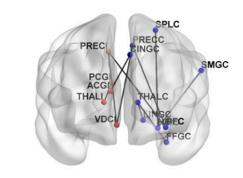


Osher Lifelong Learning Institute Our Brains: An Operator's Manual – 11/9/2021

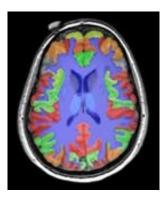


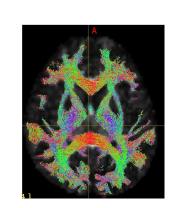
Brain Fingerprints with MRI

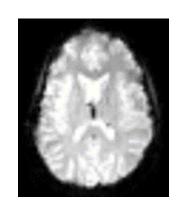
Victoria Morgan, PhD

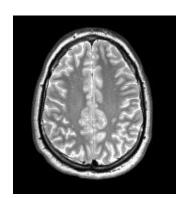
Professor

Vanderbilt University Institute of Imaging Science















MEDICAL CENTER



Imaging & Engineering:

Catie Chang, PhD Adam Anderson, PhD Baxter Rogers, PhD Bennett Landman, PhD John Gore, PhD Lucas Sainburg Kurt Schilling, PhD Hernan Gonzalez Sarah Goodale Graham Johnson Andrew Jansen, PhD

Neurology:

Bassel Abou-Khalil, MD Monica Jacobs, PhD

Neurosurgery: Dario Englot, MD, PhD

VUIIS Human Imaging Core

Funding:

NIH R01 NS075270 NIH R01 NS108445 NIH R01 NS110130 NIH R01 NS097618 (DJE)



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School of Engineering



National Institute of **Neurological Disorders** and Stroke

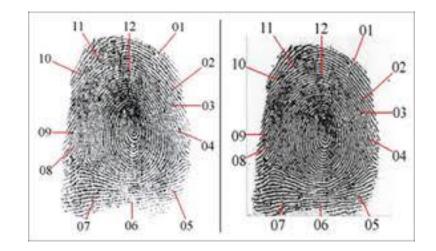


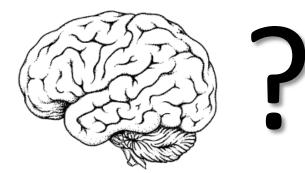
Why FINGERPRINT?

pattern



unique

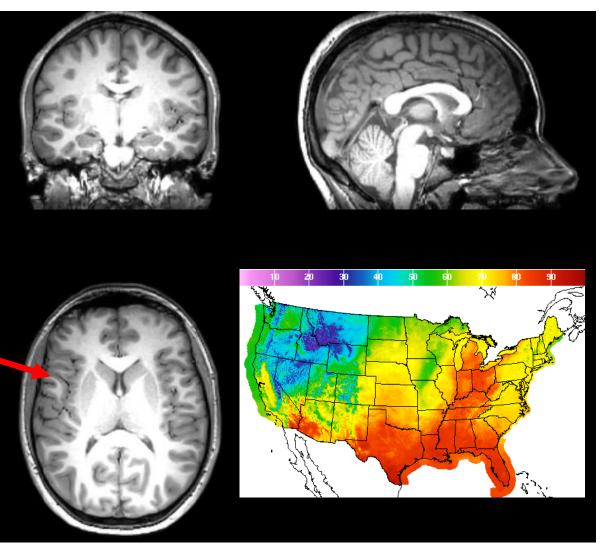




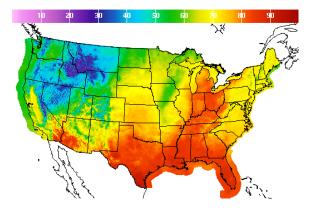
Magnetic Resonance Imaging - MRI



Makes an image from *tissue properties* parametric maps

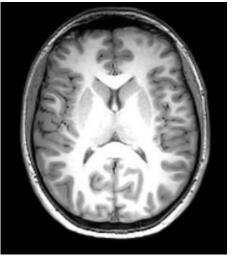


Magnetic Resonance Imaging - MRI

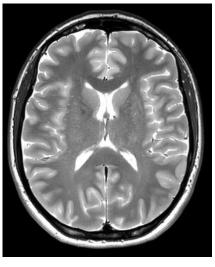


Makes an image from *tissue properties*

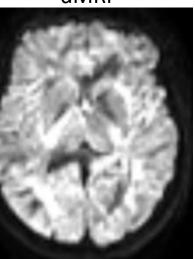
T1 weighted



T2 weighted



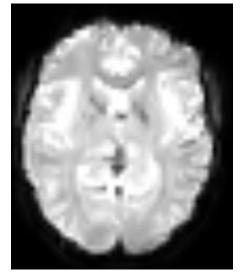
dMRI



Diffusion Weighted MRI

Intensity = how much water diffuses along a specific axis

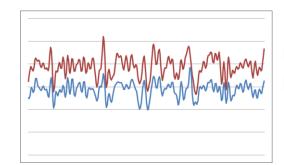
fMRI



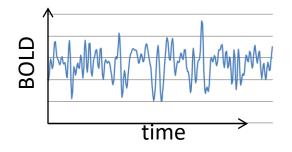
Functional MRI

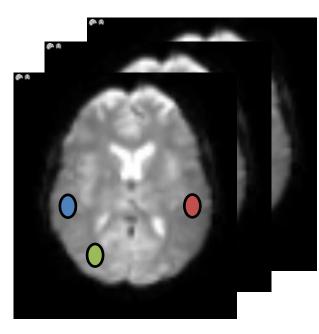
Intensity = Oxyhemoglobin to deoxyhemoglobin ratio (BOLD) = Brain activity

