

#### **Precision Medicine for Health Systems**

### Enabling the Transformation of Healthcare Systems

November 14th, 2016



### Key Messages

- 1. A variety of data types are needed to enable precision medicine
- 2. These data types include:
  - a. Clinical data
  - b. Lab and genomics data
  - c. Imaging data
  - d. Sensor data
  - e. Patient reported data
- 3. The amount and size of these data sets will require them to be collected in a cloud computing environment
- 4. Hospital systems need a well thought out, systematic approach to developing the infrastructure to collect and analyze this data
- If done properly, this comprehensive data set can be used to drive insights and better clinical outcomes and improve drug development through both traditional analytics and machine learning

#### **Enterprises are experiencing a Digital Transformation**



Data on premise, hard to access, analyze and use

Productivity tools built for individual, local usage

IT focusing on **where** it computes



Data stored in cloud, simple to query

Machine learning drives deep, actionable insights

Collaborative, cloud based productivity applications

IT changing **how** it computes.

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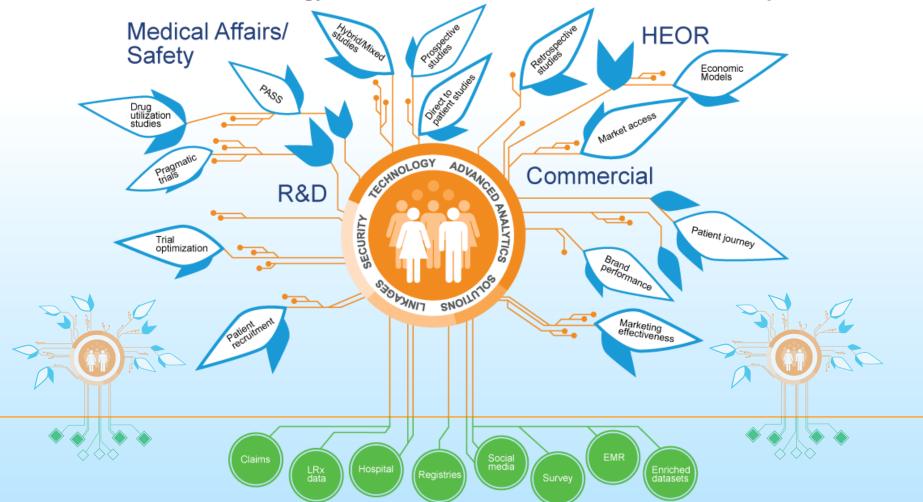
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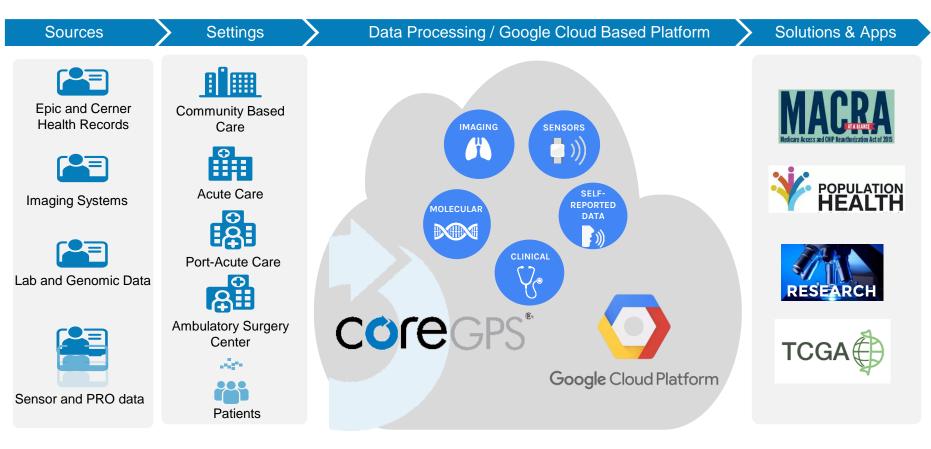
Collaborative, cloud based productivity applications

IT changing **how** it computes.

