

## **Clinical Trial Evaluation Report**

Name of the Trial:		-
Study Number:		
Phase of CT:		
<b>Study Population:</b>		
Designation Number of MOII.		
Registration Number at MOH:		
Date of Submission to MOH:	Received	on:
C	Received	on:
Date of Submission to MOH:	Received	on:
Date of Submission to MOH:  Sponsor:	Received	on:
Date of Submission to MOH:  Sponsor:  Country of Origin:	Received	on:
Date of Submission to MOH:  Sponsor:  Country of Origin:  Name of Applicant:		on:

Name of IMP:		
Type of IMP:		
Therapeutic Benefit:		
Objective of the Study:		
Market Authorization of IMP:	<b>Commercial Name:</b>	
Other Countries Authorizing the Trial:		
Not authorizing the trial:		
Importation of Other Products/Drugs:		
<b>Documents Submitted:</b>		
√ Importation Request		
√ Invoice		
√ Certificate of Analysis		
√ Certificate of Release		
$\sqrt{\text{Principle Investigator Letter}}$		
√Pharmacy Request		
√ Protocol		
$\sqrt{ m IRBs}$		
√ Investigator's Brochure		

√GMP Certificate

 $\sqrt{\text{Registration of CT in Country of Origin: }}\sqrt{\text{Annex 4}}$ 

 $\sqrt{10}$  Page Summery on: Pharmacology & Therapeutic Benefits, Pharmacokinetics, Adverse events, Results from previous studies

**Comments:** 

Rasha Hamra