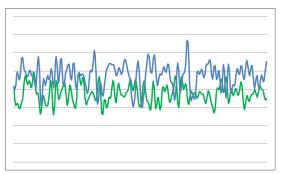
Functional MRI Connectivity



High linear correlation = High functional connectivity



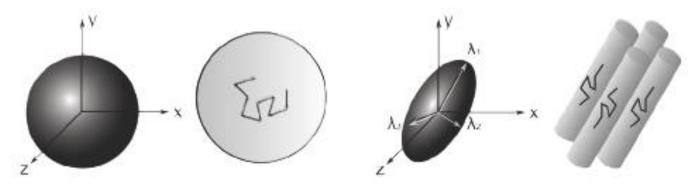




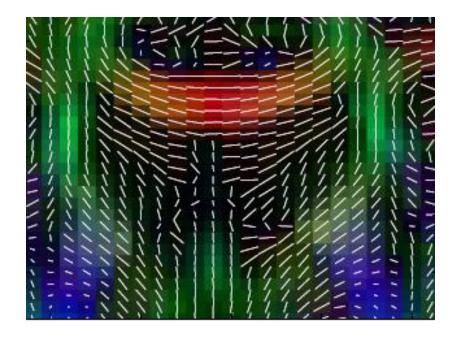
Low linear correlation = Low functional connectivity

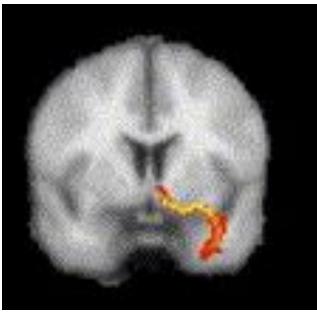
[Biswal 1995]

Diffusion MRI Connectivity



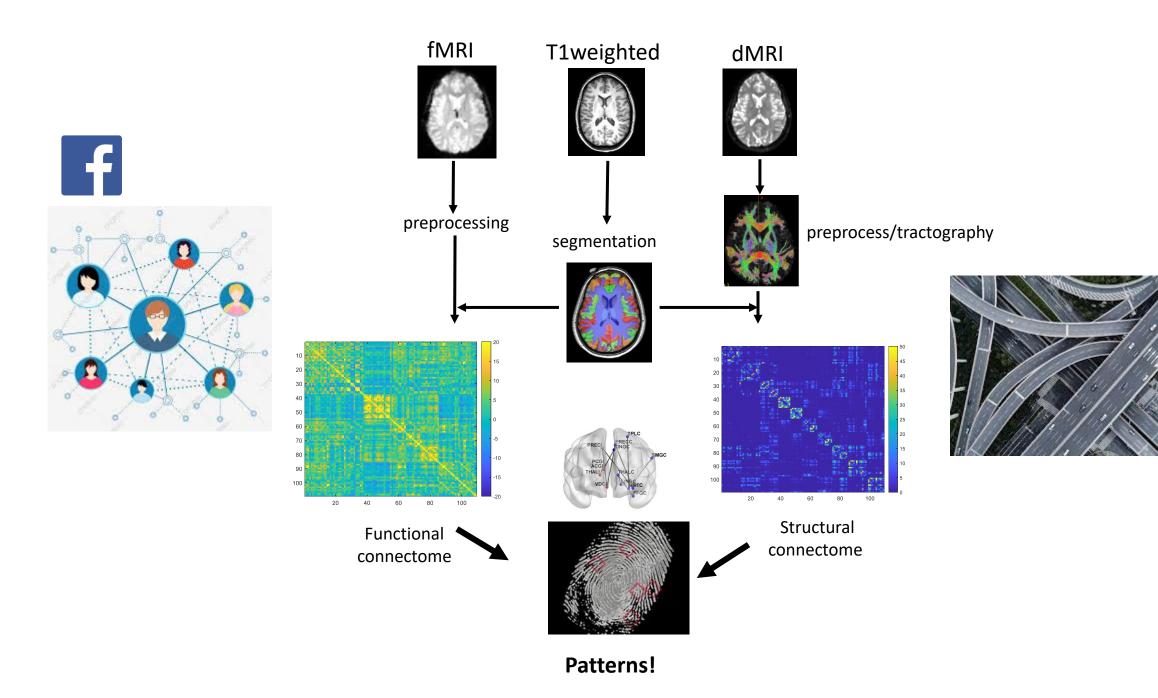
[http://qims.amegroups.com/article/view/1315/1771]





[Behrens 2007]

Structural connectivity = "trackability" of streamlines



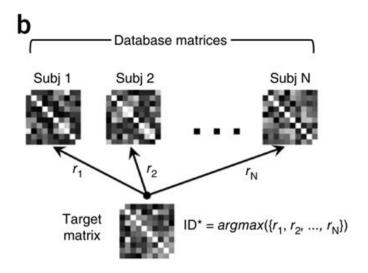
Are the patterns UNIQUE to individuals?

