SACRA Clinical Trials Capacity Building Workshop in collaboration with FARMOVS, SOCM and the Free State DOH

06 November 2019 FARMOVS Clinical Research Centre, UFS

	Topic	Presenter
09:30	Registration	FARMOVS
10:00	Opening and Welcome	Dr Rita Nathan, DOH Prof Nathaniel Mofolo, UFS Dr Vathi Papu-Zamxaka
10:05	SACRA Introduction and Aim of Workshop	Dr Nyda Fourie SACRA EXCO
10:15	 Clinical Trials Landscape – Setting the Stage Clinical Research vs Clinical Trials Stages of Drug Development and Importance or RCT Clinical Trials – Global Activities vs African, National, Provincial 	Dr Michelle Middle SACRA, FARMOVS
10:45	 How do Clinical Trials Work? Stakeholders, Approval Process, Monitoring and Auditing Key Aspects to Site Selection Pharmacovigilance, Post Trial Access and Insurance 	Ms Sue Bailey, IQVIA Country Manager
11:15	 The Patient Factor What happens to a Patient in a Clinical Trial? Patient Benefits and Risks Handling of Adverse Events, Insurance Recruitment 	Prof Makoali Makotoko, DOH, UFS Dr Thabiso Mofokeng, DOH, UFS Dr Riana van Zyl, DOH, UFS
12:00	Ethics in Clinical Trials	Dr Nyda Fourie SACRA, IATROS
12:30	Q & A	SACRA EXCO
12:45	Lunch	
13:15	Economic Aspects of Clinical Trials - Benefit to Health Sector - Opportunities for South Africa - Learning from Others	Ms Bicky Mthombeni SACRA, SANOFI
13:45	3Cs in Clinical ResearchCollaboration, Capacity Building and Career Opportunities	Dr Vathi Papu-Zamxaka SACRA, SYNEXUS
14:15	Q & A and Wrap-up	Dr Michelle Middle SACRA, FARMOVS
14:45	Refreshment Break	
15:00	Working Group Breakaway - Collaboration in Clinical Trials in the Free State, Create the Opportunity and Uncap our Potential, Including Risk Analysis	Prof Nathaniel Mofolo, UFS Dr Kwazi Ndlovu, DOH, UFS Dr Lizzy Tabane, DOH Dr Claire Barrett, UFS
16:20- 17:00	Working Group Feedback	Rapporteurs

Please RSVP to Michelle Middle Michelle.Middle@farmovs.com