The same data and technology can be used for both clinical research and patient care

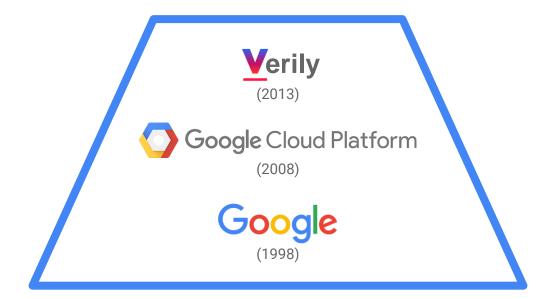


#### **Creating the Infrastructure to Support Precision Medicine**



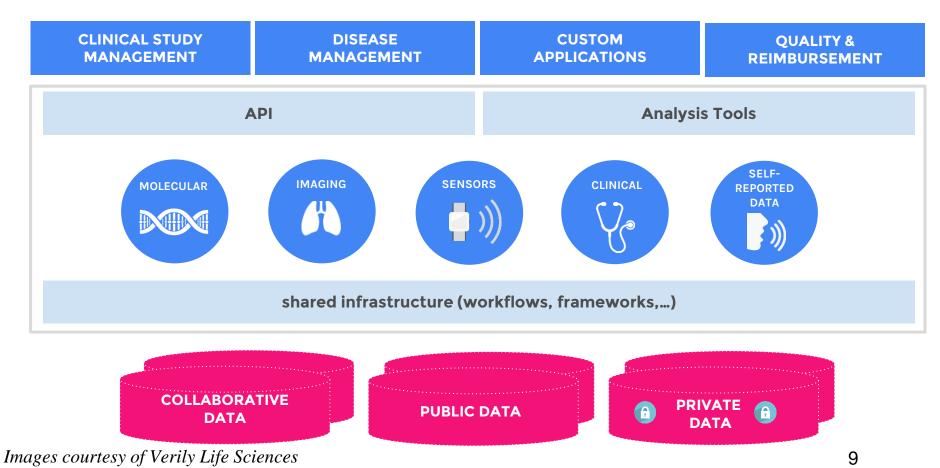
### Healthcare Capabilities

#### Standing on the shoulders of the Web

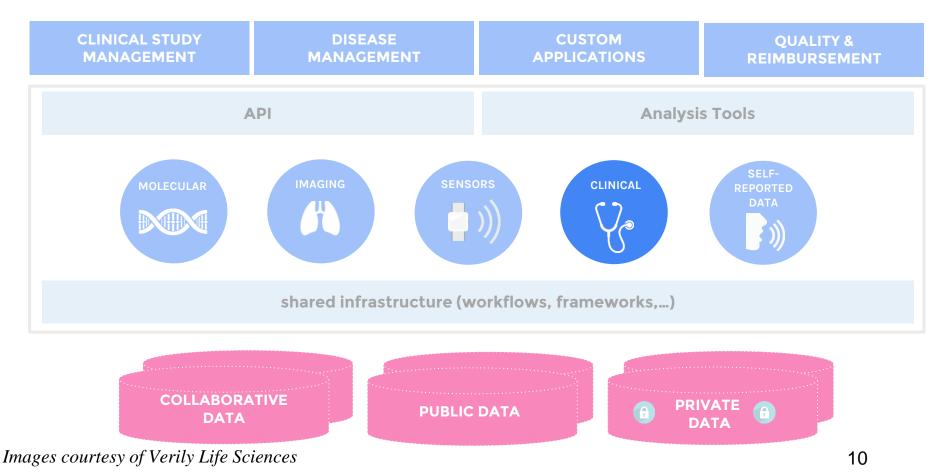


Building on Google's core infrastructure, data analytics, and machine learning.