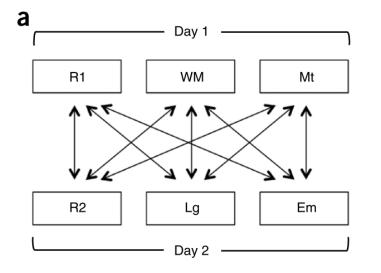


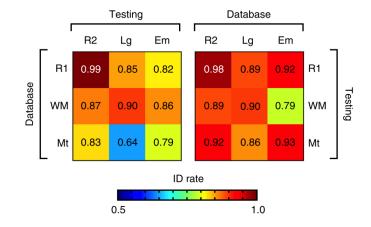
Functional connectome fingerprinting: identifying individuals using patterns of brain connectivity

Emily S Finn^{1,7}, Xilin Shen^{2,7}, Dustin Scheinost², Monica D Rosenberg³, Jessica Huang², Marvin M Chun^{1,3,4}, Xenophon Papademetris^{2,5} & R Todd Constable^{1,2,6}

- 126 subjects
- Different types of scans
- Different subsets of regions



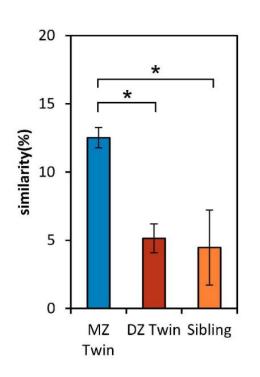




Here we show that an individual's functional brain connectivity profile is both unique and reliable, similarly to a fingerprint. We demonstrate that it is possible, with near-perfect accuracy in many cases, to identify an individual from a large group of subjects solely on the basis of his or her connectivity matrix. Although inter-individual RESEARCH ARTICLE

Quantifying Differences and Similarities in Whole-Brain White Matter Architecture Using Local Connectome Fingerprints

Fang-Cheng Yeh¹*, Jean M. Vettel^{2,3,4}, Aarti Singh⁵, Barnabas Poczos⁵, Scott T. Grafton³, Kirk I. Erickson⁶, Wen-Yih I. Tseng⁷, Timothy D. Verstynen⁸*



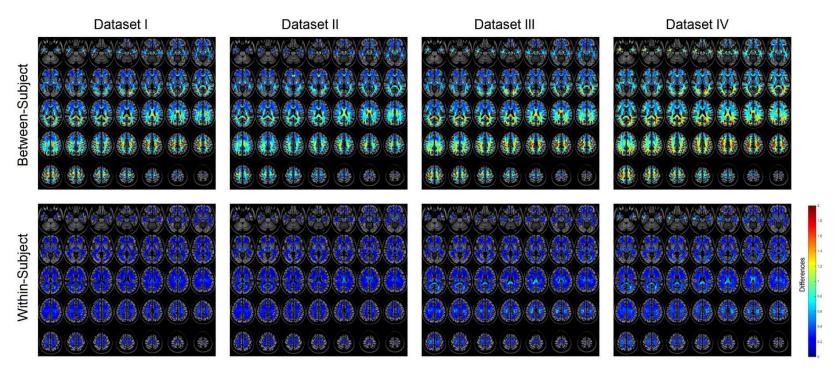
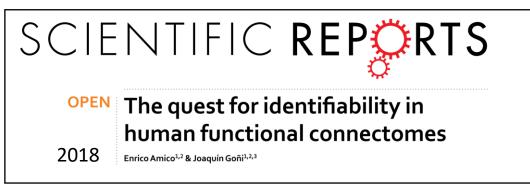


Fig 4. The spatial mapping of between-subject (first row) and within-subject (second row). Dataset I was acquired within 16 days, whereas dataset II (1~3 months), dataset III (6 months) and dataset IV (a year) were acquired at longer time intervals. High between-subject differences can be observed in white matter tissue, especially the corpus callosum and central semiovale. The within-subject differences are much smaller, and repeat scans with longer time intervals show higher within-subject differences.

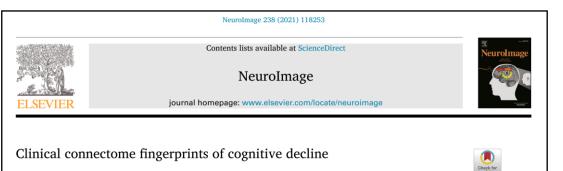


RESEARCH ARTICLE

2021 WILEY

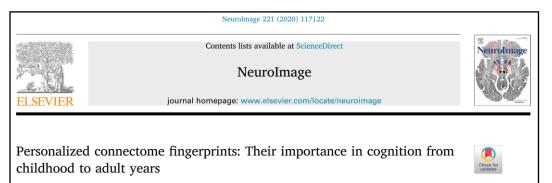
Functional connectome fingerprinting: Identifying individuals and predicting cognitive functions via autoencoder

