ATL-SSSMS 3.0.0

Stability Sample Schedule Management System
With analytics

USFDA-UKMHRA-GMP certified product

www.anveshatechlab.com

TangentTech
Infosoft Private Limited

Modules

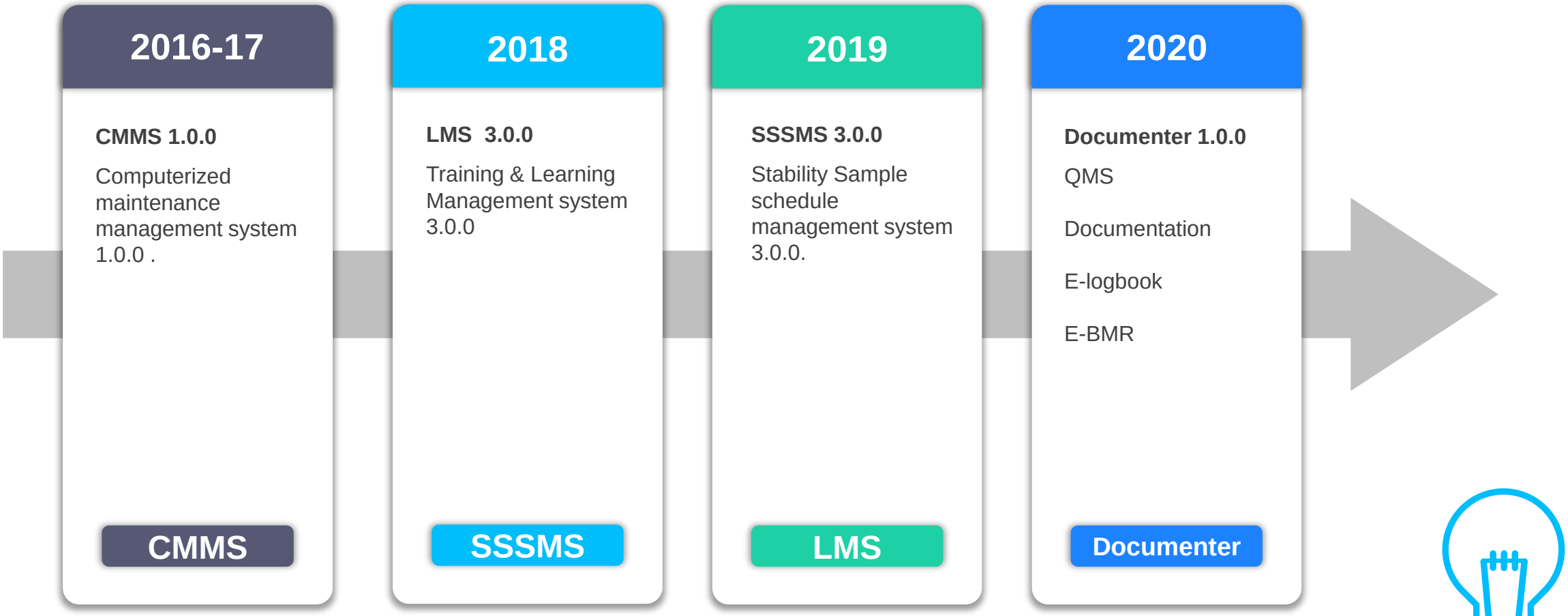
01 Protocol Preparation

02 Sample Charging

03 Sample Withdrawal

04 Lab Test Entry & COA

TechLabPharma -Timeline



Meet Our Team

Excellence Behind the product

Anvesha Tech Lab Private Limited

Develop & Conceptualized by

Tangenttech
Infosoft Private Limited

Marketed & Validated By

Specifically designed keeping in mind the stringent rules and regulations that all the stake-holders in the Pharmaceutical Industry have to adhere to.

- Create Plans for Stability Study
- Sample charging against plans
- Timely withdrawal of samples charged for study
- All test results should be recorded and kept in an organized way so that results and cross-referencing reports can be called up at any given point in time for regulatory approvals and COA generation
- Sample Destruction to maintain protocol

- Barcode Stickers generation for samples while charging to chamber
- Trending & Chart of the Stability Study

