

REVIEW ARTICLE

Dissecting the uncinate fasciculus: disorders, controversies and a hypothesis

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The uncinate fasciculus is a bidirectional, long-range white matter tract that connects lateral orbitofrontal cortex and Brodmann area 10 with the anterior temporal lobes. Although abnormalities in the uncinate fasciculus have been associated with several psychiatric disorders and previous studies suggest it plays a putative role in episodic memory, language and social emotional processing, its exact function is not well understood. In this review we summarize what is currently known about the anatomy of the uncinate, we review its role in psychiatric and neurological illnesses, and we evaluate evidence related to its putative functions. We propose that an overarching role of the uncinate fasciculus is to allow temporal lobe-based mnemonic associations (e.g. an individual's name + face + voice) to modify behaviour through interactions with the lateral orbitofrontal cortex, which provides valence-based biasing of decisions. The bidirectionality of the uncinate fasciculus information flow allows orbital frontal cortex-based reward and punishment history to rapidly modulate temporal lobe-based mnemonic representations. According to this view, disruption of the uncinate may cause problems in the expression of memory to guide decisions and in the acquisition of certain types of learning and memory. Moreover, uncinate perturbation should cause problems that extend beyond memory to include social-emotional problems owing to people and objects being stripped of personal value and emotional history and lacking in higher-level motivational value.

Keywords: diffusion tensor imaging; episodic memory; orbitofrontal cortex; schizophrenia; anterior temporal lobe

Abbreviations: DTI = diffusion tensor imaging; FTD = frontotemporal dementia

Introduction

The uncinate fasciculus is one of several long-range white matter association fibre tracts in the human brain. It connects orbitofrontal cortex to the anterior temporal lobes through a direct, bidirectional monosynaptic pathway. Understanding the function and dysfunction of the uncinate fasciculus is of particular interest because of its clinical relevance. The uncinate fasciculus has been

implicated in several developmental and psychiatric disorders. Its location makes it particularly susceptible to immediate impact and shearing injuries in head trauma cases and it is commonly implicated in white matter damage associated with traumatic brain injury (Ewing-Cobbs and Prasad, 2011; Johnson *et al.*, 2011; Seo *et al.*, 2012). In addition, the uncinate fasciculus is frequently damaged in surgical treatment of epilepsy (Kucukyuruk *et al.*, 2012) and it is also one of a small group of white matter

association tracts to mature, not reaching its developmental peak until the third decade of life (Lebel *et al.*, 2008, 2012; Lebel and Beaulieu, 2011), which perhaps leaves it more susceptible to factors causing psychiatric illness during adolescence and young adulthood (Paus *et al.*, 2008).

It is believed that the information transmission properties of any given white matter tract can be predicted by the function of the regions that it connects. At the same time, functions of cortical regions are determined by their pattern of white matter input and output (Van Essen and Maunsell, 1983; Passingham *et al.*, 2002). By virtue of its geographic placement and connectivity, the uncinate fasciculus is typically associated with the limbic system. As a result, it is assumed that its functions should align with emotion and episodic memory. In actuality, however, the function(s) and information transmission properties of this white matter tract are largely unknown. Advancements in neuroimaging techniques, such as diffusion tensor imaging (DTI), offer the potential to fill in these critical gaps in knowledge. However, to interpret the findings from DTI, we must gain a better understanding of functional tractography. In the absence of a roadmap to guide our understanding of the functional significance of white matter tracts, there is a limit to the new information these endeavours can provide.

The purpose of this review is to summarize what is currently known about the uncinate fasciculus in an effort to move towards a better understanding of its function(s) and dysfunction(s). Inferences have often been made about the function of the uncinate fasciculus based on its anatomical connectivity. Therefore, this review begins with an overview of its physical connections, highlighting current misunderstandings about its connectivity and identifying current gaps in knowledge. Our understanding of the uncinate fasciculus's anatomy comes from different techniques: autoradiography, gross dissection and DTI. It is widely accepted that radiological tracer studies offer better precision compared with dissection and DTI; however, tracer studies cannot be conducted in humans.

Next, we critically evaluate the literature associating abnormalities in the uncinate fasciculus with a range of neurological and psychiatric disorders. The specific goal is to assess the strength of the evidence linking structural abnormalities of the uncinate fasciculus to specific clinical differences (e.g. differences in symptoms, cognitive function, and treatment outcomes) and to examine how these links contribute to a current understanding of uncinate fasciculus function. The review concludes with an examination of evidence that implicates the uncinate fasciculus in memory, language and socio-emotional processing and a proposed model that more specifically associates the uncinate fasciculus with these cognitive functions.

Anatomy

The anatomy of the uncinate fasciculus has been described in detail (Horel and Misantone, 1976; Ebeling and von Cramon, 1992; Ghashghaei and Barbas, 2002; Kier *et al.*, 2004; Kringelbach and Rolls, 2004; Catani and ffytche, 2005; Schmahmann *et al.*, 2008; Choi *et al.*, 2010; Noonan *et al.*, 2010; Peltier *et al.*, 2010; Lehman *et al.*, 2011; Thiebaut de Schotten *et al.*, 2012). Before summarizing these findings, it is important to comment on several

common misunderstandings about the uncinate fasciculus. First, in DTI studies, the uncinate fasciculus is often confused with the inferior frontal occipital fasciculus, as they are near each other in the region of the insula (Curran, 1909; Kier *et al.*, 2004). This confusion does not exist in tracer studies because this method relies on axonal transport mechanisms, though it is problematic in the techniques commonly used in humans (dissection and DTI). Thus, some clinically important features of the inferior frontal occipital fasciculus, such as spread of tumour growth in humans, may be incorrectly attributed to the uncinate fasciculus due to its close proximity (Kier *et al.*, 2004). Second, it is often mistakenly stated that the uncinate fasciculus connects the hippocampus to the frontal lobe (Saur *et al.*, 2008), when in fact, most anatomy studies to date do not show any extension into the hippocampus. Last, although the anatomy of the uncinate fasciculus appears to be highly conserved across non-human primates and humans (Fig. 1; Thiebaut de Schotten *et al.*, 2012), there are differences (Croxon *et al.*, 2005). Some caution in extending function from monkeys to humans is in order given the putative role of the human anterior temporal lobe and the uncinate fasciculus in one aspect of language processing, semantic retrieval.

