

Diffusion Tractography



Tract-Density Imaging

[Calamante Neurolmage 2010]

Single HCP subject TDI @ 0.2mm





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Single HCP subject TDI @ 0.2mm





Diffusion Tractography





Overview

- Goal of tractography
- Estimating Fibre Orientations BEDPOSTX
- Probabilistic Tractography PROBTRACKX
- ProbtrackX outputs
- Tractography limitations





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What is Tractography?



Post-mortem dissection of some white matter fibre bundles (tracts)

Williams, Gluhbegovic, and Jew, "The Human Brain`; Dissections of the Real brain", Virtual Hopstital, Universitoy of Iowa, 1997

Tractography

The post-imaging reconstruction of fibre bundles/ anatomical connections in the brain using a set of DW images. (in-vivo virtual dissection)





v₁ map Principal Diffusion Direction



Principal Diffusion Direction



Assumption:

Direction of maximum diffusivity (in anisotropic voxels) is an <u>estimate</u> of the major fibre orientation.



v₁ map Principal Diffusion Direction



Principal Diffusion Direction





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Uncinate

Inferior fronto-occipital

Problems of scale



Ohno et al. 2013

- White matter (dys)connectivity is thought to form the substrate for many different neurological and psychiatric disorders.



 Tractography provides non-invasive localisation and semi-quantitative biomarkers



- Tractography provides non-invasive localisation and semi-quantitative biomarkers



- Connections constrain function

- Different regions have distinct connectivity fingerprints







Passingham et al. 2002

Tractography outputs

Known white matter tracts









What does tractography offer?

- + non-invasive
- + in-vivo
- + whole brain
- + can address new questions



Lawes et al. 2008

...But

- low resolution (large bundles)
- indirect (diffusion paths)
- error prone (MRI is noisy)
- difficult to interpret quantitatively



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But is WM always coherently organised within a voxel?



Unfortunately not, complex fibre patterns (e.g. crossings) are very common at the voxel scale.

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Predictions from the tensor model no crossing fibres









How good is the DTI Model in regions with crossing fibres?

- In voxels containing two crossing bundles, the tensor ellipsoid is pancake-shaped (oblate, planar tensor).
- In voxels containing three crossing bundles, the tensor ellipsoid is spherical.
- In these areas, DTI \mathbf{v}_1 is meaningless.





Uncertainty on DTI Fibre Orientation Estimates

Repeat an acquisition many times and obtain the variability in v_1 from the different datasets.



Cones of uncertainty on DTI v_1

Jones, 2002



DTI model (dtifit)

Do we have to use the DTI model to estimate orientations? Not really, many models exist

Ball & sticks model (bedpostx)



$$s_j = s_0 \left[(1 - f) \exp(-b_j d) + f \exp(-b_j d(x_j T v)^2) \right]$$



Do we have to use the DTI model to estimate orientations? Not really, many models exist

DTI model (dtifit)

Ball & sticks model (bedpostx)





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DTI model (dtifit)

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Do we have to use the DTI model to estimate orientations? Not really, many models exist





- Anisotropic tensors (sticks) with isotropic background (ball)
- Fibre Orientations modelled explicitly and separated from isotropic partial volumes

$$Max number of sticks (user-defined)$$

$$s_j = s_0 \left[(1 - \sum f_n) \exp(-b_j d) + \sum f_n \exp(-b_j d(x_j^T v_n)^2) \right]$$



- Anisotropic tensors (sticks) with isotropic background (ball)
- Fibre Orientations modelled explicitly and separated from isotropic partial volumes

Max number
of sticks
(user-defined)
$$s_j = s_0 \left[(1 - \Sigma f_n) \exp(-b_j d) + \Sigma f_n \exp(-b_j d(x_j^T v_n)^2) \right]$$
Signal

Measured Signa for Gradient *j*



- Anisotropic tensors (sticks) with isotropic background (ball)
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Max number
of sticks
(user-defined)
$$s_{j} = s_{0} \left[(1 - \sum f_{n}) \exp(-b_{j} d) + \sum f_{n} \exp(-b_{j} d(x_{j} \nabla v_{n})^{2}) \right]$$
Measured Signal
for Gradient *j*
b-value for gradient *j*
(known)
Unit vector representing the direction of
gradient *j* (known)



