

Toxigenic *Corynebacterium diphtheriae* Disease in South Africa, week 26 2026

Report date: 2 July 2026

Reporting period: 29 December 2025 to 28 June 2026

Date of data extraction: 2 July 2026

Data are provisional as of the date data were extracted. Case counts are reported and analysed by date of first specimen collection, or by date of first clinical presentation where the specimen collection date is unavailable. Data cleaning and validation are ongoing; therefore, the figures presented in this report are provisional and may be subject to changes in subsequent reports.

Highlights

- Since the last situational report (week 25, 2026), the following updates are included in this report
 - One new laboratory-confirmed case of toxigenic diphtheria from the Western Cape.
 - No new asymptomatic carriers of toxigenic *C. diphtheriae*.
- Appropriate public health responses were initiated for all suspected and confirmed cases.

Table 1: Number of suspected, probable and confirmed cases of toxigenic respiratory and cutaneous diphtheria in South Africa, 29 December 2025 to 28 June 2026.

Case definition	Number	Provincial distribution
Laboratory-confirmed toxigenic respiratory diphtheria	33	Gauteng (1/33, 3%) Limpopo (2/33, 6%) Western Cape (30/33, 91%)
Probable diphtheria cases	5	Limpopo (5/5, 100%)
Laboratory-confirmed toxigenic cutaneous diphtheria	1	Gauteng (1/1, 100%)
Suspected diphtheria cases with specimens sent to exclude diphtheria and tested negative	72	Free State (3/72, 4%) Gauteng (2/72, 3%) KwaZulu-Natal (5/72, 7%) Mpumalanga (2/72, 3%) Western Cape (60/72, 83%)
Asymptomatic carriers of toxigenic <i>C. diphtheriae</i> identified during contact tracing	7	Western Cape (7/7, 100%)
Deaths in probable and laboratory-confirmed toxigenic respiratory diphtheria cases	8	Gauteng (1/8, 12%) Limpopo (2/8, 25%) Western Cape (5/8, 62%)

Epidemiology of respiratory diphtheria cases and asymptomatic carriers, 29 December 2025 – 28 June 2026

Between 29 December 2025 and 28 June 2026, 33 confirmed cases of respiratory diphtheria and seven asymptomatic carriers of toxigenic *C. diphtheriae* have been identified in South Africa. The majority of confirmed cases (91%; 30/33) and all carriers were from the Western Cape, while two of the remaining confirmed cases (6%, 2/33) were from Limpopo and one (3%, 1/33) was from Gauteng (Figure 1, Table 1). Five probable cases were reported from Limpopo. The median age for cases of confirmed respiratory diphtheria was 23 years (range: 5-43 years), with 70% (22/33) being 18 years and older. The overall case-fatality ratio (CFR) among probable and confirmed respiratory diphtheria cases was 21% (8/38). The highest CFR was observed among individuals aged 30-39 years (57%; 4/7), although this estimate should be interpreted with caution given the very small number of cases in this age group (Table 2).

