

# Data Science, ML, and AI for Medical Students

Hour 1: Linking AutoAnalyzer with Colab Notebooks for Clinical Insight

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Data Science



**Machine Learning** 



**Artificial Intelligence** 

# **Relevant Disclosures**

David Liebovitz, MD

### Financial Disclosures

Advisor to Dendritic Health AI (Using AI for education)

# **Bias Mitigation**

- I have used the Feinberg bias checklist to review this lecture to mitigate bias.
- I have reviewed this lecture for use of stereotypical or offensive terms, language, etc.
- I have, to the best of my abilities, used inclusive language, pronouns, photos and images.

Despite best efforts and intentions, if there is concerning content, please feel free to contact me or report via the bias reporting system.



# **Digital Health & Data Science Training Series**

# **Course Topics**

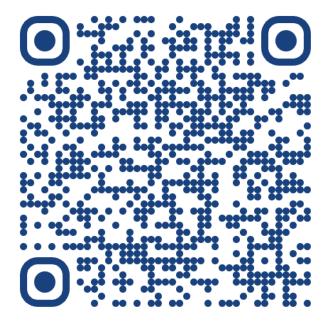
- Core Concepts in Digital Health and Data Science
- Al within CVD
- Data Exploration and Visualization ALC ← Today
- Machine Learning and Al Models ALC
- Video-based Gait Analysis using Al
- Telemedicine in Psychiatry
- Wearables and Reproductive Health
- LLMs in Healthcare
- Ethical Considerations for AI in Healthcare

# **©**\* Key Competencies

- 1 Healthcare Delivery Science: Evaluate digital health tools impact
- 2 Health Data Ecosystem: EHRs, PACS, device data, omics
- 3 Health IT Regulatory: Mandates, interoperability, compliance
- 4 Data Science Methods: ML/Al for diagnosis, precision medicine
- 5 Clinical Decision Support: CDS system principles
- 6 Modeling Applications: Apply predictive models to clinical problems
- Bias, Ethics, Health Equity: Algorithm bias, ethical implications
- 8 Sociotechnical Context: Implementation science principles

https://github.com/DrDavidL/learning-dhds/blob/main/README.md

http://bit.ly/4mxwklv





# Why Digital Health & Al Skills Are Essential



# **Current Healthcare Reality**



#### **Data-Driven Medicine**

EHRs generate 2.3 exabytes of healthcare data annually. Your patients' care increasingly relies on algorithms for diagnosis, treatment planning, and risk stratification.



#### **Al Integration Accelerating**

FDA has approved 500+ AI medical devices. By 2030, AI will be standard in radiology, pathology, drug discovery, and clinical decision support.



#### **Precision Medicine**

Genomics, proteomics, and personalized treatment protocols require computational skills to interpret and apply effectively.





#### **Clinical Practice**

You'll need to interpret AI recommendations, understand model limitations, identify bias, and communicate uncertainty to patients and colleagues.



#### **Critical Thinking**

As shown in today's data quality example, you must critically evaluate the data and algorithms that inform your medical decisions.



#### Leadership & Innovation

Whether in research, administration, or clinical practice, you'll shape how AI is developed, validated, and implemented in healthcare.



#### **The Bottom Line**

Digital health literacy isn't optional—it's as fundamental to modern medicine as anatomy or physiology. Today's session gives you practical tools to understand, evaluate, and use data-driven insights in your future practice.

# **Model Performance: AUROC vs Calibration**

### Neurological Example: Migraine Prediction

#### Two Al Models for Migraine Risk (Hypothetical Example)

#### Model A (Good AUROC, Poor Calibration):

- · AUROC: 0.87 (excellent discrimination)
- Says "40% migraine risk" for patients with 15% actual risk
- Problem: Over-prescribing preventive medications

#### Model B (Good AUROC, Good Calibration):

- AUROC: 0.85 (still excellent discrimination)
- Says "40% migraine risk" for patients with 38% actual risk
- · Benefit: Accurate shared decision-making

#### **Patient Conversation Impact**

Patient: "Doctor, the AI says I have 40% migraine risk. What does that mean?"

With good calibration: "Out of 100 patients like you, about 40 will develop migraines in the next year."

# **III** Understanding Model Metrics

AUROC (Discrimination)

"Can the model rank patients from low to high risk?"

- 0.5 = random guessing 0.8+ = good 0.9+ = excellent
- Calibration

"Are the predicted probabilities accurate?" If model predicts 30% risk, do ~30% of patients actually have the outcome?

**Population Variation** 

Performance can vary dramatically across age, sex, ethnicity, and comorbidities Always test subgroups separately!



