# **Gynaecological cancer (clinical) DSS**

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# Gynaecological cancer (clinical) DSS

### Identifying and definitional attributes

Metadata item type:	Data Set Specification		
METEOR identifier:	421105		
Registration status:	Health!, Superseded 14/05/2015		
DSS type:	Data Set Specification (DSS)		
Scope:	The purpose of the Gynaecological cancer (clinical) data set specification (DSS) is to define data standards for the national collection of gynaecological cancer data so that data collected is consistent and reliable. The data set specification is not mandated for collection but is recommended as best practice if gynaecological cancer data is to be collected. It enables individual treatment centres or health service areas to develop collection methods and policies appropriate for their service.		
	The Gynaecological cancer (clinical) data set specification is used in conjunction with the Cancer (clinical) data set specification (CCDSS). The data elements with obligations described as mandatory or conditional for collection are recommended as best practice, while the data items described as optional are for collection at the discretion of the treating centre and may be contingent, for example, on the availability of resources.		
	The scope for the Gynaecological cancer (clinical) DSS is to collect comprehensive data encompassing the time a person is first referred for the investigation of symptoms and for the entire duration of their illness so that treatment and outcomes are captured.		
	The definitions used in this data set specification are designed to capture the provision of cancer care on a day-to-day level. They relate to the realities of cancer care and the need to optimise care by correctly diagnosing, evaluating and managing patients with gynaecological cancer.		
	The data elements specified provide a framework for:		
	<ul> <li>providing a systematic foundation and promoting the delivery of evidence- based care to patients with gynaecological cancer</li> <li>informing treatment guidelines and professional education</li> <li>informing quality assurance</li> <li>guiding resource planning and the evaluation of cancer control activities</li> </ul>		
	Many of the data elements in this data set specification may also be used in the collection of data for other types of cancer.		
	This data set specification is primarily directed at the clinical and clinical epidemiological use of cancer data. Treatment centres such as hospitals, radiotherapy centres and cancer specialist practices are the settings in which implementation of the Gynaecological cancer (clinical) data set specification should be considered. The data set specification can also be used by a wider range of health and health-related establishments that create, use or maintain records on health-care clients.		
Source and reference attributes			
Submitting organisation:	Cancer Australia		
Relational attributes			

Related metadata	Has been superseded by <u>Gynaecological cancer (clinical) NBPDS</u>
references:	Health!, Standard 14/05/2015

### Metadata items in this Data Set Specification

Seq No.	Metadata item	Obligation	Max occurs
1	Person with cancer—cytopathology result, code N	Mandatory	29
2	Person with cancer—lymphovascular invasion indicator, yes/no code N	Mandatory	1
3	Person with cancer—location of lymphovascular invasion of cervix, code N	Conditional	5
	Conditional obligation:		
	This data element is only to be recorded for patients with cervical cancer, as indicated by <u>Person with cancer</u> <u>primary site of cancer</u> , topography code (ICD-O-3) ANN.N, and when <u>Person with cancer</u> <u>lymphovascular invasion</u> indicator, yes/no code N indicates the presence of lymphovascular invasion.		
4	Person with cancer—location of lymphovascular invasion of corpus uteri, code N	Conditional	5
	Conditional obligation:		
	This data element is only to be recorded for patients with endometrial cancer, as indicated by <u>Person with cancer</u> —primary site of cancer, topography code (ICD-O-3) ANN.N, and when <u>Person with cancer</u> —lymphovascular invasion indicator, yes/no code N indicates the presence of lymphovascular invasion.		
5	Person with cancer—extent of primary cancer, cervical cancer staging (FIGO) code $N[N]$	Conditional	1
	Conditional obligation:		
	This data element is only to be recorded for patients with cervical cancer, as indicated by <u>Person with cancer—primary site of cancer, topography code</u> (ICD-O-3) ANN.N.		
6	Person with cancer—extent of primary cancer, endometrial cancer staging (FIGO) code N[N]	Conditional	1
	Conditional obligation:		
	This data element is only to be recorded for patients with endometrial cancer, as indicated by <u>Person with cancer—primary site of cancer, topography code</u> (ICD-O-3) ANN.N.		
7	Person with cancer—extent of primary cancer, ovarian cancer staging (FIGO) code N[N]	Conditional	1
	Conditional obligation:		
	This data element is only to be recorded for patients with ovarian cancer, as indicated by <u>Person with cancer—primary site of cancer, topography code</u> (ICD-O-3) ANN.N.		
8	Person with cancer—distant metastatic cancer indicator, yes/no/not stated/inadequately described code N	Mandatory	1
9	Person with cancer—distant metastatic site(s) at diagnosis, code N[N]	Conditional	20
	Conditional obligation:		
	This data element is to be completed if <u>Person with cancer—distant</u> metastatic cancer indicator, yes/no/not stated/inadequately described code N indicates the presence of metastatic cancer.		