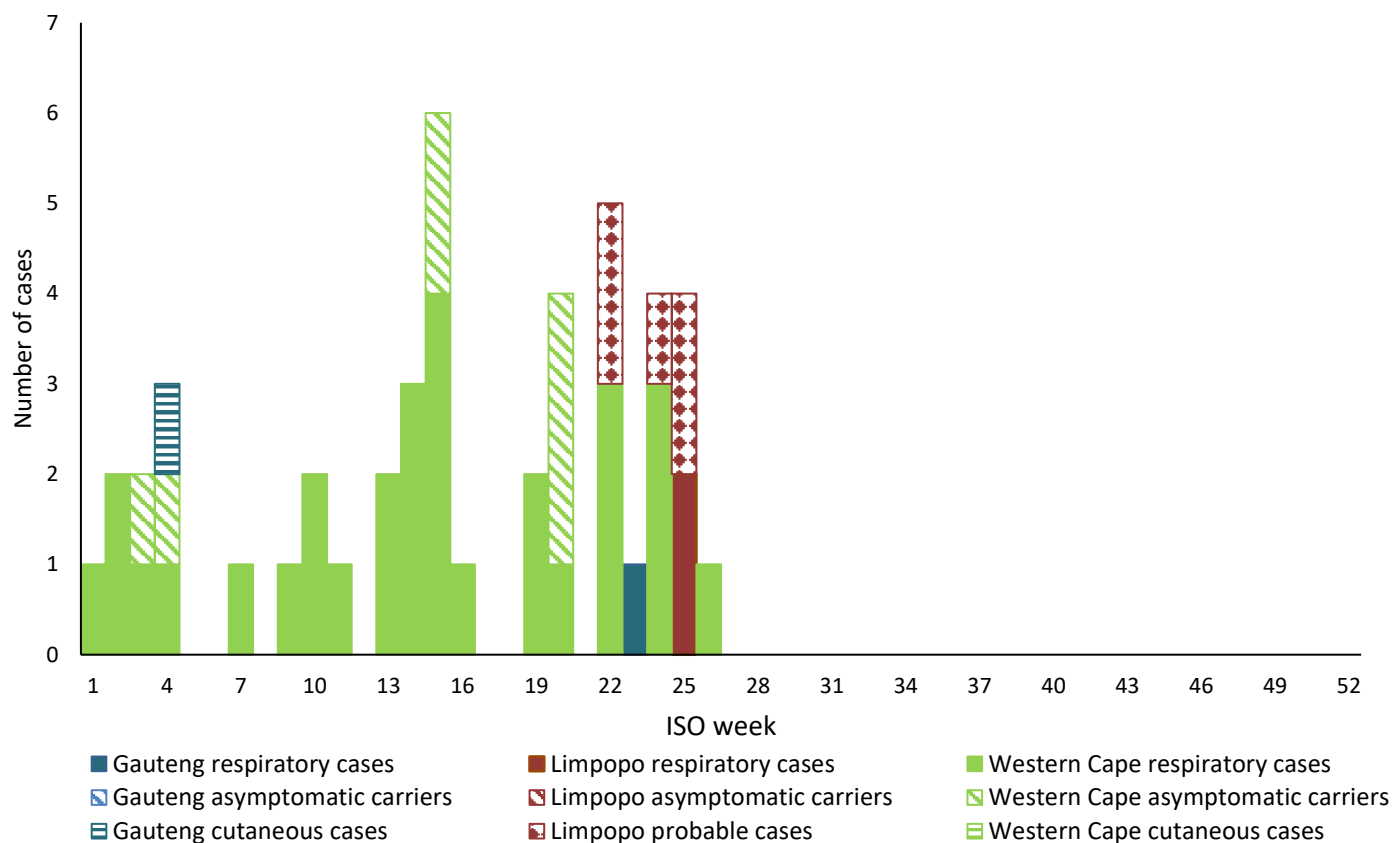


Figure 1: Number of individuals testing positive for toxigenic *C. diphtheriae* (respiratory diphtheria cases, asymptomatic carriers, and cutaneous cases) and probable cases, South Africa, 29 December 2025 – 28 June 2026.

***Note:** Case counts are reported and analysed by date of first specimen collection, or by date of symptom onset where the specimen collection date is unavailable. Data cleaning and validation are ongoing; therefore, the figures presented in this report are provisional and may be subject to changes in subsequent reports. ISO: International Organization for Standardization.

Table 2: Age distribution of laboratory-confirmed and probable respiratory diphtheria cases, diphtheria-related deaths, and case-fatality ratio, 29 December 2025 to 28 June 2026.

Age category (years)	Confirmed cases	Probable cases	Diphtheria-related deaths	Case fatality ratio (%)
0-9	5	1	1	17
10-19	7	3	3	30
20-29	12	1	0	0
30-39	7	0	4	57
40-49	2	0	0	0
50-59	0	0	0	0
≥60	0	0	0	0
Total	33	5	8	21

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Clusters and sporadic respiratory diphtheria cases by province

Between 29 December 2025 and 28 June 2026, **six diphtheria clusters** were identified, comprising 22 individuals. These included 10 confirmed respiratory cases (8 in the Western Cape and 2 in Limpopo), 7 asymptomatic carriers (all in the Western Cape), and 5 probable cases (all from Limpopo). Outside these clusters, 23 **sporadic respiratory cases** with no known epidemiologic links were reported, 22 in the **Western Cape** and 1 in **Gauteng** (Table 3).

