







KB Medical Group Inc.
9650 Research Drive, Suite A
Irvine CA 92618
P: 949-333-3785
info@kbmed.com

ENFit[®] Supply and Demand for the U.S

Dear Manufacturers, Patients, Caregivers, Healthcare and Supply Chain Professionals,

The Global Enteral Device Supplier Association (GEDSA) previously announced that member manufacturers will phase out legacy feeding devices and transition adaptors starting July 1st, 2020.

KB Medical Group would like to support the guidance from the U.S. Food and Drug Administration (FDA) to reduce medical tubing misconnections and improve patient safety. However, given the increased demand for enteral feeding caused by the pandemic, and to improve the flow of supply to remove the obstacles, KB Medical will provide ENFit supplies and legacy adaptors during the pandemic:

The Y port is used to add nutrition during the delivery application	
The drip chamber is used to allow passage during application	
The transition adaptor is used to transform the legacy syringe (disposable with a luer tip) into a syringe compatible with the ENFit [®] feeding system.	
Transfer connector plus female connector is used as a transfer from ENFit [®] enteral female connector to the legacy feeding tube.	



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To minimize any disruption to patient feeding and care, GEDSA members will monitor and commit to supply ENFit in a tool-box format to adjust needs accordingly to the hospitals.

KB Medical is fully committed of transition to ENFit as our ISO-compliant option. However, since COVID-19 created an unprecedented time, our internal team would like to support the organizations that has converted and also in the process offer them ENFit or legacy adaptors.

As a manufacturer of FDA approved ENFit products we are preparing for the increased demand.

To Ensure about product availability of ENFit, legacy adaptors, or contract manufacturing opportunities please work directly with our team and keep them updated about your demands and we will ensure you are connected with a representative.