The uncinate fasciculus is a long-range association fibre connecting the frontal and temporal lobes. In older anatomy texts, the uncinate fasciculus is considered to be part of the 'temporal stem', which also contains the anterior commissure, the inferior frontal occipital fasciculus, and Meyer's loop of the optic tract. The uncinate fasciculus is traditionally considered to be part of the limbic system. The uncinate fasciculus has a distinctive hook shape, arcing around the Sylvian fissure (Schmahmann and Pandya, 2006). For ease of description, the uncinate fasciculus is frequently broken into three parts: a dorsal/temporal segment, a middle/insular segment, and a ventral/frontal extension (Ebeling and von Cramon, 1992). The temporal segment originates from the uncus [Brodmann area (BA) 35], entorhinal and perirhinal cortices (also referred to as cortical nuclei of the amygdala; BA 28, 34 and 36), and the temporal pole/anterior temporal lobe (BA 20 and 38). The uncus is part of the olfactory cortex; entorhinal cortex is closely associated with episodic memory functions of the hippocampus; and perirhinal cortex has a controversial function in high-level object perception and object memory (Murray *et al.*, 2005). The temporal pole and nearby tissue constitute part of the anterior temporal lobe. Portions of the anterior temporal lobe (but not the temporal pole) are believed to play a role in certain types of semantic memory (Patterson *et al.*, 2007) and also in the encoding and storage of social and emotional concepts (Olson *et al.*, 2007, 2013; Zahn *et al.*, 2007; Simmons *et al.*, 2010). Within the greater anterior temporal lobe, there is a sensory-based segregation of function and intraregion connectivity: dorsal = auditory, ventral = visual, medial = olfactory, and polar = multisensory (Grossman *et al.*, 2004; Skipper *et al.*, 2011).

There is disagreement as to whether the temporal segment of the uncinate fasciculus extends into the amygdala proper (Croxon *et al.*, 2005; Petrides and Pandya, 2007; Thiebaut de Schotten *et al.*, 2012) and at least one tracer study provided clear evidence that it did not (Ungerleider *et al.*, 1989). However, several studies have provided clear evidence of monosynaptic connections between the orbital frontal cortex, anterior temporal lobe and

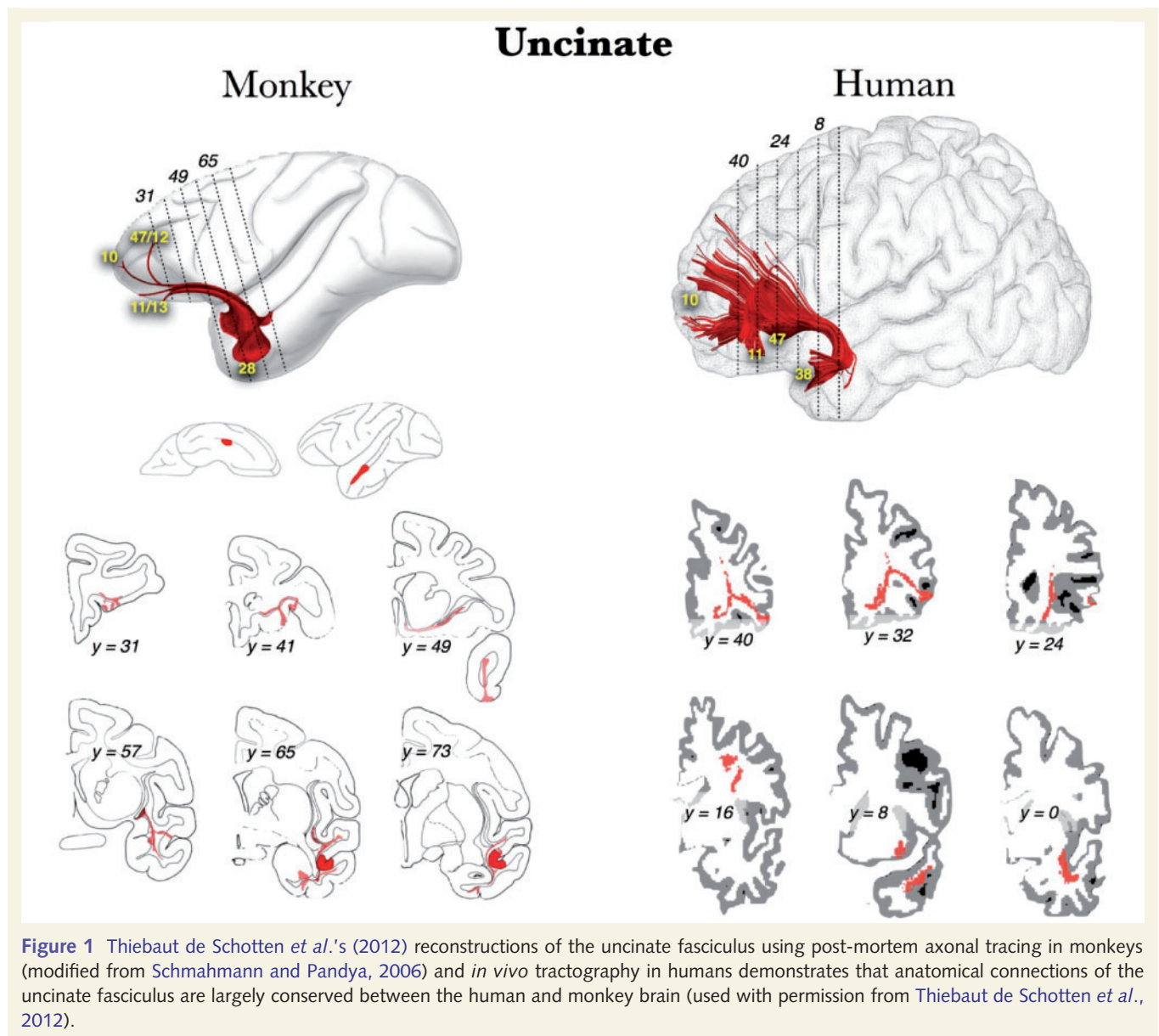


Figure 1 Thiebaut de Schotten *et al.*'s (2012) reconstructions of the uncinate fasciculus using post-mortem axonal tracing in monkeys (modified from Schmahmann and Pandya, 2006) and *in vivo* tractography in humans demonstrates that anatomical connections of the uncinate fasciculus are largely conserved between the human and monkey brain (used with permission from Thiebaut de Schotten *et al.*, 2012).

