

## CLINICAL INVESTIGATION TRAINING PROGRAM (CITP)II

2<sup>ND</sup> OF OCTOBER, 2010  
AMMAN MARRIOTT HOTEL, JORDAN

Time	Subject	Speaker
8:30 - 9:00	Registration & morning Coffee	
9:00 - 9:30	Opening Speech & Welcoming Remarks	Acting Director General Dr. Laila Ghazi Jarrar
		USAID
		Hanan Sboul
9:30 - 10:00	Clinical Studies In Jordan	Dr. Wafa Al-Khateeb
10:00 - 10:30	Good Clinical Practice & parties involved	Dr. M. Amer Al-Khatib
10:30 - 11:00	The Roles and Responsibilities of CORS in Clinical Studies or Bio-studies	Dr. Mu'taz Sheikh Salem
11:00 - 11:30	Principal Investigator	Dr. Yacoub Irshaid
11:30 - 11:45	Coffee Break	
11:45 - 12:15	Source Data & Source Documents	Dr. Muatsem Al-Ghazawi
12:15 - 12:45	Informed Consent Form & Ethical consideration	Dr. Abla Al-Bsoul
12:45 - 1:15	Investigational Product	Dr. Saleem Al-Mahrouq
1:15 - 1:45	Adverse Drug Reactions	Dr. Yacoub Irshaid
1:45 - 2:45	Lunch	
2:45 - 3:15	Institute Review Board : Roles & Responsibilities	Dr. Abla Al-Bsoul
3:15 - 3:45	Compliance with Good Clinical Practice & Regulatory Authority	Dr. Wafa Al-Khateeb
3:45 - 4:15	Trial Design & Statistical Aspects	Dr. Muatsem Al-Ghazawi
4:15 - 4:45	Quality Control & Quality Assurance in Clinical Studies	Dr. Esam Al-Shoubaki
4:45 - 5:00	Discussion & closing	