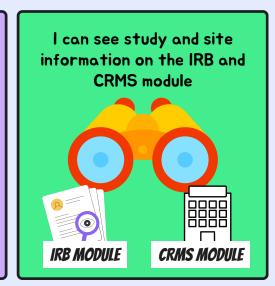
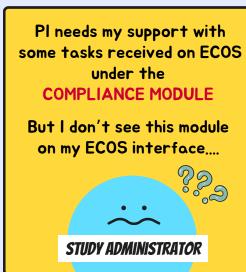
National Healthcare Group

View and Edit Access Rights

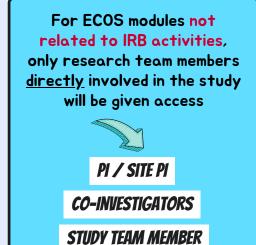
ECOS - ATTENTION TEAM MEMBERS!













CRMS ROLES

STUDY TEAM MEMBER

Site personnel directly involved in the research conduct e.g. CRCs, Study Nurses, Pharmacists, etc

STUDY ADMINISTRATOR

Site personnel **NOT directly involved** in the research but provides administrative support only, e.g. Dept Manager providing research administrative support

STUDY SPONSOR

Sponsor/CRO personnel, e.g. CRAs

View & Edit Rights

Modules Supporting IRB Submissions

User Role	IRB Module					
	Forms ¹	Study Information	User Authorisation List (UAL) ²	Site Information ²	Milestones ²	Participants ²
Study Team Member (STM)	View & Edit	View & Edit	View & Edit	View & Edit	View & Edit	View & Edit
Study Administrator (SA)	View & Edit	View & Edit	View & Edit	View & Edit	View & Edit	View & Edit
Study Sponsor (SS)	View & Edit	View & Edit	View & Edit	No Access	No Access	No Access

¹ Initial Application, Amendments, Study Status Report (SSR), Study Deviation/ Non-Compliance Report (DNC), Serious Adverse Event Report (SAE), Unanticipated Problems Involving Risks to Subjects or Others Report Form (UPIRTSO), Other Study Notifications Report Form (OSN).

Modules Supporting non-IRB related Activities

User Role	Compliance Mod	Compliance Module (PISAF)	Standing Database Module	Monitoring Module	Audit Module		
	Study Deviation/ Non- Compliance Report (DNC)	Serious Adverse Event Report (SAE)	Modules To Be Launched On ECOS				
Study Team Member (STM)	View & Edit ³	View & Edit ³		KIN	N		
Study Administrator (SA)	No Access	No Access		PROGRESS			
Study Sponsor (SS)	No Access	No Access					

³ Access to edit is granted by RI Secretariat only

ACCESS THE ECOS MODULES USER GUIDES HERE





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² Access is site-specific