

Augusta 2.0 Overview

How does AI ML benefit health systems and hospitals?

- Diagnose disease more accurately
- Improve care delivery and personalized medicine
- Align hospital reporting and administration to VBC incentives
- Effectively plan treatments



Cost to bring a single drug to market

POTENTIAL BENEFITS 30 TO 40% IMPROVED DIAGNOSIS

- Millions of Americans are misdiagnosed annually, e.g. ~12 million with incorrect diagnoses in outpatient clinics in 2014 (CBS News)
- Potential to improve diagnostic outcomes by 30% to 40%.
- Potential to reduce hospital stays, unnecessary readmissions and testing, and health care costs.

Source: Frost & Sullivan, "Al & Cognitive Computing Systems in Healthcare" (December 2016). TM Capital Industry Spotlight, "The Next Generation of Medicine: Al and Machine Learning" (2017).

Estimated potential annual benefit for an application by 2026. Source: Accenture analysis, "Al: Healthcare's New Nervous Systems" (2017).

Current State of Al Adoption - Heavily Verticalized



For what stages of drug discovery does your organization currently use artifical intelligence?



A recent survey of the Drug Discovery industry found that most current applications of AI are in areas you might expect (target identification, safety, discovery)

Most identified "Speed of Discovery" as the greatest motivating factor for using AI, however, **45% thought their application of AI would remain the same next year**

https://blog.benchsci.com/6-things-we-learned-about-artificial-intelligence-in-drug-discovery-fro m-330-scientists



Source: BenchSci and The Science Advisory survey conducted in December 2017 amongst 330 scientists who work on drug discovery, drug development, or related biomedical research Note: Respondents could select multiple answers

The Problem With Verticalized Applications of Al

• Our initial model showed a marked difference between control and disease groups, however this was only due to an undetected bias in survey site



Integrating Data for Better Accuracy



Combining MRI with genomic features allowed **better prediction performance** than with either alone - Alzheimer's Disease Neuroimaging Initiative (ADNI) data

This combination approach becomes more accurate as additional data types are included, and is effective in precision medicine & drug discovery applications

Evaluation Metric	MRI	Metabolomics	Genetics	All combined
True Positive Rate	0.73	0.73	0.77	0.83
False Positive Rate	0.66	0.63	0.5	0.39
AUC	0.56	0.55	0.68	0.73



What Limits the Widespread Application of AI in Biomedicine?





Data Variety/Heterogeneity

Lack of Standards

Different Data Types: EHR/EMR, MRI/fMRI, EEG, EKG, chemistry No standards for processing or interpreting medical data

Lack of Scalability

Challenges in pre-processing and machine learning using massive data

Single framework for integration of diverse data types Dynamic, optimized pre-processing

Deploy anywhere, on any architecture

Introducing Augusta



Actionable insight for quality of care

Personalized diagnostic models

Population-scale health analysis

Standardized data features and R&D

What Makes Augusta Unique?

Parameter optimization is typically performed after selecting an ML model





What Makes Augusta Unique?



Substantial bias can be introduced in pre-processing, optimization should begin much sooner



What Makes Augusta Unique?



Augusta is a single language that you can use to process and interpret biomedical data, allowing you to build automated AI workflows



Augusta 2.0 - A Biomedical AI Language

All operations are treated as optimizable blocks:



Source - Data source (file, directory of files)



Transform - Data transformation, such as a preprocessing step for a specific datatype (e.g. skull subtraction for an MRI) or an operation (e.g. calculating a mean)



Model - Any number of machine learning models



Performance - Examine impact of transformation & model parameters

How the Augusta[™] Platform Works

Augusta[™] can start with siloed, complex, and raw data of multiple types:



- Integration of diverse and large-scale data types
- Data of any type, size, and dimensionality explored and modeled
- Biological, clinical, genomics, precision medicine, metabolomic, lab testing, drug compound data



- Accurate results and automated distribution
- Built for massive data architecture
- End-to-end machine learning
- Post-analysis & data visualization



Our Partners and Market Environment





DRUG DISCOVERY

Reduction of Chemical Survey Space Prediction of Molecular Mechanism Experimental Data Integration



PRECISION MEDICINE

Genomic Risk Assessment Improved Disease Diagnosis Patient Data Integration



VALUE-BASED CARE

Patient stratification Outcome & Event Prediction Operational Analysis



Use Cases: Outcomes Prediction

- Continuously updated calculation of severity scores (including SOFA)
- Generate dynamic risk scoring for patients using time-based models
- Symptoms extracted from clinical notes
- Prescription names mapped to relevant MeSH terms, allowing systematic analysis of prescriptions by type

MIIMC Patient Dashboard Age: 49.0 (deceased icity: BLACK/AFRICAN AMERICAN Marital status: SINGLE a nerwider: Madica Mal hep kid w er i 2 3 End stage renal class 3 4 Schizzohrenia NOS-unep 4 5 Anomia in chr kidney dia lected Tonic: (Slide to adjust relevance metric-9 0.6 $\lambda = 1$ 0.0 0.2 0.4 0.0 Intertopic Distance Map (via multidimensional scaling) Top-30 Most Relevant Terms for Topic 2 (15.1% of tokens) 0.9% sodium chloride stoproiol tartrat Marginal topic distributio

Overall term frequency



Use Cases: Value Based Care



- GE Healthcare (US) major initiative in value based care, predicting outcomes for patients with chronic conditions (NLP-based)
- OrbCare (Canada) analysis of clinical billing to identify diagnostic code over/under use
- Intacare (UK) homogenization of billing codes





Use Cases: Precision Medicine



- Automatable, dynamic processing of medical images & streaming data from virtually any format
- Easy integration of multiple lines of clinical data
 - Scalable model generation using cloud and conventional computing









Use Cases: Drug Discovery & Building an Automated Platform



- Initial models focused on predicting zebrafish phenotypes
- Prioritized compounds showed correlates in clinical application
- In a collaborative effort, we included genomic, proteomic, and transcriptional data, generating predictions of molecular mechanism underlying phenotypes
- Currently working to automate this process:
 - Begin with prediction of phenotype
 - Automated phenotypic barcoding
 - Predict molecular mechanism & validate
 - Refine process



Our Team



Anthony lacovone Co-Founder & Board Chairman



Gabriel Musso, PhD

Chief Scientific Officer



Wendy Tsai, MBA VP Business Development



Babak Afshin-Pour, PhD VP of Technology



Victoria Catterson, PhD Principal Data Scientist



Cindy Lopes Application Developer

The BioSymetrics Strategic Advisory Board

Dr. Calum MacRae Harvard University & Brigham and Women's Hospital

Dr. Robert DeVita Icahn School of Medicine at Mount Sinai (New York), Merck

Mr. Chinnappa Kodira GE Healthcare, Broad Institute, Roche

Dr. Paul Boutros

Ontario Institute for Cancer Research, University of Toronto

Dr. Gene Salkind Holy Redeemer Hospital, Albert Einstein Medical Center





www.biosymetrics.com | Twitter @biosymetrics

CONTACT

Wendy Tsai | VP Business Development | wendy@biosymetrics.com Gabe Musso | Chief Scientific Officer | gabe@biosymetrics.com