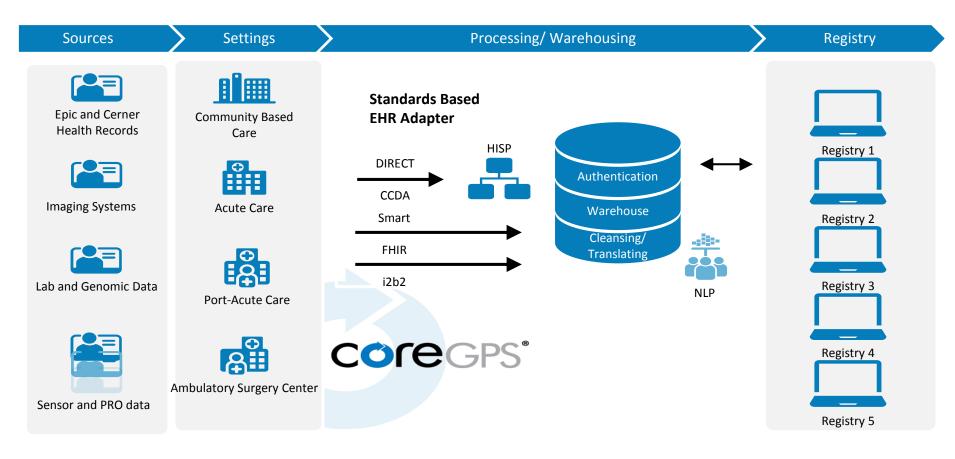
#### **Platform Vision**



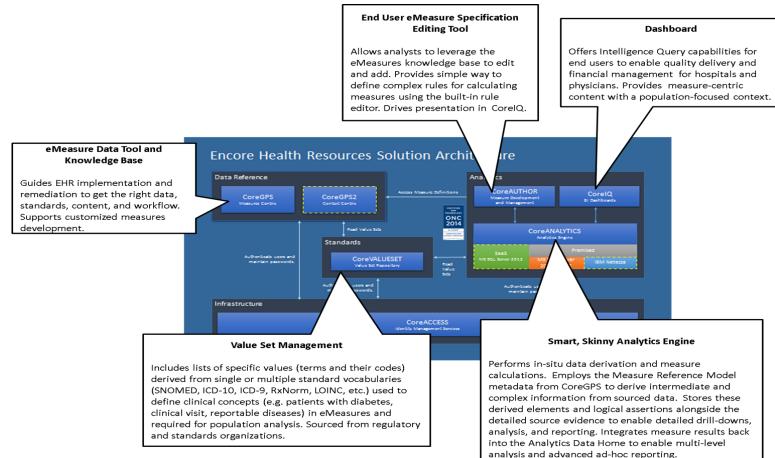
#### **Clinical Capabilities**



#### **Mapping Clinical Data**



#### **Getting Data Mapping Right - Core Suite Tools**



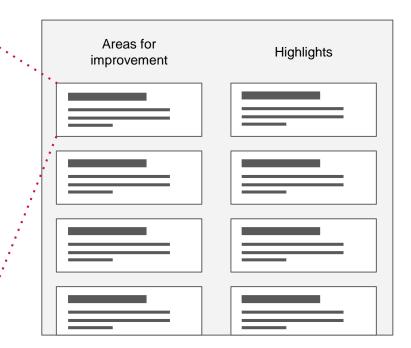
#### **Getting Data Mapping Right - Risk Assessment Process**

Risk Assessment Score Report														
Source Content from CMS					Detailed Content from Risk Assessment									
Program	Version	Measure Category	Measure Name	Measure Description	Risk Priority Code	Risk Priority Sequence	Functionality / Certification	Workflow	Adoption	Content	Data	Reporting	Total	
EH MU Stage 1	Not Applicable	CQM - EH	ED-1 Subset-1	Emergency Department Throughput – Median Time from ED Arrival to ED Departure for Admitted Patients	Not Prioritized	1	0.6	o	1.2	0.3	0.6	0.9	3.6	
EH MU Stage 1	Not Applicable	CQM - EH	ED-1 Subset-2	Emergency Department Throughput – Median Time from ED Arrival to ED Departure for Admitted Patients	Not Prioritized		0	0	0	0	0	0	0	
EH MU Stage 1	Not Applicable	CQM - EH	ED-1 Subset-3	Emergency Department Throughput – Median Time from ED Arrival to ED Departure for Admitted Patients	Not Prioritized		0	0	0	0	0	0	0	
EH MU Stage 1	Not Applicable	CQM - EH	ED-2 Subset-1	Emergency Department Throughput – Admit Decision Time to ED Departure Time for Admitted Patients	Not Prioritized	1	1.2	0.3	0.6	0.3	0.6	O	з	
EH MU Stage 1	Not Applicable	CQM - EH	ED-2 Subset-2	Emergency Department Throughput – Admit Decision Time to ED Departure Time for Admitted Patients	Not Prioritized		0	0	0	0	0	0	o	
EH MU Stage 1	Not Applicable	CQM - EH	ED-2 Subset-3	Emergency Department Throughput – Admit Decision Time to ED Departure Time for Admitted Patients	Not Prioritized		0	0	0	0	0	0	0	
EH MU Stage 1	Not Applicable	CQM - EH	Stroke-2	Ischemic Stroke – Discharge on Antithrombotics	Not Prioritized	1	1.2	0.3	0.6	0.3	0.3	0	2.7	
EH MU Stage 1	Not Applicable	CQM - EH	Stroke-3	Ischemic stroke – Anticoagulation for A-fib/flutter	Not Prioritized	1	1.2	0.6	0	0.3	0.3	0.9	3.3	
EH MU Stage 1	Not Applicable	CQM - EH	Stroke-4	Ischemic stroke – Thrombolytic therapy for patients arriving within 2 hours of symptom onset	Not Prioritized	1	1.2	0.3	1.2	0	0.3	0.9	3.9	
EH MU Stage 1	Not Applicable	CQM - EH	Stroke-5	Ischemic Stroke – Antithrombotic Therapy by End of Hospital Day 2	Not Prioritized	1	1.2	0.6	1.2	0.6	0.3	0.9	4.8	
EH MU Stage 1	Not Applicable	CQM - EH	Stroke-6	Ischemic stroke – Discharge on statins	Outstanding	1	1.2	0	1.2	0.3	0.3	0.9	3.9	
EH MU Stage 1	Not Applicable	CQM - EH	Stroke-8	Ischemic or hemorrhagic stroke – Stroke education	Not Prioritized	1	1.2	0.3	1.2	0.6	0.3	0.9	4.5	
EH MU Stage 1	Not Applicable	CQM - EH	Stroke-10	Ischemic or hemorrhagic stroke – Rehabilitation assessment	Not Prioritized	1	1.2	0.6	0.6	0.6	0.3	0.9	4.2	

