



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

**Public Summary**

<b>Summary for ARTG Entry:</b>	19427	10% DEXTRAN 40 in 0.9% SODIUM CHLORIDE 500mL injection bag FKB5043G
<b>ARTG entry for</b>	Medicine Registered	
<b>Sponsor</b>	Baxter Healthcare Pty Ltd	
<b>Postal Address</b>	PO Box 88, TOONGABBIE, NSW, 2146 Australia	
<b>ARTG Start Date</b>	30/09/1991	
<b>Product Category</b>	Medicine	
<b>Status</b>	Active	
<b>Approval Area</b>	Drug Safety Evaluation Branch	

**Conditions**

Conditions applicable to the relevant category and class of therapeutic goods as specified in the document "Conditions Applying to Therapeutic Goods Accepted for Registration or Listing as Goods Currently Supplied at the Commencement of the Therapeutic Goods Act 1989"

Conditions applicable to all therapeutic goods as specified in the document "Standard Conditions Applying to Registered or Listed Therapeutic Goods Under Section 28 of the Therapeutic Goods Act 1989" effective 1 February 1992.

Conditions applicable to the relevant category and class of therapeutic goods as specified in the document "Standard Conditions Applying to Registered or Listed Therapeutic Goods Under Section 28 of the Therapeutic Goods Act 1989" effective 1 February 1992.

**Products**

**1 . 10% Dextran 40 in 0.9% Sodium chloride 500mL injection bag FKB5043G**

<b>Product Type</b>	Single Medicine Product	<b>Effective Date</b>	3/07/2002
---------------------	-------------------------	-----------------------	-----------

**Permitted Indications**

No Permitted Indications included on Record

**Indication Requirements**

No Indication Requirements included on Record

**Standard Indications**

This product accepted for registration/listing as 'currently supplied' at the time of commencement of the Act. Indications held in ARTG paper records. (Old code)

**Specific Indications**

No Specific Indications included on Record

**Warnings**

See Product Information and Consumer Medicine Information for this product

**Additional Product information**

**Container information**

Type	Material	Life Time	Temperature	Closure	Conditions
Bag	Not recorded	Not recorded	Not recorded	Not recorded	Not recorded

**Pack Size/Poison information**

<b>Pack Size</b>	<b>Poison Schedule</b>
500mL x 15	Not scheduled. Not considered by committee

**Components**

**1 . Medicine Component**

<b>Dosage Form</b>	Injection, solution
<b>Route of Administration</b>	Intravenous
<b>Visual Identification</b>	Clear solution

**Active Ingredients**

dextran 40	100 g/L
sodium chloride	9 g/L

Public Summary



**Australian Government**

**Department of Health**  
Therapeutic Goods Administration

**Other Ingredients (Excipients)**

hydrochloric acid  
sodium hydroxide  
water for injections

© Commonwealth of Australia. This work is copyright. You are not permitted to re-transmit, distribute or commercialise the material without obtaining prior written approval from the Commonwealth. Further details can be found at <http://www.tga.gov.au/about/website-copyright.htm>.

Public Summary