Title:

Euclidean distance as a measure to distinguish ventral and dorsal white matter connectivity in the human brain

Authors:

Philipp Kellmeyer[1,2], Magnus-Sebastian Vry[3]

Affiliations

1 Translational Neurotechnology Lab, Department of Neurosurgery, Medical Center – University of Freiburg, Germany

2 BrainLinks-BrainTools Cluster of Excellence, University of Freiburg, Germany

3 Department of Psychiatry, Medical Center - University of Freiburg, Germany

Corresponding author: Philipp Kellmeyer, Email: philipp.kellmeyer@uniklinik-freiburg.de

Addresses

Philipp Kellmeyer:

Translational Neurotechnology Lab (AG Ball), Universitätsklinikum Freiburg, Engelbergerstr. 21, D-79106 Freiburg im Breisgau, Germany

Magnus-Sebastian Vry:

Klink für Psychiatrie und Psychotherapie, Universitätsklinikum Freiburg, Hauptstr. 5, D-79104 Freiburg im Breisgau, Germany

Corresponding Author

Philipp Kellmeyer

E-mail: philipp.kellmeyer@uniklinik-freiburg.de

Phone: ++49-761-270-87570

ORCID code

Philipp Kellmeyer: 0000-0001-5538-373X

Abstract

Fiber tractography based on diffusion tensor imaging (DTI) has become an important research tool for investigating the anatomical connectivity between brain regions in-vivo. Combining DTI with functional magnetic resonance imaging (fMRI) allows for the mapping of the structural and functional architecture of large-scale networks for cognitive processing. This line of research has shown that ventral and dorsal fiber pathways subserve different aspects of stimulus- and task-dependent processing in the human brain.

Here we investigate the feasibility and usefulness of Euclidean distance as a geometric measure to differentiate ventral and dorsal long-range white matter fiber pathways between parietal and inferior frontal cortex in a body of studies that used probabilistic tractography.

We show that, on average, ventral pathways between parietal and inferior cortex have a significantly longer Euclidean distance in 3D-coordinate space than dorsal pathways. Thus, Euclidean distance could provide a simple means to assess patterns of connectivity in fMRI studies in the absence of DTI data. This would allow for a much broader meta-analytic assessment of general patterns of ventral and dorsal large-scale connectivity for different cognitive operations in the brain.

Keywords: DTI; Dorsal; Euclidean Distance; Structural Connectivity; Ventral

Introduction

In recent years, fiber tracking based on diffusion tensor imaging (DTI) and functional magnetic resonance imaging (fMRI) have allowed for the anatomically constrained investigation of large-scale white matter pathways for cognitive processing in different domains. This research has demonstrated that ventral and dorsal fiber systems subserve different aspects of cognitive processing in a variety of domains like vision, attention, language and the motor system. With respect to the white matter fiber pathways supporting language processing, for example, DTI-based fiber tracking has identified a ventral pathway via the middle longitudinal fascicle and the extreme capsule fiber system that connects parietal and posterior temporal areas with inferior frontal areas - in addition to the classic dorsal fiber pathway via the arcuate fascicle and superior longitudinal fascicle fiber system. (Frey et al. 2008; Saur et al. 2008; Saur et al. 2010; Kellmeyer et al. 2013).

DTI as a research tool for the in-vivo mapping of white matter fiber systems in the human brain has become widely available about a decade ago. From before the advent of DTI, however, we have a large body of functional imaging studies in which the patterns of structural connectivity between the brain regions identified with fMRI (or PET) remain unexplored. Thus, it would be useful if the information on the human fiber system from DTI-based fiber tracking we accumulate now could be harvested to retroactively infer patterns of connectivity between brain coordinates in the absence of DTI data. This would allow for much larger meta-analyses on patterns of large-scale connectivity across a variety of cognitive domains. This, in turn, would increase the aggregated evidence on large-scale networks for cognitive processing based on functional neuroimaging.