amygdala in monkeys. Using radioactive tracers, Ghashghaei and Barbas (2002) and Ghashghaei *et al.* (2007) detected substantial projections from the caudal orbital frontal cortex and from BA 36 of the anterior temporal lobe to the basolateral and basomedial nuclei of the amygdala; projections from both regions terminated in the basomedial nucleus of the amygdala. Although the aforementioned tracts were unnamed, Schmahmann and Pandya (2006) consider these tracts to be part of the uncinate fasciculus. Direct connections between the orbital frontal cortex and the anterior temporal lobe in monkeys are also well documented (Petrides and Pandya, 1988; Rempel-Clower and Barbas, 2000; Saleem *et al.*, 2008). The caudal orbital frontal cortex projects to entorhinal and perirhinal cortex, the rostral orbital frontal cortex to inferior ventral anterior temporal lobe and to dorsal anterior temporal lobe (Rempel-Clower and Barbas, 2000).

Uncinate fasciculus cell bodies are found in the temporal segment. From there, the uncinate fasciculus passes upward over the

lateral nucleus of the amygdala, through the limen insula, and either near or through two smaller white matter tracts, the external capsule and extreme capsule (Catani *et al.*, 2002; Mori *et al.*, 2002). At this point, the uncinate fasciculus is inferior to the inferior frontal occipital fasciculus. From there, it passes into the orbital regions of the frontal lobe (BA 11 and 47), where it has a horizontally-oriented fan shape. This fan splits into two branches, a larger ventro-lateral branch and a smaller medial branch. The ventral branch terminates in the lateral orbitofrontal cortex while the medial branch terminates in the frontal pole (BA 10; Dejerine, 1895; Klingler and Gloor, 1960; Crosby *et al.*, 1962; Thiebaut de Schotten *et al.*, 2012). In adult humans, the uncinate fasciculus has a width of 3–7 mm, a height of 2–5 mm, and a volume of $\sim 140 \text{ mm}^3$ (Ebeling and von Cramon, 1992; Schmahmann and Pandya, 2006).

Findings of hemispheric differences in the uncinate fasciculus volume are discordant. Some authors using DTI methods have

reported a leftward bias for uncinate fasciculus volume and fractional anisotropy values (Kubicki *et al.*, 2002; Hervé *et al.*, 2006; Rodrigo *et al.*, 2007; Hasan *et al.*, 2009), whereas other studies using post-mortem dissection methods have reported a rightward bias (Highley *et al.*, 2002; Park *et al.*, 2004). For example, a post-mortem study by Highley *et al.* (2002) found the right uncinate to be 27% larger in ~80% of participants studied. The failure to find laterality differences in all subjects could potentially be due to handedness differences, which were not examined by Highley *et al.* (2002). It is also possible that asymmetry exists in some parts of the uncinate fasciculus but not others (Park *et al.*, 2004; Rodrigo *et al.*, 2007).

Clinical disorders and the uncinate fasciculus

In this section, we discuss evidence for uncinate fasciculus involvement in five disorders: anxiety, schizophrenia, psychopathy, epilepsy and frontotemporal dementia. These disorders were chosen for their prevalence in the uncinate fasciculus literature. Although we do not discuss developmental trajectories or disorders of the uncinate fasciculus in this review, it should be noted that several authors have implicated uncinate fasciculus dysfunction in developmental disorders such as autism and conduct disorder (Sarkar *et al.*, 2012; Travers *et al.*, 2012).

Anxiety and schizophrenia

Several studies have reported altered functional activity in frontal regions, along with increased activations in limbic/paralimbic regions in generalized anxiety disorder and social anxiety disorder (Cooney *et al.*, 2006; Evans *et al.*, 2008; Bishop, 2009). One would expect then, that white matter tracts connecting limbic regions to orbitofrontal cortices, like the uncinate fasciculus, might be structurally impaired in anxiety disorders. However, our review of the DTI literature indicates that the uncinate fasciculus *per se* plays either a small role, or no role, in anxiety disorders (Table 1). Although several recent studies have provided evidence of a correlation between fractional anisotropy measures in the uncinate fasciculus and various measures of anxiety (Phan *et al.*, 2009; Tröstl *et al.*, 2011; Baur *et al.*, 2012; Hettrema *et al.*, 2012), other studies have reported negative correlations (Baur *et al.*, 2012) or no correlation (Han *et al.*, 2008; Liao *et al.*, 2011). It seems more likely that global white matter plays a role in anxiety disorders (Westlye *et al.*, 2011).

Similarly, the disconnection hypothesis of schizophrenia (Friston and Frith, 1995) has generated interest in discovering whether abnormalities exist in white matter tracts connecting the prefrontal cortex to other brain regions. Overall, the results of DTI studies measuring changes in uncinate fractional anisotropy values in schizophrenics are contradictory. Several studies have reported reduced fractional anisotropy values in the uncinate fasciculus of schizophrenics (Burns *et al.*, 2003; Kubicki *et al.*, 2005; McIntosh *et al.*, 2008; Price *et al.*, 2008; Kawashima *et al.*, 2009; Voineskos *et al.*, 2010; Kitis *et al.*, 2012). This association tends to be more

apparent in older and chronic populations of schizophrenics, or in populations with generally greater symptoms (Mandl *et al.*, 2013). However, other studies have reported increased, or no change in fractional anisotropy values (Highley *et al.*, 2002; Kubicki *et al.*, 2002; Jones *et al.*, 2006; Karlsgodt *et al.*, 2009) relative to healthy control subjects. It is possible that uncinate fasciculus dysfunction correlates with specific symptoms of schizophrenia such as flattened affect and lack of social engagement (Kitis *et al.*, 2012); however, few studies have taken this experimental approach.

Several design flaws, such as small sample sizes and samples that are clinically heterogeneous, are endemic in the DTI literature on schizophrenia, so it is difficult to know whether a true association between the left uncinate fasciculus and schizophrenic symptoms exists. Even if such an association exists, changes in left uncinate fractional anisotropy values do not appear to be particular to schizophrenia, as similar findings have been reported in bipolar disorder (McIntosh *et al.*, 2008; Sussman *et al.*, 2009; Lin *et al.*, 2011) and various anxiety disorders.

