Details of Treatment and Outcome: *Use separate appropriate form.***18. If Dead,****18.1 Cause of Death**

- (1) As a result of cancer ☐ (2) Most probably due to cancer ☐ (3) Intercurrent Death ☐
- (4) Treatment related Death ☐ (8) Others (specify)..... ☐ (9) Unknown ☐

18.2 Date of Death

dd	mm	yy

19. Remarks (add additional sheet(s) if necessary)
