

For the effective nutritional care of infants with malabsorption problems, the answer may be elemental.

Pepti-Junior is a balanced elemental feed, which is easily digested and well absorbed. It has been specially developed for infants and young children with digestive or malabsorption problems and/or severe food intolerances.

Pepti-Junior is well tolerated, with a highly palatable taste.

The formulation comprises hydrolysed whey protein, glucose syrup, MCT oil, maize oil, minerals and vitamins.

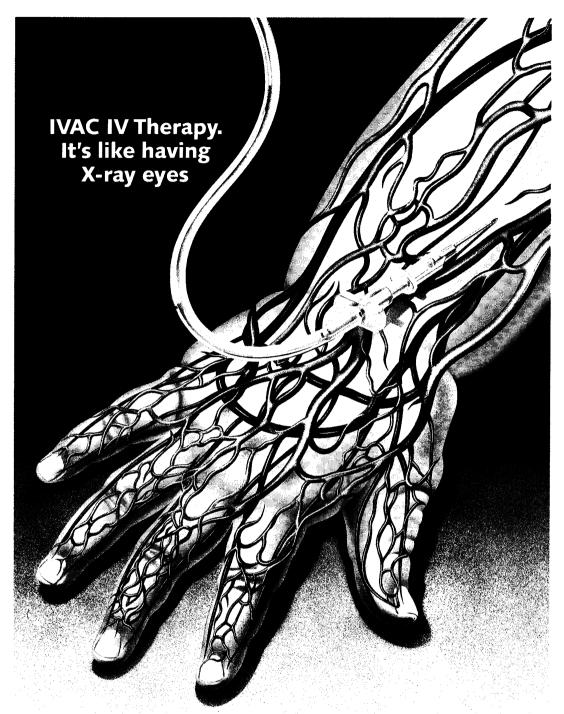
Osmolality is low at 210 mosmol/kg H₂0.



Pepti-Junior is ACBS prescribable, and the latest addition to the extensive Cow & Gate range of Specialised Formula Foods.

The balanced elemental formula for infant malabsorption

References 1. Taylor CJ, Jenkins P, Manning D Clinical Nutrition, 1988; 7: 183-190. 2. Keohane PP, Grimble GK, Brown B et al. Gut, 1985; 26: 907-913. 3. Silk DBA et al. JPEN, 1980; 4: 548-553. 4. Silk DBA et al. Gut 1979; 20: 291-299. Breast milk is the best milk for babies. PEPTI-JUNIOR has been prepared for infants with impaired gastrointestinal function and/or severe food intolerances. PEPTI-JUNIOR should only be used under strict medical supervision. For further information contact Cow & Gate, Medical Department, Trowbridge, Wiltshire BA14 8YX. (0225 768381).













Unfortunately, the most common causes of IV complications are not always visible to the naked eye, so you have to trust the IV instrument to help you.

Imagine a blocked IV, the patient stops receiving critical medication, or imagine a vesicant solution, such as potassium which is being infused interstitially due to an infiltration.

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the patient is at risk.