Biao Cai¹[©] | Gemeng Zhang¹ | Aiying Zhang¹[©] | Li Xiao¹ | Wenxing Hu¹ | Julia M. Stephen² | Tony W. Wilson³[©] | Vince D. Calhoun⁴ | Yu-Ping Wang¹



Pierpaolo Sorrentino^{a,1}, Rosaria Rucco^{b,c,1}, Anna Lardone^d, Marianna Liparoti^c, Emahnuel Troisi Lopez^c, Carlo Cavaliere^e, Andrea Soricelli^{c,e}, Viktor Jirsa^a, Giuseppe Sorrentino^{b,c,f,e}, Enrico Amico^{g,h,e*}

Original Article	CHRONIC STRESS
Pretreatment Brain Connectome Fingerprint Predicts Treatment Response in Major Depressive Disorder	Chronic Stress Volume 4: 1–8 © The Author(s) 2020 Article reuse guidelines: sagepub.com/journals-permissions DOI: 10.1177/2470547020984726 journals.sagepub.com/home/css ©SAGE
Siyan Fan ^{1,2} , Samaneh Nemati ^{1,2} , Teddy J. Akiki ^{1,2,3} , Jeremy Roscoe ^{1,2} , Christopher L. Averill ^{4,5} , Samar Fouda ^{1,2} , Lynnette A. Averill ^{1,2,4,5} , and Chadi G. Abdallah ^{1,2,4,5}	2020



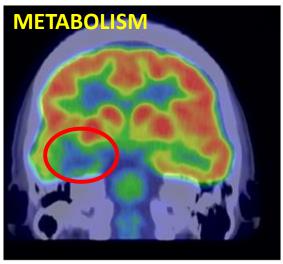
B.C. Munsell^{d,e,*}, E. Gleichgerrcht^a, E. Hofesmann^c, J. Delgaizo^a, C.R. McDonald^b, B. Marebwa^a, M.A. Styner^{d,e}, J. Fridriksson^f, C. Rorden^g, N.K. Focke^h, J.H. Gilmore^d, L. Bonilha^a

Why am I interested in brain fingerprinting?



temporal lobe epilepsy (TLE)







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ELECTROPHYSIOLOGY

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SEIZURE SYMPTOMS





Improvements in identifying the focus have not led to better outcomes

200 A 10	Contents lists available at ScienceDirect	
5-5-2-2	Epilepsy & Behavior	
ELSEVIER	journal homepage: www.elsevier.com/locate/yebeh	

Review

History of surgery for temporal lobe epilepsy 2017 Ali A. Asadi-Pooya^{a,b,*}, Cyrus Rostami^a

markable and revolutionary progress. This progress has resulted in tremendous advancements in understanding the underlying causes and pathophysiology of epilepsies. With the help of these technologies and advancements, we may now offer surgery as a safer therapeutic option to more patients who are suffering from drug-resistant temporal lobe epilepsy. However, the degree of improvement in surgery outcome has not been proportionate to the technological progress.

SUPPLEMENT ARTICLE

Lpilepsia

Algorithms in clinical epilepsy practice: Can they really help us predict epilepsy outcomes?

Lara Jehi 💿 🛛 2021

best. The "stepwise process" we follow to get to this end can be simplified into the following steps: (1) identify the surgical candidate, (2) localize the epilepsy, (3) resect the epileptogenic zone, and (4) the end to reach: the patient will be seizure-free. Over the past few decades, steps 1 through 3 have witnessed significant progress, yet our end has not improved proportionally. Heaps of academic literature has

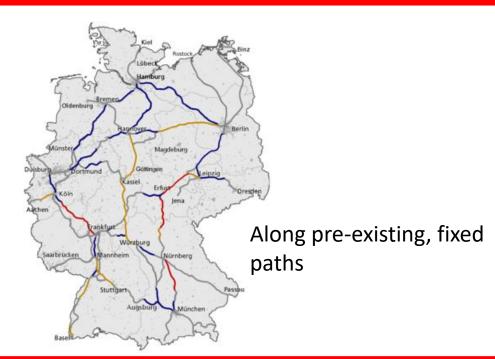
How does a focal seizure affect the rest of the brain?



Equal to all regions, transient



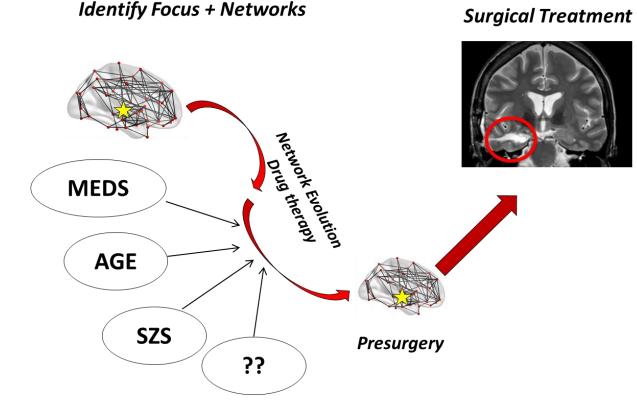
specific, along unique path, transient





Unique paths, progressive

Networks evolve prior to surgery



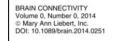
Functional connectivity homogeneity correlates with duration of temporal lobe epilepsy