- Anisotropic tensors (sticks) with isotropic background (ball)
- Fibre Orientations modelled explicitly and separated from isotropic partial volumes





Predictions from the ball and sticks model crossing fibres



Markov Chain - Monte Carlo (MCMC) Sampling




Output in Each voxel = Distributions of Parameters







Ball & Sticks Model Selection

- Model selection problem: One, two or more fibres within a voxel?
- Automatic Relevance Determination: Only estimate complexity that is supported by the data







ARD1

Measured Signal



 No benefit from including a 2nd fibre => 2nd volume fraction goes to zero

 Measured signal is explained better by more complex model => 2nd volume fraction is non-zero





 No benefit from including a 2nd fibre => 2nd volume fraction goes to zero

 Measured signal is explained better by more complex model => 2nd volume fraction is non-zero





- After running BedpostX all voxels will have estimated parameters for the maximum number of sticks requested.
- But due to ARD, the sticks that are not supported in a voxel will have an almost zero volume fraction.
- We use a threshold (e.g. >5%) to **exclude sticks with tiny volume fraction**.





Ball & Sticks Orientations

All sticks, with secondary ones thresholded (*f*_n>5%)





Ball & Sticks Orientations

All sticks, with secondary ones thresholded (*f_n*>5%)





DTI vs Ball & Sticks Orientations

DTI









A large portion of the WM supports crossing fibres

Coherence in orientations shows that we are not over-fitting (the ARD works)



Multi-Shell Diffusion Acquisitions Why bother?



Higher b value gives us more angular contrast!!!





Multi-Shell Diffusion Acquisitions Why bother?







Generalised Ball & sticks Model Gets best of both worlds

- Multi-shell model (or model=2) in Bedpostx options.
- Allows representation of multiple diffusivities within a voxel (rather than just one).
- More accurate model for multi-shell data & partial volume effects.



Human Connectome Project Data

*Jbabdi, Sotiropoulos et al, MRM 2012 * Sotiropoulos, Jbabdi et al, NeuroImage 2013

Faster bedpostx on GPUs



Hernandez et al, Plos One 2013



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DTI Streamline Tractography



Formally, we solve numerically the differential equation:



Mori S, Neuron 2006



DTI Streamline Tractography

But When to Stop? Heuristics to avoid error propagation. + Knowledge of the anatomy

Curvature Change Threshold: To avoid crossings of boundaries and very bended trajectories, impose a smoothness criterion.

Anatomical criteria (e.g. reach grey matter)





Streamline tractography can dissect major bundles



arcuate fasciculus



corpus callosum



uncinate fasciculus



cingulum bundle



inferior fronto-occipital



corona radiata



inferior longitudinal fasciculus



fornix



cerebellar tracts



- Use the major axis of the DTI ellipsoid as a fibre orientation estimate.

- Propagate curves within this vector field until empirical thresholds are exceeded.

- Major fibre bundles can be reconstructed.



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Streamlining reproducibility

Repeat an acquisition many times and repeat streamline tracking.

Due to uncertainty in v_1 , curves will not perfectly overlap

Create a map that shows the degree of overlap across the trials.

Streamlines from a single dataset





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Streamlines from a single dataset



Map that shows where results across datasets overlap



Low Reproducibility

High Reproducibility



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Streamlines from a single dataset



Map that shows where results across datasets overlap



Low Reproducibility

High Reproducibility



Probabilistic Tractography

- We normally have one dataset per subject, not many.
- Probabilistic Tractography as a two-step process:

a) Use DWI data and a model to infer a fibre orientation **and its uncertainty** in each voxel.

b) Use the estimates and the uncertainty to build a path probability map to a seed.



Probabilistic tractography

 But now, we no longer have a single direction at each voxel. How can we do tractography?





'Streamlining'

Probabilistic tractography Behrens et al, 2003, Parker et al. 2003, Hagmann et al 2003, Jones et al. 2004



Probabilistic tractography

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'Streamlining'

Probabilistic tractography Behrens et al, 2003, Parker et al. 2003, Hagmann et al 2003, Jones et al. 2004





Behrens et al, 2003 Parker et al, 2003

- Propagate N streamlines from a seed, but for each propagation step choose randomly an orientation from the underlying distribution.