#### **Clinical Takeaway**

Both metrics matter: AUROC for identifying high-risk patients, calibration for making treatment decisions. Test performance across all relevant patient populations.



# **Statistical Reasoning: Beyond P-Values**

# Neurological Example: Coffee and Stroke Risk

#### Study Findings (Hypothetical Example)

#### Observational study results:

- · Heavy coffee drinkers: 15% stroke rate
- Non-coffee drinkers: 10% stroke rate
- p < 0.001 (highly significant!)
- Conclusion: Coffee causes strokes?

#### The Hidden Confounders

Heavy coffee drinkers also had:

- Higher smoking rates (60% vs 10%)
- More sedentary jobs (desk work)
- · Higher stress levels
- Poor sleep habits

The association disappeared after controlling for these factors!

# ✓ Statistical Reasoning Principles

- Association ≠ Causation: Strong statistical associations can be misleading without considering confounders
- P-value Limitations: Statistical significance ≠ clinical significance. A tiny effect can be "significant" with large sample sizes
- Effect Sizes Matter: A 50% reduction from 0.02% to 0.01% risk is statistically dramatic but clinically minimal
- Confidence Intervals: Wide CIs suggest uncertainty, regardless of statistical significance

# ♥ Clinical Takeaway

Always ask: "Is this association clinically meaningful?" and "What else could explain this relationship?" before making treatment decisions.

### **©** For Your Practice

Focus on effect sizes, confidence intervals, and biological plausibility—not just p-values. Consider confounders and alternative explanations.



# The Evolving Story of Statins: A Data Science Case Study

# **Timeline of Discovery**



First Statin Approved: Lovastatin introduced for cholesterol reduction



WOSCOPS Trial Published: Demonstrated cardiovascular benefits



Unexpected Finding: WOSCOPS post-hoc analysis revealed diabetes association



Pattern Confirmed: Multiple trials confirmed increased diabetes risk

# Neurologic Connection

Dual Purpose: Statins prevent both coronary artery disease (CAD) and stroke through cholesterol reduction and plaque stabilization, making them essential in neurology practice.

### ▲ The Unexpected Association

#### WOSCOPS (2001): The First Signal

- Originally showed 30% reduction in diabetes cases
- 5,974 participants, 139 developed diabetes
- · Used non-standardized criteria for diabetes diagnosis
- Initially appeared protective

#### **Subsequent Studies: Pattern Reversal**

- JUPITER trial: Higher diabetes incidence with rosuvastatin
- Meta-analyses show 9-12% increased risk
- Risk greater with high-intensity statins
- Consistent pattern across multiple studies



#### **Key Clinical Insight**

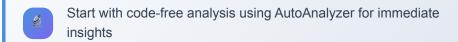
Benefits still outweigh risks: statins prevent ~5 cardiovascular events for every new diabetes

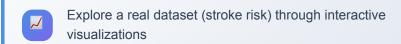


#### **Why This Matters for Your Future Practice**

In our rapidly evolving healthcare environment, unexpected associations like this are discovered continuously. The tools and analytical approaches you're learning today—data exploration, statistical testing, and machine learning—enable much earlier detection of these patterns, potentially improving patient safety and outcomes.

# **Session Goals**





Dive deeper with Colab notebooks (don't worry - Al will help!)

Build predictive models and prepare for future sessions





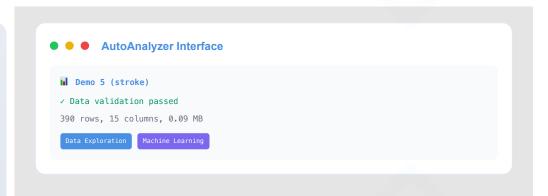
# Part 1: Code-Free Analysis with AutoAnalyzer



### **Getting Started**

#### Access URLs:

- autoanalyze.azurewebsites.net
- autoanalyze-beta.azurewebsites.net
- ▲ Work in small groups server capacity limited
- Public data only hosted on Northwestern's Azure for privacy





### **Future Option**

Available on GitHub for local deployment

→ Consider informatics elective for local use with sensitive data

# Our Analysis Plan



1. Clean & Filter

Prepare dataset for analysis



2. Explore & Visualize

Charts, plots, patterns



3. Predict Stroke

Build ML model



# Part 2: Deeper Dive with Colab Notebooks - Use GitHub links



Don't Fear the Code! Al is Your Assistant

When we move to code, remember: Al tools like ChatGPT, Claude, and GitHub Copilot can help you understand, debug, and write code. You're not expected to memorize syntax!

We are going to use just 2 notebooks today that are pre-made for this session to stay in our time slot! Use links to these on the GitHub page!