Zulfi Haneef ^{a,b,*,1}, Sharon Chiang ^{c,1}, Hsiang J. Yeh ^d, Jerome Engel Jr. ^{d,e}, John M. Stern ^d *Z. Haneef et al. / Epilepsy & Behavior 46 (2015) 227–233*

Multivariate white matter alterations are associated with epilepsy duration

Thomas W. Owen¹, Jane de Tisi³, Sjoerd B. Vos^{3,4,5}, Gavin P. Winston^{3,5,6}, John S Duncan^{3,5}, Yujiang Wang^{1,2,3}, Peter N. Taylor^{1,2,3}

bioRxiv 2020



ORIGINAL ARTICLE

Evolution of Functional Connectivity of Brain Networks and Their Dynamic Interaction in Temporal Lobe Epilepsy

Victoria L. Morgan,¹ Bassel Abou-Khalil,² and Baxter P. Rogers¹

Epilepsia, 52(9):1741–1749, 2011 doi: 10.1111/j.1528-1167.2011.03196.x

FULL-LENGTH ORIGINAL RESEARCH

Cross hippocampal influence in mesial temporal lobe epilepsy measured with high temporal resolution functional magnetic resonance imaging

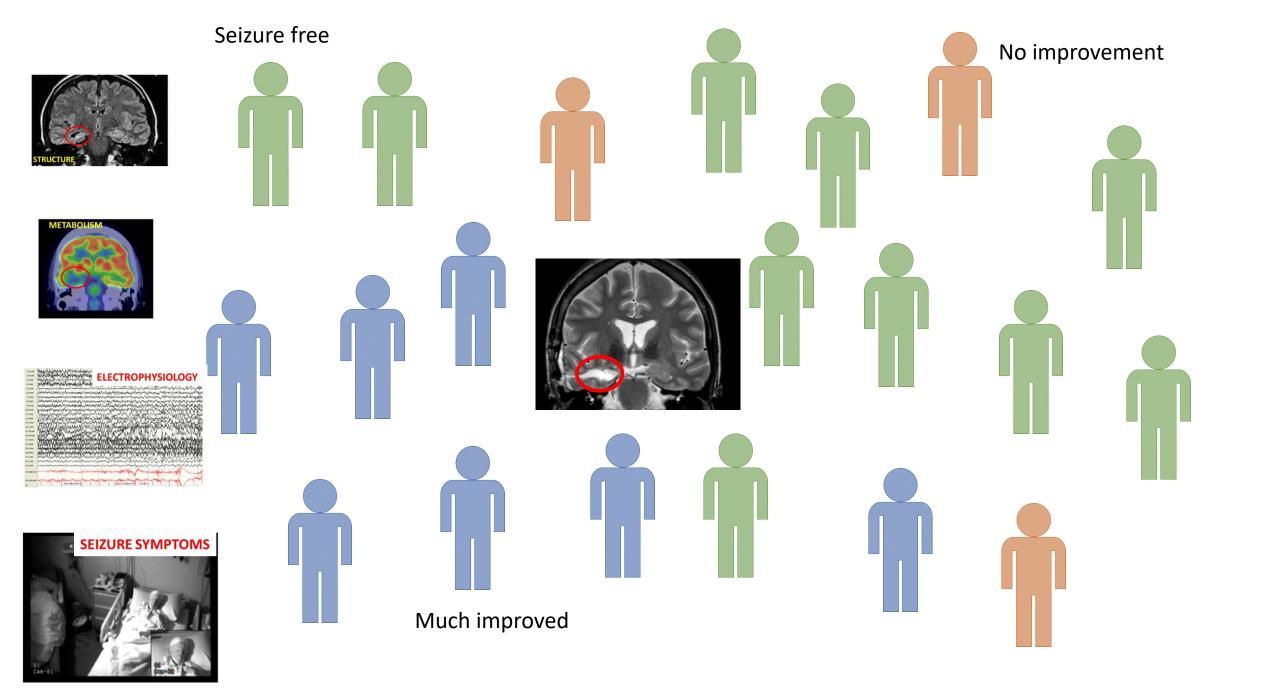
*Victoria L. Morgan, *Baxter P. Rogers, †Hasan H. Sonmezturk, *John C. Gore, and †Bassel Abou-Khalil **Our hypothesis – The temporal lobe epilepsy FINGERPRINT**

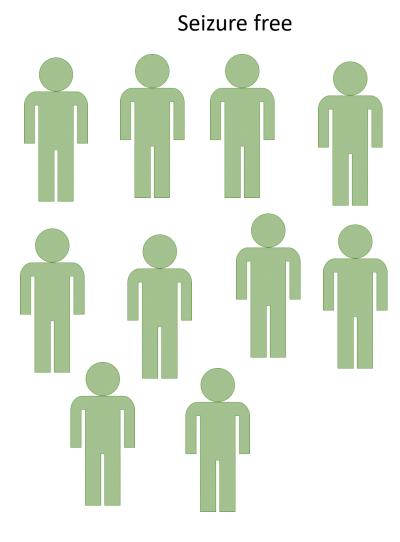
The focus may be the same, but the connections are different



