Quality Policy

Wismed is committed to the implementation of robust processes for effective medical device management over the entire life cycle of our products. We employ clinically proven and well-controlled quality practices during product development, manufacturing, verification, validation and packaging, both at the company and our 3-rd party manufacturing partners.

Wismed Executives, Management and Staff strive to achieve this quality commitment through

- Management of organization, along with established quality objectives and defined responsibilities for their fulfilment
- Establishing, applying, maintaining and continual improvement of effectiveness of Quality Management System according to ISO 13485:2016
- Directing the business to satisfy clinical requirements and expectations
- Complying with applicable regulatory and governing body requirements
- Proactive risk management of medical devices and ensuring highest standards of device safety
- Continual enhancement of customers' satisfaction and patient safety
- Training and educating of staff on new technologies
- Careful selection of suppliers and subcontractors

The framework for setting quality objectives is defined in the Quality Manual.

I, the undersigned Chief Executive Officer will ensure that the Quality Policy is communicated to all persons working for or on behalf of the organization and making it available to the public.

Pawel Wisniewski, Chief Executive Officer

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