- 1. Stroke Al Generated Analysis (Completed) Al-powered stroke data analysis with LLM integration
- 2. Stroke Analysis Your Turn Hands-on stroke analysis practice with LLM tools

Notebooks below - also on GitHub - use for more in-depth learning!



### Notebook 1: Data Exploration

- · Load and inspect the stroke dataset
- · Identify variable types and missingness
- Visualize distributions (histogram, bar chart)
- Form clinical questions from observed patterns



### Notebook 2: Statistical Analysis

- · Compare groups using t-tests and chi-square
- Calculate effect sizes: risk difference and ratio
- · Interpret p-values, confidence intervals
- Recognize limits of observational associations



### Notebook 3: Prediction Model

- · Split data into train/test for generalization
- · Build logistic regression with preprocessing
- Evaluate discrimination (AUROC) and calibration
- · Discuss model limitations: leakage, bias, fairness



### **P** AutoAnalyzer → Colab Bridge

Use AutoAnalyzer insights to guide your code-level learning in Colab notebooks



# **Data Quality & Integrity: A Growing Concern**



# **Recent Findings: Federal Data Manipulation**

Freilich J, Kesselheim AS. The Lancet (2025)

Study of 232 U.S. public health datasets revealed concerning patterns



49% of Datasets Altered

114 of 232 datasets changed without public logging



93% Changed Gender → Sex

Most common alteration: terminology changes



Only 13% Logged

Vast majority of changes unrecorded

# **©** Clinical Implications

- Research Validity: Unlogged changes make studies irreproducible
- Variable Meaning: 'Sex' ≠ 'Gender' different biological vs. social constructs
- Clinical Decisions: Policy changes based on manipulated data
- Public Trust: Transparency essential for scientific credibility



### **For Your Practice**

#### Always verify data provenance

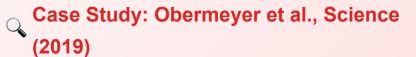
- Check data source documentation
- · Look for version control and change logs
- Understand variable definitions
- Document your own data processing steps



Remember: Good data science starts with trustworthy, well-documented data



# **Subgroup Analysis: Why Fairness Matters**



Analysis of a commercial algorithm used by health systems to identify patients for care management programs



The Algorithm's Goal

Identify patients who would benefit from extra care management



What It Actually Did

Used healthcare costs as a proxy for health needs



The Bias

Selected less sick white patients over sicker Black patients

# **Root Cause Analysis**

- Flawed Proxy: Healthcare costs ≠ health needs
- Access Disparities: Black patients had less access to expensive care
- 3 Historical Bias: Algorithm learned from biased historical data
- 4 No Subgroup Testing: Algorithm wasn't tested across racial groups

### **▼** The Solution

**Mandatory subgroup analysis** revealed the bias. When researchers tested performance by race, they found:

- · Black patients were significantly sicker at same risk scores
- · Algorithm needed recalibration for equitable care



Key Takeaway: Always test Al models across different patient populations before deployment



# Calibration vs Discrimination: Understanding Model Performance

# **III** Discrimination (AUROC)

"Can the model separate high-risk from low-risk patients?"

- AUROC = 0.5: Random guessing
- AUROC = 0.7: Acceptable
- AUROC = 0.8: Good
- AUROC = 0.9: Excellent



"Are the predicted probabilities accurate?"

If model says "30% stroke risk," do 30 out of 100 similar patients actually have strokes?

# Clinical Example: Stroke Risk

#### Model A: Good discrimination, poor calibration

- AUROC: 0.85 (Excellent at ranking patients)
- But predicts 20% risk for patients with 5% actual risk
- · Problem: Over-treatment, unnecessary procedures

#### Model B: Good discrimination, good calibration

- AUROC: 0.82 (Still excellent at ranking)
- Predicts 20% risk for patients with 19% actual risk
- Benefit: Accurate shared decision-making

## **∇** Clinical Impact

#### Patient conversation:

"Doctor, you said I have a 20% stroke risk. What does that mean for me?"

With good calibration: "Out of 100 patients like you, about 20 will have a stroke in the next year."



Both matter: Discrimination for identifying high-risk patients, calibration for making treatment decisions



# **Essential Facts — What to Remember**



Garbage in → garbage out: prioritize data quality



Association ≠ causation; confounding is common



Calibration is as important as discrimination



Always check subgroup performance for fairness



Document and share methods for reproducibility

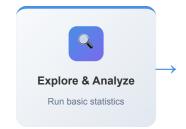


Al tools are your coding companions, not replacements



# **AutoAnalyzer Integration**











# **Next Steps — Future Sessions**