Seq No.	Metadata item	Obligation	Max occurs
10	Person with cancer—depth of myometrial invasion, total millimetres N[N]	Conditional	1
	Conditional obligation:		
	This data element is only to be recorded for patients with endometrial cancer, as indicated by <u>Person with cancer</u> <u>primary site of cancer</u> , topography code (ICD-O-3) ANN.N.		
11	Person with cancer—myometrial thickness, total millimetres N[N]	Conditional	1
	Conditional obligation:		
	This data element is only to be recorded for patients with endometrial cancer, as indicated by <u>Person with cancer</u> <u>primary site of cancer, topography code</u> (ICD-O-3) ANN.N.		
12	Person with cancer-depth of cervical cancer invasion, total millimetres N[N]	Conditional	1
	Conditional obligation:		
	This data element is only to be recorded for patients with cervical cancer, as indicated by <u>Person with cancer—primary site of cancer, topography code</u> (ICD-O-3) ANN.N.		
13	Person with cancer—tumour outside primary site indicator, yes/no/not stated/inadequately described code N	Mandatory	1
14	Person with cancer—tumour size outside primary site, code N	Mandatory	1
15	Person with cancer—multiple primary tumours indicator, yes/no code N	Mandatory	1
16	Person with cancer—multiple primary tumours descriptor, code N	Conditional	1
	Conditional obligation:		
	This data element is to be recorded if <u>Person with cancer—multiple primary</u> <u>tumours indicator, yes/no code N</u> indicates the presence of multiple primary tumours.		
17	Person—tissue sample collected indicator, yes/no code N	Mandatory	30
18	Organisation—organisation name, text X[X(199)]	Conditional	10
	Conditional obligation:		
	This data element is to be recorded when the data element <u>Person—tissue</u> <u>sample collected indicator, yes/no code N</u> indicates that a tissue sample has been collected.		
	DSS specific information:		
	Use this data element to record the name of the laboratory or biobank in which a tissue sample is stored. Collect this data element in conjunction with <u>Person—tissue sample collected indicator, yes/no code N</u> .		
19	Medical specialist—surgical specialty, initial gynaecological surgical speciality code N[N]	Conditional	10
	Conditional obligation:		
	This data element is only to be recorded for patients who have undergone		

surgery relating to their initial course of treatment for gynaecological cancer.

Seq No.	Metadata item	Obligation	Max occurs
20	Cancer treatment—post-initial surgery residual tumour size category, code N	Conditional	20
	Conditional obligation:		
	This data element is to be recorded then the data element <u>Cancer treatment</u> <u>—residual (R) tumour indicator, yes/no code N</u> indicates the presence of residual tumour after surgery.		
21	Cancer treatment—residual (R) tumour indicator, yes/no code N	Conditional	1
	Conditional obligation:		
	This data element is to be recorded for patients with ovarian cancer and stage IV endometrial cancer when surgical treatment for gynaecological cancer has been completed.		
22	Cancer treatment—primary surgical treatment complication indicator, yes/no/unknown code N	Mandatory	1
23	Cancer treatment—treatment complication type, cancer-related primary surgery complication type code N[N]	Conditional	10
	Conditional obligation:		
	This data element is to be recorded when <u>Cancer treatment—primary surgical</u> <u>treatment complication indicator, yes/no/unknown code N</u> indicates the presence of a treatment complication.		
24	Cancer treatment—gynaecological cancer post-radiotherapy complication indicator, yes/no/unknown code N	Conditional	1
	Conditional obligation:		
	This data element should be recorded in relation to the primary course of treatment for gynaecological cancer.		
	DSS specific information:		
	This relates to the primary course of treatment for gynaecological cancer.		
25	Cancer treatment—treatment complication type, gynaecological cancer-related radiotherapy code N	Conditional	10
	Conditional obligation:		
	This data element should be recorded when <u>Cancer treatment</u> <u>gynaecological cancer post-radiotherapy complication indicator</u> , <u>yes/no/unknown code N</u> indicates the presence of a radiotherapy related treatment complication.		
26	Cancer treatment—treatment complication type, text X[X(149)]	Conditional	10
	Conditional obligation:		
	This data element is to be recorded when either <u>Cancer treatment</u> <u>treatment</u> <u>complication type, gynaecological cancer-related radiotherapy code N</u> or <u>Cancer treatment</u> <u>treatment complication type, cancer-related primary</u> <u>surgery complication type code N[N]</u> indicates an 'Other' type of treatment complication.		

