DATA SHEET
GASTROLYTE

Name of Medicine
Gastrolyte

Active ingredients:
Sachet: Sodium chloride 470mg, potassium chloride 300mg, sodium acid citrate 530mg, glucose 3.56g.
Tablet: Sodium chloride 117mg, potassium chloride 186mg, anhydrous citric acid 384mg, sodium bicarbonate 336mg, glucose 1.62g.

Excipients:
Oral glucose electrolyte preparations
Sachet Silicon dioxide, saccharin sodium or aspartame, orange flavour, pineapple flavour.
Tablet Saccharin sodium, blackcurrant flavour.

Presentation
Sachet 1 L of made up solution (5 sachets, 5x200mL) contains sodium (Na⁺) 60mmol, potassium (K⁺) 20mmol, chloride (Cl⁻) 60mmol, citrate 10mmol and glucose 90mmol.
Tablet 1L of made-up solution (10 tablets, 5x200mL quantities) contains sodium (Na⁺) 60mmol, potassium (K⁺) 25mmol, chloride (Cl⁻) 45mmol, citrate 20mmol and glucose 90mmol.

Uses

Actions
Compound sodium chloride and glucose oral powder and effervescent tablet. Total osmolarity is 240mOsm/L.

Indications
Oral correction of fluid and electrolyte loss in infants, children and adults. The product has been formulated to treat fluid and electrolyte loss associated with acute dehydration in infantile diarrhoea, but is also appropriate for the treatment of older children and adults. Gastrolyte is also indicated for the correction of dehydration due to traveller's gastroenteritis and vigorous and prolonged exercise.

Dosage and Administration
For oral administration only.

Reconstitution: The contents of one sachet or two effervescent tablets should be made up to 200ml with fresh drinking water. (For infants, the water should be freshly boiled and cooled before mixing with GASTROLYTE). An infant's feeding bottle is a convenient measure of this volume. The solution should be made up immediately prior to feeding and any solution remaining an hour after reconstitution should be discarded. However, the solution may be used for up to 24 hours if stored in a refrigerator immediately after reconstitution. The reconstituted solution must not be boiled.
Dosage: The volume of reconstituted GASTROLYTE needed per day should be determined by the physician, taking into account the weight of the patient and severity of the condition.

**Toddlers and older children:**

For toddlers and older children, GASTROLYTE may be given freely until their thirst is satisfied or frequently while diarrhoea persists.

**Infants:**

A number of different regimens are recommended for the treatment of infantile diarrhoea, but the basic principle is to provide enough fluid to rehydrate the infant. Further fluid is then given to allow for ongoing losses until the diarrhoea settles. During the first 12-24 hours of rehydration, milk and solid feeds can be omitted, but these should be rapidly reintroduced to avoid malnutrition. Milk feeds may be half-strength initially, but full-strength feeds should be given within 24 hours. If the diarrhoea persists or is worsened by the milk, then the possibility of a temporary intolerance to lactose should be considered. If confirmed, a low lactose formula can then be offered.

Infantile diarrhoea is uncommon in breast-fed infants. However, if treatment with GASTROLYTE becomes necessary, it is suggested that for each feed the chosen regimen is followed such that the infant is given the appropriate volume of GASTROLYTE for that feed and then put to the breast until satisfied. Expression of residual milk from the breasts may be necessary during this period.

In those patients who are vomiting at the start of treatment, it may be advisable to offer very small volumes initially until vomiting is under control. If the vomiting and diarrhoea show no sign of moderating, the patient should be reassessed.

The dosage and regrading schemes are only a general guide and the volume of GASTROLYTE given and the speed of reintroduction of the normal feeds is at the discretion of the physician.

**Mild Cases (no dehydration):**

During the first 12-24 hours the infant is offered GASTROLYTE solution in the same quantities as are used for the usual feeds. For the next 12 to 24 hours, the infant can be given full strength feeds of the usual formula mixed with an equal volume of water. Following this, full strength feeds should be recommenced. Ordinary solid feeds should be continued throughout.

**Dosage: Volume of fluid required over 6 hours for the prevention of dehydration**

<table>
<thead>
<tr>
<th>Patient’s weight (kg)</th>
<th>Volume of Gastrolyte (mL) in 6 hours</th>
</tr>
</thead>
<tbody>
<tr>
<td>5</td>
<td>150 – 210</td>
</tr>
<tr>
<td>10</td>
<td>300 – 420</td>
</tr>
<tr>
<td>15</td>
<td>450 – 630</td>
</tr>
<tr>
<td>20</td>
<td>600 – 840</td>
</tr>
<tr>
<td>30</td>
<td>900 – 1260</td>
</tr>
</tbody>
</table>

For patients over 30kg, fluid replacement should be 5-7mL/kg/hr plus replacement of any additional losses.

