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 Act, 1965 (101 of 1965) in June 2017, to include the regulatory oversight of Medical Devices; the Department of Health intends to host a Medical Device Workshop.

The workshop will address the licence application process for wholesalers and is scheduled as follows:

Date: 19 February 2018

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

Time: Please note that three sessions will be held:

Session 1:	**09h00 - 10h00**	**Q & A Session:**	**10h00- 10:45**
Session 2:	**11h15- 12h15**	**Q & A Session:**	**12h15 – 13h00**
Session 3:	**14h00- 15h00**	**Q & A Session:**	**15h00- 15h45**

The information presented in each of the sessions is the same. Please be advised that you are therefore invited to attend **one** of the three sessions.

Please note that participation from stakeholders is limited to a maximum of one (1) representative from each organisation/institution and the invitation is targeted at Authorised Representatives and personnel responsible for Quality Assurance/Quality Control.

Kindly confirm your attendance and selected session number via e-mail to Ms Puleng Nkosi at Puleng.Nkosi@health.gov.za by no later than12 February 2018.

Enquiries:
Tel: +27 12 395 9402

MS P NKAMBULE
Acting Cluster Manager: Food Control, Pharmaceutical Trade & Product Regulation