To this end, we explore systematic differences in Euclidean distance (ED) as a geometric measure to distinguish between ventral and dorsal fiber pathways in studies that have used DTI-based probabilistic fiber tracking. Our aim is to investigate, whether the anatomical segregation into ventral and dorsal fiber pathways, demonstrated by probabilistic fiber tracking, can be captured by a simple measure of spatial distance. Differentiating ventral and dorsal connectivity between brain coordinates by Euclidean distance would provide a simple tool for assessing patterns of connectivity in the brain even in the absence of DTI data.

Euclidean distance has been previously shown to relate to functional connectivity between brain regions in resting state fMRI in the context of psychiatric disorders. In one study in patients with childhood-onset schizophrenia, the differences between subjects in the topology of

functional networks in the resting-state was reflected by variation in ED between connected regions (Alexander-Bloch et al. 2013). Another study which compared resting-state fMRI between elderly patients with depression and controls showed increased ED between connected brain regions in the depressed group (Bohr et al. 2013). With respect to long-range anatomical connectivity via white matter fiber pathways in the brain, however, no geometrical relationships, power laws or allometric principles based on in-vivo DTI imaging data is available so far to our knowledge.

Here we analyzed Euclidean distances between coordinates in parietal and inferior frontal brain regions for which the pattern of ventral or dorsal connectivity has been demonstrated previously by DTI-based probabilistic fiber tracking. We chose probabilistic fiber tracking because we have substantial experience in developing and applying this fiber tracking method (Kreher et al. 2008). In this previous work, probabilistic fiber-tracking investigated ventral and dorsal fronto-parietal fiber pathways in the context of language processing (Saur et al. 2008; Saur et al. 2010), motor imagery (Hamzei et al. 2015; Vry et al. 2012), attention (Suchan et al. 2012; Umarova et al. 2009) and mental arithmetic (Klein et al. 2013) we observed that ventral fronto-parietal fiber pathways seem to travel much longer distances in the brain than dorsal fronto-parietal pathways. However, experiments on perception show that visual estimation of geometric distance between two points often results in an overestimation of the distance, with relative errors increasing with the length of curvilinear paths (that were used as visual cues, called the "detour effect") (Faineteau et al. 2003). Based on our observation from these tractography studies, we hypothesize that the ventral pathway connecting parietal and inferior frontal cortex is on average longer than the dorsal pathway. To test this hypothesis, we analyze systematic geometric differences in ventral and dorsal connectivity between parietal and inferior frontal regions quantitatively by measuring Euclidean distance in 3D coordinate space in these studies.

Methods

To test this approach, we confined the analysis to two well characterized regions in terms of fiber connectivity, the parietal lobe and the inferior frontal gyrus (opercular and triangular parts). As we have studied white matter fiber connectivity using probabilistic fiber-tracking in variety of contexts and have a good understanding of the methodology of these measurements,

we confined this investigation to studies using probabilistic fiber tracking (Kellmeyer et al. 2013; Suchan et al. 2012; Umarova et al. 2009; Vry et al. 2012).

Identification of coordinates

For the list of studies that were used for identifying the coordinates and their respective connectivity pattern from the fiber tracking see Table 1. All studies that were included in the analysis reported brain coordinates in Montreal Neurological Institute (MNI) space.

Measuring geometrical distance

We used Euclidean distance (ED) $d = \sqrt{(x_2 - x_1)^2 + (y_2 - y_1)^2 + (z_2 - z_1)^2}$ to measure the distance between two coordinates, located in the inferior parietal and inferior frontal cortex respectively, in three-dimensional MNI space.

We measured ED between two coordinates for which connectivity has been demonstrated previously by DTI-based probabilistic fiber tracking. Fiber pathways from the tracking results were either classified as dorsal or ventral in accordance with the classification from the respective studies (**Fig. 1**).

