

CLINICAL TRIAL – ON-SITE MONITORING

On the request of the **PI/Head of Unit at CRCO**, the **On-site Monitor** and **Quality Assurance Manager of CRCO** examine whether all the Protocol requirements are satisfied at the Site to initiate the trial.



During the conduction of the trial, the **On-site Monitor** administrates all the information from the incoming **follow-up letters**.

During the conduction of the trial, the **On-site Monitor** supervises the delegated study coordinators' work.

During the conduction of the trial, the **Quality Assurance Manager** makes sure the **SOP's** are appropriate and in use, and are related to the protocol.



If the **PI/Head of Unit at CRCO** requires, the **On-site Monitor** supports the **preparation** of the Site for an Audit/Inspection.



Quality Assurance Manager supports the **PI** during the close-out procedure and helps to archive all related documents of the trial.