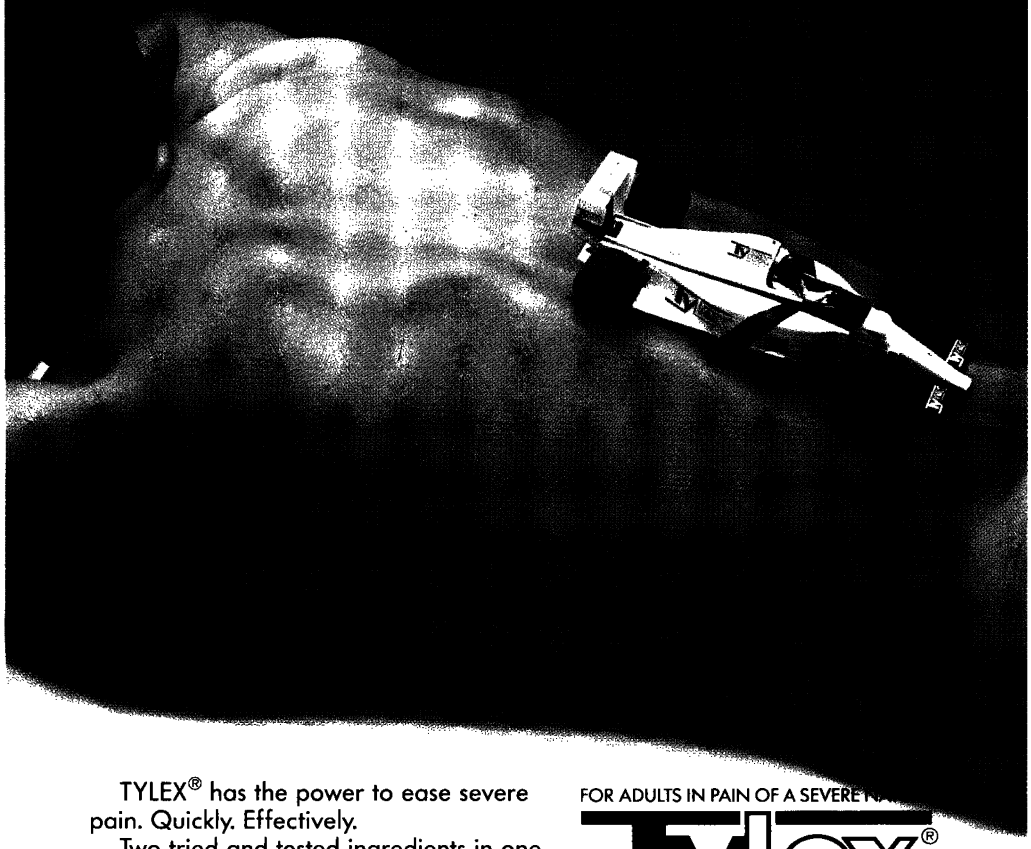
WHEN PAINFUL BACKS NEED POWERFUL RELIEF FROM SEVERE PAIN



TYLEX® has the power to ease severe pain. Quickly. Effectively.

Two tried and tested ingredients in one capsule with a flexibility of dose to treat even the severest back pain.

TYLEX® Because it works.

FOR ADULTS IN PAIN OF A SEVERE N...

Tylex®
CAPSULES

Paracetamol Ph.Eur.500mg/
Codeine phosphate Ph.Eur.30mg

ONE OR TWO CAPSULES Q.D.S.