Psychopathy and antisocial personality disorder

Craig *et al.* (2009) measured fractional anisotropy in adult psychopaths and matched control subjects and found significantly reduced mean fractional anisotropy in the right uncinate of psychopaths compared with control subjects (Table 1 and Fig. 2). The volume of the tract, which was measured by the tractography algorithm as the number of trajectories, correlated negatively with antisocial behaviour in both the left and right uncinate. Given the correlation between white matter volume and antisocial behaviour, the authors suggest that their fractional anisotropy findings reflect antisocial behaviour rather than psychopathy. To test this hypothesis, the same group later measured fractional anisotropy in individuals with antisocial personality disorder and in healthy control subjects (Sundram *et al.*, 2012). Participants in the antisocial personality disorder group had significantly reduced fractional anisotropy and significantly increased mean diffusivity compared with control subjects in the right uncinate fasciculus. However, as the antisocial personality disorder group also had significantly reduced fractional anisotropy in several other white matter tracts, this study provides no evidence for an uncinate-specific impairment. An uncinate fasciculus-specific impairment in psychopathic individuals was, however, reported by Motzkin *et al.* (2011).

Interestingly, it has been reported that neural regions linked by the uncinate fasciculus—the orbital frontal cortex and temporal pole—are relatively thinner or have decreased volume in psychopaths and subjects with antisocial personality disorder relative to control subjects (Raine *et al.*, 2000; Yang *et al.*, 2009; Gregory *et al.*, 2012).

In summary, evidence for uncinate fasciculus dysfunction in psychopathy and antisocial personality disorder is limited, as researchers have only recently begun to investigate the matter. Nevertheless, compared with the other clinical disorders reviewed in this manuscript, we find the evidence for the uncinate fasciculus playing a role in these disorders to be more compelling. The cognitive function(s) disrupted by abnormalities of the right uncinate and how this relates to psychopathy is unclear, although it has

Table 1 Studies providing evidence for or against a particular role of the uncinate fasciculus in various psychiatric disorders

Evidence type	Summary of findings
Anxiety DTI (Han <i>et al.</i> , 2008; Kim and Whalen, 2009; Phan <i>et al.</i> , 2009; Liao <i>et al.</i> , 2011; Tröstl <i>et al.</i> , 2011; Westlye <i>et al.</i> , 2011; Baur <i>et al.</i> , 2012; Hetteima <i>et al.</i> , 2012)	Findings are mixed. The uncinate fasciculus appears to play either a small role, or no role, in this disorder.
Schizophrenia DTI (Burns <i>et al.</i> , 2003; Kubicki <i>et al.</i> , 2005; McIntosh <i>et al.</i> , 2008; Price <i>et al.</i> , 2008; Kawashima <i>et al.</i> , 2009; Voineskos <i>et al.</i> , 2010; Kitis <i>et al.</i> , 2012; reviewed by Kubicki <i>et al.</i> , 2007).	Findings are mixed. The uncinate fasciculus appears to play either a small role, or no role, in this disorder.
Psychopathy DTI (Craig <i>et al.</i> , 2009; Motzkin <i>et al.</i> , 2011; Sundram <i>et al.</i> , 2012). Volumetric MRI (Raine <i>et al.</i> , 2000; Yang <i>et al.</i> , 2009; Gregory <i>et al.</i> , 2012).	All DTI studies reported reduced fractional anisotropy values in the right uncinate fasciculus. Volumetric studies show bilateral volume losses in regions connected by uncinate fasciculus.

