

Clostridium difficile Infection (CDI) Surveillance

CDI case definition

A CDI episode is defined as a laboratory confirmed *C. difficile* infection. Diagnosis is by a PCR-based toxin test on faeces from a symptomatic patient. Record only positive results from **diarrhoeal** stools, i.e. do not report positive results from screening specimens or asymptomatic patients.

Inclusions

- > Cases from all patients attending an acute care facility while symptomatic (including inpatients, HITH patients and patients attending Emergency departments, outpatient departments or haemodialysis units etc.).

Exclusions

- > Cases where a known previous positive specimen has been reported within the previous 8 weeks (an isolate obtained from a patient more than 8 weeks since the last positive test is regarded as a new episode)
- > Patients less than 2 years old at date of attendance/admission.

CDI exposure classification

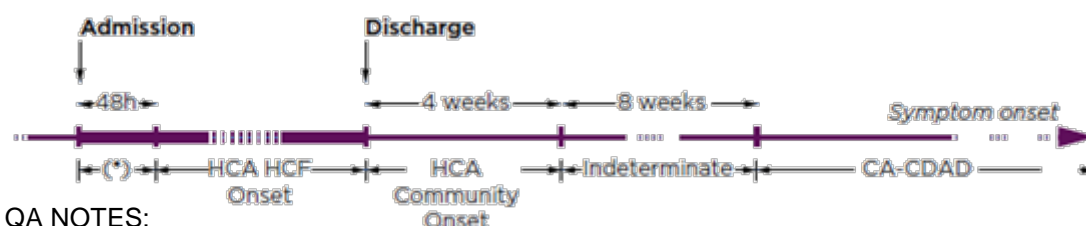
Currently the national data collection for CDI only includes those episodes that are first identified in a specimen collected in an acute care facility. However, the Australian Commission for Safety and Quality in Health Care has developed an Implementation Guide for CDI surveillance that describes sub-categorisation of episodes according to their most likely place of acquisition into 5 separate categories. Available from: <http://www.safetyandquality.gov.au/our-work/healthcare-associated-infection/national-hai-surveillance-initiative/>.

At the present time, South Australian contributors are requested to only apply the Category A (HCA-HCF) exposure classification (see Figure 1 below) as a sub-set of Hospital identified.

Category A: Healthcare associated – Healthcare facility onset (HCA-HCF)

Date of CDI symptom onset more than 48 hours after admission to a health care facility and prior to discharge from the facility. If date of symptom onset is not available, the date/time of specimen collection is used as a proxy.

Figure 1



QA NOTES:

- Outpatient and Emergency records – Attendance date should match specimen/symptom onset date*
- Category A records – ensure reported specimen/onset date is > 48hrs after admission.
- Specimen/onset date must be between admission and discharge dates

*Sources for identifying onset date may include (but are not limited to) medical notes, lab request date, specimen request form or the patient.

Data Element Table

Field Name	Description	Details
UR or Postcode	Unique record identification number	<ul style="list-style-type: none"> This is the patient's medical record number (MRN) or postcode for Private hospitals that do not supply MRN Mandatory field, cannot be null
Gender	Sex of the patient	<ul style="list-style-type: none"> Mandatory field, cannot be null
Date of Birth	The patients full year of birth, including day and month	<ul style="list-style-type: none"> If date of birth is not known or cannot be provided, provision of a generic estimate is acceptable (the first day of the appropriate month or 01/01/ of the appropriate year Format date as dd/mm/yyyy Mandatory field, cannot be null
Date of Admission / Attendance	The date the patient attended the Emergency or Outpatient Dept or date the patient was admitted	<ul style="list-style-type: none"> Format date as dd/mm/yyyy Mandatory field, cannot be null
Attendance Type	Identifies if the type of hospital attendance/admission	<ul style="list-style-type: none"> Permissible values: Inpatient, Outpatient or Emergency Mandatory field, cannot be null
Ward	The ward where patient was located at time specimen was collected.	<ul style="list-style-type: none"> This is NOT the acquisition ward , only the ward at the time the specimen was taken Field should not be Null if "Attendance Type" = Inpatient
Specimen/Onset Date[#]	Identifies the date of symptom onset or the date the specimen was taken	<ul style="list-style-type: none"> Format date as dd/mm/yyyy Must be within the reporting month Mandatory field, cannot be null
LAB Name	Identifies the laboratory organisation that processed the specimen	<ul style="list-style-type: none"> Mandatory field, cannot be null
Specimen Number	Positive specimen's unique identification number	<ul style="list-style-type: none"> Identifier allocated by the laboratory to the pathology result Mandatory field, cannot be null
Category	Identifies record as healthcare associated where applicable	<ul style="list-style-type: none"> Permissible values: Cat A or N/A Field should not be null
Comment	Record any relevant additional information	