Table 3. Summary of diphtheria clusters reported in South Africa from 29 December 2025 to 28 June 2026

Cluster No.	Week (week start date) ¹	Province (City)	Details	Cluster type
1	2 (5 January 2026)	Western Cape (Cape Town)	2 respiratory cases 1 asymptomatic carrier	Household/family
2	4 (19 January 2026)	Western Cape (Cape Town)	1 respiratory case 1 asymptomatic carrier	Household/family
3	15 (6 April 2026)	Western Cape (Cape Town)	2 respiratory cases 1 asymptomatic carrier	Household/family
4	15 (6 April 2026)	Western Cape (Cape Town)	2 respiratory cases 1 asymptomatic carrier	Household/family
5	19 – 20 (4 - 11 May 2026)	Western Cape (Cape Town)	1 respiratory case 3 asymptomatic carriers	Correctional facility
6	22-25 (29 May- 21 June)	Limpopo (Tzaneen)	2 respiratory cases 5 probable cases	Household/family

¹Period between the first recorded infection and the last observed infection within each cluster, expressed in weeks. Each week starts on a Monday and is labelled by its start date, following the ISO 8601 standard. Based on the date of clinical presentation or sample collection.

Notified suspected cases of diphtheria

From 29 December 2025 to 28 June 2026, there were 72 reports of suspected diphtheria cases who tested negative for *C. diphtheriae*. Alternative diagnoses became available for some individuals with suspected diphtheria, including *Streptococcus pyogenes*, *Streptococcus dysgalactiae* and respiratory viruses such as influenza.

Cutaneous toxigenic diphtheria cases

One cutaneous toxigenic diphtheria case was reported from Gauteng.

Non-toxigenic diphtheria

From 29 December 2025 to 28 June 2026, 26 individuals with non-toxigenic *C. diphtheriae* (14 asymptomatic, 11 cutaneous, one bacteraemia) have been detected (Table 4).

Table 4: Number of non-toxigenic *Corynebacterium* spp. infections by species and clinical presentation, South Africa, 29 December 2025 to 28 June 2026.

Clinical presentation	<i>C. diphtheriae</i>	<i>C. belfantii</i>	<i>C. ulcerans</i>	Total
Asymptomatic*	14	0	0	14
Cutaneous	11	0	0	11
Respiratory	0	0	0	0
Other**	1	0	0	1
Total	26	0	0	26

*Asymptomatic carriers identified as part of contact tracing before confirmation of non-toxigenic *C. diphtheriae*

** Bacteraemia

Microbiology

Of the 41 toxigenic *C. diphtheriae* infections identified, 31 were culture-confirmed; of these, four isolates lost viability, leaving 27 isolates for further phenotypic and genotypic characterisation.

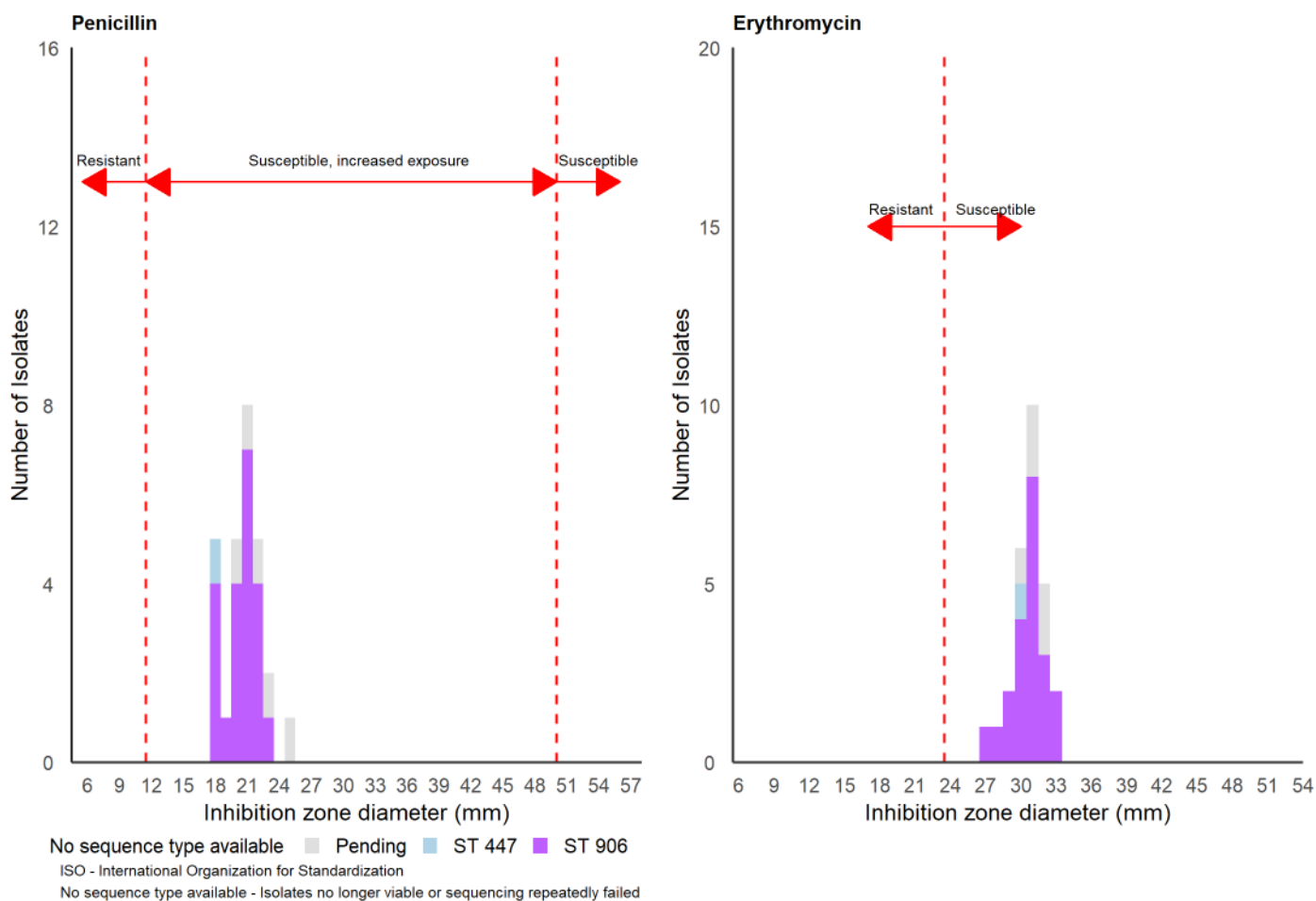
The phenotypic Elek test to confirm toxin production showed 100% agreement with the toxin-gene PCR among isolates tested by both methods to date (n = 27). No non-toxigenic toxin-gene-bearing (NTTB) isolates have been identified; such isolates would be Elek-negative but PCR-positive for the toxin gene.