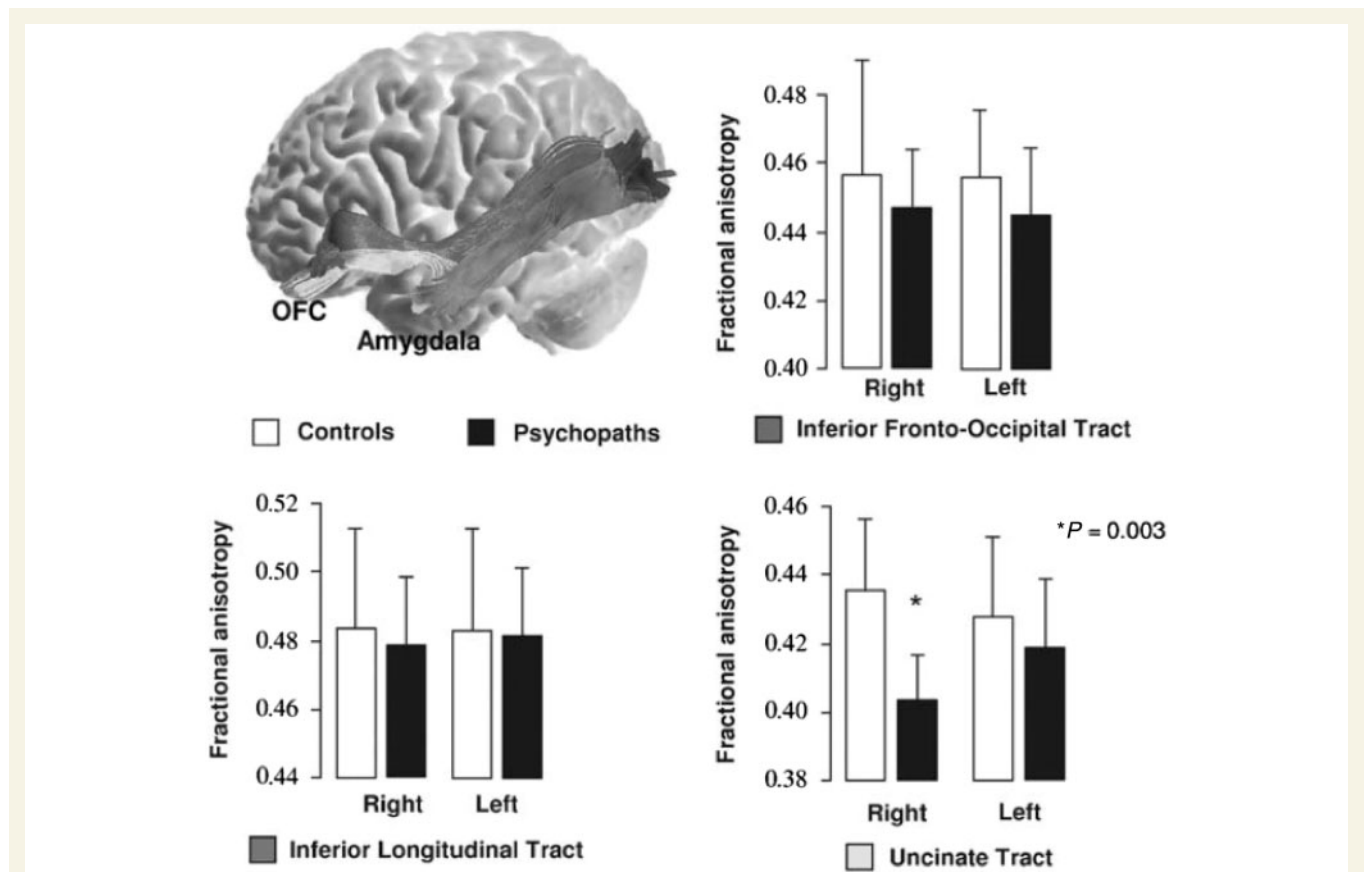


Figure 2 Studies of antisocial behaviour and psychopathy consistently report abnormalities of the right uncinate fasciculus. For example, Craig *et al.* (2009) reported significantly reduced mean fractional anisotropy in the right uncinate fasciculus ($P = 0.003$) for psychopaths compared with age- and IQ-matched control subjects and no significant differences in fractional anisotropy in the left uncinate fasciculus ($P = 0.448$) or in two control tracts: the inferior longitudinal tract and the inferior frontal-occipital tract (used with permission from Craig *et al.*, 2009). This pattern of findings is unique and noteworthy because studies of patients with other psychiatric disorders consistently report abnormalities in the left or bilateral uncinate fasciculus rather than right uncinate fasciculus alone.

been proposed that it may relate to particular types of learning and memory.

Epilepsy

Unilateral resection of the anterior temporal lobe has long been used as a treatment for intractable temporal lobe epilepsy

(Penfield and Baldwin, 1952; Jensen, 1975, 1976; Jensen and Vaernet, 1977). Although it is rarely noted or commented on, the uncinate fasciculus is frequently severed in these surgeries. Despite the frequency of surgical severance of the uncinate fasciculus, a recent review of the temporal lobe epilepsy surgery literature noted that little could be said about the function of this white matter tract (Kucukyuruk *et al.*, 2012).

Even before resection surgery, temporal lobe epilepsy, especially in the left hemisphere, is strongly associated with decreased white matter integrity in the uncinate fasciculus (Rodrigo *et al.*, 2007; Diehl *et al.*, 2008; McDonald *et al.*, 2008; Riley *et al.*, 2010). It is likely that changes in white matter properties of the uncinate fasciculus are the result of aberrant neuronal morphology or firing and thus epiphenomenal to the disorder because most seizures are localized in the medial and anterior temporal lobe regions. Although it is unlikely that alterations in the uncinate fasciculus cause epilepsy, uncinate fasciculus abnormalities may contribute to certain cognitive deficits commonly associated with temporal lobe epilepsy. Abnormal fractional anisotropy values in the bilateral uncinate fasciculus are correlated with deficits in immediate verbal memory (Diehl *et al.*, 2008; McDonald *et al.*, 2008), delayed memory (Diehl *et al.*, 2008; McDonald *et al.*, 2008; Riley *et al.*, 2010), and confrontational naming/semantic memory as tested with the Boston Naming Test (McDonald *et al.*, 2008) in patients with temporal lobe epilepsy. Similar deficits are reported after unilateral left temporal resection surgery, especially proper naming deficits (Lu *et al.*, 2002; Hamberger and Drake, 2006).

The epilepsy literature also hints that the uncinate fasciculus may have non-mnemonic functions. There is a subgroup of grand mal temporal lobe seizures known as 'uncinate fits', so named because the seizure focus is in the uncus, on the medial surface of the anterior temporal lobe at the anterior end of the parahippocampal gyrus. An early case study (Jackson and Colman, 1898) identified an epileptic patient who experienced 'dreamy states' and unusual tastes immediately before seizing; post-mortem examination found damage only in the left uncinate fasciculus and the left uncus. Later research found that uncinate fits were also associated with emotional and sexual arousal, olfactory and gustatory hallucinations, and involuntary movement of the face and mouth, including spitting (Penfield and Baldwin, 1952), olfactory hallucination (Efron, 1957; West and Doty, 1995), depressive symptoms (Weil, 1955) and intense anxiety (Parrino, 1971). The olfactory symptoms that accompany uncinate fits do not appear to arise from the uncinate fasciculus, but rather tissue damage to the insula or olfactory cortex (Daly, 1958; Howe and Gibson, 1982). The remaining symptoms are reminiscent of the gustatory, sexual and emotional symptoms of Kluver-Bucy disorder, which can be caused by temporal lobectomy surgery (for a review see Olson *et al.*, 2007).

In general, research on the uncinate fasciculus from the perspective of epilepsy and surgical treatment of epilepsy is surprisingly weak although there is a clinical consensus that it must be important for disease progression and cognitive deficits following temporal resection. Based on this literature, it appears that retrieval of proper names and the retention and/or retrieval of verbal material are disproportionately affected by severance of the left uncinate fasciculus. The proper naming findings are compelling, since several laboratories using different patient groups and different techniques have reported similar findings (Damasio *et al.*, 1996; Tranel *et al.*, 1997; Grabowski *et al.*, 2001; Papagno, 2011). There are also older findings on uncinate fits linking the dominant hemisphere uncinate fasciculus to emotional, sexual and gustatory symptoms. However, our understanding of the uncinate fasciculus from the perspective of epilepsy is at least in part driven by the

kinds of tests performed on these patients—typically verbal memory and lexical retrieval—with few studies testing non-verbal memory or emotional processing.

