# MEDICINES CONTROL COUNCIL

DEPARTMENT OF HEALTH
Republic of South Africa

MEDICINES CONTROL COUNCIL

## WORKSHOP BETWEEN THE REGULATOR AND INDUSTRY ON MEDICAL DEVICES

### TO ALL APPLICANTS

Following the promulgation of the Medicines and Related Substances Amendment Act (2015), Act 14 of 2015, the Department of Health intends to host a workshop on transitional arrangements for medical devices.

The workshop will address issues pertaining to the Roadmap for the Regulation of Medical Devices, Section 21 Authorisation, QMS requirements, Licensing and Vigilance.

The workshop is scheduled as follows:

**Date:** 13 July 2017 / 14 July 2017

**Venue**: Impilo Boardroom, Podium level, Civitas Building, 42 Thabo Sehume Street, Pretoria

**Time:** 09h00 – 16h00

You are invited to attend on one of the scheduled dates.

Please note that participation from stakeholders is limited to a maximum of two (2) representatives, from each organisation/institution and the invitation is targeted at Authorised Representatives and personnel responsible for Quality Assurance/Quality Control.

Kindly confirm your availability to participate via e-mail to Ms Puleng Nkosi at Puleng.Nkosi@health.gov.za by no later than 01 July 2017.

*Enquiries:*

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**DR JC GOUWS**
**REGISTRAR OF MEDICINES**