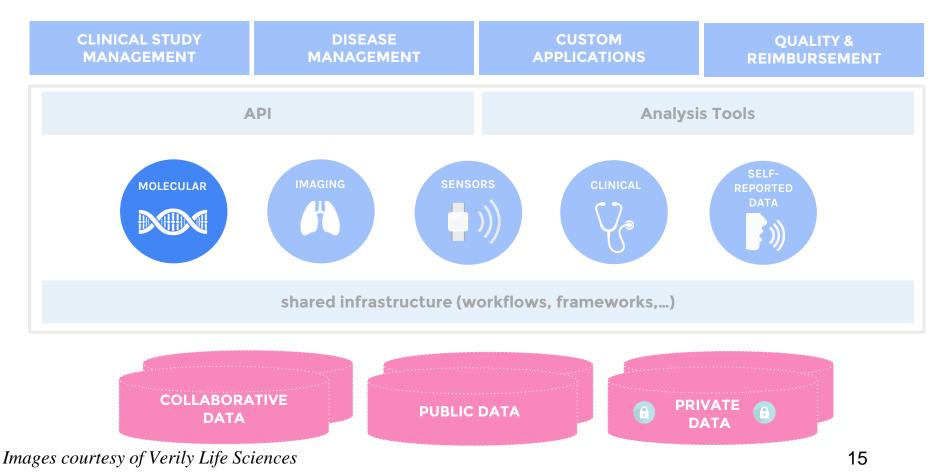
Functionality/CertificationContentWorkflowDataAdoptionReporting

#### **Provider analytics**

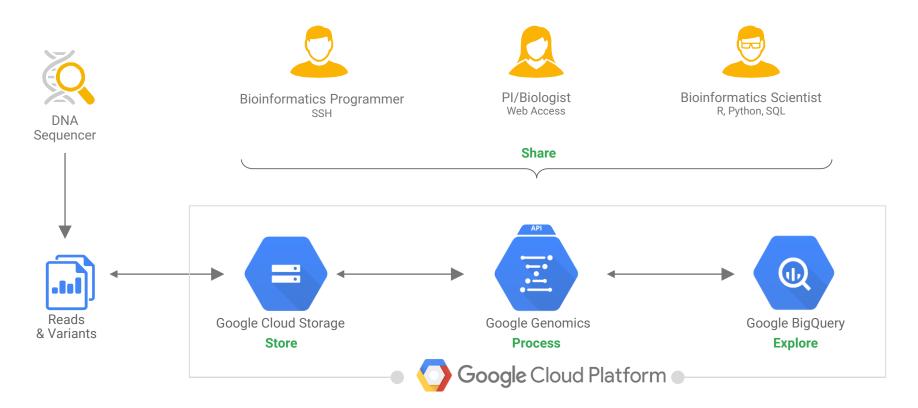
Your hospital had a 10.1% longer length-of-stay for Knee Joint Replacement (127 bed-days which costs \$549,004.20). This change may be driven by severity 2 cases, which are higher by 10.3%. The longer stays in severity 2 could account for 49.5% (63 bed-days) of the total increase.



