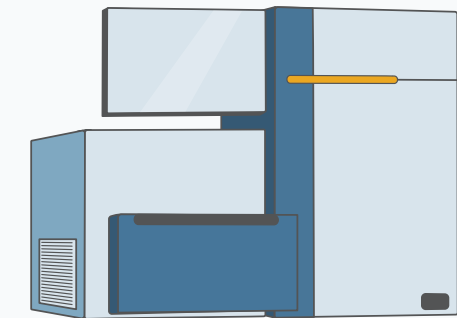


## Standardized Method

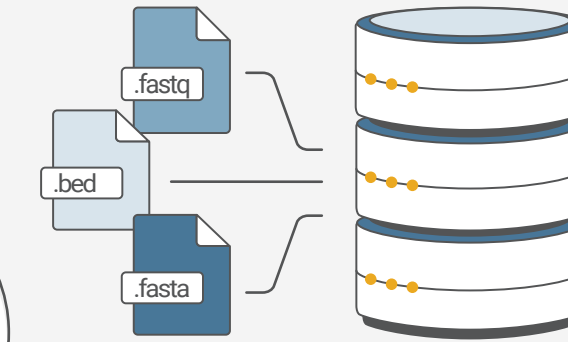
The NGS-based viral safety assays developed in-house need standardization to be successfully validated.



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## End-to-End Data Management

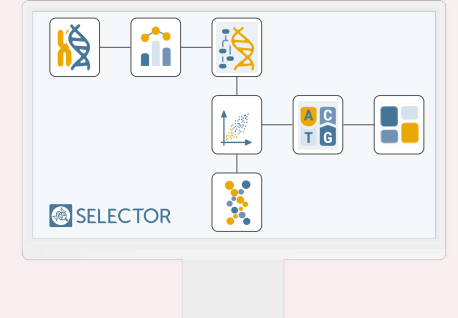
A validated platform for tracking samples across different instruments and assays is essential in a GMP environment.



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## Automated Workflows

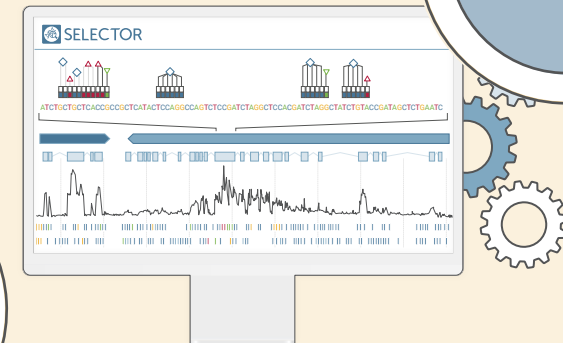
Automating standardized tasks increases productivity, reduces error rates, and is crucial in GMP environments.



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## Computer System Validation (CSV)

CSV enables the recording of traceable data trails that are securely stored and accessible at any time.



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## Genedata Selector

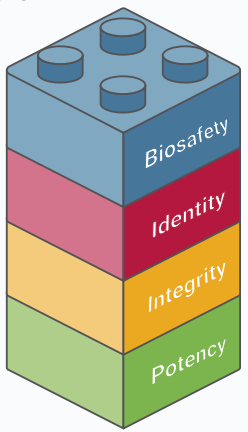
Connect the dots across R&D workflows.



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## Establishing Multi-Attribute Method

Monitoring multiple critical quality attributes (CQAs) is essential to ensure the safety and efficacy of bioterapeutics.



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## Breaking Down Silos

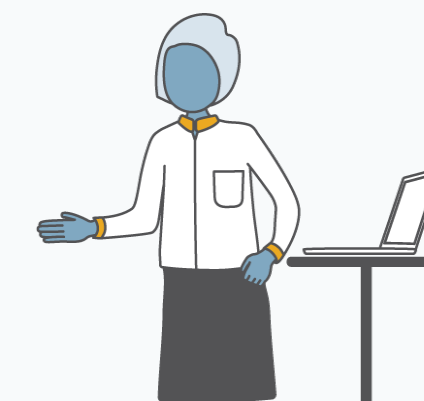
Establishing connectivity between sites and groups is instrumental in optimizing workflows across the organization.



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## User Training

The dynamic regulatory landscape and pressure to deliver life-saving drugs on time require quick and efficient user training.



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## Regulatory Compliance

There is an increasing pressure on companies to ensure compliance with EMA and FDA regulations regarding the use of NGS.



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