The frontotemporal dementias

Frontotemporal dementia (FTD) is a progressive neurological disease characterized by degeneration of frontal and/or anterior temporal lobe tissue with more medial regions, such as the hippocampus, remaining intact at early stages of the disease. It is a heterogeneous disorder with three prominent subtypes: semantic dementia, behavioural variant FTD, and progressive non-fluent aphasia (Weder *et al.*, 2007). Semantic dementia is a subtype in which general naming deficits are observed, typically in association with deterioration of the temporal lobes, prominently in the left anterior regions (Snowden *et al.*, 1989). Naming deficits are a diagnostic symptom of this disease (Snowden *et al.*, 1989; Hodges *et al.*, 1992; Neary *et al.*, 1998), although patients also usually present with personality changes and are reported by family members to be emotionally distant and lacking in empathy.

Like epilepsy, uncinate fasciculus dysfunction clearly does not cause FTD. However, because the anterior temporal lobe and orbital portions of the frontal lobe are among the earliest regions to experience cell loss in this disorder, the integrity of the uncinate fasciculus is necessarily affected. Several studies have reported an association between decreased fractional anisotropy values/radial diffusivity in the left uncinate fasciculus and semantic dementia (Matsuo *et al.*, 2008; Agosta *et al.*, 2010; Whitwell *et al.*, 2010; Acosta-Cabronero *et al.*, 2011; Galantucci *et al.*, 2011) with a smaller number reporting changes in both the left and right uncinate fasciculus (Matsuo *et al.*, 2008; Galantucci *et al.*, 2011). One study reported that patients with semantic dementia had lower fractional anisotropy values in bilateral uncinate fasciculus compared with both non-fluent aphasics, who exhibit agrammatism and motor speech errors, and healthy control subjects, suggesting that the uncinate fasciculus is particularly associated with semantic retrieval rather than speech output (Galantucci *et al.*, 2011).

Findings from another subtype of frontotemporal dementia, behavioural variant FTD, complicate this picture. The symptoms of behavioural variant FTD tend to centre around deficits in emotional regulation, social cognition, motivation and decision making, although language deficits typically appear later in the disease progression (Piguet *et al.*, 2011). Patients exhibit apathy, impulsivity, inappropriate sexual behaviours and hoarding (Piguet *et al.*, 2011). Several studies have reported decreased fractional anisotropy values in both the left and right uncinate fasciculus in patients with behavioural variant FTD (Zhang *et al.*, 2009; Piguet *et al.*, 2011; Mahoney *et al.*, 2012). Another study compared patients diagnosed with behavioural variant FTD and Alzheimer's disease, and found no significant differences between the two groups in fractional anisotropy measures of the uncinate fasciculus, although patients with behavioural variant FTD did, on average, have smaller fractional anisotropy values in both the left and right uncinate fasciculus (Tartaglia *et al.*, 2012). Furthermore, patients with behavioural variant FTD had significant degeneration of the grey matter connected directly to the uncinate fasciculus in the

right hemisphere, as compared with the patients with Alzheimer's disease (Tartaglia *et al.*, 2012). Alternatively, Agosta *et al.* (2012) found that the damage to the left uncinate fasciculus, as measured by DTI, was the best predictor of behavioural variant FTD diagnosis, as opposed to progressive aphasia, and damage to the left uncinate fasciculus was the best predictor of overall degeneration in these patients. These reports clearly indicate that the uncinate fasciculus is affected in behavioural variant FTD, but the differential effects of the left, right or bilateral uncinate fasciculus degradation remains unclear.

Summary: adult clinical disorders and the uncinate fasciculus

In the preceding paragraphs we reviewed evidence linking uncinate fasciculus dysfunction to three psychiatric and two neurological disorders. Existing evidence fails to support the contention that the uncinate fasciculus has a primary role in anxiety disorders or schizophrenia. Future studies should focus on more distributed networks of white matter as they relate to these disorders, rather than on the uncinate fasciculus alone.

We found stronger evidence for uncinate fasciculus dysfunction in psychopathy and epilepsy. With regards to epilepsy, further studies of temporal lobe epilepsy in which patients with affected and unaffected uncinate fasciculus can be compared may be critical to elucidating not only the function of the uncinate fasciculus, but also how to preserve emotional processing and verbal memory in treatment of temporal lobe epilepsy. Of note, the epilepsy literature typically includes only a small battery of tests, thus the potential range of deficits has not been fully explored. There is also strong evidence for involvement of uncinate fasciculus degeneration in the clinical diagnosis of FTD, however, it is unclear whether it is more related to semantic retrieval deficits or social-emotional abnormalities.

Recommendations for future researchers interested in investigating the role of the uncinate fasciculus in various psychiatric and neurological disorders include: testing larger and more homogeneous clinical samples; correlating specific symptoms and/or theory-driven test performance, rather than disease presence or absence, with various DTI values; and using more rigorous statistical thresholding procedures. Future studies should also take care to investigate other white matter tracts, as well as measures of global white matter, alongside their investigations of the uncinate fasciculus. In doing so, care must be taken to dissociate the uncinate fasciculus from other nearby tracts. Greater experimental control and rigor will help prevent false positives and also help to elucidate the specific functional involvement of the uncinate fasciculus in psychiatric and neurological disease.

Putative functions of the uncinate fasciculus

The literature refers to three functions of the uncinate fasciculus: associative and episodic memory functions; linguistic functions; and social-emotional functions.

Episodic memory and the uncinate fasciculus

Episodic memory is a record of a person's experience that holds date information and spatio-temporal relations (Tulving, 1983). Episodic memory formation relies on the function of medial temporal lobe structures, especially the hippocampus and portions of the frontal lobe, such as dorsolateral prefrontal cortex. There is long-standing interest in understanding how fibre pathways that connect these regions function in episodic memory.

Markowitsch (1982) proposed that retrograde amnesia was caused by the disconnection of temporal limbic structures from the frontal lobe 'The task of the uncinate fascicle will be to guide and channel this information flow to the prefrontal cortex and to transmit preprocessed information back to the temporal cortex for the final act of representation' (Markowitsch, 1982). His proposal was based on a review of neuropsychological case studies of selective retrograde amnesia, in which the majority of cases involved damage to the ventral frontal lobe and the anterior temporal lobe with presumed involvement of the uncinate fasciculus. Along these lines, Levine *et al.* (1998) reported the case of Patient ML, who suffered traumatic brain injury after a bicycle accident that resulted in damage to portions of the visual cortex, the hypothalamus, right and left prefrontal cortex, and the uncinate fasciculus. Several years after this incident, the patient continued to suffer from social impairments and severe retrograde memory loss, but only mild anterograde memory loss. Although this case is striking, it is important to note that Patient ML's brain damage was not limited to the uncinate fasciculus. Indeed, there are no instances of retrograde or anterograde amnesia in which the uncinate alone was destroyed.