Seq No.	Metadata item	Obligation	Max occurs
27	Cancer treatment—systemic therapy treatment modification indicator, yes/no/unknown code N	Conditional	1
	Conditional obligation:		
	This data element is to be recorded for patients who have undergone systemic therapy as part of their cancer treatment. This includes chemotherapy, hormone therapy and immunotherapy.		
28	Cancer treatment—treatment modification type for cancer-related systemic therapy, code N[N]	Conditional	10
	Conditional obligation:		
	This data element is to be recorded when <u>Cancer treatment—systemic</u> <u>therapy treatment modification indicator, yes/no/unknown code N</u> indicates a modification to planned systemic therapy treatment.		
29	Cancer treatment—treatment plan modification, text X[X(149)]	Conditional	10
	Conditional obligation:		
	This data element is to be recorded when <u>Cancer treatment—treatment</u> <u>modification type for cancer-related systemic therapy, code N[N]</u> indicates an 'Other' type of treatment modification.		
30	Cancer treatment—primary course of chemotherapy delay indicator, yes/no/unknown code N	Conditional	5
	Conditional obligation:		
	This data element is to be recorded for patients who have undergone chemotherapy as part of their cancer treatment.		
31	Cancer treatment—primary course of chemotherapy delay reason, code N	Conditional	10
	Conditional obligation:		
	This data element is to be recorded when <u>Cancer treatment—chemotherapy</u> <u>delay indicator, yes/no/unknown code N</u> indicates a delay in planned chemotherapy treatment.		
32	Cancer treatment—cancer treatment type, code N[N]	Conditional	5
	Conditional obligation:		
	This data element is to be recorded for a patient having a first recurrence of cancer. All treatments administered to the patient during the first recurrence of cancer should be recorded.		
	DSS specific information:		
	This data element is to be recorded separately for the primary course of treatment and treatment for the first recurrence of cancer. All treatments administered to the patient as part of the primary course of treatment for the first recurrence of cancer should be recorded.		

Seq No.	Metadata item	Obligation	Max occurs
33	Cancer treatment—other cancer treatment, text X[X(149)]	Conditional	10
	Conditional obligation:		
	This data element is to be recorded for a patient having treatment that cannot be defined as surgery, radiotherapy or systemic therapy according to the definitions of those data items in this data set specification.		
	DSS specific information:		
	This data element is to be used to describe treatment, other than surgery, radiotherapy or systemic therapy, used to treat a first recurrence of gynaecological cancer.		
34	Cancer treatment—outcome of treatment, code N.N	Conditional	5
	Conditional obligation:		
	This data element is conditional on a patient completing treatment for their first recurrence of cancer.		
	DSS specific information:		
	This data element is to be recorded for patients who have completed their primary course of treatment or treatment for the first recurrence of cancer. For patients who have completed treatment for their first recurrence of cancer this should be recorded multiple times, once in relation to their primary course of treatment and once in relation to treatment for the first recurrence of cancer.		
35	Cancer (clinical) DSS	Mandatory	1
	DSS specific information:		
	The Cancer (clinical) data set specification is intended to only describe data collected in relation to the initial course of cancer treatment. The initial course of treatment includes all treatments administered to the patient from diagnosis and before disease progression or recurrence.		
-	Chemotherapy for cancer cluster	Conditional	1
	Conditional obligation:		
	Conditional on patient receiving chemotherapy.		
	- Cancer treatment—chemotherapy completion date, DDMMYYYY	Mandatory	1
	- <u>Cancer treatment—chemotherapy cycles administered, number of cycles</u> <u>N[NN]</u>	Mandatory	99
	- <u>Cancer treatment—chemotherapy start date, DDMMYYYY</u>	Mandatory	1
	- <u>Cancer treatment—systemic therapy agent or protocol, eviQ protocol</u> <u>identifier NNNNN</u>	Conditional	3
	Conditional obligation:		
	Conditional on the administration of systemic therapy agents according to a prespecified regimen or protocol, and on the availability of the protocol number on the eviQ website.		
	- Cancer treatment—systemic therapy agent or protocol, text X[X(149)]	Mandatory	99
-	Hormone therapy for cancer cluster	Conditional	1
	Conditional obligation:		

Conditional on patient receiving hormone therapy.