Antimicrobial susceptibility testing was performed on 27 isolates using the disc diffusion method in accordance with European Committee on Antimicrobial Susceptibility Testing (EUCAST) guidelines. All tested isolates were susceptible to penicillin, with increased exposure (zone diameter [ZD] range 18–23 mm) and susceptible to erythromycin (ZD range 27–33 mm) (Figure 2) (1,2).

To date, sequence data are available for 22 of 27 viable toxigenic *C. diphtheriae* isolates. The remaining 5 isolates have pending sequence results. All Western Cape Province isolates are sequence type (ST) 906 (n = 21), the same lineage first detected in 2023 and currently appearing to be localised within this province. One cutaneous diphtheria isolate from Gauteng is ST447.

Table 5: Toxigenic *C. diphtheriae* infections by sequence type and province, South Africa, 29 December 2025 to 28 June 2026.

Province	ST 906	ST 447	Pending	Total
Gauteng	0	1	0	1
Limpopo	0	0	1	1
Western Cape	21	0	4	25
Total	21	1	5	27



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Figure 2: Antimicrobial susceptibility testing for toxigenic *C. diphtheriae* isolates from respiratory diphtheria cases, asymptomatic carriers, and cutaneous cases, South Africa, 2026 (N = 27). The red vertical dashed lines indicate the clinical breakpoints defined by the European Committee on Antimicrobial Susceptibility Testing (EUCAST). Arrows represent the inhibition zone diameter ranges at which *C. diphtheriae* is classified as susceptible, “susceptible at increased exposure”, or resistant to each antibiotic.

