

## INSTITUTIONAL RESEARCH – ETHICS COMMITTEE (IR-EC)

## PROGRESS REPORT FORM

Sponsor Protocol Number:		
*Protocol Title:		
*Principal Investigator/s:		
Contact Details: Telephone Number:	Fax:	Email
Type of Research: [ ] Clinical Trial	[ ] Clinical Research [ ] Sociobehavioural	[ ] Public Health [ ] Laboratory Research
Descritor and Chatra	to	
Total number of patients screened: Total number of patients randomized, if application Total number of screen failures: Total number of patients discontinued: Total number of patients completed the study:	prov	
Total number of SAEs:  Total number of Unexpected Event/ Unanticipa	•	eport, if necessary.
Total number of Protocol Deviation/ Violation:	provide re	eport, if necessary.
Remarks:	provide re	eport, if necessary.
Data		

To be filled-up by the IR-EC:
Reviewer/s Recommendations:
Changes to the protocol recommended? No Yes Comments :
Changes to the Informed Consent Form and/or other documents recommended? No Yes Comments :
Reviewer's Name : Signature: Date :
IR-EC Final Action:
** Request an amendment to the protocol or the Informed Consent Form Request further information Suspend or terminate the study Take note and no further action is needed Others:
Type of review: Expedited review Full board review  Date of meeting:
Name of Member/Secretary: Signature: Date: