CRF Design Changes

Royal Papworth Hospital
NHS Foundation Trust

This is for changes that are made after the study has been approved to go into production.

Please send completed form to your Trial Coordinator via email or via safehaven fax 01480 364356.

Site Name (if applicable): _____ Study Title: _____ Study Number: _____

CRF Name	Change Requested	Reason for Change	Notes
Example	*Add options for DI,OM and IM to lesion site.*	*Had not expected patients of this type, not specified in protocol.*	*Added, made version 1.3, moved previous subjects to this version.*

Requested by: Signature: Date:

Approved by (CPM): Signature: Date:

Amended by: Signature: Date: