## Get ready for the new ENFit connector Transition Checklist for Nurses and Clinicians

A new design standard for medical device tubing connectors is on its way. Starting with enteral feeding, application-specific standards will help ensure that connectors do not fit into ports other than the type for which they are intended, reducing the incidence of misconnections.

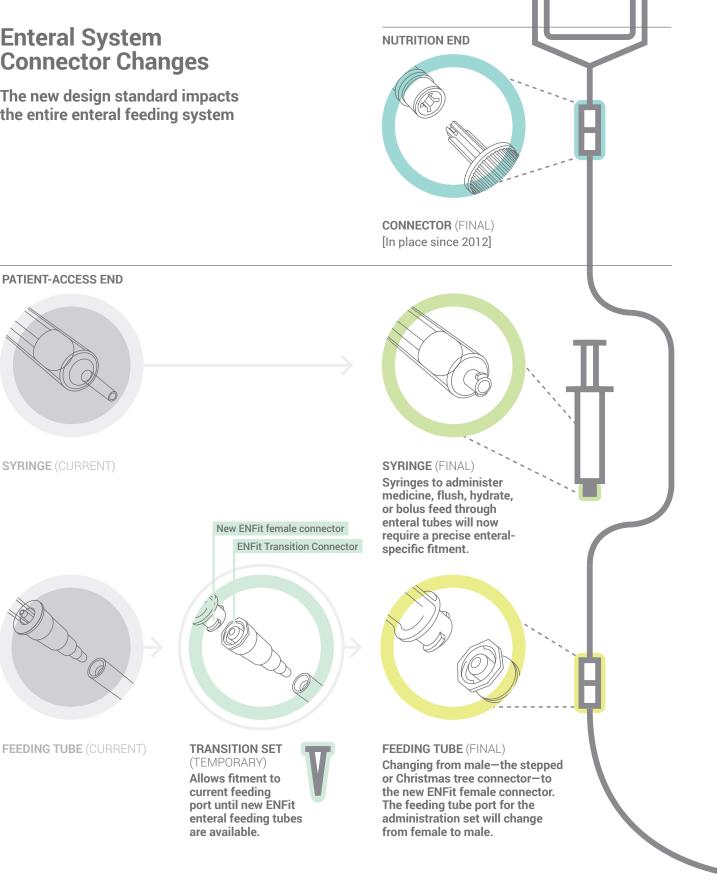
This global patient safety initiative starts in the US, Canada, and Puerto Rico, with the goal of completion in these markets by 2016.

More than anyone else, nurses and clinicians will be handling and using the new ENFit connectors, so your preparedness is key. This is not intended to be a complete list, but use the STEPS below to get started:

S	Supplier communication	<ul> <li>Familiarize yourself with all the product-specific changes</li> <li>Practice new connections with all products including feeding sets, enteral-specific syringes, and feeding tubes</li> <li>Understand anticipated timing of the transition</li> </ul>
Т	Training	<ul> <li>Engage your nursing education team</li> <li>Communicate importance of connector changes to enhance patient safety</li> <li>Explain how new feeding sets will change and demonstrate how they will connect</li> <li>Reinforce locking feature and safety benefits of the new ENFit connector</li> <li>Identify a super user for each shift and seek hands-on training opportunities</li> </ul>
Ε	Education	<ul> <li>Patient/Caregiver</li> <li>Discuss implications of impending connector changes, including timing and transition process</li> <li>Plan opportunities for patients/caregivers going home on enteral feeding to learn about: <ul> <li>The difference between current and new devices, and how they work</li> <li>Administering enteral formula and medications if prescribed</li> <li>Flushing the tube and checking residuals as appropriate</li> <li>Providing additional hydration</li> </ul> </li> <li>Give checklist to patient/caregivers discharged on enteral feeding</li> <li>Staff</li> <li>Direct product-specific questions to the manufacturer/supplier</li> <li>Direct procedural questions to a multidisciplinary transition team</li> </ul>
Ρ	Process	<ul> <li>Assemble a multidisciplinary transition team to evaluate current procedures and protocols and adjust to include the new ENFit connectors</li> <li>Update medication and enteral delivery order sets for use of new products</li> <li>Review pharmacy medication preparation and delivery processes to incorporate new enteral-specific syringes</li> <li>Develop communication mechanisms between nursing and pharmacy to identify patients who need medication through feeding tube</li> <li>Inform prescribers and pharmacists that medication orders must specify route—enteral (tube) or oral (mouth), and not say "po" for both. Until ENFit connectors are fully transitioned, the order must also indicate which connection the patient is using</li> </ul>
S	Supply management	<ul> <li>Audit and consolidate all storage areas for enteral devices (feeding sets, enteral-specific syringes, feeding tubes) in preparation for new inventory</li> <li>Reduce excess inventory of enteral devices with current connectors</li> <li>Let products with ENFit Transition Connectors and ENFit feeding tubes flow through inventory</li> <li>Avoid overstocking any enteral feeding products</li> </ul>

## Enteral System Connector Changes

The new design standard impacts the entire enteral feeding system





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