Clinical Informatics for Varied EHR Systems

State of Science and Gaps

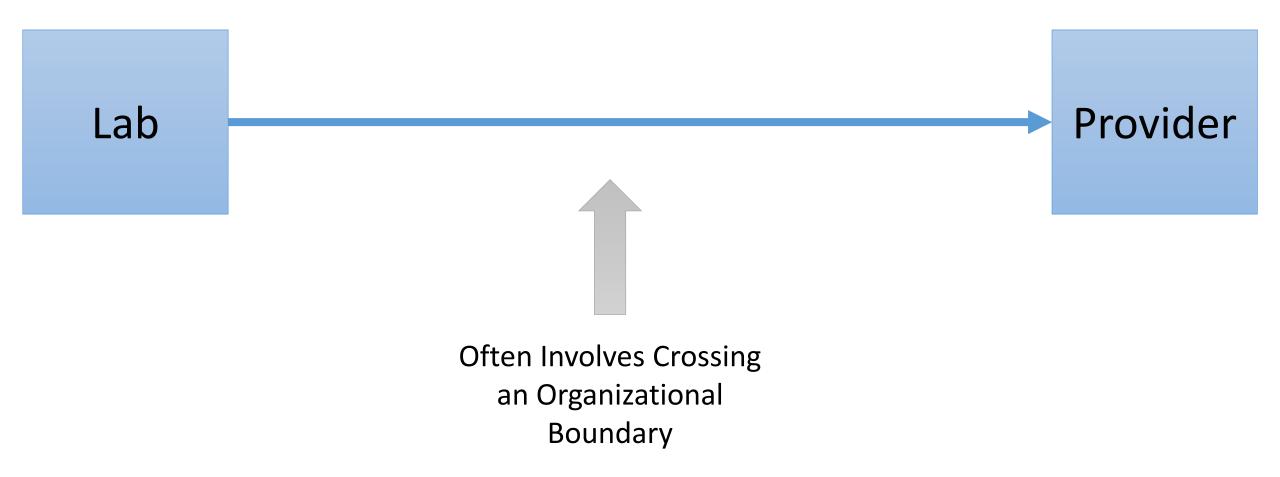
Clinical Decision Support

Federate Display Data Through Networks and Repositories

Clinical Decision Support

Federate Display Data Through Networks and Repositories

Genetic Results Transmission



Amount of Structure Patient Demographics

Types of Tests

Panels

Genotypes Variants (HGVS | Genomic Coor)

PDF

Interpretations (Variant|Gene|Test Level)

Exome Results Test Coverage

Genome Results Meta Data | Annotations Scope

Reported data

All Variants

Raw Data

Lab

Transfer Method Onotogies | Codes **Security**

Secure Conduit HL7 v2 | 3 Push Variants | Genes

BAA Required? Diseases **FHIR**

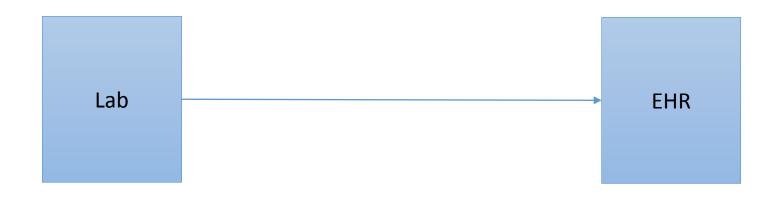
Authentication Batch File Transfer **Tests**

