Auditor

For some studies, notably clinical trials, a monitoring role may be required. Study monitors are responsible for reviewing and verifying data at the study sites. They only require the ability to review data and raise queries. They do normally need to edit patient data so should have the following access:

- Data export - Full Data Set
- Add/Edit Reports
- Stats & Charts
- Open, close and respond to queries. (Note that Monitors can raise queries even if they cannot edit records).
- Data Entry Rights - Read only access to all forms