

Quality Policy

Our objective is to ensure that all devices, meets customers and regulatory expectations for quality, safety and efficacy. Our commitment is to create an advanced Quality System applicable across the life cycle of each device through a continuous, in-depth and forward-looking monitoring of the medical devices regulatory environment.


Quality is fully integrated into our processes; it is seen as a joint responsibility of all personnel and it is achieved by combining the collective ambition and individual contribution at all levels of the organization.

The quality of our products is assured by the cumulative effect of the premises, equipment, materials, services and personnel involved directly and indirectly in the processes.

Our main target is to distribute and produce high quality medical devices and meet our customer's expectations. Therefore, we:

- Ensure patient safety through risk management, quality assurance of medical devices and compliance with In MDR 2017/745
- Seek for the continuously enhancement of customers' satisfaction
- Track and apply new innovative technologies
- Maintain and continuously improve the effectiveness of our Quality Management System according to the standards of ISO 9001:2015, ISO 13485:2016/A11:2021, ISO 14971:2019 and ISO 37001:2016 in every aspect of business processes.
- Invest on the continuous improvement of our personnel through continuous training.

The management and employees of KeV Medical Imaging are committed to a continuous effort to improve quality with the ultimate goal of maximizing customer satisfaction.

General Manager 
KeV Medical Imaging
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