


Doc Number SAHPP-QA-01	Title <b>Quality Policy</b>	
Revision 01	Effective Date: 14 February 2021	

## SAHPRA QUALITY POLICY STATEMENT

The South African Health Products Regulatory Authority (SAHPRA) is responsible for the:

- Regulation of health products intended for human and animal use;
- Licensing of manufacturers, wholesalers and distributors of medicines, medical devices, radiation emitting devices and radioactive nuclides; and
- Conduct of clinical trials.

Our organisation is committed to providing the highest standard of regulatory services to all customers. This will be achieved through the use of guidelines, policies, processes and procedures, which guarantee an effective quality management system that conforms to the requirements of ISO 9001:2015 and other applicable requirements.

Continual improvement, reliable service and meeting customer requirements inspire all our efforts in ensuring quality, safety and efficacy of all regulated health products used in South Africa.

SAHPRA shall therefore commit adequate required resources for implementing, maintaining and continually improving the quality management system to achieve set objectives; and maintain an adequate workforce that is trained, motivated and empowered to achieve results.

Actions required for Top Management includes:

- Supporting the QMS and actively promoting the agenda by defining and communicating the Quality Policy
- Encouraging the establishment and monitoring of Quality objectives.
- Ensuring appropriate resources are available.

This quality policy shall be communicated to all SAHPRA employees and shall be reviewed annually for continuing suitability.

### Signed by the Executive of SAHPRA

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