Seq No.	Metadata item	Obligation	Max occurs
	- Cancer treatment—hormone therapy completion date, DDMMYYYY	Mandatory	1
	- <u>Cancer treatment—hormone therapy start date, DDMMYYYY</u>	Mandatory	1
	- <u>Cancer treatment—systemic therapy agent or protocol, eviQ protocol</u> identifier NNNNN	Conditional	3
	Conditional obligation:		
	Conditional on the administration of systemic therapy agents according to a prespecified regimen or protocol, and on the availability of the protocol number on the eviQ website.		
	- <u>Cancer treatment—systemic therapy agent or protocol, text X[X(149)]</u>	Mandatory	99
-	Immunotherapy for cancer cluster	Conditional	1
	Conditional obligation:		
	Conditional on patient receiving immunotherapy.		
	- <u>Cancer treatment—immunotherapy completion date, DDMMYYYY</u>	Mandatory	1
	- <u>Cancer treatment—immunotherapy start date, DDMMYYYY</u>	Mandatory	1
	- <u>Cancer treatment—systemic therapy agent or protocol, eviQ protocol</u> identifier NNNNN	Conditional	3
	Conditional obligation:		
	Conditional on the administration of systemic therapy agents according to a prespecified regimen or protocol, and on the availability of the protocol number on the eviQ website.		
	- <u>Cancer treatment—systemic therapy agent or protocol, text X[X(149)]</u>	Conditional	99
-	Radiotherapy for cancer cluster	Conditional	1
	Conditional obligation:		
	Conditional on the patient receiving radiotherapy.		
	- Cancer treatment—radiation dose administered, total Gray N[NN.NN]	Mandatory	99
	- <u>Cancer treatment—radiotherapy completion date, DDMMYYYY</u>	Mandatory	99
	- Cancer treatment—radiotherapy fractions administered, total fractions N[N]	Mandatory	99
	- <u>Cancer treatment—radiotherapy start date, DDMMYYYY</u>	Mandatory	99
	- <u>Cancer treatment—radiotherapy target site, code N[N]</u>	Mandatory	99
	- <u>Cancer treatment—radiotherapy treatment type, code N[N]</u>	Mandatory	99
-	Surgery for cancer cluster	Conditional	1
	Conditional obligation:		
	Conditional on the patient receiving cancer-directed surgery.		
	- Cancer treatment—surgery target site, topography code (ICD-O-3) ANN.N	Mandatory	99
	- <u>Cancer treatment—surgical procedure date, DDMMYYYY</u>	Mandatory	99
	- <u>Cancer treatment—surgical procedure for cancer, procedure code (ACHI</u> 8th edn) NNNN-NN	Mandatory	99
-	Systemic therapy procedure for cancer cluster	Conditional	1
	Conditional abligation		

#### Conditional obligation:

Conditional on the patient receiving a systemic therapy procedure.