#### **Genomics Capabilities**



#### **Genomics workflow**



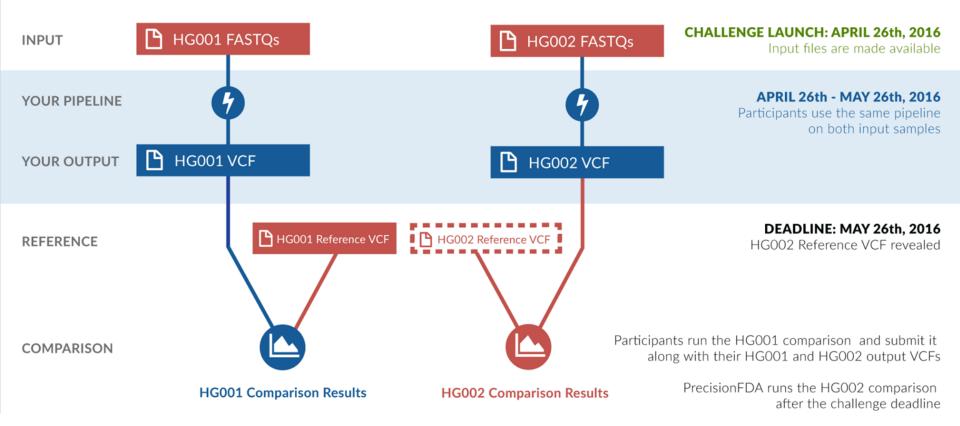
## PrecisionFDA Truth Challenge



President Obama's Precision Medicine Initiative envisions a day when an individual's medical care will be tailored in part based on their unique characteristics and genetic makeup. The goal of the FDA's second precisionFDA challenge, similarly to the first challenge, is to continue engaging the genomics community in advancing the quality standards in order to achieve more accurate and consistent results in the context of genetic tests (related to whole human genome sequencing), advancing the goal of better personalized care. PrecisionFDA invites all innovators to take the challenge and assess their (or their favorite!) software on the supplied human datasets. Participation is voluntary, but instrumental in helping the community prepare for the coming genomic data revolution.

#### **PrecisionFDA Truth Challenge**

April 26, 2016 through May 26, 2016

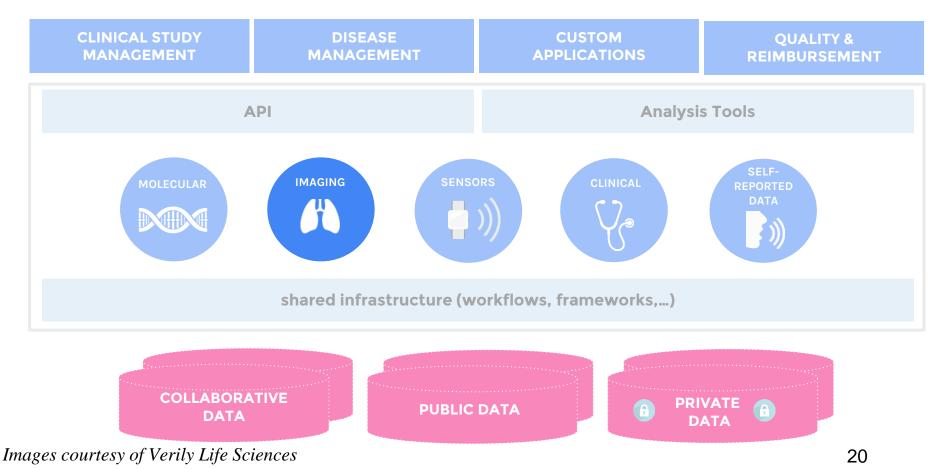


#### **PrecisionFDA Truth Challenge**



						F-SCORE				TRUTH	TRUTH	QUERY	QUERY		
ENTRY	TYPE	SUBTYPE	SUBSET	GENOTYF	PE	<b>↑</b>	RECALL	PRECISION	FRAC_NA	ТР	FN	ТР	FP	FP GT	% FP MA
<b> \$</b>	SNP	*	◆ *	\$ *	\$										
rpoplin-dv42	SNP	•	•	•		99.9587 %	<b>99.9447</b> %	<b>99.9728</b> %	19.0681 %	3052930	1689	3052766	832	433	52.0433 %

#### **Imaging Capabilities**





#### Other research possibilities ...

#### **RETINA**



#### EYE DISEASES

Glaucoma Age-related macular degeneration

#### SYSTEMIC DISEASES

Stroke & heart attack risk Diabetic nephropathy, neuropathy Vascular dementia, Alzheimer's Mortality? Hospitalizations?

