

# Benefits of ISO Standardization to the ENFit® Enteral Feeding Connector



## Improved patient safety

While there are many supply chain, clinical, and user benefits associated with the standardized ENFit connector, improved patient safety is the key benefit and top priority.



## Less supply disruptions / backorders

- Split production lines (legacy product & ENFit product) can now both run ENFit, doubling output.
- Warehousing capacities and safety stock levels can increase, since stock isn't split between legacy & ENFit.
- Raw materials are no longer being competed over by split product lines.
- Alternative suppliers are available because the standardized ENFit products all work together.
- Unlike ENFit, proprietary systems, often have no alternative supplier when a supply disruption occurs.



## Patient Portability

With patients and facilities standardized to one system, there are no product incompatibilities when a patient is transferred, no matter whether that transfer is across the street or halfway across the globe.



## Product Innovations

- With standardized connectors, new technologies emerge because the tech doesn't have to accommodate numerous connection types.
- Several new innovations, that improve patient outcomes and experiences, have already emerged around ENFit.
- Manufacturers are also able to allocate more resources into R&D, instead of having those resources managing split product lines.



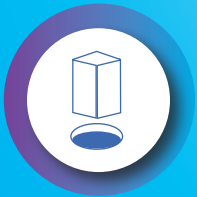
## Reduced Training

Clinicians are not having to constantly learn new product systems and connections because of ENFit's robust supply chain and multitude of backup suppliers.



## More product options

With nearly all manufacturers standardizing to one connector, an enormous array of compatible products becomes available. Clinicians then have a greater ability to select the different features and benefits of products they want from different manufacturers, with the confidence that they will all be compatible with one another.



## No more need for adaptors

Dissimilar systems require additional components to adapt/couple one system to the other. This creates extra SKUs and adaptors can have many negative benefits, as pointed out by ISMP<sup>1</sup>, and they are also discouraged by the FDA<sup>2</sup>.

<sup>1</sup> ISMP. Transition adapters for ENFit syringes can defeat the purpose of ENFit itself. Institute For Safe Medication Practices. <https://www.ismp.org/resources/transition-adapters-enfit-syringes-can-defeat-purpose-enfit-itself>. Published September 21, 2017. Accessed July 15, 2022.

<sup>2</sup> Maisel WH. September 7, 2018 to: Manufacturers of Enteral Feeding tubes. FDA. <https://www.fda.gov/media/115846/download>. Published September 7, 2018. Accessed July 15, 2022.