Figure 1 Patterns of dorsal and ventral connectivity from the studies used for measuring Euclidean distance: Panels (A)-(D) show the different Studies with each panel including the

seeds from all studies. (A) Umarova et al. 2009 (yellow spheres), (B) Vry et al. 2012 (blue spheres), (C) Kellmeyer et al. 2013 (green spheres), (D) Klein et al. 2013 (orange spheres). Blue dotted line = dorsal pathway; red dotted line = ventral pathway. For MNI coordinates (see Table 1).

Comparing mean Euclidean distance between dorsal and ventral connections

The null-hypothesis, that Euclidean distance between dorsal and ventral pathways connecting parietal and inferior frontal cortex does not significantly differ, was addressed in a two-sample t-test comparing the mean ED between pathways classified as dorsal or ventral by DTI-based probabilistic fiber tracking.

Results

Euclidean distance between ventral and dorsal fronto-parietal pathways

Table 1 shows the frontal and parietal MNI coordinates (x,y,z) from the 4 studies that were used in our analysis and the respective Euclidean distance between them.

Study	Parietal region	Parietal MNI coordinates			Frontal region	Frontal MNI coordinates			Euclidean distance	Ventral / dorsal
Umarova et al. 2009	R IPS	27	-51	51	R IFGpo	54	12	12	78.86	dorsal
	R IPS	27	-51	51	R IFG pt	51	21	-3	93.15	ventral
Vry et al. 2012	LIPSa/hIP2	-48	-42	51	L PMv	-54	3	39	58.56	dorsal
	L PSm/hIP2	-42	-51	51	L PMv	-54	3	39	56.60	dorsal
	L IPSp/hIP1	-33	-54	42	L IFG pt	-45	24	-3	90.85	ventral
	L IPSp/hIP1	-33	-54	42	L IFG pt	-39	45	3	106.57	ventral
Kellmeyer et al. 2013	L IPLm	-39	-42	39	L IFG po	-45	12	27	55.64	dorsal
	L IPLa	-39	-42	45	L IFG po	-48	12	27	57.63	dorsal
	R IPLa	36	-42	45	R IFG po	45	12	24	58.63	dorsal
	L IPLp	-27	-60	36	L IFG po	-48	12	27	75.54	dorsal
	R IPLa	36	-42	45	R IFG pt	45	36	12	85.17	ventral
	L IPLa	-39	-42	45	L IFG pt	-45	39	9	88.84	ventral
	L IPLp	-27	-60	36	L IFG pt	-45	39	9	104.18	ventral
Klein et al. 2013	L IPSa	-45	-38	45	L IFG po	-49	11	26	52.71	dorsal
	L IPSp	-30	-71	49	L IFG pt	-53	19	26	95.70	ventral
	R IPSp	41	-49	45	R IFG pt	53	34	19	87.80	ventral
	R SPLp	19	-68	57	R IFG pt	53	34	19	114.04	ventral

Table 1 Studies using DTI-based probabilistic fiber tracking with coordinates in parietal and inferior frontal cortex
Abbr.: L=left hemisphere. R=right hemisphere; a=anterior. p=posterior. m=medial; IPS=intraparietal suclcus; hIP=intraparietal sulcus (1.2.3); IPL=inferior parietal lobule; IPS= intraparietal sulcus; PMv=ventral premotor cortex; IFG=inferior frontal gyrus. po=pars opercularis. pt=pars triangularis.

The two-sample t-test comparing median Euclidean distance (ED) between brain coordinates in parietal and inferior frontal cortex showed that ED for coordinates connected by ventral pathways (*N*=9, mean 96.26, standard deviation 9.84, standard error 3.28) are significantly (p<0.0001) longer than for dorsal pathways (*N*=8, mean 61.77, standard deviation 9.75, standard error 2.43) (**Figure 2**).

Figure 2 Plot showing mean Euclidean distance for dorsal and ventral pathways between the parietal and inferior frontal coordinates from **Table 1**.