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Define the degree of overlap at each location B, as:

M:number of streamlines that go through B N: total streamlines generated from A

This is the probability of a curve starting at A and going through B.





Behrens et al, 2003 Parker et al, 2003

- Can now propagate through isotropic regions (e.g. GM).
- Do not need to stop when anisotropy is low, as in deterministic tracking.
 - The high uncertainty will be reflected in the probability map.







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Probabilistic Tractography in Multi-Fibre Fields







Parker & Alexander 2003, Behrens et al, 2007

When multiple fibre orientations exist in a voxel, choose the one that is most compatible with the incoming trajectory.

Probabilistic Tractography in Multi-Fibre Fields Examples



Cortico-spinal tracts. 9 subjects 냥 Internal capsule ----- Primary motor cortex

Behrens et al, 2007



one fibre

two fibres

* If one fibre is modelled and we track through a crossing, a) we may not make it through the crossing, b) if we make it, the connectivity index will be relatively low.

Probabilistic Tractography in Multi-Fibre Fields Examples



Acoustic radiations.

9 subjects

Behrens et al, 2007





one fibre

two fibres



Path Probability Map

- Recall that it assesses how <u>reproducible</u> results are

- Often called "connection probability", "connectivity index", "connectivity strength". But it does not quantify how strong a connection is...

- Rather, how robust it is against noise/uncertainty







Probabilistic Tractography

Low Probability

High Probability



- Needs apart from orientation estimates, an estimate of their uncertainty. Does not need to be the ball and stick model, the DTI model can be used instead!

- Propagate streamlines repeatedly from a seed, but the orientation field is no longer deterministic. In each propagation step choose randomly an orientation from the underlying distribution.

- A connection probability value>=0 can be obtained from a seed A to any voxel in the brain B. This assesses the reproducibility of the path from A to B, along which water molecules preferably diffuse.





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ProbtrackX outputs

Known white matter tracts









 Because of the uncertainty propagation, the spatial distribution of paths is often very wide.



Low Probability	High Probability



Fdt GUI:



Once a seed is specified, prior anatomical knowledge can be imposed to assist the dissection of a specific tract.

Waypoint ROIs If a curve does not go through, it is discarded.

Exclusion ROI If a curve goes through, it is discarded.

Termination ROI If a curve goes through, it is terminated.



Cortico-spinal tract

Seed: M1, hand area





No ROIs



Cortico-spinal tract

Seed: M1, hand area





Exclusion: Mid-Sagittal plane



Cortico-spinal tract

Seed: M1, hand area





Waypoint: Internal Capsule



Corpus Callosum

Seed: dorsal PMC





No ROIs



Corpus Callosum

Seed: dorsal PMC





Waypoint: Corpus Callosum

Surfaces as constraints



No surface constraint



Surface as termination mask











- Register to standard space
 - b0 or FA -> TIw -> standard TIw
 - FA -> standard FA





- Register to standard space
 - b0 or FA -> TIw -> standard TIw
 - FA -> standard FA
- Don't transform masks -> diffusion space





- Register to standard space
 - b0 or FA -> TIw -> standard TIw
 - FA -> standard FA
- Don't transform masks -> diffusion space
- Don't transform diffusion -> standard space





- Register to standard space
 - b0 or FA -> TIw -> standard TIw
 - FA -> standard FA
- Don't transform masks -> diffusion space
- Don't transform diffusion -> standard space

Tell probtrackX about transform:

PROBTRACKX Probabilistic track	king 💻	
Data Options		
BEDPOSTX directory		
Single mask =		
Seed Image/Surface:		
E Seed space is not diffusion		
🗖 nonlinear		
Select Seed to diff transform		<u></u>
Select diff to Seed transform		<u>S</u>
□ surface		
 Optional Targets Waypoints masks Exclusion mask Termination mask Classification targets 		
Output directory:		
Go	Exit	Help

Connectivity - Why do we care?

 Tractography provides non-invasive localisation and semi-quantitative biomarkers



Connectivity - Why do we care?

- Tractography provides non-invasive localisation and semi-quantitative biomarkers



XTRACT: generating tracts for you



5 cm

Warrington et al. (2020) NeuroImage Warrington et al. (2022) Sci. Adv. Assimopoulos et al. (2024) Brain Struct. Func.