Seq No.	Metadata item	Obligation	Max occurs
-	Cancer treatment—systemic therapy procedure date, DDMMYYYY	Mandatory	99
-	Cancer treatment—systemic therapy procedure, code N[N]	Mandatory	99
-	Cancer staging—cancer staging scheme source edition number, code N[N]	Mandatory	1
-	Cancer staging—cancer staging scheme source, code N[N]	Mandatory	1
-	Cancer staging—staging basis of cancer, code A	Mandatory	1
-	Cancer treatment—cancer treatment type, code N[N]	Mandatory	1
-	Cancer treatment—other cancer treatment, text X[X(149)]	Conditional	99
	Conditional obligation:		
	Conditional on the patient having treatment that cannot be defined as surgery, radiotherapy or systemic therapy according to the definitions of those data items in this data set specification.		
-	Cancer treatment—outcome of treatment, code N.N	Mandatory	1
-	Date—accuracy indicator, code AAA	Mandatory	1
-	Establishment—organisation identifier (state/territory), NNNNN	Mandatory	1
-	Healthcare provider—organisation identifier, N(16)	Mandatory	1
-	Patient—cancer status, code N	Mandatory	1
-	Patient—date of last contact, DDMMYYYY	Mandatory	1
-	Patient-diagnosis date of cancer, DDMMYYYY	Mandatory	1
-	Patient—diagnosis date of first recurrence as distant metastasis, DDMMYYYY	Conditional	1
	Conditional obligation:		
	Conditional on the patient being diagnosed with recurrence involving a distant metastasis.		
-	Patient—diagnosis date of first recurrence as locoregional cancer, DDMMYYYY	Conditional	1
	Conditional obligation:		
	Conditional on the patient being diagnosed with recurrence of locoregional cancer.		
-	Person (address)—address line, text X[X(179)]	Mandatory	1
-	Person (name)—family name, text X[X(39)]	Mandatory	1
-	Person (name)—given name, text X[X(39)]	Mandatory	1
-	<u>Person with cancer—distant metastasis status, M stage (UICC TNM</u> Classification of Malignant Tumours, 7th ed) code X[XX]	Mandatory	1
-	Person with cancer—extent of primary cancer, stage grouping other, code X[XXXXX]	Mandatory	1
-	Person with cancer—extent of primary cancer, TNM stage (UICC TNM Classification of Malignant Tumours, 7th ed) code X[XX]	Mandatory	1
-	Person with cancer—histopathological grade, code N	Mandatory	1
-	Person with cancer—laterality of primary cancer, code A	Mandatory	1
-	Person with cancer—morphology of cancer, code (ICD-O-3) NNNN/N	Mandatory	1
-	Person with cancer—most valid basis of diagnosis of a cancer, code N	Mandatory	1

Seq No.	Metadata item	Obligation	Max occurs
-	Person with cancer—most valid basis of diagnosis of the first recurrence, code N	Conditional	1
	Conditional obligation:		
	Conditional on the return, reappearance or metastasis of cancer of the same histology after a disease-free intermission or remission.		
-	Person with cancer—number of positive regional lymph nodes, total N[N]	Conditional	1
	Conditional obligation:		
	Conditional on the regional lymph nodes being excised and examined by a pathologist and demonstrated to be positive for malignancy.		
-	Person with cancer—number of regional lymph nodes examined, total N[N]	Conditional	1
	Conditional obligation:		
	Conditional on the regional lymph nodes being excised and examined by a pathologist.		
-	Person with cancer—primary site of cancer, topography code (ICD-O-3) ANN.N	Mandatory	1
-	<u>Person with cancer—primary tumour status, T stage (UICC TNM Classification of Malignant Tumours, 7th ed) code X[XXX]</u>	Mandatory	1
-	Person with cancer—region of first recurrence as distant metastasis, topography code (ICD-O-3) ANN.N	Conditional	99
	Conditional obligation:		
	Conditional on the patient being diagnosed with recurrence of distant metastasis.		
-	Person with cancer—region of first recurrence as locoregional cancer, topography code (ICD-O-3) ANN.N	Conditional	99
	Conditional obligation:		
	Conditional on the patient being diagnosed with recurrence of locoregional cancer.		
-	Person with cancer—regional lymph node metastasis status, N stage (UICC TNM Classification of Malignant Tumours, 7th ed) code X[XX]	Mandatory	1
-	Person with cancer—solid tumour size (at diagnosis), total millimetres NNN	Conditional	1
	Conditional obligation:		
	Conditional on the histopathological examination of the tumour, and excludes Phyllodes tumours, sarcomas or lymphomas.		
-	Person-date of birth, DDMMYYYY	Mandatory	1
-	Person-date of death, DDMMYYYY	Conditional	1
	Conditional obligation:		
	Recorded when the patient has died.		
	DSS specific information:		
	This field must be greater than or equal to Date of diagnosis of primary cancer.		

### Seq Metadata item No.

## Obligation Max occurs

Mandatory 1

Mandatory 1

Mandatory 1

Mandatory 1

Conditional 1

- Person—government funding identifier, Medicare card number N(11)
- Person-Indigenous status, code N
- Person—person identifier, XXXXXX[X(14)]
- Person—sex, code N
- Person-underlying cause of death, code (ICD-10 2nd edn) ANN-ANN

Conditional obligation:

Recorded when the patient has died.