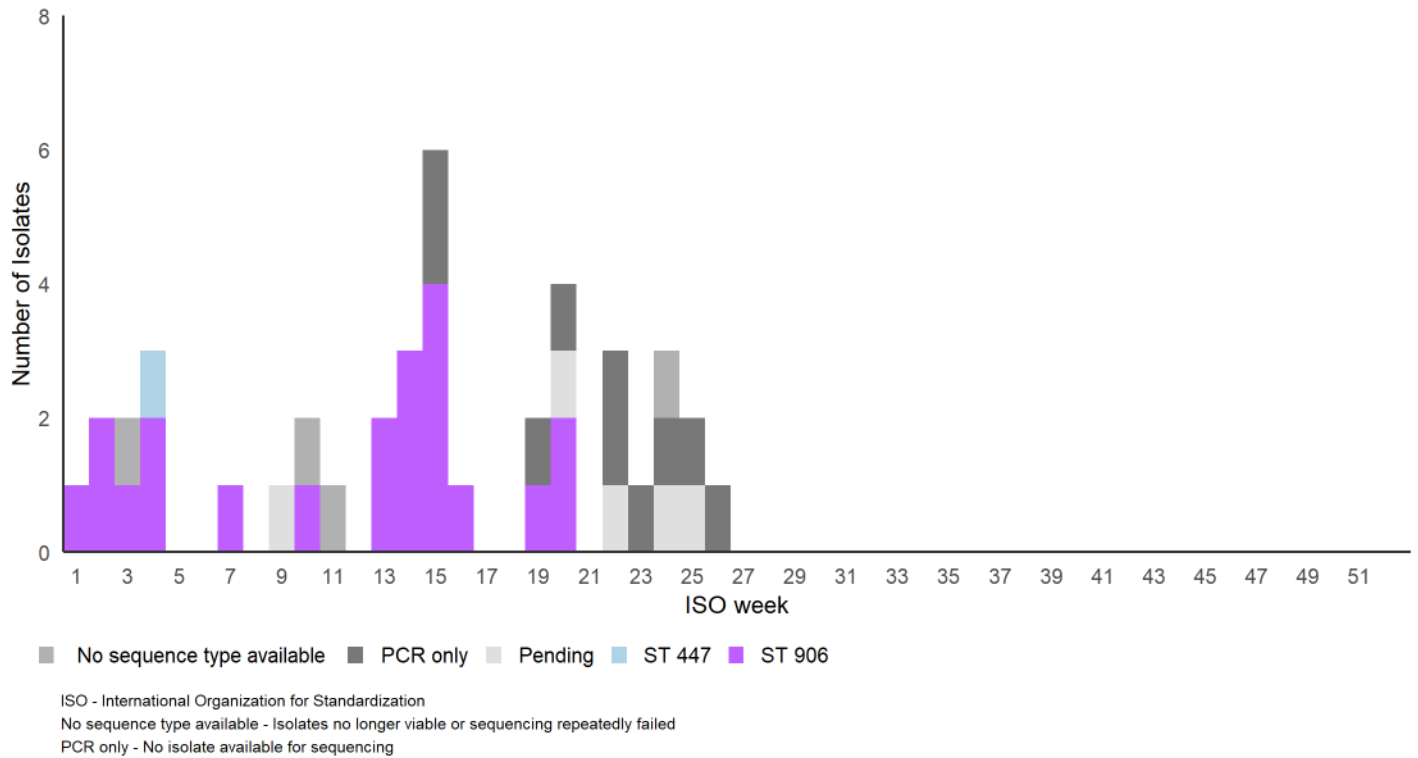


Figure 3: Number of individuals testing positive for toxigenic *C. diphtheriae* (respiratory diphtheria cases and asymptomatic carriers and cutaneous cases) (N=41) by sequence type, South Africa, 29 December 2025 to 28 June 2026.

Additional resources

Further information on diphtheria case definitions, laboratory testing, clinical management and outbreak response can be found on the National Institute for Communicable Diseases website: <https://www.nicd.ac.za/diseases-a-z-index/diphtheria/>

References

1. EUCAST. Antimicrobial susceptibility testing: EUCAST disk diffusion method. Version 11.0. 2023. http://www.eucast.org/ast_of_bacteria/disk_diffusion_methodology/
2. Berger A, Badell E, Åhman J, Matuschek E, Zidane N, Kahlmeter G, Sing A, Brisse S. *Corynebacterium diphtheriae* and *Corynebacterium ulcerans*: development of EUCAST methods and generation of data on which to determine breakpoints. *J Antimicrob Chemother.* 2024 May 2;79(5):968-976. doi: 10.1093/jac/dkae056. PMID: 38497937.
3. Williams MM, Waller JL, Aneke JS, Weigand MR, Diaz MH, Bowden KE, Simon AK, Peng Y, Xiaoli L, Cassidy PK, Winchell J, Tondella ML. Detection and Characterization of Diphtheria Toxin Gene-Bearing *Corynebacterium* Species through a New Real-Time PCR Assay. *J Clin Microbiol.* 2020 Sep 22;58(10):e00639-20. doi: 10.1128/JCM.00639-20. PMID: 32727830; PMCID: PMC7512153.

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4. Engler KH, Glushkevich T, Mazurova IK, George RC, Efstratiou A. A modified Elek test for detection of toxigenic corynebacteria in the diagnostic laboratory. *J Clin Microbiol.* 1997 Feb;35(2):495-8. doi: 10.1128/jcm.35.2.495-498.1997. PMID: 9003626; PMCID: PMC229610.