• Libraries of tractography protocols



- Libraries of tractography protocols
- Human (adult and neonate), macaque within FSL



- Libraries of tractography protocols
- Human (adult and neonate), macaque within FSL
- Many more available online



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- Easy...
- tract visualisation: xtract_viewer


XTRACT

- Libraries of tractography protocols
- Human (adult and neonate), macaque within FSL
- Many more available online
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XTRACT

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- tract visualisation: xtract_viewer
- QC: xtract_qc
- tract-wise summary statistics: xtract_stats



XTRACT: connectivity blueprints

 Easy connectivity blueprints: xtract_blueprint



Mars et al. (2018) eLife Warrington et al. (2022) Sci. Adv.





Warrington et al. (2022) Sci. Adv. Assimopoulos et al. (2024) Brain Struct. Func.



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Vertex A_v Vertex N_a Vertex A_x Vertex M; slf1 Vertex A, Cross-species divergence: mdlf mdlf mdlf fma xtract divergence fma Connections to vof WM tracts • Uses connectivity Macaque Neonate Adult blueprints to compare brains without **Predicted Macaque** Myelin map geometrical alignment I. Similarity maps Human 2. Parcellation/ROI Myelin Map compare Common predictions Space (XTRACT **Bundles**) Measured (Original) Macaque Myelin Map **High Cortical** Low Cortical Myelin Myelin

fma

Vertex A_v Vertex N_a Vertex A_x Vertex M; slf1 Vertex A, Cross-species divergence: mdlf mdlf mdlf xtract divergence fma Connections to vof WM tracts • Uses connectivity Macaque Neonate Adult blueprints to compare brains without **Predicted Macaque** Myelin map geometrical alignment Similarity maps Human 2. Parcellation/ROI Myelin Map compare Common predictions Space (XTRACT 3. Translate cortical maps **Bundles**) Measured (Original) Macaque Myelin Map **High Cortical** Low Cortical

Myelin

Myelin

ProbtrackX outputs

Known white matter tracts









Resulting matrix:

	?	?	?
?		?	?
?	?		?
?	?	?	



- Seed from blue
- Other ROIs are waypoints
- Fill first row of matrix Resulting matrix:

(0)		?	?	?
ROIs	?		?	?
Seed ROIs	?	?		?
	?	?	?	



- Seed from yellow
- Other ROIs are waypoints
- Fill first row of matrix Resulting matrix:





- Seed from green
- Other ROIs are waypoints
- Fill first row of matrix Resulting matrix:





- Seed from copper
- Other ROIs are waypoints
- Fill first row of matrix Resulting matrix:

















Fdt GUI: PROBTRACKX Probabilistic tracking 📃 Data Options BEDPOSTX directory /Users/ndcn0236/Work/projects/fsl_course/ni -Seed Space Multiple masks 😐 -Masks list-ROI 1 **ROI 3** ROI 4 Remove Image Load List Save List Add Image Seed space is not diffusion Optional Targets Waypoints masks Exclusion mask Termination mask Output directory: <u></u> Go Exit Help



Fdt GUI: PROBTRACKX Probabilistic tracking 📃 Data Options BEDPOSTX directory /Users/ndcn0236/Work/projects/fsl_course/n/ 🔄 Seed Space Multiple masks -Masks list-ROI 1 **ROI 3** ROI 4 Remove Image Load List Save List Add Image Seed space is not diffusion Optional Targets Waypoints masks Exclusion mask Termination mask Output directory: <u></u> Go Exit Help





Seed voxels

Resulting matrix:







Seed voxels

Resulting matrix:





Seed voxels



Resulting matrix:







Seed voxels

Resulting matrix:

Target ROIs



etc...





no contrast on conventional MRI

VL -> M1



MD -> PFC



Behrens et al, 2003 (probabilistic tractography)





RT VLc MD VPLo VPM CM VPI VPMpc

Rouiller et al, 1998 (BDA anterograde tracing)