Results

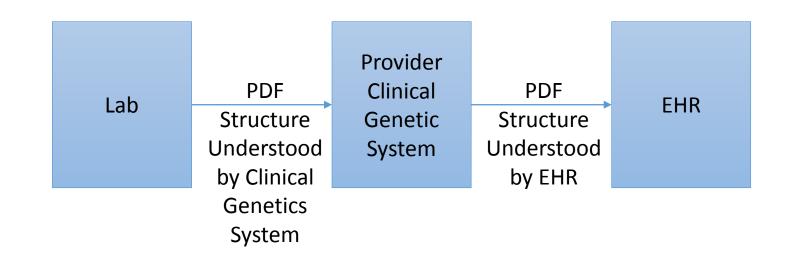
Provider

Provider EHR Ecosystem Options

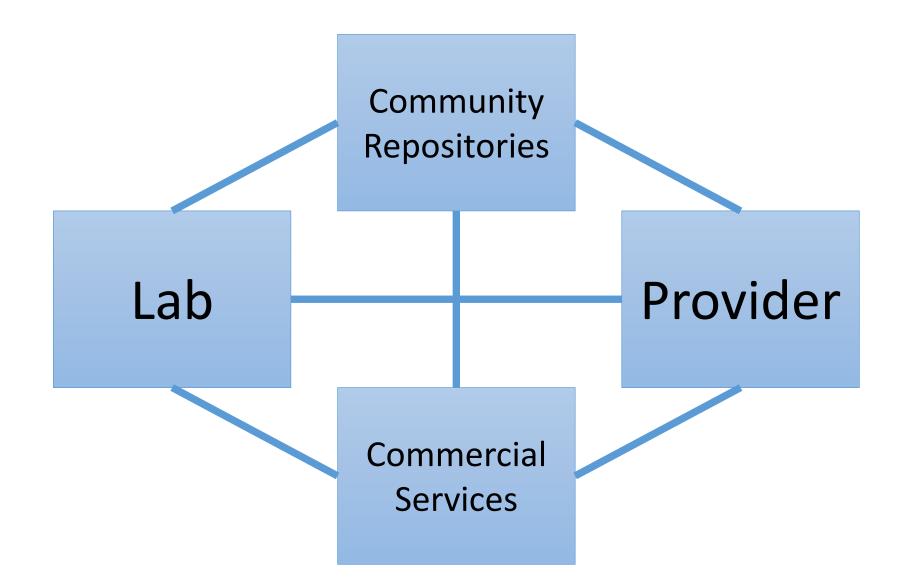
Direct to EHR



With Specialized Clinical Genetic System Intermediary



Potential Interpretation Locations

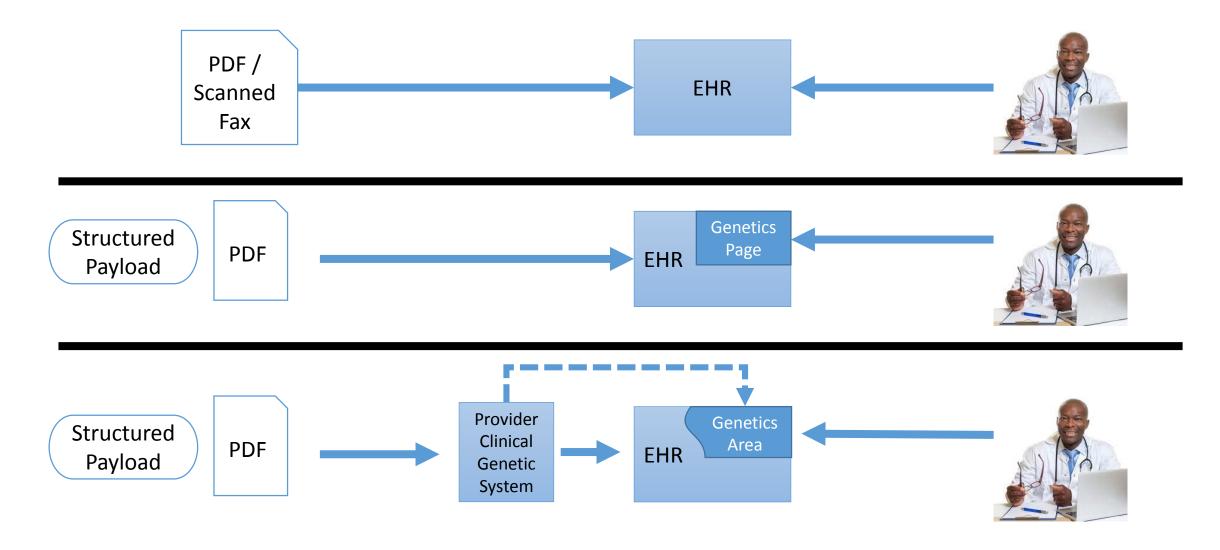


Clinical Decision Support

Display Data

Federate
Through Networks
and Repositories

Display Mechanisms



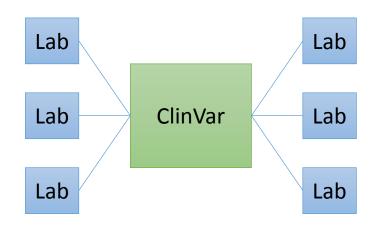
Ux Design Elements

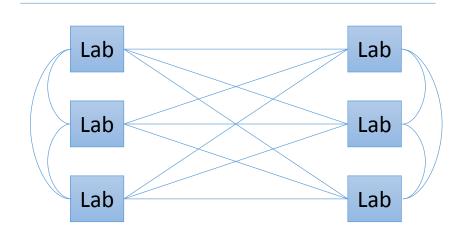
- Graphic Design
- Information Layout | Cognitive Optimization
- Screen Construction
- User Interface Evaluation

Clinical Decision Support

Federate Display Data Through Networks and Repositories

Federate Through Networks





Content

- Genetic Knowledge
- Aggregate Case Data
- Individual Case Data
 - Genetic Results
 - Phenotypes
 - Indications
 - OUTCOMES
 - De-identified / Identified

Clinical Decision Support

Federate Display Data Through Networks and Repositories

Clinical Decision Support

- Passive On Screen Information and Options
 - eResources and Infobuttons
 - Annotations
 - Calculators
 - Population Based Analytics
- Active Proactive Alerts
 - Pharmacogenomic Interventions
 - Knowledge Update Alerts
 - Genetic risk driven reminders