

## NEW ZEALAND DATA SHEET

### 1. PRODUCT NAME

Hyalase® 1500 I.U. Powder for Solution for Injection / Infusion

### 2. QUALITATIVE AND QUANTITATIVE COMPOSITION

Each ampoule contains 1,500 international units of Hyaluronidase.

For excipients, see section 6.1.

### 3. PHARMACEUTICAL FORM

Powder for solution for injection / infusion.

A white, sterile freeze-dried powder for solution for injection or infusion.

### 4. CLINICAL PARTICULARS

#### 4.1 Therapeutic Indications

Hyaluronidase can be used to enhance permeation of subcutaneous or intramuscular injections, local anaesthetics and subcutaneous infusions and to promote resorption of excess fluids and blood in the tissues.

#### 4.2 Dose and method of administration

*Adults, children and the elderly:*

**With subcutaneous infusion (hypodermoclysis):** 1500iu of Hyaluronidase dissolved in 1ml of water for injections or normal saline injected into the site, before the infusion is set up, or injected into the tubing of the infusion set, about 2cm back from the needle, at the start of the infusion. 1500iu is sufficient for administration of 500-1000ml of most fluids. Refer to Section 4.4 for information on solutions for hypodermoclysis. Care should be taken in young children and the elderly to control the speed and total volume of fluid administered and to avoid over-hydration, especially in renal impairment.

**With subcutaneous or intramuscular injections:** 1500iu of Hyaluronidase dissolved directly in the solution to be injected.

**With local anaesthetics:** 1500iu Hyaluronidase is mixed with the quantity of local anaesthetic solution to be used. In ophthalmology, 15iu of Hyaluronidase per ml is recommended.

**Extravasation:** Where dispersal rather than localisation is indicated, 1500iu of Hyaluronidase in 1ml water for injections or normal saline infiltrated into the affected area as soon as possible after the extravasation is noted.

**Haematoma:** 1500iu of Hyaluronidase dissolved in 1ml water for injections or normal saline infiltrated into the affected area.

Immediately before use dissolve the freeze-dried powder in approximately 1ml of water for injections or directly in the solution with which Hyaluronidase is to be combined.

### **4.3 Contraindications**

Hypersensitivity to hyaluronidase.

Not to be used for intravenous injections.

Not to be used to reduce the swelling of bites or stings or at sites where infection or malignancy is present.

Not to be used for anaesthetic procedures in cases of unexplained premature labour.

### **4.4 Special warnings and precautions for use**

Do not apply directly to the cornea.

Hyaluronidase should not be used to enhance the absorption and dispersion of dopamine and/or alpha agonist drugs.

Solutions for subcutaneous administration should be isotonic with extracellular fluid. Hyaluronidase is physically compatible with the commonly used infusion fluids. Use in hypodermoclysis has been reported with 0.9% sodium chloride, 0.18% sodium chloride with 4% glucose, 0.45% sodium chloride with 2.5% glucose and 5% glucose.

Potassium 34mmol/litre has been administered by hypodermoclysis in isotonic glucose or saline with 1500 I.U./litre hyaluronidase.

Electrolyte-free fluids are less preferable than those containing electrolytes and should not be given too rapidly. Hyaluronidase has also been mixed with morphine, diamorphine, hydromorphone, chlorpromazine, metoclopramide, promazine, dexamethasone, local anaesthetics and adrenaline (see 6.2. Incompatibilities).

### **4.5 Interactions with other medicines and other forms of interaction**

None stated.

### **4.6 Fertility, pregnancy and lactation**

It is not known whether the drug enters breast milk although it is unlikely to harm the breast-fed infant. Caution should be exercised in administering it to nursing mothers.

There is no evidence on the drug's safety in human pregnancy nor is there evidence from animal work that it is free from hazard. Avoid use in pregnancy unless there is no safer alternative.

#### **4.7 Effects on ability to drive and use machines**

None known.

#### **4.8 Undesirable effects**

Oedema has been reported in association with hypodermoclysis. Allergic reactions have included rare reports of periorbital oedema occurring with the use of hyaluronidase in conjunction with local anaesthetics in ophthalmology.

Severe allergic reactions including anaphylaxis have been reported rarely. Local irritation, infection, bleeding and bruising occur rarely.

#### Reporting of suspected adverse reactions

Reporting suspected adverse reactions after authorisation of the medicine is important. It allows continued monitoring of the benefit/risk balance of the medicine. Healthcare professionals are asked to report any suspected adverse reactions <https://nzphvc.otago.ac.nz/reporting/>

#### **4.9 Overdose**

No cases of overdose appear to have been reported.

For advice on the management of overdose please contact the National Poisons Centre on 0800 POISON (0800 764766).

### **5. PHARMACOLOGICAL PROPERTIES**

#### **5.1 Pharmacodynamic properties**

Hyaluronidase is an enzyme that has a temporary and reversible depolymerising effect on the polysaccharide hyaluronic acid, which is present in the intercellular matrix of connective tissue.

#### **5.2 Pharmacokinetic properties**

Not applicable

#### **5.3 Preclinical safety data**

There are no additional pre-clinical data of relevance to the prescriber.

## **6. PHARMACEUTICAL PARTICULARS**

### **6.1 List of excipients**

None.

### **6.2 Incompatibilities**

Physical incompatibility has been reported with heparin and adrenaline, although in clinical practice very low concentrations of adrenaline are combined with hyaluronidase without problems. Furosemide, the benzodiazepines and phenytoin have been found to be incompatible with hyaluronidase.

### **6.3 Shelf life**

Unopened: 3 years.

Once opened use immediately and discard any unused contents.

### **6.4 Special precautions for storage**

Do not store above 25°C.

### **6.5 Nature and contents of container**

1ml neutral glass ampoule containing a plug of white freeze-dried powder.

Pack size: 10 ampoules.

### **6.6 Special precautions for disposal**

The solution should be used immediately after preparation. The appearance of the solution is clear and not more than faintly yellow.

For detailed instructions on preparation and administration, see section 4.2.

For single use only. Discard any unused contents.

## **7. MEDICINE SCHEDULE**

Prescription Medicine

## **8. SPONSOR**

Artex Ltd  
PO Box 249  
15 Ruataniwha St  
Waipukurau

Ph: 021 730344

**9. DATE OF FIRST APPROVAL**

31 Dec 1969

**10. DATE OF REVISION OF THE TEXT**

Aug 2025

Summary Table of Changes

<b>Section Changed</b>	<b>Summary of New Information</b>
7	Change from General Sale to Prescription Medicine
8	Update to Sponsor's contact details