

|  |
| --- |
| Cancer and Screening Registers16517 |
| Outcome objective: Victorians are healthy and well  Output group: Public Health  Output: Health Protection |

**OFFICIAL**

## 1. Service objective

To record data about cancer and screening activity.

## 2. Description of the service

To maintain a register to record data about cancers and screening results for Victorians (as prescribed by legislation where applicable).

## 3. Client group

The client group is Victorians who have screened for cancer.

## 4. Obligations specific to this activity

In addition to the obligations listed in the Service Agreement, organisations funded to deliver this activity must comply with the following:

### 4a. Registration and Accreditation

N/A

### 4b. Program requirements and other policy guidelines

* [Cancer Control Registers Operating Guidelines](file:/F:/Oracle/Middleware/Oracle_Home/user_projects/domains/bi2)

<>

* [Improving Cancer Outcomes Act 2014 (VIC) – Instrument of Delegation](https://www.legislation.vic.gov.au/in-force/acts/improving-cancer-outcomes-act-2014/002)

<https://www.legislation.vic.gov.au/in-force/acts/improving-cancer-outcomes-act-2014/002>

* [Improving Cancer Outcomes Act 2014 (VIC)](https://www.legislation.vic.gov.au/in-force/acts/improving-cancer-outcomes-act-2014/002)

<https://www.legislation.vic.gov.au/in-force/acts/improving-cancer-outcomes-act-2014/002>

## 5. Performance

Funding is subject to achieving the performance targets specified in Schedule 2 of the Service Agreement.

Performance is measured as follows:

### Key performance measure 1: Statistical Report within an agreed timeline and publicly available

|  |  |
| --- | --- |
| **Aim/objective** | To maintain cancer and screening databases |
| **Target** | The performance measure target is provided in the Service Agreement |
| **Type of count** | Cumulative |
| **Counting rule** | Submit performance reports |
| **Data source(s) collection** | * Victorian Cervical Cytology Registry Data Collection * Cancer Council Victoria - Cancer Registry Data Collection |
| **Definition of terms** | The terms are defined in the Funding and Service Agreements |

## 6. Data collection

The reporting requirements for this service are:

| **Data collection name** | **Data system** | **Data set** | **Reporting cycle** |
| --- | --- | --- | --- |
| Cancer Council Victoria - Cancer Registry Data Collection | Manual Data Collection | Defined in Funding and Service Agreement | Quarterly |
| Victorian Cervical Cytology Registry Data Collection | Manual Data Collection | Defined in Funding and Service Agreement | Quarterly |

|  |
| --- |
| To receive this publication in an accessible format, email Monitoring Framework Helpdesk <monitoringframework.helpdesk@dffh.vic.gov.au>  Authorised and published by the Victorian Government, 1 Treasury Place, Melbourne. © State of Victoria, Department of Health, July 2019.  In this document, ‘Aboriginal’ refers to both Aboriginal and Torres Strait Islander people. ‘Indigenous’ or ‘Koori/Koorie' is retained when part of the title of a report, program or quotation.  ISSN 2653-0937 – Online (PDF/Word)  Available at Department of Families, Fairness and Housing and Department of Health activity search  [<http://providers.dffh.vic.gov.au/families-fairness-housing-health-activity-search](http://providers.dffh.vic.gov.au/families-fairness-housing-health-activity-search)> |