

# TIXAGEVIMAB AND CILGAVIMAB

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|-------------------------|---|
| BRAND NAME              | EVUSHELD  |
| DRUG CLASS              | Immunoglobulin (IgG1) monoclonal antibody (human), antiviral (SARS-CoV-2)   |
| AVAILABILITY            | There are <b>two vials</b> - use both vials to prepare the dose.<br>one vial contains 150 mg/1.5 mL of tixagevimab (100 mg/mL)<br>one vial contains 150 mg/1.5 mL of cilgavimab (100 mg/mL). <sup>1</sup><br>Also contain histidine, histidine hydrochloride monohydrate, sucrose and polysorbate-80. <sup>1</sup><br>The solutions are clear to opalescent and colourless to slightly yellow. <sup>1</sup>   |
| WARNING                 | <div style="border: 2px solid red; padding: 5px;"><p>The occupational hazard of intermittent low dose exposure to tixagevimab and cilgavimab and is not known. Wear a mask and gloves when preparing the dose to minimise exposure.</p><p>Hypersensitivity reactions including anaphylaxis may occur. Resuscitation facilities must be readily available.<sup>1</sup></p></div>   |
| pH                      | 6 <sup>1</sup>  |
| PREPARATION             | To prepare a 300 mg dose (150 mg of tixagevimab and 150 mg of cilgavimab) for pre-exposure prophylaxis: draw up 150 mg/1.5 mL of tixagevimab and 150 mg/1.5 mL of cilgavimab into separate syringes suitable for IM use. <sup>1,2</sup>   |
| STABILITY               | Vials: store at 2 to 8 °C. Do not freeze. Protect from light. <b>Do not shake.</b> <sup>1</sup><br>The solutions are stable in a syringe for 4 hours at 25 °C or at 2 to 8 °C. <sup>1</sup>   |
| ADMINISTRATION          |   |
| <b>IM injection</b>     | Inject the tixagevimab dose into the gluteal muscle, then inject the cilgavimab dose into the opposite gluteal muscle. <sup>1</sup><br>The ventrogluteal site is preferred.   |
| <b>SUBCUT injection</b> | Not recommended   |
| <b>IV injection</b>     | Not recommended   |
| <b>IV infusion</b>      | Not recommended   |
| COMPATIBILITY           | Do not mix with other medicines   |
| INCOMPATIBILITY         | No information  |
| SPECIAL NOTES           | Hypersensitivity reactions including rash and urticaria may occur. <sup>1</sup><br>Check your local guidelines for monitoring requirements. Clinical trial participants were monitored for adverse effects for at least 1 hour after the injections. <sup>3</sup><br>Anaphylactic reactions are rare but are a medical emergency. Commence treatment immediately. <sup>1</sup><br>Pain, redness and itching at the injection sites are common. <sup>1</sup> |

## REFERENCES

1. Product information. Available from [www.tga.gov.au](http://www.tga.gov.au). Accessed 07/04/2022.
2. National COVID-19 Clinical Evidence Taskforce. Australian guidelines for the clinical care of people with COVID-19: 6.1.9.1 Tixagevimab plus cilgavimab (Evusheld). Available from [www.covid19evidence.net.au](http://www.covid19evidence.net.au). Accessed 08/04/2022.
3. Levin MJ, Ustianowski A, De Wit S, Launay O, Avila A, Templeton A, et al. Intramuscular AZD7442 (Tixagevimab- Cilgavimab) for prevention of COVID-19. *N Engl J Med* 2022; Apr 20. DOI: 10.1056/NEJMoa2116620.