#### **OTHER IMAGING**





#### **SKIN CONDITIONS**

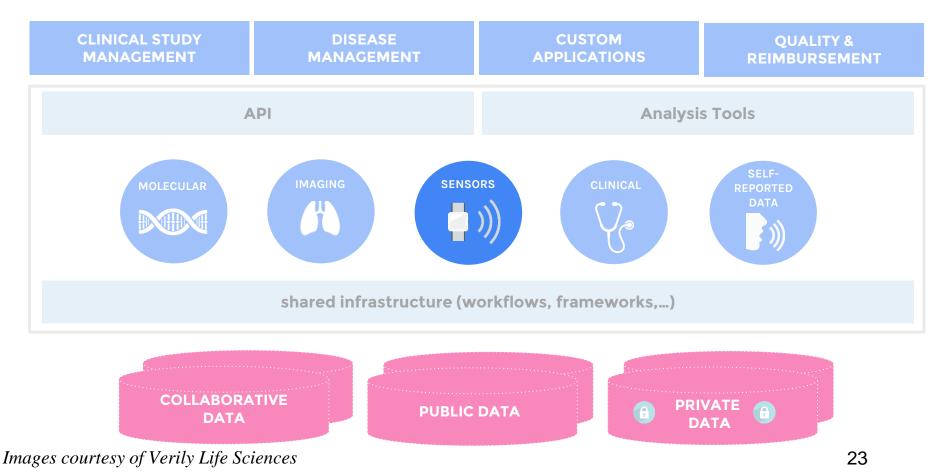
Moles Skin cancer Infections Acne/rosacea Dermatitis Hair/nail