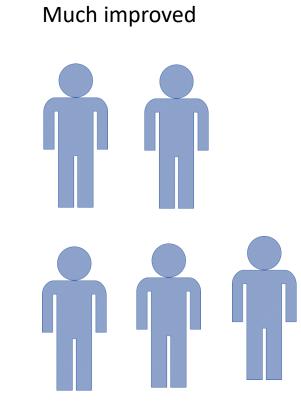
A pattern of network connections in the brain at the time of surgery that will result in seizure free outcome

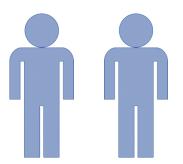




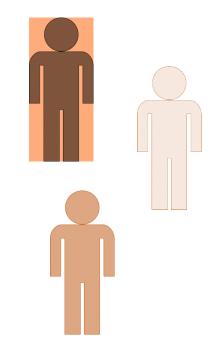








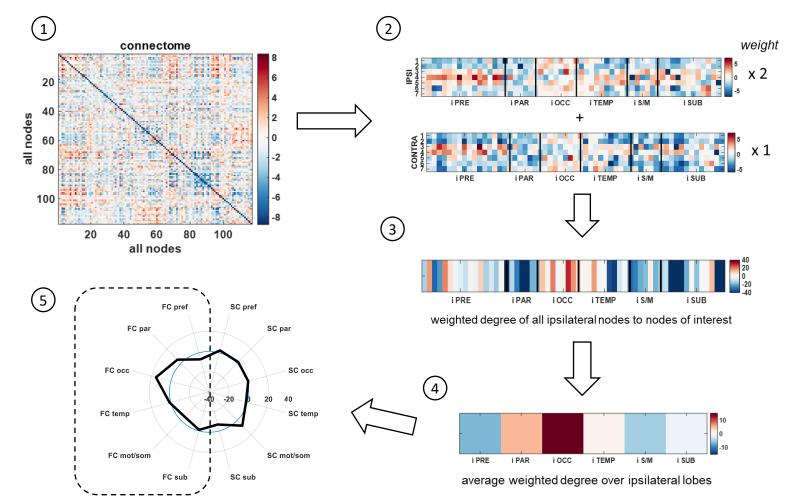
No improvement

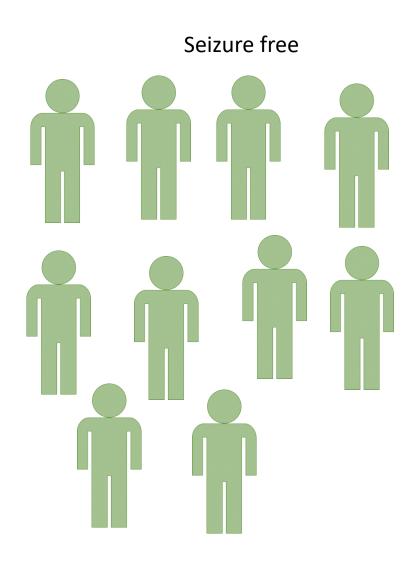


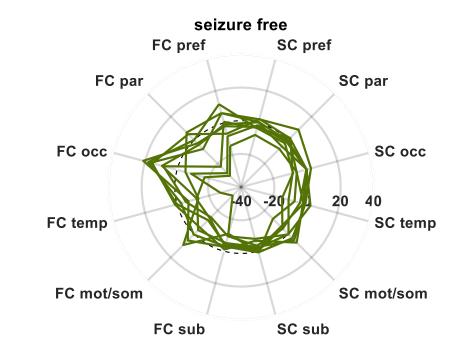
Different from pattern in various ways.

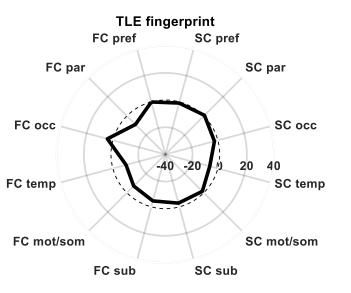
> Multi focus? Bilateral?



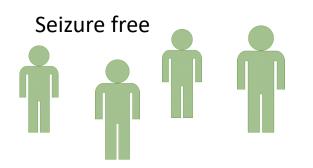


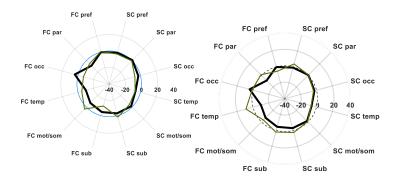






Now to test this!