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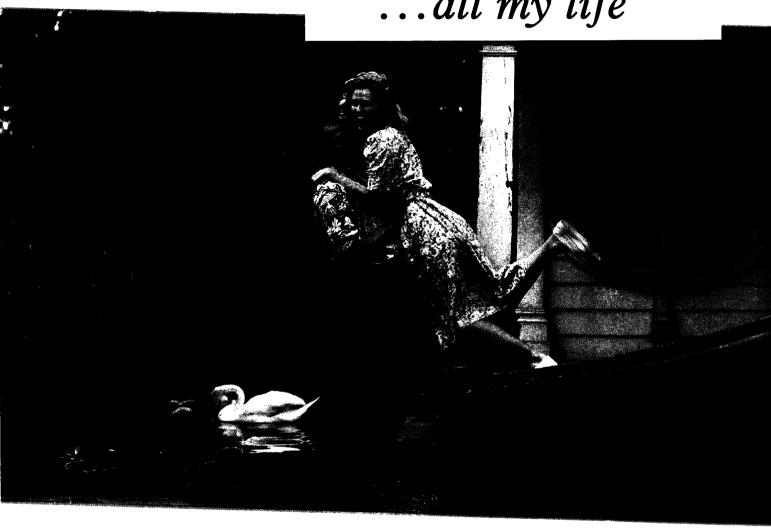
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Abbreviated Prescribing Information Presentation: Human Mixtard 30/70 (Bi-phasic Insulin Injection 30/70, Human Insulin (emp). John visia containing 100 IU/ml. Uses: Treatment of insulin-requiring diabetic patients. Dosage and Administration: The dosage is determined by the physician. Human Mixtard 30/70 may be given subcutaneously or intramuscularly. The suspension should be well mixed by gently inverting the vial several times. Care should be taken to avoid accidental intravascular injection. The injection site should be changed regularly. Contra-indications: Hypoglycaemia Precautions: Some patients previously treated with insulin of beef or pork origin may require a dosage adjustment on transfer to highly purified human insulin (emp). This is more likely in patients previously treated with beef or mixed beef/pork insulin. Reduction of early warning symptoms for hypoglycaemia may be seen upon tightening control and has been reported by a few patients on transfer from animal source to human insulin. Hyperglycaemia can be enhanced by drugs including triodothyronine; thyroxine; various natural and synthetic steroids, diuretics and cyclophosphamide. Certain beta-blockers, may affect insulin requirements and mask signs of hypoglycaemia. Monoamine oxidase inhibitors MAOI) may potentiate insulin action. Side Effects: Local reactions at the injection site are rare and may include transient crythema, induration, urticaria and ocedma. True generalised hypersensitivity reactions are very rare. Localised lipodystrophy is rare but cannot be totally excluded. Use in Pregnancy and Lactation: It is important to maintain continuous good glycaemic control during conception and throughout pregnancy. Use in the elderly: Insulin may have a more prolonged action, dose requirements should be regularly reviewed. Product Licence Number: Human Mixtard 30/70 31320037. Basic NHS Price: 10ml glass vial 18.45. Product Licence Holder: Novo Nordisk House, Broadfield Park, Brighton Road, Pease Pottage, Crawley, West Sussex Rf111 PRIC, Englan

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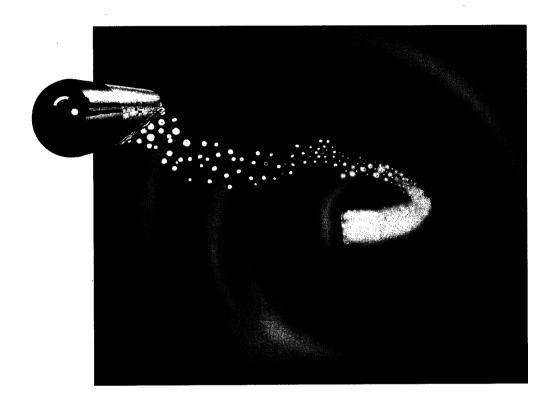
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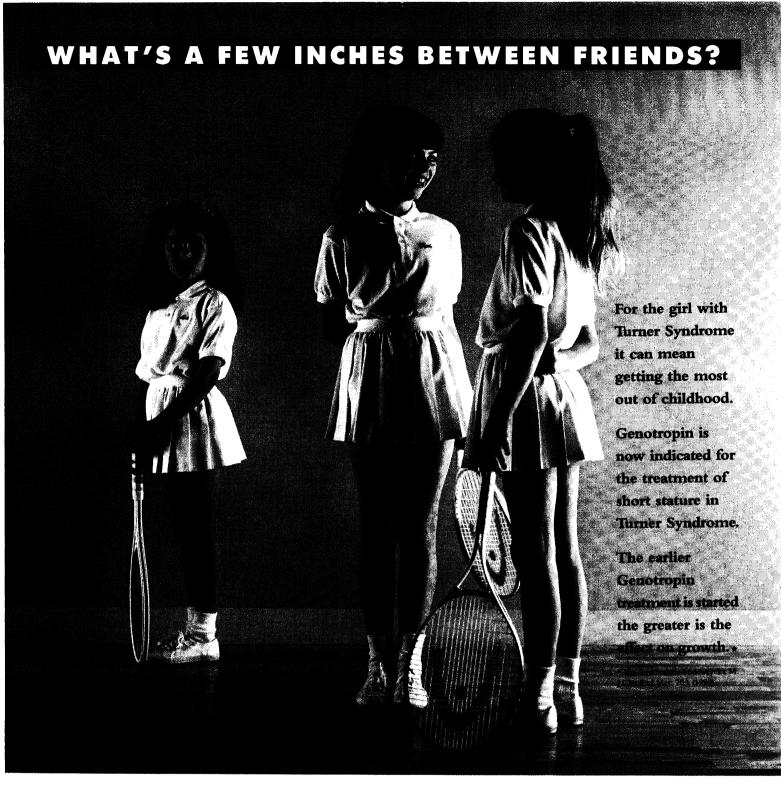