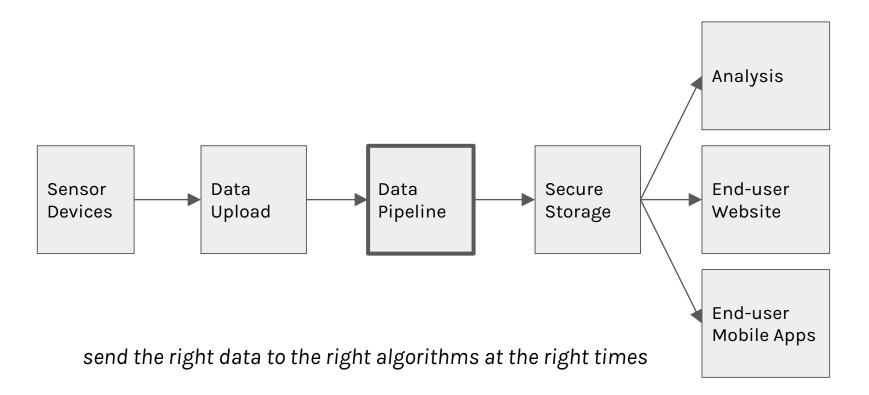
EAR, NOSE, THROAT Ear infections

Sore throat

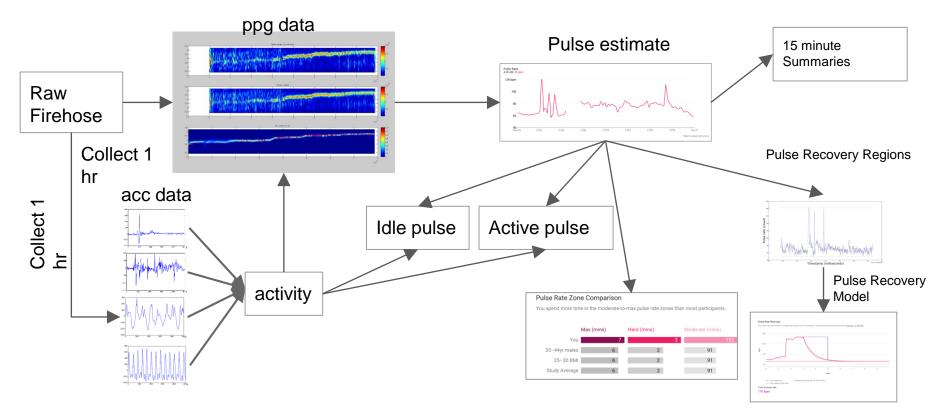
#### **Sensor Data Capabilities**

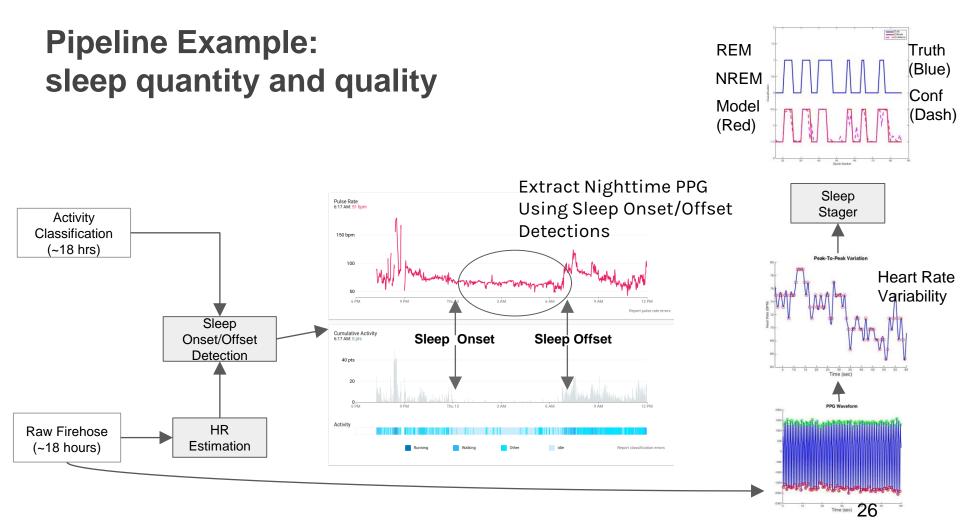


#### **Architecture Overview**

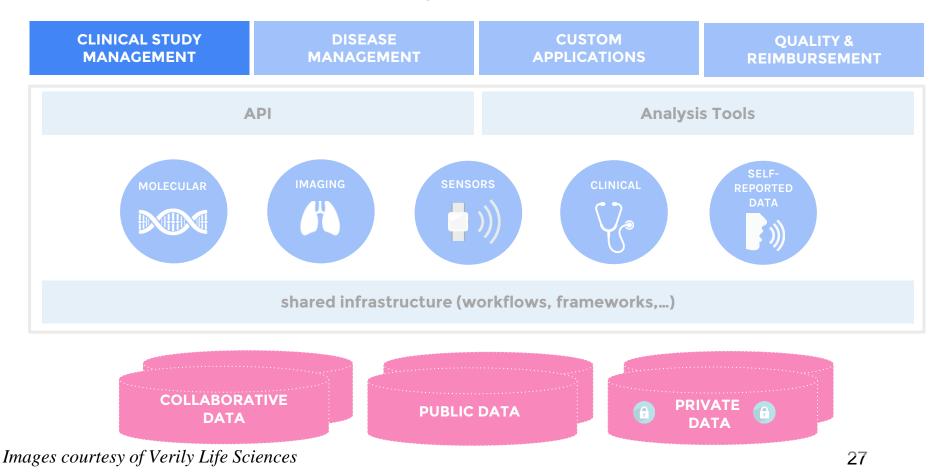


#### **Pipeline Example: pulse data computation**





#### **Solutions: Baseline Study**



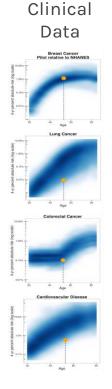
## "Google has embarked on what may be its most ambitious and difficult science project ever: a quest inside the human body."

Wall Street Journal | July 2014

#### **Broad and Deep Molecular, Device, and Clinical Phenotyping Data for Each Participant**

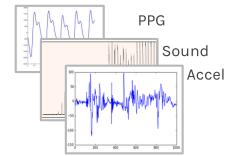
Imaging

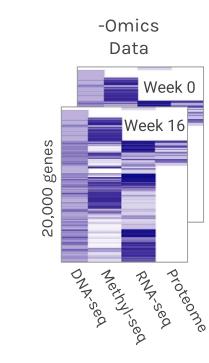
Data



Device Data







Immunoprofiling Data Neek 0 Week 16 Monocytes B cells

CD4 T cells
 CD8 T cells

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### **Example: Supported Verily Analyses**



**Ingestion, preprocessing, QC:** Import data at LIMS-level. Automatically survey data quality and highlight areas of concern. Determine pre-analytical, analytical, and biological variability.



