

<i>Product Licence Number</i>	<i>Company Name</i>	<i>Product Name</i>	<i>Active Ingredients</i>	<i>Indications (Non-Prescription Items)</i>
PL0109/0041R	Roussel Laboratories Limited	Hydrotalcite Suspension (formerly Altacite)	Hydrotalcite Light 500.00mg	Use as an antacid. Hydrotalcite is indicated for symptomatic relief in the following conditions: peptic ulceration, dyspepsia, hyperacidity; gastritis; heartburn; especially when associated with reflux oesophagitis or hiatus hernia, and heartburn in pregnancy. <i>Route of Administration:</i> Taken Orally <i>Legal Category:</i> Pharmacy Medicine (500 ml size) General Sale List (250ml and 100ml size) <i>Date of Authorisation:</i> 25th March 1991
PL0113/5054R	Fisons Plc — Pharmaceutical Division	Dextraven 150 6% in 5% Dextrose	Dextran of weight average molecular weight — 150,000 6.0% w/v	<i>Legal Category:</i> Prescription Only Medicine <i>Date of Authorisation:</i> 30th October 1991
PL0113/5060R	Fisons plc — Pharmaceutical Division	Lomodex 70 in 5% Dextrose	Dextran of weight average molecular weight — 70,000 6.0% w/v	<i>Legal Category:</i> Prescription Only Medicine <i>Date of Authorisation:</i> 30th October 1991
PL0113/5061R	Fisons Plc — Pharmaceutical Division	Lomodex 70 in 0.9% Sodium Chloride	Dextran of weight average molecular weight — 70,000 6.0% w/v	<i>Legal Category:</i> Prescription Only Medicine <i>Date of Authorisation:</i> 21st October 1991
PL0113/5062R	Fisons Plc — Pharmaceutical Division	Lomodex 40 in 5% Dextrose	Dextran of weight average molecular weight — 40,000 10.0% w/v	<i>Legal Category:</i> Prescription Only Medicine <i>Date of Authorisation:</i> 29th October 1991
PL0113/5063R	Fisons Plc — Pharmaceutical Division	Lomodex 40 in 0.9% Sodium Chloride	Dextran of weight average molecular weight — 40,000 10.0% w/v	<i>Legal Category:</i> Prescription Only Medicine <i>Date of Authorisation:</i> 22nd October 1991
PL0113/5085R	Fisons Plc — Pharmaceutical Division	Dextran Powder 70	Dextran 70 100.00% w/w	Not applicable. The material is an ingredient in Dextran Injection <i>Date of Authorisation:</i> 20th February 1991
PL0113/5086R	Fisons Plc — Pharmaceutical Division	Dextran Powder 75	Dextran 75 100.00% w/w	No applicable. The material is an ingredient in Dextran Injection <i>Date of Authorisation:</i> 20th February 1991
PL0113/5083R	Fisons Plc — Pharmaceutical Division	Dextran Powder 40	Dextran 40 100.00% w/w	Not Applicable. The material is an ingredient in Dextran Injection <i>Date of Authorisation:</i> 20th February 1991