Stability Modules

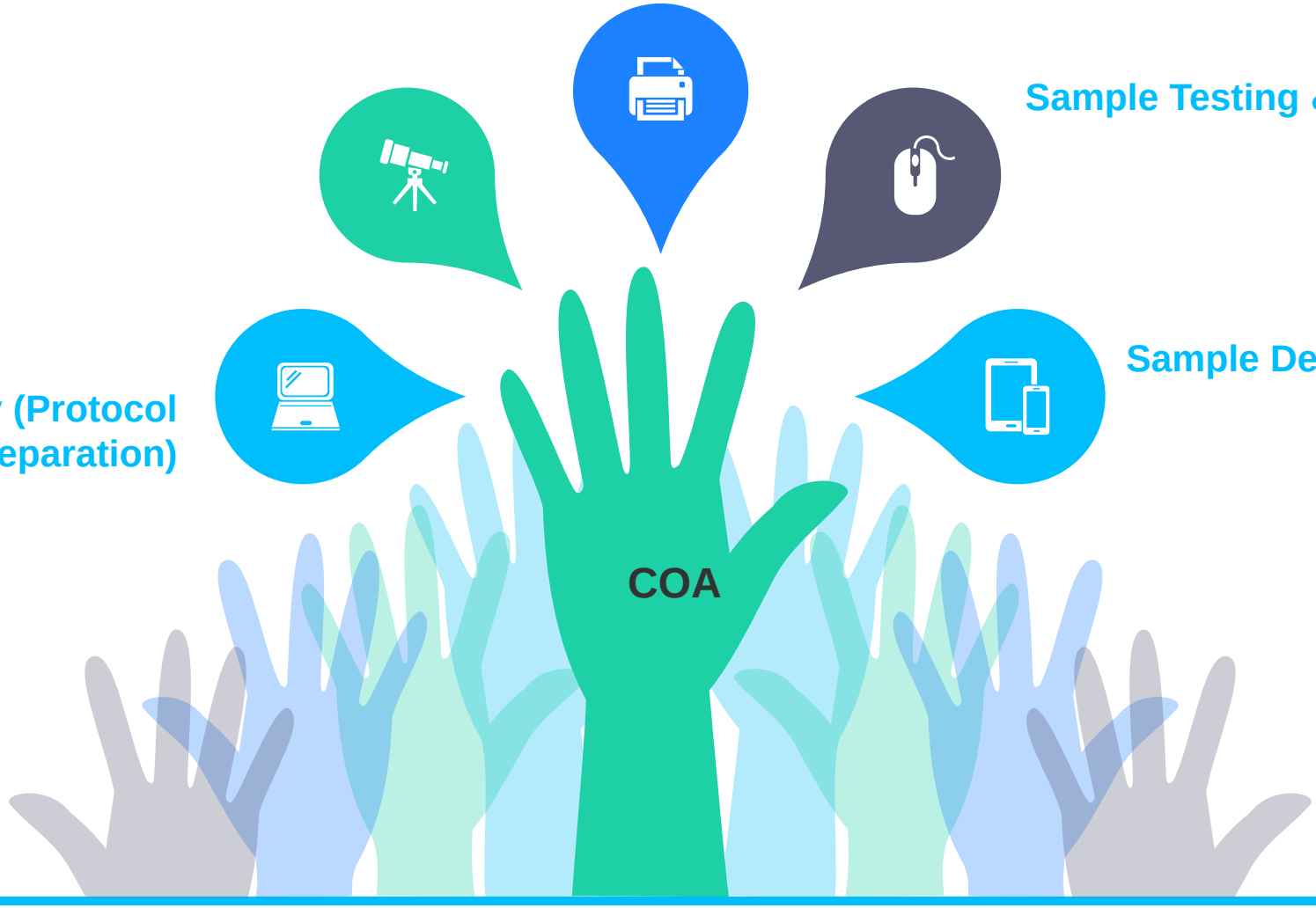
Sample Withdrawal

Sample Charging

Sample Testing & Entry

Planning of Study (Protocol Preparation)

Sample Destruction



Features of SSSMS 3.0.0

Dashboard

- Status dashboards
- Extensive Reporting
- Real – time data Trending

Alerts & Notification

- Authorization
- Daily –Weekly Withdrawals
- Out of range lab test results
- Many more

MIS

- COA
- Pending Withdrawals
- Sample Inventory
- Chambers Inventory
- Many more

Question Banks:

A huge set of questions with the relevant answers can be stored in the system to serve as a central paper-setter. Each employee will get a unique set of questions/the same set of questions in a scrambled order so that no malpractice takes place while conducting tests.

Benefits to the organization

- Completely automated solution prevents manual errors
- Handles sensitive data on a large number of samples efficiently.
- Keeps a log of each entry in the system which facilitates a transparent audit.
- The system is reliable, easily manageable and scalable according to your requirements.
- Provides extensive results, summary reports, exception reports and triggers alarms and predefined procedures to be undertaken at times of emergencies.
- Authorized access to data prevents breach of secrets and relevant test data on products.