Prior cortical parcelaltion





Resulting matrix: Target ROIs

M1 PMC PFC





Prior cortical parcelaltion





Resulting matrix: Target ROIs





Prior cortical parcelaltion





Resulting matrix: Target ROIs

M1 PMC PFC





Prior cortical parcelaltion





Resulting matrix: Target ROIs





Prior cortical parcelaltion





Resulting matrix: Target ROIs

M1 PMC PFC





Prior cortical parcelaltion





Resulting matrix: Target ROIs





Prior cortical parcelaltion





Hard thalamic parcellation





DBS for treatment of tremor in Parkinsons





Pouratian et al. JNS 2011



PROBTRACKX Probabilistic tracking -
BEDPOSTX directory /Users/ndcn0236/Work/projects/fsl_course/ni Seed Space Single mask Seed Image/Surface: Seed space is not diffusion Optional Targets Vaypoints masks Exclusion mask Classification targets Targets list
Add Image Remove Image Load List Save List Output directory:
Go Exit Help



Fdt GUI: PROBTRACKX Probabilistic tracking -Options Data BEDPOSTX directory /Users/ndcn0236/Work/projects/fsl_course/ni Seed Space Single mask 😐 Seed Image/Surface: ROI 4 Seed space is not diffusion Optional Targets Waypoints masks Exclusion mask Termination mask Classification targets Targets list-ROI 1 ROI 3 Remove Image Load List Save List Add Image Output directory: 3 Go Exit Help


ROI 1 voxels





Resulting matrix: ROI 1 voxels ? ? ? ? ? ? ? ? ? ? ? ? ? ? ? ?



ROI 2 voxels



	Mask2
Maskl	Matrix3



Data	Options				
⊢Ba	sic Options				
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	■ Matriv3:	Mask1 x Ma	ask2 Matriv		

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📕 Loopcheck	
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→ Matrix Options	
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Matrix2: Seed x Mask2 Matrix	
Matrix3: Mask1 x Mask2 Matrix	

Exit

Help

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PROE	Fdt GUI: BTRACKX Probabilistic tracking
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	Atrix Options
	Matrix1: Seed x Seed Matrix
	Matrix2: Seed x Mask2 Matrix
	Matrix3: Mask1 x Mask2 Matrix
	Go Exit Help

Dense connectome

Cortical seed (matrix1)



WM seed (matrix3)



Dense connectome

Cortical seed (matrix1)



WM seed (matrix3)



Cortical vertices



Dense connectome

Cortical seed (matrix1)



WM seed (matrix3)



Cortical vertices







Medial area 6 contains two distinct regions with very different connectivity: SMA and Pre-SMA





Medial area 6 contains two distinct regions with very different connectivity: SMA and Pre-SMA





Medial area 6 contains two distinct regions with very different connectivity: SMA and Pre-SMA





Medial area 6 contains two distinct regions with very different connectivity: SMA and Pre-SMA

Can we define a border based on a change in connectivity profile?