RIGHT ON TARGET - RIGHT FROM THE START

Prescribing Information — Presentation: Brown-yellow capsules containing enteric coated granules of pancreatin equivalent to: 9,000 BP units of amylase, 8,000 BP units of lipase, 210 BP units of protease. Available in packs of 100. Basic NHS price £13.33. Indication: Pancreatic exocrine insufficiency. Dosage and administration: Adults and children: Initially one or two capsules with meals, then adjust according to response. The capsules can be swallowed whole, or for ease of administration they may be opened and the granules taken with fluid or soft food, but without chewing. If the granules are mixed with food, it is important that they are taken immediately, otherwise dissolution of the enteric coating may result.

Contra-indications, Warnings, etc: Contra-indications: Substitution with pancreatic enzymes is contra-indicated in the early stages of acute pancreatitis. Warnings: Use in pregnancy: There is inadequate evidence of safety in use during pregnancy. The product is of porcine origin. Rarely cases of hyper-uricosuria and hyper-uricaemia have been reported with high doses of pancreatin. Overdosage could precipitate meconium ileus equivalent. Perianal irritation could occur, and, rarely, inflammation when large doses are used. Product Licence Number: 5727/0001. Name and address of Licence Holder: Kali Chemie Pharma GmbH, Postfach 220, D-3000, Hannover 1, West Germany.



Genotropin (rbe)



ABBREVIATED PRESCRIBING INFORMATION Presentation Genotropin 121U Multidose A vial of sterile lyophilised powder containing 12 IU somatropin (tbe) supplied with 1 ml of Water for Injections. Indications The treatment of short stature caused by decreased or absent secretion of pituitary growth hormone. The diagnosis should be verified by a specialist medical practitioner. Short stature in gonadal dysgenesis (Turner Syndrome). Reconstitution Add 3 ml of Water for Injections with 0.25% m-cresol to the 12 IU and 1 ml Water for Injections on the 4 IU in the vial. Dissolve the drug gently without shaking vigorously. Dosage and Administration Administer by subscutaneous injection. Generally a dose of 0.5 – 0.7 IU/kg body weight per week is recommended. Contra-Indications Only patients with unfused epiphyses should be treated. Precautions Patients with diabetes mellitus may require adjustment of their antidiabetic therapy. Patients treated with Genotropin should be regularly assessed by a child growth specialist. The injection site should be rotated to prevent lipoatrophy. Pregnancy and Lactation In the event of pregnancy occurring during Genotropin therapy, treatment should be discontinued is available as to whether peptide hormones pass into breast milk. Side-Efficets A few children have developed transient local reactions. In Turner Syndrome temporary exacerbation of lymphoedema has been reported. Antibodies towards growth hormone may be formed in some patients but these antibodies rarely have clinical significance. Pharmaceutical Precautions 12 IU Multidose may be used over a period of 21 days if kept in the refrigerator. Genotropin if 10 Vivo pack sizes of Genotropin 1 IU are available: a combined pack containing one vial of somatropin (rbe) 4 IU and one ampoule of 1 ml Water for Injections, and a combined pack containing to 10 Vivo pack sizes of Genotropin 1 IU are available: a combined pack containing one vial of somatropin (rbe) 4 IU and one ampoule of 1 ml Water for Injections, and a combined pack containing to





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The types of Fellowship, which are open to men or women are

- A To enable paediatricians from any part of the Commonwealth overseas to spend up to twelve weeks in the United Kingdom, meeting British Paediatricians and seeing something of their work. Preference will be given to those recently established in an academic career who can arrange their visit to allow attendance at the Annual Meeting of the British Paediatric Association in April 1992.
- To enable paediatricians from the United Kingdom of Registrar, Senior Registrar, or Consultant status, or non-medically qualified members of the Association, in the early years of professional life to make a short working visit (up to three months) to a centre in a developing country, teaching or conducting research so as to benefit both fellow and hosts.

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Applications for C Fellowships can be accepted by 31st January, or 31st July 1991.

The conditions for the Fellowships and applications forms may be obtained from the British Paediatric Association, 5 St Andrew's Place, Regent's Park, London NW1 4LB.