**More Severe Cases (dehydration):**

Prompt medical attention should be sought if dehydration is suspected. Signs of dehydration include documented weight loss, reduced urine output, and diminished skin turgor. A suggested regimen for the treatment of infantile diarrhoea with marked dehydration based on bodyweight in kilograms is given.
Dosage: Volume of fluid required over 6 hours for the treatment of dehydration

<table>
<thead>
<tr>
<th>Patient’s weight (kg)</th>
<th>Volume of Gastrolyte (mL) in 6 hours</th>
</tr>
</thead>
<tbody>
<tr>
<td>5</td>
<td>200 – 400</td>
</tr>
<tr>
<td>10</td>
<td>400 – 800</td>
</tr>
<tr>
<td>15</td>
<td>800 – 1000</td>
</tr>
<tr>
<td>20</td>
<td>900 – 1500</td>
</tr>
<tr>
<td>30</td>
<td>1000 – 2000</td>
</tr>
</tbody>
</table>

After 6 hours, fluid replacement should be 5 – 7 mL/kg/hour plus replacement of any additional losses.

If the condition worsens, or the diarrhoea has not stopped within 6 hours in infants under 6 months, within 12 hours for children under 3 years, 24 hours in children 3-6 years of age or 48 hours in children over 6 years of age, or the child has decreased urinary output, the clinician should ensure patients and parents are aware of the risk and early warning signs of dehydration in young children and infants.

If the condition further worsens or fails to improve, IV fluid replacement is required.

Dosage: Treatment of dehydration in older children and adults

<table>
<thead>
<tr>
<th>Approx. number of sachets</th>
<th>Child</th>
<th>Adult</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>1 sachet after each loose motion or 6-12 sachets per 24 hours</td>
<td>1 or 2 sachets after each loose motion or 8-16 sachets per 24 hours</td>
</tr>
<tr>
<td>Diet Day 1</td>
<td>Limited food if desired</td>
<td></td>
</tr>
<tr>
<td>Diet Day 2</td>
<td>Gradually return to normal diet by slowly reducing the amount of Gastrolyte and increasing the amount of food (Light solids at first, e.g. cereals, toast, etc.)</td>
<td></td>
</tr>
</tbody>
</table>

Contraindications

There are no known contraindications to GASTROLYTE. However, there may be a number of conditions where treatment with GASTROLYTE will be inappropriate, e.g. intestinal obstruction requiring surgical intervention.

Warnings and Precautions

Use water ONLY to mix Gastrolyte powder or effervescent tablets. Never dilute with lemonade, soft drinks, cordials or any other fluid than water.

Gastrolyte powder is available with either sodium saccharin or aspartame as a sweetening agent. Phenylketonurics are warned that this product, in some forms, contains phenylalanine. Please refer to the packaging for information relating to the sweetening agent used.

For oral administration only. The contents of one sachet or two effervescent tablets should always be made up to 200ml with water (the product must not be reconstituted in diluents other than water, e.g. must not be included in milk solutions). A weaker solution than recommended will fail to provide adequate sugar and electrolytes and a stronger solution than recommended may give rise to hypernatraemia. Intravenous rehydration is required for dehydrated children with shock, very large stool losses (>10mL/kg/hour), severe vomiting that interferes with oral fluid replacement, or glucose malabsorption. Patients with gastroenteritis should be monitored carefully to ensure that their condition does not deteriorate. In particular, young infants may become severely dehydrated in a short time.
Patients and/or parents should be advised to seek medical advice if the condition worsens. Clinicians should particularly ensure patients are aware of the risk of dehydration in young children and infants. Early warning signs of impending dehydration should be discussed. Seek medical advice if diarrhoea persists for more than 6 hours in infants under 6 months, 12 hours in children aged 6 months to 3 years, 24 hours in children aged 3 to 6 years and 48 hours in children over 6 years of age.

**Pregnancy and Lactation**

There is no contraindication to the use of Gastrolyte in pregnancy and lactation. However, pregnant or lactating women with gastroenteritis should seek specialist medical advice.

**Adverse Effects**

None reported.

**Medicine Classification**

General Sale

**Package Quantities**

<table>
<thead>
<tr>
<th>Sachets</th>
<th>Saccharin: Orange flavour: 5.2g, 10’s</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Aspartame: Natural and Fruit flavour: 4.9g, 10’s</td>
</tr>
<tr>
<td>Tablets, effervescent</td>
<td>Blackcurrant: 20’s</td>
</tr>
</tbody>
</table>

**Name and Address**

Distributed by:
sanofi-aventis new zealand limited
Auckland
Ph: (09) 580 1810

**Date of Preparation**

May 2008