**Clustering:** unbiased or hypothesis-weighted clustering of multi-omics data to reveal unique patterns.



**Regression:** supervised or semi-supervised methods that import known biological information.



Longitudinal: analysis and sequence prediction in longitudinal data.



eQTL/mQTL: integrative analysis combining multiple genetic data types.

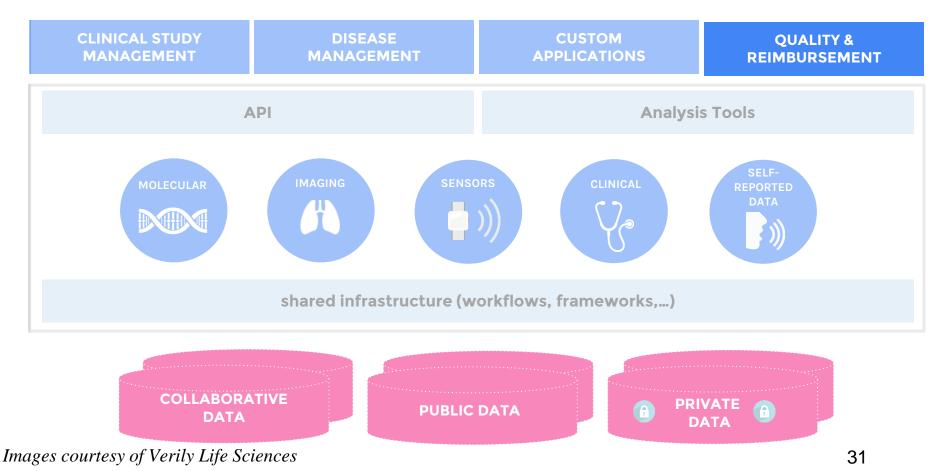


**GRS:** genomic predisposition; advanced modeling across multiple population data.

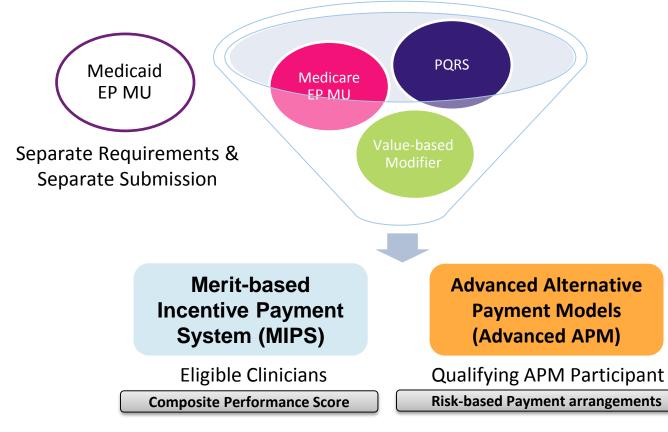


**Advanced machine learning:** for integrative pathway discovery & analysis, data annotation, quality control, and phenotype-\*omics associations.

#### **Solution: Quality Improvement / MACRA**



# The MACRA Quality Payment Program Consolidates key aspects of three existing physician-based programs



### **Data Submission Mechanisms for Groups**

Merit-based Incentive Payment System (MIPS)

<ul> <li>Qualify</li> <li>Qualified Clinical Data Registry (QCDR)</li> <li>Qualified registry</li> <li>EHR</li> <li>CMS Web Interface (groups of 25 or more)</li> <li>CMS-approved survey vendor for CAHPS for MIPS (must be reported in conjunction with another data submission mechanism</li> </ul>	<ul> <li>CPIA</li> <li>Attestation</li> <li>QCDR</li> <li>Qualified registry</li> <li>EHR</li> <li>CMS Web Interface (groups of Administrative claims (if techn submission required)</li> </ul>	
Resource Use <ul> <li>Administrative claims (no submission required)</li> </ul>	<ul> <li>Advancing Care Information</li> <li>Attestation</li> <li>QCDR</li> <li>Qualified registry</li> <li>EHR</li> <li>CMS Web Interface (groups of the second se</li></ul>	

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