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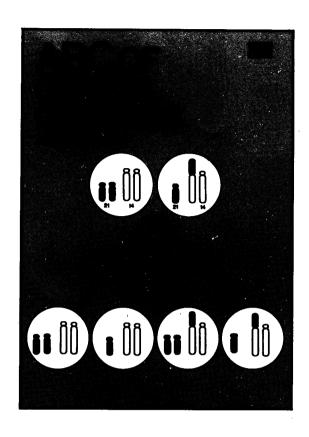
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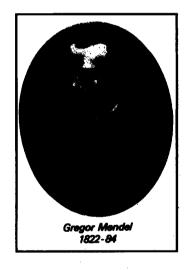
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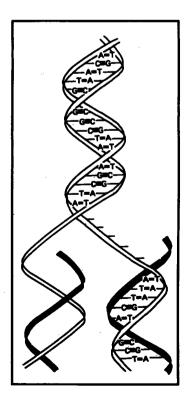
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Genetics is a subject fundamental to medicine, but rapid advances in recent years may make the subject complex and confusing to clinicians and paramedical staff outside the specialty. Nevertheless, the demand for genetic investigation and counselling is increasing, and these services need to be offered by clinicians other than clinical geneticists. In the ABC of Clinical Genetics Helen Kingston, consultant clinical geneticist at St. Mary's Hospital, Manchester, reviews basic concepts of inheritance in mendelian, chromosomal, and multifactorial disorders; outlines the investigation and management of genetic conditions; and illustrates the application of recombinant DNA technology to clinical practice. Implications of genetic diseases for relatives are also considered, together with the ethical and human dilemmas that face doctors and patients and their families.

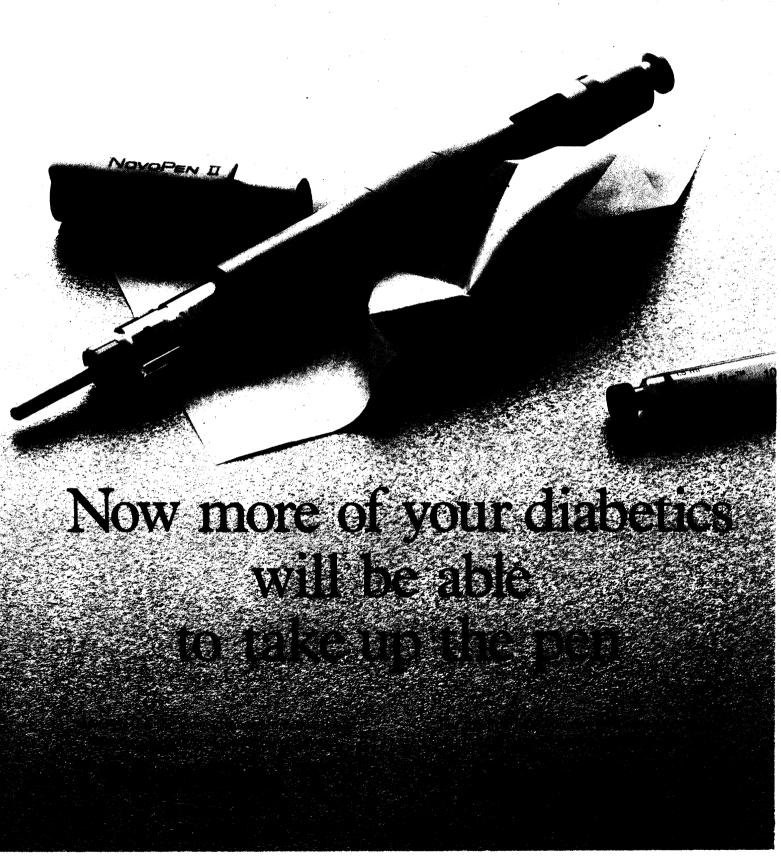


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requirements. Use in Pregnancy: Insulin requirements usually fall during first trimester and increase during second and third trimester. Side-effects: Lipodystrophy insulin resistance and hypersensitivity are rarely reported with human insulin. Product Licence Number and Basic NHS Price: PL 4668/0020, 5 × 1.5ml cartridges £8.10. Product Licence Holder: Novo Nordisk A/S, Novo Alle, DK-2880 Bagsvaerd, Denmark. Sole Distributor: Farillon Limited, Ashton Road, Harold Hill, Romford, Essex RM3 8UE. Further Information is available on request from: Novo Nordisk Pharmaceuticals Ltd, Novo Nordisk House, Broadfield Park Brighton Road, Pease Pottage, Crawley, West Sussex RH11 9RT. Tel: 0293 613555.

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