Prescribing information. Presentation. TYLEX® Capsules are size 0 hard gelatin capsules imprinted C30 with a white body and red cap. Each capsule contains 500mg of Paracetamol and 30mg Codeine Phosphate. Uses: For the relief of severe pain. Dosage and administration: TYLEX® Capsules are given orally. The usual adult dose is one or two capsules every four hours as required. The total daily dose should not exceed 240mg Codeine Phosphate. Contra-indications, warning, etc. Precautions: Patients receiving other central nervous system depressants concomitantly with this drug may exhibit an additive depressant effect. Use of TYLEX® Capsules is not recommended during pregnancy, lactation or in children since safety in these groups has not been established. Warning: Codeine can produce drug dependence of the morphine type and therefore has the potential for being abused. Side effects: The most frequently observed adverse reactions include light-headedness, dizziness, sedation, nausea and vomiting. Overdosage: Serious overdosage of codeine can result in respiratory depression, bradycardia and hypotension. Opioid antagonists may be employed. Hepatic toxicity can result from paracetamol over-dosage. Pharmaceutical precautions: Protect from light. Legal category: POM. Package quantities: Containers of 100 capsules. Further information: Nil. Product licence number: PL0076/0109. Basic N.H.S. Price: £8.60/100, £43/500 capsules. Further information available from: Cilag Limited, Saunderton, High Wycombe, Buckinghamshire, HP14 4HJ, England. Date of preparation January 1993. ® Denotes Registered Trademark

Cilag

Haemaccel®

polygeline

The flexible solution, with more than a decade of British experience

Presentation: Each 500ml bottle contains 17.5g polygeline. **Cations** (mmol/500ml): sodium 72.50, potassium 2.55, calcium 3.13. **Anions** (mmol/500ml): chloride 72.50, phosphate – traces, sulphate – traces. Sterile distilled water to 500ml. **Uses: 1.** As a plasma volume substitute in the initial treatment of hypovolaemic shock due to a) Haemorrhage (visible or concealed), b) Burns, peritonitis, pancreatitis, crush injuries. **2.** Fluid replacement in plasma exchange. **3.** Extra-corporeal circulation. **4.** Isolated organ perfusion. **5.** Carrier solution for insulin. **Dosage and administration:** Haemaccel should be administered intravenously in a volume approximately equal to the estimated blood loss. Normally 500ml will be infused in not less than 60 minutes but in emergencies Haemaccel can be infused rapidly. **Hypovolaemic shock:** 500-1,000ml Haemaccel intravenously initially. Up to 1.5 litres blood loss can be replaced entirely by Haemaccel. For between 1.5 and 4 litres blood loss, fluid replacement should be with equal volumes of Haemaccel and blood, given separately. (See Pharmaceutical Precautions). For losses over 4 litres the separate infusions should be in the ratio two parts blood to one part Haemaccel. **Burns:** At least 1ml Haemaccel per kg body weight, multiplied by the percent of body surface burned, should be infused in each 24 hours for 2 days. **Plasma exchange:** Haemaccel should be given either alone or in combination with other replacement fluids in a volume adequate to replace the plasma removed. Up to 2 litres have been given as sole replacement fluid. **Contraindications, warnings etc:** There are no absolute contraindications to the use of Haemaccel. However, caution should be used in any patient likely to develop circulatory overloading. Inappropriately rapid administration of Haemaccel, especially to normovolaemic patients, may cause the release of histamine. Histamine release may be especially hazardous in patients with known allergic conditions such as asthma. In these cases prophylaxis with H_1 and H_2 receptor antagonists is advisable. In the events of anaphylactic shock the infusion should be discontinued and adrenaline (5-10ml of 1:10,000 by slow intravenous injection or 0.5-1.0ml of 1:1,000 by intramuscular/subcutaneous injection) should be given immediately. Haemaccel contains calcium ions and caution should be observed in patients being treated with cardiac glycosides. Haemaccel should, if possible, be warmed to body temperature before use. However in emergencies it may be infused at ambient temperatures. **Pharmaceutical precautions:** As Haemaccel contains no preservatives, any unused fluid should be discarded once a bottle has been opened. Citrated blood may be infused immediately before or after Haemaccel provided there is adequate flushing of the infusion set. **PL**0086/0040. **Date of Preparation** June 1993. **Basic NHS Hospital Price** £3.43 per 500ml bottle. Behring – A division of HOECHST UK LTD, Salisbury Road, Hounslow, Middlesex TW4 6JH.

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