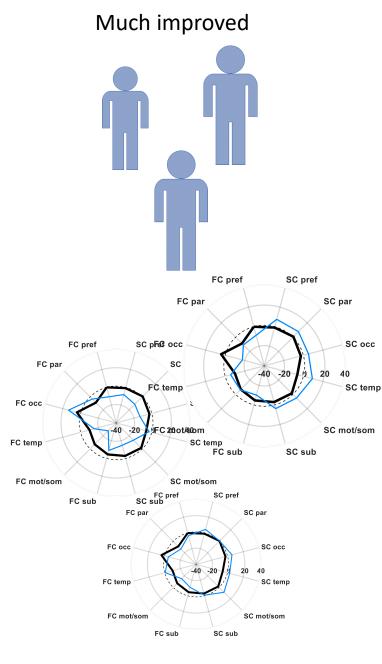
FC pref SC pref FC par SC par FC pref SC pref FC par SC par FC occ SC occ FC occ -40 -20 20 40 FC temp SC temp 20 40 .40 FC temp FC mot/som SC mot/som FC sub SC sub FC mot/som SC mot/som

FC sub

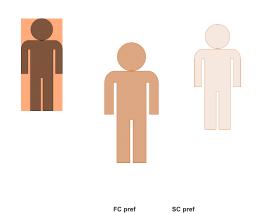
SC occ

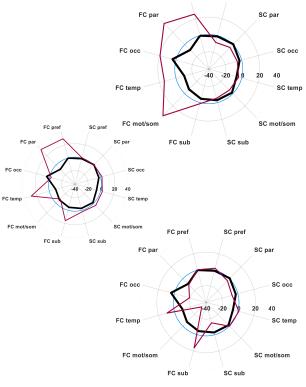
SC temp

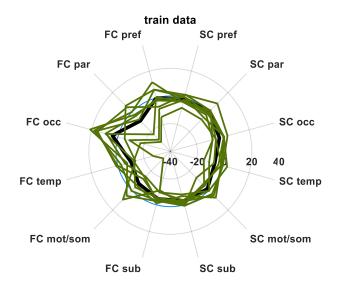
SC sub



No improvement





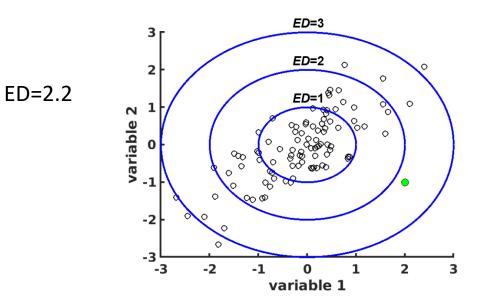


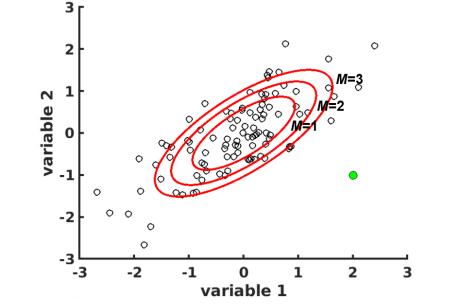
Similarity to fingerprint indicates better chance of good outcome.

Quantify similarity using 2 distances – Euclidean and Mahalanobis for magnitude and pattern

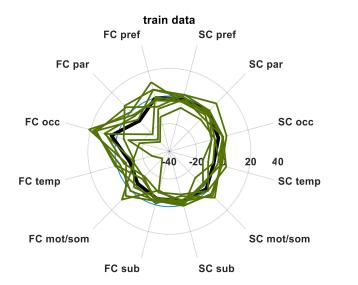
functional and structural connectivity distance

Also calculate total distance







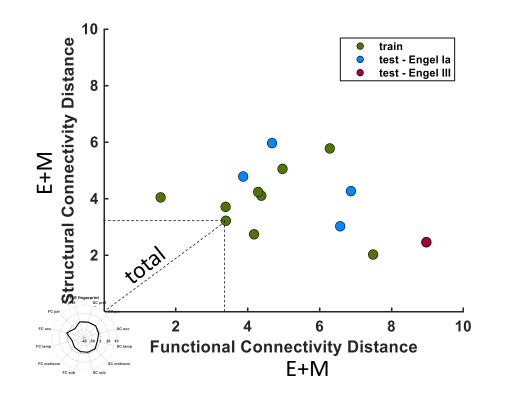


Similarity to fingerprint indicates better chance of good outcome.

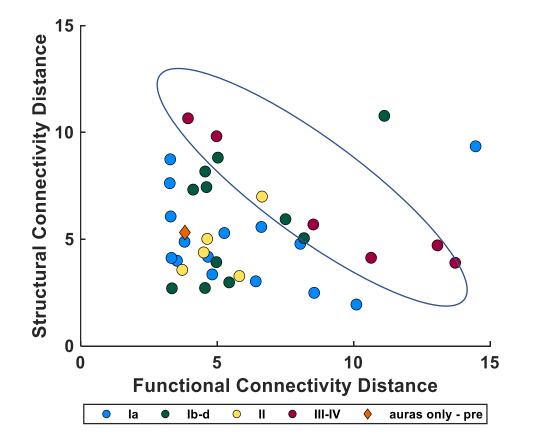
Quantify similarity using 2 distances – Euclidean and Mahalanobis for magnitude and pattern