Focal unilateral lesions of the human uncinate fasciculus occur from time to time, usually in the context of surgical removal of low-grade gliomas. Papagno *et al.* (2011) studied 18 right-handed patients before and after left uncinate fasciculus removal. The results showed that effects on memory were modest; short-term memory and visual long-term memory were preserved, while verbal memory as tested by list learning was impaired immediately after surgery but improved to normal levels after 3 months. It is possible that the effects on memory were modest because (i) resection was unilateral; (ii) the wrong type of memory was tested; or (iii) the uncinate fasciculus's role in memory is redundant with that of other white matter tracts.

The importance of the type of memory tested is highlighted by research on non-human primates. Information flow from the inferior temporal lobe to orbital frontal cortex is dramatically reduced by uncinate fasciculus disconnection (Ungerleider *et al.*, 1989), even though indirect routes between these regions exist (e.g. through the amygdala and medial dorsal nucleus of the thalamus; Eacott and Gaffan, 1992). Uncinate fasciculus disconnection does not impair visual object discrimination learning, suggesting that some associative learning processes are not reliant on the uncinate fasciculus (Parker and Gaffan, 1998; Gaffan *et al.*, 2002). Similarly, concurrent object-reward association learning, reversal learning, configural learning, and delayed matching-to-sample are unimpaired by uncinate fasciculus disconnection in

monkeys (Eacott and Gaffan, 1992; Gaffan and Eacott, 1995; Gutnikov *et al.*, 1997). Object-in-place learning is mildly impaired after uncinate fasciculus resection (Browning and Gaffan, 2008). The one type of learning that is consistently impaired after uncinate fasciculus dissection is conditional rule learning. In this task, an instructional cue at the beginning of each trial indicates what types of choices will be rewarded (Gaffan *et al.*, 1988; Gutnikov *et al.*, 1997; Parker and Gaffan, 1998; Bussey *et al.*, 2002).

The DTI literature provides relatively strong support for the role of the uncinate fasciculus in episodic memory. Thomas *et al.* (2012) followed-up the monkey studies described above; in their study, participants were required to learn face–scene associations through computerized feedback across 1000+ learning trials. Faces were used as stimuli, since portions of the ventral anterior temporal lobe are critical for person memory (Von Der Heide *et al.*, 2013). The results showed that subjects who had faster learning rates also had higher fractional anisotropy values in the left, but not right, uncinate fasciculus. There was no correlation in nearby white matter tracks. One interpretation of these findings is that the uncinate fasciculus was necessary for retrieving information and keeping count as to which face–scene associations were rewarded or punished in the past.

Other relevant DTI studies tend to use a variety of tasks drawn from the neuropsychology literature rather than the relevant monkey literature. Several DTI studies have reported correlations between left uncinate fasciculus white matter fractional anisotropy values and performance on auditory–verbal memory tasks, such as list learning (but not visual memory) by cued or free recall (Diehl *et al.*, 2008; McDonald *et al.*, 2008; Niogi *et al.*, 2008). Such a relationship between the left uncinate fasciculus and verbal memory has been reported not only in adults, but also in children and adolescents (Mabbott *et al.*, 2009). Another study reported correlations between the right uncinate fasciculus and auditory–verbal memory (Fink *et al.*, 2010). Likewise, fractional anisotropy values of the uncinate fasciculus are significantly lower in amnesic mild cognitive impairment (Fuji *et al.*, 2008) and correlate with verbal memory impairment in traumatic brain injury (Niogi *et al.*, 2008) and with age-related reductions on several declarative memory tasks (Engvig *et al.*, 2012).

It is important to note that in some of these studies (Engvig *et al.*, 2012), the uncinate fasciculus was grouped together with nearby structures—the internal capsule—so it is not clear that the uncinate fasciculus *per se* was associated with memory

performance. Although null reports are not commonly reported, Seo *et al.* (2012) reported finding no correlation between changes in DTI parameters and episodic memory performance in a group of patients with traumatic brain injury.

In summary, the reviewed findings fail to support the contention that the uncinate fasciculus has a general role in episodic memory (Markowitsch, 1982; McCauley *et al.*, 2011). A more likely candidate for this function is the fornix (Metzler-Braddeley *et al.*, 2011). Several studies in monkeys and a related study in humans (Thomas *et al.*, 2012) suggest that the uncinate fasciculus supports some types of associative learning, but perhaps not in the obvious way of forming the actual linkage between two stimuli, such as a face and a place. We draw this conclusion for two reasons. First, cells in the ventral anterior temporal lobe can rapidly form associative pairings between visual stimuli (Sakai and Miyashita, 1991; Eifuku *et al.*, 2010), therefore it is unclear what the uncinate fasciculus or the orbital frontal cortex would add to this process. Second, research in non-human primates has shown that associative learning is not abolished by uncinate fasciculus resection, but rather, that certain types of associative learning are affected. The most compelling finding in this literature is the observed deficit in conditional rule learning after uncinate fasciculus resection. One interpretation of this finding is that the uncinate fasciculus is involved in associating stimuli with various types of rewards, non-rewards, and punishments that when broadly defined, would include error monitoring (Metzler-Braddeley *et al.*, 2011) that may serve to modulate retrieval of correct pairings (Table 2).

