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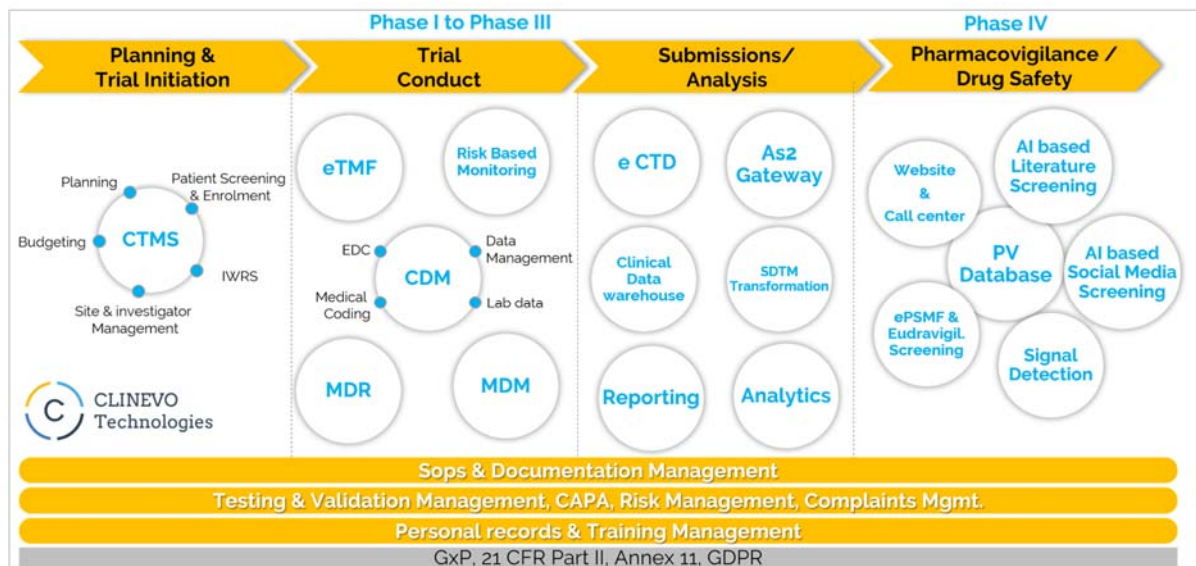
Clinevo-Cliniminds Health Sciences Technology Training Partnership Agreement

Cliniminds is pleased to announced the agreement with Clinevo Technologies to provide hands on training to Cliniminds Students on various health sciences technology platforms.

Clinevo Technologies is an ISO certified company headquartered in Bangalore, India having offices in US and Europe specialized in developing robust technology solutions for Clinical Trials and Pharmacovigilance.

Clinevo is committed to delivering the most efficient and practical end-to-end solutions with HIPAA, GXP, CSV, 21 CFR Part 11, Annex 11 and GDPR guidelines.

Below are some of the Clinevo Platforms which are used by 50+ global Pharma and Biotech companies who do submissions to global regulatory agencies including USFDA, EMA, SFDA, DCGI,etc.



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Please visit www.clinevotech.com for more information.

Electronic Trail Master File (eTMF):

Clienvo eTMF is a dynamic trial master file in electronic format for organizing and storing documents, images, and other digital content of clinical trials.

Clienvo eTMF meets the regulatory requirements including Digital content archiving, Security, Access control, Change controls, Audit trails, 21CFRPart11/Annex11, GDPR and GxP validation.

Clienvo eTMF has,

- Inbuilt and configurable DIA TMF reference model
- File Plan and Milestone setups for tracking the study progress and missing documents
- Configurable lifecycle management for documents

- Electronic signatures
- Real time dashboards and reports on Compliance, Completeness, Quality and Timeliness
- Bulk uploads, Indexing, Notifications, and many more.

