BARD® Nasogastric Sump Tubes
Instructions For Use

NG Tubes, PVC with Radiopaque Stripe

<table>
<thead>
<tr>
<th>Fr.</th>
<th>Size (mm, inch)</th>
<th>Length (cm, inch)</th>
</tr>
</thead>
<tbody>
<tr>
<td>10</td>
<td>3.2 (1/8”)</td>
<td>45 (18”)</td>
</tr>
<tr>
<td>12</td>
<td>4.0 (5/32”)</td>
<td>60 (24”)</td>
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<tr>
<td>14</td>
<td>4.7 (3/16”)</td>
<td>60 (24”)</td>
</tr>
<tr>
<td>16</td>
<td>5.3 (7/32”)</td>
<td>60 (24”)</td>
</tr>
<tr>
<td>18</td>
<td>6.0 (7/32”)</td>
<td>60 (24”)</td>
</tr>
</tbody>
</table>

Standard Nasogastric Sump Tube
- EN0042100
- EN0042120
- EN0042140
- EN0042160
- EN0042180

Nasogastric Sump Tube and PREVENT Anti-Reflux Filter with ENFit™ Connectors
- EN0046100
- EN0046120
- EN0046140
- EN0046160
- EN0046180

Nasogastric Sump Tube and PREVENT Anti-Reflux Filter with ENFit™ connectors and STATLOCK® Nasogastric Stabilization Device
- EN0046163
- EN0046183

Nasogastric Sump Tube and PREVENT Filter and Lopez Valve® with ENFit™ connectors
- EN0056120
- EN0056140
- EN0056160
- EN0056180

Nasogastric Sump Tube and PREVENT Filter and Lopez Valve® with ENFit™ Connectors and STATLOCK® Nasogastric Stabilization Device
- EN0056163
- EN0056183

Attention: All references to the NG Tube, PREVENT® Filter and Lopez Valve® hereafter are understood to be ENFit™ compatible. Read all instructions prior to use.

Indications for Use:
BARD® Nasogastric Sump Tubes with ENFit™ connector are intended to be used for:
• Decompression of stomach by suction or aspiration of gastric contents.
• Short-term administration of term tube feeding, lavage fluid and medications.

Contraindications:
• Patients with known tape or adhesive allergies.

Warnings
1. Use with caution in patients with a history of head trauma, facial trauma, esophageal diseases and patients with potential for vomiting.
2. Do not force Nasogastric Tube during insertions; damage to the nasal passage and mucosa and bleeding may occur.
3. MEASURE INSERTION LENGTH CAREFULLY- Excessive insertion length of tube into the stomach may lead to coiling and/or formation of tube-knot.
4. Lubricate the tube generously with water soluble lubricant prior to insertion. DO NOT use petroleum-based products as they may be harmful to the respiratory tract.
5. Reflux of gastric contents into the blue vent lumen indicates that the suction lumen is obstructed or suction is too low. Routinely check for reflux in the blue vent lumen and clear as per applicable directions. Failure to clear the obstruction or clear PREVENT® Filter may cause gas & fluid buildup in stomach, aspiration of gastric contents, aspiration pneumonia and other complications.
6. DO NOT inject fluid through the PREVENT® Filter as this may result in blockage and leakage of filter.

STATLOCK® Nasogastric Stabilization Device: Avoid contact with alcohol or acetone; both can weaken bonding of components and STATLOCK® Stabilization Device pad adherence.

Instructions for Nasogastric Tube insertion
1. Explain the procedure to the patient.
2. Carefully measure to find desired length of the tube using the Nasogastric Tube as a measurement aid. To determine the insertion length: measure the tube from the tip of the nose to the earlobe and from the earlobe to the tip of the xiphoid process. Mark the length of the tube to be passed with a small piece of tape.
3. Check the patient’s nostrils for patency; select the nostril with best patency.
4. Lubricate the full length of tube to be inserted.
5. Insert the tube through the nose aiming down and back. When the tube hits the pharynx, if patient is able, have him or her flex his/her head forward and swallow. Advance the tube as the patient swallows. If resistance is met, rotating the tube may facilitate placement.
6. Continue to advance the tube until the marked position on the tube is reached. DO NOT advance beyond the marked length as coiling and or knotting of the tube in the stomach may occur.
7. Confirm tube placement per hospital policy. The tube has a radiopaque stripe facilitating x-ray confirmation. If proper placement of tube within the stomach cannot be confirmed, remove the tube gently and start the procedure again.
8. Secure with a securement device or tape per hospital protocol.
9. If applicable, ensure the Lopez Valve® with ENFit™ connector is snugly inserted into suction lumen to prevent suction loss.
10. Keep blue vent lumen above the level of the patient’s stomach to prevent reflux of stomach fluids into the blue lumen.
11. DO NOT clamp air vent port while suction is being applied.

Recommended Suction Settings
Always use lowest suction setting that will effectively decompress the stomach. For intermittent suction via thermotic pump, use “High” (Gomco, 120mm Hg). For intermittent suction via central source, set at “Low” (30-40mm Hg). For continuous suction, set at “Low” (30-40mm Hg). Increase slowly until flow is observed as necessary.

Instructions for PreVent® Anti-Reflux Filter with ENFit™ connector
1. Firmly twist the white base of anti-reflux filter in blue air lumen vent of nasogastric tube.
2. If gastric reflux in vent lumen is observed, clear the obstruction in the main lumen by following your hospital’s standard protocol. Attach ENFit™ - compatible syringe to ENFit™ male fitting on anti-reflux filter and inject a minimum of 15cc of air to clear the blue air vent lumen of any gastric reflux. DO NOT inject fluid through filter.
3. To cap nasogastric tube when tube is not connected to a suction source insert tethered cap that is attached to the barb in the suction lumen of nasogastric tube.

Instructions for Lopez Valve® with ENFit™ connector (when included):
1. Attach medication port cap to side “C”.
2. Turn valve to position one and attach NG Tube to side “B”.
3. Attach suction tube to side “A” if suction is desired.
4. To administer medication, remove cap. Attach syringe to sideport, push and twist until tight and turn valve to position four. Flush valve per facility protocol following administration.
5. Return valve to position one when complete to avoid leakage.

Instructions for the STATLOCK® Nasogastric Stabilization Device (when included):
Remove oil and moisturizer from targeted skin area. Apply skin prep to targeted STATLOCK® Stabilization Device area for skin protection and enhanced pad adherence. ALLOW TO DRY COMPLETELY.

Caution: Federal (U.S.A.) laws restricts this device to sale by or on the order of a physician.

Sterile/Single Patient Use - Do not resterilize.

The STATLOCK® Stabilization Device does not contain natural rubber latex.

Note: After use, this product may be a potential biohazard. Handle and dispose of in accordance with accepted medical practice and applicable local, state and federal laws and regulations.

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Lopez Valve is a registered trademark of ICU Medical.

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