functional and structural connectivity distance

Also calculate Total distance from (0,0) in plot



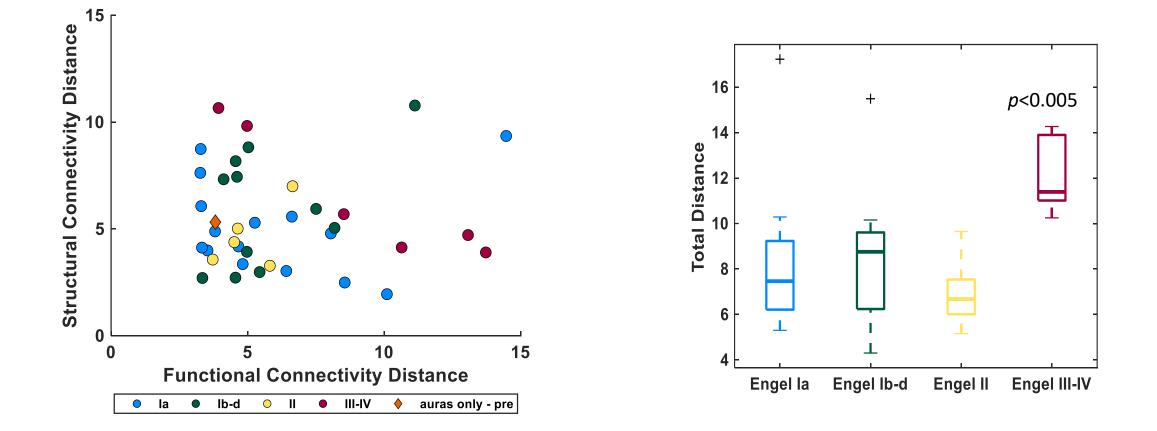
Testing data



Observations:

Functional vs. structural I and II mixed Few outliers

Class I	Free of disabling seizures
Class II	Rare disabling seizures ("almost seizure-free")
Class III	Worthwhile improvement
Class IV	No worthwhile improvement



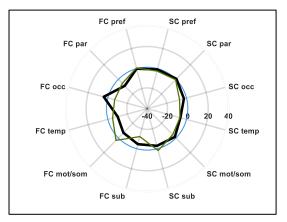
Patients with seizure recurrence have larger total distance from TLE fingerprint.

- Not in functional, structural, Euclidean or Mahalanobis only

temporal lobe epilepsy (TLE)

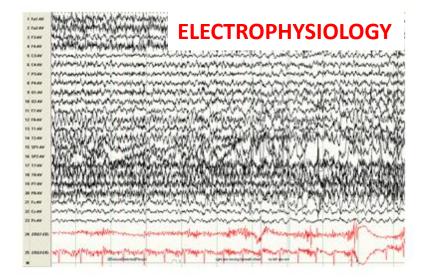






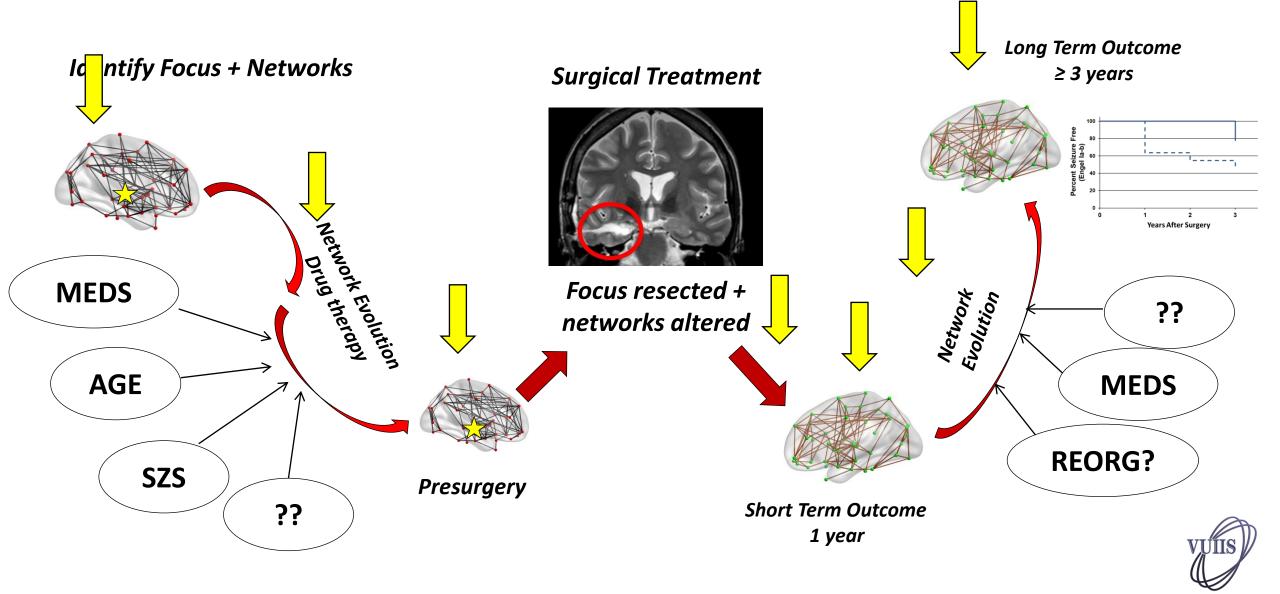




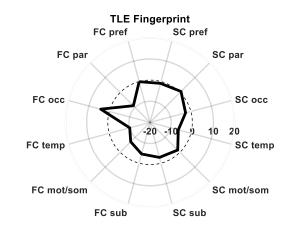




Moving forward









QUESTIONS??



