# **USING COMPUTER VISION TO DISENTANGLE** FEATURES ENABLING AI TO LEARN SELF-REPORTED RACE AND ETHNICITY FROM MEDICAL IMAGES

### **INTRODUCTION**

#### The Danger of Algorithmic Bias in Healthcare

A 2019 clinical Artificial Intelligence (AI) applied to ~200 million Americans underdiagnosed patients of color by 50%.



The global market for AI in Medical Imaging is estimated to grow >10 times by 2033.

AI-powered medical imaging tools are expanding and exacerbating inequity in clinical care for Black and Brown patients, and other vulnerable communities.

### **AI Can Learn Self-Reported Race And Ethnicity**

In 2021, AI models were trained to recognize patients' self-reported race and ethnicity from medical images, even when there are no indications of race or

## **METHODS**

#### **Discovering Hidden Signals Using CNNs**

My goal was threefold: (1) to identify key image features that could be hidden signals, (2) to extract each feature from RVMs, and (3) to train an AI model to learn race and ethnicity from the isolated feature to assess its individual significance.

After conducting 100+ experiments to identify key image features, I designed a novel approach to deconstruct an RVM into three key features: **Number of** Nonzero Pixels (NNP), Pixel Intensity Values (PIVs), and Spatial Arrangement (SA).



#### ethnicity visible to human experts. This has stumped experts worldwide.



AI *might* be learning false correlations between race or ethnicity and disease, but we're not sure because the hidden features that signal AI to race and ethnicity are UNKNOWN.

We need to discover how AI learns self-reported race and ethnicity when humans cannot.



SA Normalized







### **Convolutional Neural Network Model Training**

If the model trained on an isolated feature performs better than random in detecting self-reported race and ethnicity on a modified test set, we infer that the specific RVM feature is a hidden signal!

### RESULTS



AI can learn self-reported race and ethnicity from RVMs with AU-ROCs from 92.0 to 95.0.

BIOMARKE

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SED

SEGMENTATION

Adult Dataset

**RESEARCH GOAL** 

Discover the hidden signals in retinal images that enable algorithms to learn self-reported race and ethnicity.

NNP* Isolated	<b>UKBB:</b> 70.0, 1.14·10 <sup>-7</sup>	10000	Pixels i-ROP: $p = 2.45 \cdot 10^{-160}$ UKBB: $p = 2.14 \cdot 10^{-24}$	nonzero pixels in Black RVMs vs. White RVMs		<ul> <li>are present in Black RVMs vs. White RVMs in the central regions near the optic disc.</li> <li>The peripheral regions in Black RVMs contain more choroidal vessels/capillaries than White RVMs.</li> </ul>
PIVs* Isolated	<b>i-ROP:</b> 79.6, 3.01·10 <sup>-5</sup> <b>UKBB:</b> 58.1, 2.55·10 <sup>-2</sup>	Pixel Intensity Distribution vs. Pigmentation	Pixel Intensity Distributions <b>i-ROP:</b> z = 14.3 <b>UKBB:</b> z = 43.0	Black RVMs have more pixels of medium intensity, White RVMs have more pixels of high intensity		
SA* Extracted	<b>i-ROP:</b> 82.0 – 97.5, 3.72·10 <sup>-13</sup> – 1.16·10 <sup>-6</sup>	i-ROP Infant RVM Dataset	UK BIOBANK Adult RVM Dataset	Retinal vessels in center regions are very highly weighted in learning self- reported race		
* NNP: Number of Nonzer PIVs: Pixel Intensity Val	5.24·10 <sup>-12</sup> – 6.25·10 <sup>-6</sup> ro Pixels ues	0 Contraction of the contraction	0 425 0 40 0 40	Model artifacts in exterior regions are very highly weighted in learning self- reported race		All images created by student researcher unless stated otherwise.

SA: Spatial Information