NOTE:

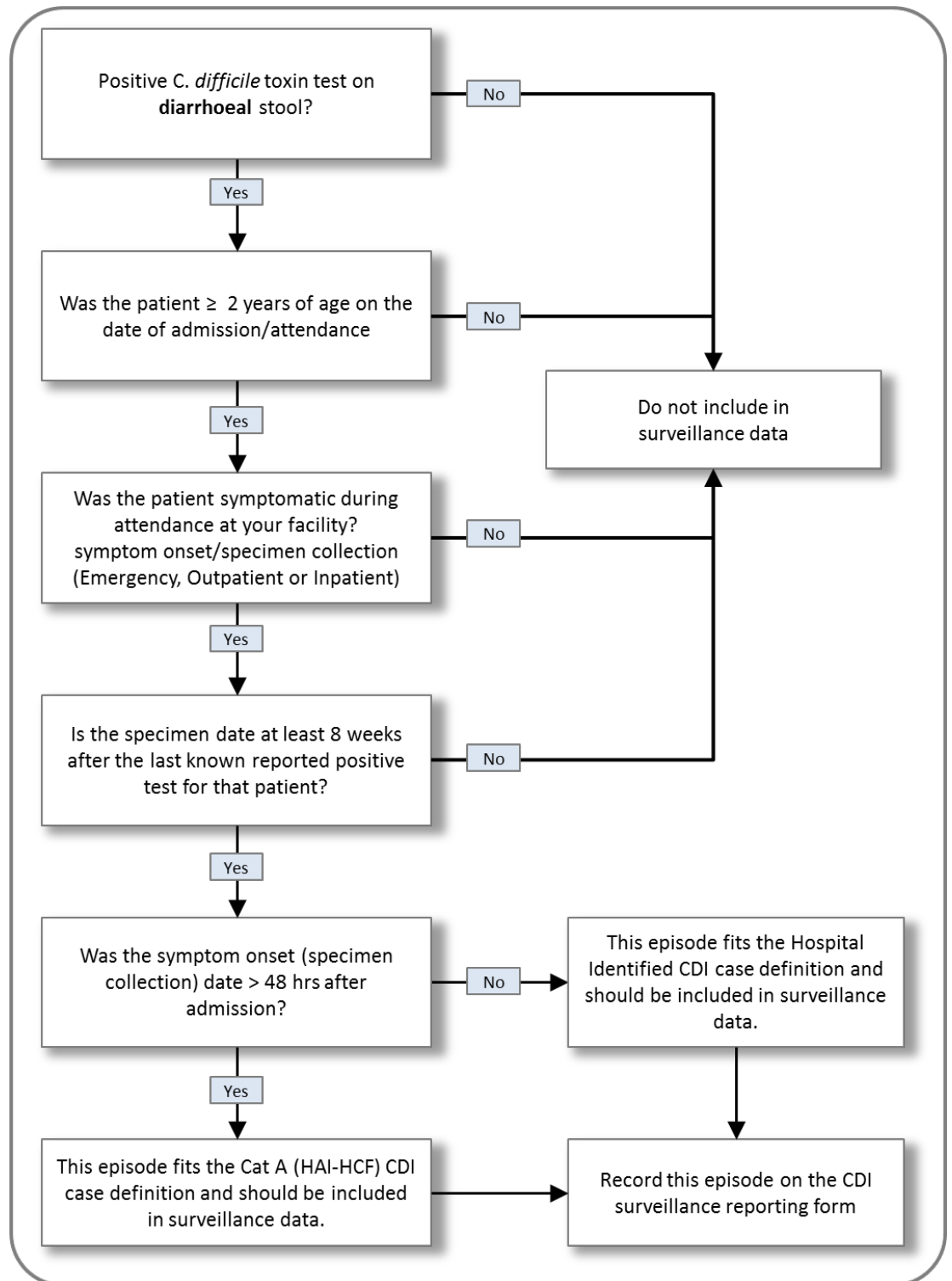
- [#]Identification/investigation of onset date is only required if the specimen collection date is >48hours after admission.
- Given state reporting currently only collects data on hospital identified and Category A (HCA-HCF) episodes as a sub-set, if a patient has a specimen taken during a current visit and it is identified as being associated with a previous admission, do not classify as a "post discharge" in the secondary acquisition, classify as NEW. Post-discharge information can be recorded as a comment in ICIMS.

Careconnect.sa infection control users (ICIMS)

To ensure record is included in the Category A field select:

- Primary Acquisition = HCA
- Secondary Acquisition = New.

Flow Chart



For more information

**Infection Control Service
Communicable Disease Control Branch
Telephone: 1300 232 272**

www.sahealth.sa.gov.au/infectionprevention

Public-I1-A2

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