National Healthcare Agreement: PI 22—Healthcare associated infections: Staphylococcus aureus bacteraemia, 2017



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National Healthcare Agreement: Pl 22—Healthcare associated infections: Staphylococcus aureus bacteraemia, 2017

Identifying and definitional attributes

Metadata item type: Indicator

Indicator type: Progress measure

Short name: PI 22–Healthcare associated infections: Staphylococcus aureus bacteraemia,

2017

METEOR identifier: 630047

Registration status: <u>Health!</u>, Superseded 30/01/2018

Description: Staphylococcus aureus bacteraemia (SAB) associated with acute care public

hospitals (excluding cases associated with private hospitals and non-hospital

care).

Indicator set: National Healthcare Agreement (2017)

Health!, Superseded 30/01/2018

Outcome area: Hospital and Related Care

National Health Performance Authority (retired), Retired 01/07/2016

Health!, Standard 07/07/2010

Data quality statement: National Healthcare Agreement: PI 22-Healthcare associated infections;

Staphylococcus aureus bacteraemia, 2017 QS

Health!, Standard 31/01/2017

Collection and usage attributes

Computation description:

Acute care public hospitals are defined as all public hospitals including those hospitals defined as public psychiatric hospitals in the Public hospital establishments National minimum data set (NMDS). All types of public hospitals are included, both those focusing on acute care, and those focusing on non-acute or subacute care, including psychiatric, rehabilitation and palliative care.

Unqualified newborns, hospital boarders and posthumous organ procurement are excluded from the indicator.

A patient-episode of SAB is defined as a positive blood culture for *Staphylococcus aureus*. For surveillance purposes, only the first isolate per patient is counted, unless at least 14 days has passed without a positive blood culture, after which an additional episode is recorded.

A *Staphylococcus aureus* bacteraemia will be considered to be healthcare-associated if: the first positive blood culture is collected more than 48 hours after hospital admission or less than 48 hours after discharge, OR, if the first positive blood culture is collected 48 hours or less after admission and one or more of the following key clinical criteria was met for the patient-episode of SAB:

- 1. SAB is a complication of the presence of an indwelling medical device (e.g. intravascular line, haemodialysis vascular access, cerebrospinal fluid (CSF) shunt, urinary catheter)
- 2. SAB occurs within 30 days of a surgical procedure where the SAB is related to the surgical site
- 3. An invasive instrumentation or incision related to the SAB was performed within 48 hours
- 4. SAB is associated with neutropenia contributed to by cytotoxic therapy. Neutropenia is defined as at least two separate calendar days with values of absolute neutrophil count (ANC) <500 cells/mm³ (0.5 × 10⁹ / L) on or within a seven-day time period which includes the date the positive blood specimen was collected (Day 1), the 3 calendar days before and the 3 calendar days after.

Exclusions:

Cases where a known previous positive test has been obtained within the last 14 days are excluded. For example: If a patient has SAB in which 4 sets of blood cultures are positive over the initial 3 days of the patient's admission only one episode of SAB is recorded. If the same patient had a further set of positive blood cultures on day 6 of the same admission, these would not be counted again, but would be considered part of the initial patient-episode.

Note: If the same patient had a further positive blood culture 20 days after admission (i.e. greater than 14 days after their last positive blood culture on day 5), then this would be considered a second patient-episode of SAB.

See <u>Establishment—number of patient days, total N[N(7)]</u> for the definition of patient days.

Analysis by state and territory is based on location of the hospital.

Presented as:

- a number, and
- per 10,000 patient days.

Coverage: Denominator ÷ Number of patient days for all public hospitals in the state or territory.

Any variation from the specifications by jurisdictions will be footnoted and described in the data quality statement.

Computation: Numerator

10,000 x (Numerator + Denominator).

Numerator: Number of SAB patient episodes (as defined above) associated with acute care public hospitals.

Numerator data elements:

-Data Element / Data Set-

Person—Staphylococcus aureus bacteraemia episode indicator

Data Source

State/territory infection surveillance data

Guide for use

Data source type: Administrative by-product data

Data Element / Data Set-

Person—person identifier

Data Source

State/territory infection surveillance data

Guide for use

Data source type: Administrative by-product data

Denominator:

Number of patient days for public acute care hospitals under surveillance (i.e. only for hospitals included in the surveillance arrangements).

Exclude unqualified newborns, posthumous organ procurement and hospital boarders.

Denominator data elements:

Data Element / Data Set-

Episode of admitted patient care—admission date

Data Source

State/territory admitted patient data

Guide for use

Data source type: Administrative by-product data

Data Element / Data Set-

Episode of admitted patient care—separation date

Data Source

State/territory admitted patient data

Guide for use

Data source type: Administrative by-product data

Data Element / Data Set-

Establishment—Staphylococcus aureus bacteraemia surveillance indicator

Data Source

State/territory admitted patient data

Guide for use

Data source type: Administrative by-product data

Data Element / Data Set-

Establishment—organisation identifier (Australian)

Data Source

State/territory admitted patient data

Guide for use

Data source type: Administrative by-product data

Disaggregation:

2010–11, 2011–12, 2012–13, 2013–14 (updated for amended denominator), 2014–15 (updated for resupplied data and amended denominator), 2015–16—State and territory, by:

 Methicillin-resistant Staphylococcus aureus (MRSA)/Methicillin-sensitive Staphylococcus aureus (MSSA)

Some disaggregation may result in numbers too small for publication.