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Clinical Trial Management System (CTMS):

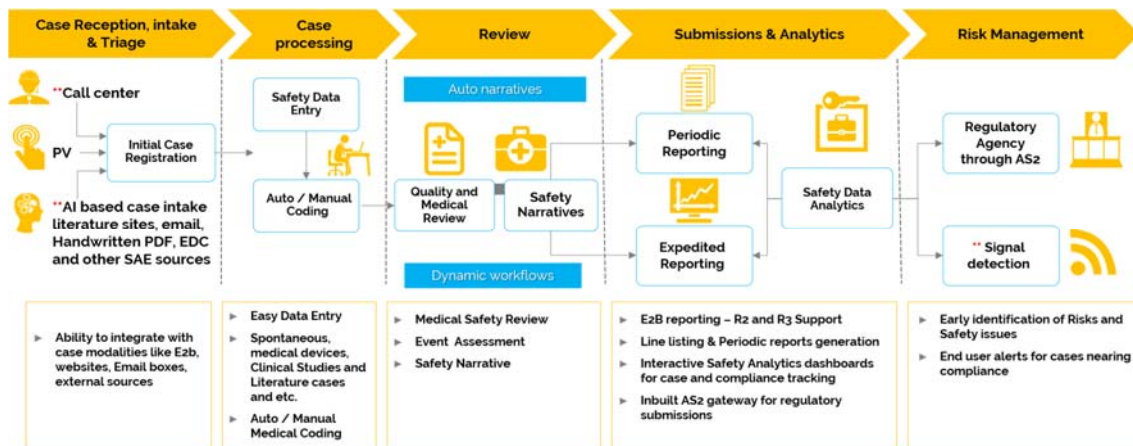
Clinevo CTMS: An end-to-end Clinical Trial Management System with below features.

- Site Monitoring Visits - Scheduling, Approvals, SV report generation, etc
- Milestones - Scheduling, Tracking and Closures
- Tasks - Scheduling, Tracking and Closures
- Trainings - Scheduling, Tracking and Closures
- Subject Visit scheduling and Tracking
- Site and Investigator Billing and Payments
- Inventory - IP
- Safety Reports Tracking
- Protocol Submissions and Deviations Tracking
- Study Documents and eTMF Integration
- Administration, Master Data Management & Many more

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Pharmacovigilance Database:

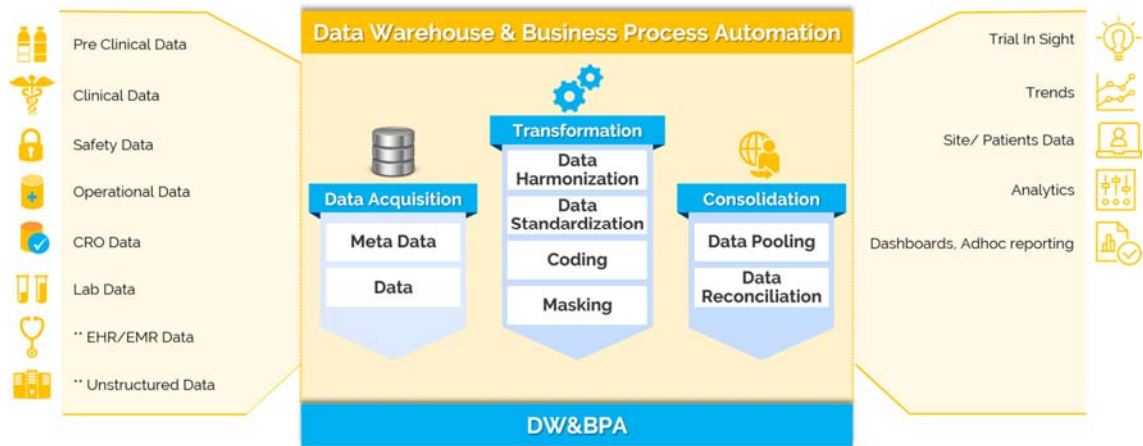
Clinevo Safety is a cloud-based, end-to-end Pharmacovigilance (PV) / Drug Safety Database which provides PV Intake, Case processing, AI, Analytics, Submissions /AS2 gateway, and Safety signals capabilities under one platform.



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Clinical Data Warehouse:

Clinevo Data Warehousing and Business Process Automation Console is a secured, regulatory compliant Clinical Trials Data Warehouse to Acquire, Store, Transform, Consolidate and Report diverse data from clinical trials in one place and AUTOMATE any of the manual, cumbersome business processes. This platform can enable companies to perform Cross Study Analysis, Data mining, Predictive Analytics, etc.



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