



**Australian Government**  
**Department of Health**  
 Therapeutic Goods Administration

**TGA use only**

This form, when completed, will be classified as 'For official use only'.

For guidance on how your information will be treated by the TGA see: Treatment of information provided to the TGA at <<https://www.tga.gov.au/treatment-information-provided-tga>>.

## Special Access Scheme – Category B

### Important information

Email completed form to [SAS@health.gov.au](mailto:SAS@health.gov.au) (preferred) or fax to 02 6232 8112.

The SAS Category B application form should be completed if guidance for use of an unapproved good will be met and the SAS Category A or SAS Category C pathways are not applicable.

### Privacy information

For general privacy information, go to <<https://www.tga.gov.au/privacy>>.

The TGA is collecting personal information in this form in order to:

- Assess the application.
- Contact the medical practitioner and discuss the application where necessary.
- The personal information of the medical practitioner may be disclosed to State and Territory authorities with responsibility for therapeutic goods or medical practitioner registration.

**Do not provide the name of the patient. Only provide the patient's initials and other information as requested on this form.**

**Please complete the form clearly and in full. Applications cannot be assessed if the form is incomplete or illegible. PLEASE PRINT IN BLOCK LETTERS.**

### Patient details (minimum of 3 (three) identifiers required)

Diagnosis(es) or Medical Condition(s): NON KETOTIC HYPERGLYCEMIA

Indication: HIGH GLYCINE

Clinical justification for use of product: (e.g. Include seriousness of condition, details of previous treatment including reasons why a therapeutic good currently listed on the ARTG cannot be used for the treatment of this patient in this circumstance)

LIFE THREATENING DISEASE - CHILD CANNOT SURVIVE WITHOUT THESE MEDICATIONS

### Product details (attach efficacy and safety data to support proposed use of the product and details of intended monitoring)

Therapeutic good type    Medicine     Biological     Medical device

Medicine/biological		Medical device	
Trade Name (if known) Amzoate	Sponsor / Supplier Medsurge Healthcare	Trade name	
Active ingredient(s) Sodium Benzoate		Product description (including variant <sup>2</sup> )	
Dosage form (e.g. tablet) Tablet	Strength (e.g., 1 mg/ml) 500mg	No of units	Sponsor / Supplier
Route of administration (e.g., IV) Oral	Dose & frequency (1 tds) 1000 to 1500mg bd	Proposed duration of treatment	Intended date of use
Quantity <sup>1</sup> required for treatment or duration 12 months			

Treating health practitioner details

Submitter details (if different)

Please send this form to the TGA only

<sup>1</sup> The health practitioner type for the treating health practitioner details above can be any of the following: Medical practitioner; ATSI health practitioner; dentist; radiographer; nurse; midwife; occupational therapist; optometrist; pharmacist; podiatrist; psychologist. Other health practitioner types can be included as the submitter.

<sup>1</sup> For substances captured by the Customs (Prohibited Imports) Regulations 1956 the quantity must be provided

<sup>2</sup> Variant means a medical device the design of which has been varied to accommodate different patient anatomical requirements (for example, relating to the shape, size, length, diameter or gauge of the device)