ATL- LMSv3.0.0 USP

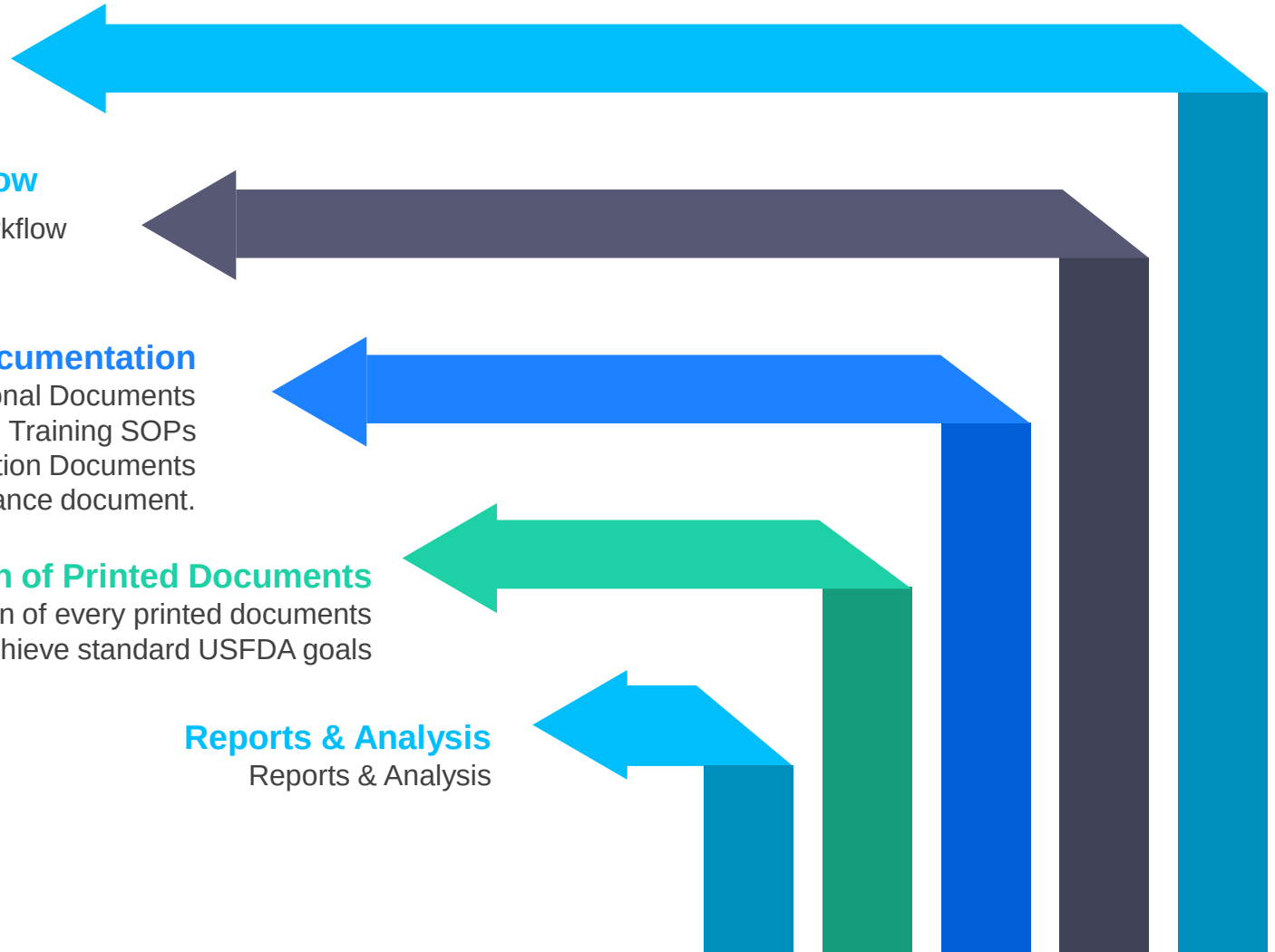
Checker-Maker System,
Inbuilt Preview-Review-Approval system

Configurable Workflow
Totally configurable workflow

Centralize Documentation
Employee Personal Documents
Training SOPs
Evaluation Documents
Classroom Attendance document.

Reconciliation of Printed Documents
Facility of reconciliation of every printed documents
help industry to achieve standard USFDA goals

Reports & Analysis
Reports & Analysis



Thank You

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