

DAY 1: 22 September 2010											
Time	Duration	Topic	Speaker	Presentation level	Room	Duration			Presentation level	Room	
08h30-08h45	10min	Welcome - Opening of conference - SACRA chair	Savi Chetty-Tulsee		OR 2						
		Using your med-bay card			OR 2:						
08h45-09h45	60min	Plenary: Keynote speaker: The Caprisa 004 trial experience	Prof Salim Abdool-Karim		OR 2						
09h45-10h15	30min	NHREC - The Ethics Committee Accreditation Process - an update: challenges, the achievements, lessons, current status in this process.	Marzelle Haskins	Core	OR 2	30min	Impact of speaking books in educating communities in clinical trials	Zane Wilson	Core	Auditorium in the International Centre:	
TEA BREAK					OR1	TEA BREAK					OR2
10h45-11h30	45min	Drug Counterfeiting: New Actions and Initiatives	Griffith Molewa	Intermediate	OR 2	45min	Informed Consent Translations - Do subjects want to sign these	Suhella Abdul Karim	Core	Auditorium	
11h30-12h15	45min	Quality of conduct of South African Clinical trials - the inspectorate's perspective	Laurette Bonthuys	Intermediate	OR 2	45min	Creating a collaborative environment at monitoring visits - a global perspective	Savi Chetty-Tulsee	Intermediate	Auditorium	
12h15-13h00	45min	Clinical Trials in Sub-Saharan Africa	Dr Yuka Manabe	Intermediate	OR 2:	45min	When the guinea pig says Yes! - Defining vulnerability in the Informed Consent Process	Dr Lawrence Reiter	Intermediate	Auditorium	
LUNCH					OR1	LUNCH					OR1
14h00-15h00	60min	Expanding South Africa's Reach into Africa: Lessons learned and Successes	Viola Mari -Raubenheimer (Quintiles), Judith Ramuthaga (Sanofi Aventis), Nathaniel Ramuthaga (Pfizer), Catherine Lund (OnQ)	Intermediate	OR 2	60min	Fraud and Misconduct in Research	Prof. Lesley Burgess	Intermediate	Auditorium	
TEA BREAK					OR1	TEA BREAK					OR1
15h30-16h30	60min	Improving clinical research productivity with strategic partnerships	Dr Bets Breedt, Dr H Hanekom	Intermediate	OR 2	60min	Writing monitoring visit reports	Nicola Main, Retha Britz	Intermediate	Auditorium	
16h30-17h15	45min	TB - recent changes in methodology for diagnosis of MTB sensitivity	Dr. Jessica Trusler	Intermediate	OR 2	45min	Supervisory responsibilities of investigators	Teresa Scanes	Intermediate	Auditorium	
17h15-18h15	60min	POSTER SESSION			OR1	60min	POSTER SESSION			OR1	
18h15-20h00	Networking evening sponsored by Pfizer									King Shaka in Domestic	
DAY 2: 23 September 2010											
08h30-09h00	Plenary session: Recognition awards		Essack Mitha		OR 2:						
09h00-10h00	60min	Legal implications of informed consent	Neil Kirby	Intermediate	OR 2	60min	FDA warning letters: Lessons Learnt when things go wrong	Leigh Howes	Intermediate	Auditorium	
TEA					OR1	TEA					Auditorium
10h30-11h15					OR 2:	45min	Biomarkers in drug development	Dr Sandy Evans	Intermediate	Auditorium	
11h15-12h00	90min	Patient recruitment workshop	Claus Mark Nielsen	Intermediate	OR 2	45min	Site Selection Process	Joan Selema	Intermediate	Auditorium	
12h00-13h00	60min	Factors that influence effective patient recruitment and the impact of provision of a recruitment budget	Dr Luthando Adams, Dr Sanet Aspinall	Intermediate	OR 2	60min	An Internal Quality Control system in clinical trials: does it make a difference?	Thelma Leopeng	Intermediate	Auditorium	
LUNCH					OR1	LUNCH					OR1
14h00-15h00	60min	Ethical issues in the conduct of Clinical Research specifically in the therapeutic area of HIV and TB in South Africa	Belinda Alport	Intermediate	OR 2	60min	Monitoring in Africa? - everything you need to know	Teresa Scanes	Core	Auditorium	
TEA					OR1	TEA					
15h30-16h30	60min	Plenary session powered by Mini Bronze sponsors: "Big Mamma and the Horse"	Vusi Thembekwayo		OR 2						
16h30	CLOSE										

Poster Presentations:

OR1

1	Interpreting FDA warning letters	Tharnija Lalbahadur
2	Duties and responsibilities of a study coordinator	Ilse de Necker, Heleen Pieterse, Christina Naude, Lesley Burgess
3	The training needs of a study coordinator	Jenny Snyman, Elfreda Nel, Lesley Burgess
4	Informed Consent - a process not an event	Marli Terblanche, Lesley Burgess
5	A framework of reference for an informed decision to a valid consent	Joan Selema,, Ashina Gokul, Cecilia Nomsobo

6	Assessment of time orientation of CRAs in the pharmaceutical industry	Koretha Ras
7	Teaching clinical research to all	Carmelita Jeftha
8	Survey of SSC fees	Zelna Vermooten
9	Research on Tuberculosis in Cameroon - A literature review	Dr Armand Nkwescheu
10	Regulatory barriers in clinical research	Kirti Narsai