Rest of brain

	이가 사람은 성적 위에 가장 할 것 같다. 말 말 같은 것 같아요. 물건 것 같아요. 물건 것 같아요. 물건 것 같아요.	승규는 영화 전 문화
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	이 소문을 많은 것 않는 것 같아? 이 말을 수 없는 것 같은 것 같아요. 것 같아요. 것 같아요. 가지 않는 것 같아요. 것 같아요. 것 같아.	이 같은 것은 것은 것은 것은 것을 가지 않는 것을 알려요. 이 가지 않는 것은 것은 것을 가지 않는 것을 하는 것을 했다. 같은 것은
는 것은 이상 상황에서 이상되어도 있다. 2015년 2월 14일 - 11일 -	이가 사람 한국가 가장 있는 것 못했다. 전 이는 것 이는 가장 가장 가지 않는 것 것 같아. 영화 것 같아. 영화 것 것 같아. 영화 것 같아. 영화 것 같아. 영화 것 같아. 영화 것 것 같아. 영화 것 같아. 영화 것 것 같아. 이 것 같아. 영화 것 같아. 영화 것 같아.	
白色的 法保证的 建分配的		法保险 化物理试验检试验 机运行机 建苯
	이 아이들은 것 같은 것	化二乙酸 化化合成基本正式化合成合金 建筑
است والداري بالحالي ويعاد الأورية والمارية ويرارية المعادية. المحادثة المحادية المحاد	는 가수는 물건에 물건을 전한 것이 가 이 이 가지 않는 것은 것이 있는 것이 가지 않는 것이 같은 것이 가지 않는 것이 있다. 가지 않는 것이 있는 것이 가지 않는 것이 있는 것이 있는 것이 있다. 이 제 에 전 1995년 에 에너지 않는 것이 같은 것이 있는 것이 같은 것이 있는 것이 있는 것이 있는 것이 있는 것이 있다. 이 가지 않는 것이 있는 것이 있는 것이 있는 것이 있는 것이 있는 것 - 이 가 것이 같은 것이 있는 것이 있	ار در این از این از این از مینیان به در این از میشود با در این از میشود. از میشود از میشود از میشود. از میشود از میشود. از میشود از م
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		남자 지수는 사람을 가지면 눈 문제 환자했다.
말 다 같은 것은 것은 것을 것을 것을 했다.	가 있는 것은 것을 가장 가장에 가장에 가장 것을 수 있는 것을 가장에 가장을 가지 않는 것을 가장을 가장을 가지 않는다. 같은 것은 것은 것은 것은 것을 것을 것을 하는 것을 것을 것을 하는 것을 것을 수 있는 것을	
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	二氢二磷酸 建制物油 医肥胖性 建磷酸酸 机石材料 化铁合合物 医分子检查力能 铁石头的过去式	
	化自己化物 医结核神经 化可能振动 化乙基乙基乙酰乙基乙烯乙酯 计分配分配数 计算法 的复数 网络马克尔斯	
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지지 것 같은 동네 방송가 봐.	1997 · 小学校、多生学校的学校和教师的主题和学校的教师,和学校、学校、学校、学校、学校、学校、教育、教育、教育、学校、学校、学校、 1998 · 1999 · 1999 · 1999 · 1997 · 1999 · 1999 · 1999 · 1999 · 1994 · 1997 · 19	يەربىي ئىلىكى بىلىرى بىلىكى بىلىك ئىلى بىلىكى بى
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Seed voxels

Crosscorrelation matrix

Seed voxels



Rest of brain

Seed voxels



algorithm





•Clusters in the re-ordered matrix represent seed voxels with similar connectivity

•Breaks between clusters represent where connectivity patterns change





•Clusters in the re-ordered matrix represent seed voxels with similar connectivity

•Breaks between clusters represent where connectivity patterns change



Johansen-Berg et al. 2004







Temporo-parietal junction Lateral Parietal Mars 2012 Mars 2011



Substantia Nigra Menke 2010





Lateral pre-motor Tomassini 2007



Striatum Tziortzi 2013



Broca's area Klein 2007



Insular cortex Cerliani 2012



Medial prefrontal Johansen-Berg 2004



Thalamus Behrens 2003



Amygdala Saygin 2011



Occipital cortex Thiebaut de Schotten 2013

ProbtrackX outputs

Known white matter tracts









Overview

- Goal of tractography
- Estimating Fibre Orientations BEDPOSTX
- Probabilistic Tractography PROBTRACKX
- ProbtrackX outputs
- Tractography limitations







• Number of axons connecting 2 areas?



- Number of axons connecting 2 areas?
- Proportion of axons from a seed that reach a target?



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- "Integrity" of the connecting white matter ... –Effective conductivity?
 - -Degree of myelination?
 - -Packing density?



- Number of axons connecting 2 areas?
- Proportion of axons from a seed that reach a target?
- "Integrity" of the connecting white matter ... –Effective conductivity?
 - -Degree of myelination?
 - -Packing density?
- What are we measuring?
 - -The probability that the **dominant** path through the <u>diffusion field</u> passes through this region.



- They may reflect "Connection Strength"
- But they do also reflect other uninteresting factors, such as:

<u>Connection length</u>: Longer connections have smaller probability than shorter ones

<u>Geometric complexity</u>: Probabilities of connections that go through regions of complex structure will be smaller than connections than go through more coherent regions

- Cross-subject comparison of the same tract is more meaningful than comparing different tracts



Can we trust tractography?

Is the direction of least hindrance to diffusion a good proxy for fibre orientation?



mapping between axon geometry and diffusion profile can be ambiguous

White matter organisation can be surprising



Whole mouse brain Electron Microscopy! Mikula Binding Denk, Nature Methods 2012

Validation: comparison with classical chemical tracing





point of entry within the CB



DBS for treatment of tremor in Parkinsons





Pouratian et al. JNS 2011



The Human Connectome Project www.humanconnectome.org





That's all folks



That's all folks





Diffusion Tractography