A-CTO functions available

Application/approval
Health Authorities
Ethics Committees
Data Protection Agencies
Amendments

Pharmacovigilance
SAE reports
SUSAR reporting
Annual safety reports

Monitoring
Risk Assessment
Monitoring Plan
Organisation
Contact to monitors

Trial organisation
Participating centers
Contracts with sites
Investigator site files
Investigator meetings etc.

Study drug
Organisation of delivery
Ordering

Economy
Budget
Handle payments

Data Management
CRF
Database
Queries

Patient inclusion
Registration
Randomisation

Biobanking
Collection of samples
Coordination of correlative studies