Disaggregation data elements:

Data Element / Data Set

Establishment—Australian state/territory identifier

Data Source

State/territory infection surveillance data

Guide for use

Data source type: Administrative by-product data

Data Element / Data Set-

Methicillin-resistant *Staphylococcus aureus* MRSA)/Methicillin-sensitive *Staphylococcus aureus* (MSSA) indicator

Data Source

State/territory infection surveillance data

Guide for use

Data source type: Administrative by-product data

Comments:

Most recent data available for 2017 National Healthcare Agreement performance reporting: 2015–16.

The number of SAB patient episodes associated with acute public hospitals under surveillance includes SAB patient episodes associated with all public hospitals, and the number of patient days for public acute care hospitals under surveillance includes the number of patient days for all public hospitals under surveillance.

For some states and territories there is less than 100 per cent coverage of hospitals. This may impact on the reported rate. For those jurisdictions with incomplete coverage of acute care public hospitals (in the numerator), only patient days for those hospitals that contribute data are included (in the denominator). Specifically, if a hospital was not included in the SAB surveillance arrangements for part of the year, then the patient days for that part of the year are excluded. If part of the hospital was not included in the SAB surveillance arrangements (e.g. children's wards, psychiatric wards), then patient days for that part of the hospital are excluded. Patient days for 'non-acute' hospitals (such as rehabilitation and psychiatric hospitals) are included if the hospital was included in the SAB surveillance arrangements, but not otherwise. However, all these patient days are included in the coverage rate denominator measure of total number of patient days for all public hospitals in a state or territory.

Some states operate a 'signal surveillance' arrangement for smaller hospitals whereby the hospital notifies the appropriate authority if a SAB case is identified, but the hospital is not considered to have formal SAB surveillance as per larger hospitals. Where this arrangement is in place, these hospitals should be included as part of the indicator. That is, SAB patient episodes and patient days should be included as 'under surveillance'.

Only episodes associated with acute public hospital care in each jurisdiction should be counted. If a case is associated with care provided in another jurisdiction (cross border flows) then it is reported (where known) by the jurisdiction where the care associated with the SAB occurred.

There may be patient episodes of SAB identified by a hospital which did not originate in the identifying hospital (as determined by the definition of a patient episode of SAB), but in another public hospital. If the originating hospital is under SAB surveillance, then the patient episode of SAB should be attributed to the originating hospital and should be included as part of the indicator. If the originating hospital is not under SAB surveillance, then the patient episode is unable to be included in the indicator.

Patient episodes associated with care provided by private hospitals and non-hospital health care are excluded.

Patient days for unqualified newborns, hospital boarders and posthumous organ procurement are excluded.

Almost all patient episodes of SAB will be diagnosed when the patient is an admitted patient. However, the intention is that cases are reported whether they were associated with admitted patient care or non-admitted patient care in public acute care hospitals.

Where there is significant variation, for example non-coverage of cases diagnosed less than 48 hours after admission, in the data collection arrangements it will affect the calculation of values across states and territories.

Variation in admission practices across jurisdictions will influence the denominator for this indicator impacting on comparability of rates.

Jurisdictional manuals should be referred to for full details of definitions used in infection control surveillance.

Note that the definition of a healthcare-associated SAB was revised by the Australian Commission on Safety and Quality in Health Care in 2016. In particular, the clinical criterion for SAB associated with neutropenia was revised. Data for 2010–11, 2011–12, 2012–13, 2013–14 and 2014–15 are provided according to the previous neutropenia criterion:

 SAB is associated with neutropenia (<1 × 10⁹) contributed to by cytotoxic therapy

Data for 2015–16 are provided according to the new neutropenia criterion:

 SAB is associated with neutropenia contributed to by cytotoxic therapy. Neutropenia is defined as at least two separate calendar days with values of absolute neutrophil count (ANC) <500 cell/mm³ (0.5 × 10⁹/L) on or within a seven-day time period which includes the date the positive blood specimen was collected (Day 1), the 3 calendar days before and the 3 calendar days after.

Note that patient episodes of SAB are just one type of healthcare associated infection. Hence, this performance indicator is not a complete measure of healthcare associated infections for the outcome area of Hospital and Related Care.

Representational attributes

Representation class: Rate

Data type: Real

Unit of measure: Episode

Format: NN[N]

Indicator conceptual framework

Framework and dimensions:

Safety

Data source attributes

Data sources:

Data Source

State/territory admitted patient data

Frequency

Annual

Data custodian

State/territory health authorities

Data Source

State/territory infection surveillance data

Frequency

Annual

Data custodian

State/territory health authorities

Accountability attributes

Reporting requirements: National Healthcare Agreement

Organisation responsible for providing data:

Australian Institute of Health and Welfare

Benchmark: National Healthcare Agreement: PB g-Better health services: the rate of

> Staphylococcus aureus (including MRSA) bacteraemia is no more than 2.0 per 10,000 occupied bed days for acute care public hospitals by 2011–12 in each

state and territory, 2017

collection required:

Further data development / Specification: Substantial work required, the measure requires significant work to

be undertaken.

Relational attributes

Related metadata references:

Supersedes National Healthcare Agreement: PI 22-Healthcare associated infections: Staphylococcus aureus bacteraemia, 2016

Health!, Superseded 04/08/2016

Has been superseded by National Healthcare Agreement: PI 22-Healthcare associated infections: Staphylococcus aureus bacteraemia, 2018

Health!, Superseded 19/06/2019

See also National Healthcare Agreement: PB g-Better health services: the rate of Staphylococcus aureus (including MRSA) bacteraemia is no more than 2.0 per 10,000 occupied bed days for acute care public hospitals by 2011–12 in each state and territory, 2017

Health!, Superseded 30/01/2018

See also National Healthcare Agreement: PI 23-Unplanned hospital readmission rates, 2017

Health!, Superseded 30/01/2018