Discussion

In this sample of DTI-based probabilistic fiber tracking studies, we found a significant difference for Euclidean distance to distinguish ventral from dorsal pathways connecting inferior frontal and parietal areas. One explanation for this pattern could be that ventral parieto-frontal connections tend to connect to more anterior-inferior subregions in IFG (pars triangularis, pars orbitalis), whereas dorsal pathways tend to connect to more anterior-superior (pars opercularis). This raises the question, whether there is an anatomical divide within IFG that determines ventral or dorsal connectivity to posterior brain regions (like parietal and temporal cortex)?

To replicate and validate our findings and to address this question, we propose the following steps for further analyses: First, a voxel-by-voxel connectivity analysis between defined volumes in parietal cortex (comprising, for example, inferior and superior parietal lobule) and inferior frontal cortex (comprising inferior frontal gyrus, pars opercularis triangularis and orbitalis) could be performed to map the complete ventral/dorsal connectome between these two volumes. Furthermore, the results could be cross-validated on studies using different fiber-tracking methods. If these studies corroborate our initial finding here, we would propose to use a measure based on Euclidean distance (for example: minus one standard deviation of the mean ED of ventral connections) to distinguish ventral from dorsal connectivity. In the example of our study this would mean, that any given pair of parietal and inferior frontal coordinates that has an ED > 83 (mean ED for ventral connections – 1 STD: 91.84-8.956 = 82.89) is connected via a ventral pathway with high probability (95%).

Previous research with DTI-related connectivity-based parcellation (Johansen-Berg and Rushworth, 2009) approach has already shown the pattern of connectivity for inferior frontal cortex (Anwander et al. 2007), orbitofrontal cortex (Kahnt et al. 2012) (using fMRI) and inferior parietal cortex (Ruschel et al. 2014) separately. However, no voxel-to-voxel whole connectome between parieto-frontal regions is available thus far. If such a whole region-to-region connectome was available, the ED between all coordinates for which either ventral and/or dorsal connectivity was found by the fiber tracking could be measured. Then the mean ED could be compared between ventral and dorsal pathways and the boundary ED value that reliably (>95% of cases) distinguishes between ventral and dorsal connectivity could be computed. In addition, this data could then be used to delineate the anatomical divide at which fronto-parietal connectivity segregates between ventral and dorsal pathways.

This measure would be a particularly useful tool for large-scale meta-analysis of fMRI studies in which no DTI data is available. In such a meta-analysis, one could address the question whether the network involved in particular cognitive operations, say speech repetition based on semantic versus phonological features, can be assigned to different pathways according to the MNI coordinates that were identified in fMRI studies on the topic. Information on the structural connectivity of "active" brain regions in neuroimaging studies is of interest for both cognitive and clinical neuroscientists.

If ventral fiber pathways from posterior-superior to anterior-inferior brain regions were indeed systematically longer than dorsal pathways, we might ask what the neurocomputational effects of this difference, for example on cognition, could be?

For cognitive operations within widely distributed perisylvian networks connected via dorsal and ventral fiber tracks, the fiber length can indeed make a difference with respect to computational efficiency. It has been shown, for example, that the average conduction delay in axonal transmission scales with the average length of long-range connections and average axon diameter (Herculano-Houzel et al. 2010). In the healthy brain, differences in domain-independent cognitive processing speed seem to correspond to the length of white matter pathways connecting regions involved in the tasks used (Behrman-Lay et al. 2014). Synthesizing findings from functional and structural imaging studies, dorsal and ventral pathways seem to support separable computations within (and across) cognitive domains. In the case of language, for example, the ventral temporo- and parieto-frontal pathways subserve "higher-order" aspects of language like syntactic or lexical-semantic processing (Rijntjes et al. 2012; Saur et al. 2008a). The same seems to be the case for higher-order aspects of the cognitive motor system in the context of motor action, for example the imitation of gestures or pantomime (Hamzei et al. 2015; Vry et al. 2012). More generally, it has been proposed that the ventral pathway might subserve processing that is time-independent and related to semantic aspects and that the dorsal pathway is involved in sequential processing in a time-sensitive manner for rapid sensorimotor integration (Hamzei et al. 2015; Kellmeyer et al. 2013; Rijntjes et al. 2012; Weiller et al. 2011).

From a clinical perspective, whether a particular pathway is significantly longer than another pathway and/or may traverse regions that are more prone to damage (e.g. due to differences in territorial vascular supply) may also entail differences in neurological vulnerability. These differences in vulnerability between ventral and dorsal pathways, in turn, may partly explain dissociable clinical phenotypes of cognitive deficits after brain injury, for example the different aphasia syndromes (Kümmerer et al. 2013).

A clear limitation of our exploratory work here is that we used only a very selected body of studies to explore our hypothesis. The motivation behind this restriction, however, was that we wanted to explore differences in geometric distances between ventral and dorsal fiber pathways in DTI studies that clearly demonstrate the existence of white matter pathways between brain regions. As probabilistic fiber tracking in DTI data is more robust both against detecting false positive connections and false negatives (e.g. through crossing fibers) than deterministic fiber tracking (Kreher et al. 2008), we restricted the exploration to DTI studies using probabilistic tracking.

Furthermore, the precise relationship between ED and actual fiber length cannot be inferred from the analysis presented here. It may well be the case, for example, that the given pathways are optimal in terms of their distance under the given physical and anatomical constraints on the actual brain's global shape, local curvature, folding and gyration, Moreover, as the pathways are not necessarily monosynaptic connections, the ventral pathway is relayed in insular cortex for instance, we do not know how ED relates to whether a particular pathway is mono- or polysynaptic. In this respect, it could be of interest to investigate whether the relationship between Euclidean distance and patterns of ventral and dorsal connectivity may reflect some kind of underlying allometric scaling law for governing principles of long-range white matter fiber connectivity (Bush and Allman, 2003; Changizi, 2001; Gould, 1975; Herculano-Houzel et al. 2010; Zhang and Sejnowski, 2000). For example, it may be the case that the need for long-range connections follows from the fact, that the volume of white matter increases at 4/3 power of the volume of gray matter (Zhang and Sejnowski, 2000).

From a comparative and evolutionary perspective, it might be interesting to investigate how geometric measures of connectivity, whether Euclidean distance or other measures, are useful metrics to differentiate patterns of fiber connectivity in other mammalian species, for example non-human primates for which neuroanatomical data in 3D coordinate space is available (Markov et al. 2013). This may give further insight into cross-species principles of white matter formation in nervous systems, particularly links between global principles like minimal wiring and more local organizing principles of neuronal circuitry and topographical maps (Perin et al. 2011; Rothschild and Mizrahi, 2015). Such information would also be important to link the global geometry of the white matter fiber system to the hierarchical organization of multimodal cortical areas (Zhang and Sejnowski, 2000).

Another potentially promising approach could be to explore geometric measures based on manually or automatedly retracing the course of fiber pathways in the raw data of the individuals' DTI. We speculate that this would most likely result in a non-euclidean, curvilinear measure of geometric distance between given coordinates (Styner et al. 1999). In terms of the underlying geometrical principles that may best describe long-range fiber connectivity, algorithms for DTI analysis based on Riemannian and geodesic distance measures have been used that could have advantages over the currently used voxel-based analyses (Astola et al. 2007; Hao et al. 2011; Wang et al. 2014).

Further studies should also address, whether differences in Euclidean distance related to ventral and dorsal connectivity can also be found in connections between other brain regions (for

instance fronto-temporal, occipito-temporal etc.). DTI-based connectivity studies that use different tracking methods such as deterministic fiber tracking should also be examined in order to facilitate large-scale meta-analyses on the structure and function of dorso-ventral pathway systems based on fMRI studies.

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