Language and the uncinate fasciculus

The left uncinate fasciculus has frequently been associated with language function because it connects regions of the brain that have putative functions in language: the anterior temporal lobes and portions of the frontal lobes, both of which have been proposed to encode, store and retrieve semantic knowledge (Grossman *et al.*, 2004; Catani and Mesulam, 2008). Based on this evidence, it was hypothesized that the uncinate fasciculus, particularly on the left, was part of the ‘ventral language pathway’ and that it supported semantic naming by relaying sensory information about objects (presumably in ventral temporal cortex) to language supporting regions (in the lateral frontal lobe; Parker *et al.*, 2005). The problem with this logic is that it is based on a

Table 2 A list of psychological functions that do and do not appear to be associated with information transmission through the uncinate fasciculus

	Functions linked to the uncinate fasciculus	Functions not linked to the uncinate fasciculus
Episodic memory	Reversal learning; learning from feedback/rewards/punishments; formation of associations that motivate behaviour; value-based updating of stored representations	Encoding and consolidation of common episodic memories including autobiographical memory
Language	Retrieval of proper names for people; possibly some aspects of semantic memory retrieval	General linguistic functions, e.g. speech production, speech comprehension, syntax, most aspects of semantic memory
Social–emotional processing	Valuation of stimuli; social reward processing; higher-level emotional meaning of concepts	Generation of emotions; personality; motivation; anxiety

rather crude assessment of anatomy. The uncinate does not connect the specific region of the frontal lobe known to be critical for language production i.e. Broca's area/inferior frontal gyrus, to the anterior temporal lobe; instead, it connects regions in the ventral and medial frontal lobe (which are not commonly associated with linguistic function) to the anterior temporal lobe and surrounding structures.

Beyond the flawed geography evidence, there is a gaping lack of evidence to support the hypothesis that the uncinate fasciculus plays a general role in language. General language function, excepting verbal memory, is typically unimpaired after unilateral, temporal lobectomy surgery for epilepsy (Parker *et al.*, 2005), and left uncinate fasciculus stimulation during neurosurgery is not associated with general language impairments (Duffau *et al.*, 2009). Also the uncinate fasciculus is not implicated in primary progressive aphasia (Galantucci *et al.*, 2011). Because of this, more recent accounts have discarded the notion that the uncinate fasciculus is part of the ventral language pathway in favour of the extreme capsule, which connects the middle temporal lobe and the ventrolateral prefrontal cortex (Saur *et al.*, 2008).

Although the left uncinate fasciculus does not appear to play a general or exclusive role in language, there is some evidence that it plays a minor supporting role, particularly in lexical retrieval of semantic knowledge. For example, a study of memory in healthy older adults found that performance on a variety of semantic memory tasks, typically involving naming (such as Pyramids and Palm Trees), correlated positively with fractional anisotropy values in the left uncinate fasciculus (de Zubicaray *et al.*, 2011). Temporal lobe resection surgery for epilepsy commonly leads to confrontation naming deficits whereas general language functions remain preserved (Lu *et al.*, 2002; Hamberger and Drake, 2006). The most commonly observed confrontation naming deficit is in proper naming. Papagno *et al.* (2011) assessed patients with left uncinate fasciculus removal due to surgery for low-grade gliomas and found significant deficits in naming famous faces immediately after surgery and 3 months later. In contrast, other deficits that were present immediately after surgery, such as difficulties in remembering lists of words, and deficits in verbal fluency, improved to normal levels 3 months post-surgery. These findings indicate that the left uncinate fasciculus has redundant functions with other fibre pathways for some aspects of lexical retrieval and verbal memory, with the exception of proper naming.

The association between proper naming and the left uncinate fasciculus was indirectly studied by Damasio *et al.* (1996, 2004) in their milestone studies of lexical retrieval and the anterior temporal lobes (Tranel *et al.*, 1997). These studies explored the deficits in patients with stable temporal lobe lesions, commonly caused by temporal lobe resection surgery. Patients with lesions restricted to the left anterior temporal lobe, and possibly including damage to the uncinate, had a specific deficit for retrieving the names of persons, and at times, landmarks. However, these patients performed normally when naming animals and objects. It is important to bear in mind that one cannot determine whether the observed deficits were due to uncinate damage *per se*, or due to other surgical or epilepsy-related damage.

More global semantic retrieval deficits are observed in semantic dementia and there is some evidence, reviewed above, linking

semantic dementia to uncinate fasciculus deterioration. However, the literature on semantic dementia is new and typically fails to correlate specific symptoms to the uncinate fasciculus.

These findings strongly indicate that the uncinate fasciculus is not dedicated to general linguistic functions. This should not be taken to imply that the uncinate fasciculus has no role in language; instead, the left uncinate fasciculus appears to have an important, though non-critical, role in semantic retrieval (Table 2). However, it is also likely to be involved in social-emotional processing. The marriage of these two processes is evident when one considers that language is inherently social; you speak to transfer ideas and emotions to another person. The conceptual knowledge that underlies our expression of ideas is coloured by our preferences and emotional history to the degree that emotion is part-and-parcel of our conceptual representations.

Social-emotional processing and the uncinate fasciculus

There is reason to believe that the uncinate fasciculus plays a central role in certain types of social and emotional processing that are instrumental in decision making and shaping behaviour (Table 2). The first line of evidence is anatomical: the uncinate fasciculus connects limbic and paralimbic brain regions. The orbitofrontal cortex has long been considered essential for reward-based decision making (Adolphs *et al.*, 1994; Angrilli *et al.*, 1999; Grabenhorst and Rolls, 2011) and there is evidence that portions of the anterior temporal lobe are involved in the storage and retrieval of social memories including person memory, as well as theory of mind (Olson *et al.*, 2007; Zahn *et al.*, 2007; Simmons *et al.*, 2010).

Second, there is also evidence from some clinical populations suggesting a social-emotional role for the uncinate. For instance, patients with behavioural variant FTD have deficits that are primarily related to social and emotional processing (Piguet *et al.*, 2011), as opposed to other variants of FTD, whose symptoms are more linguistic and mnemonic (Hodges, 2001; Whitwell *et al.*, 2010). Findings of uncinate abnormalities are quite common in this population (Matsuo *et al.*, 2008; Agosta *et al.*, 2010; Whitwell *et al.*, 2010; Acosta-Cabronero *et al.*, 2011; Galantucci *et al.*, 2011). One study reported that patients with semantic dementia and progressive non-fluent aphasia had decreased fractional anisotropy values in the left and right uncinate, respectively, compared with healthy control subjects, but only patients with behavioural variant FTD showed bilateral uncinate fasciculus abnormality and that decreased fractional anisotropy values in the uncinate specifically correlated with behavioural inhibition (Hornberger *et al.*, 2011). Studies have shown grey matter loss in two regions connected by the uncinate fasciculus, the anterior temporal lobe and orbitofrontal cortex, as well as decreased fractional anisotropy values in the right uncinate fasciculus.

Related to this, literature on traumatic brain injury provides some interesting case studies to support our hypothesis. A patient with moderate traumatic brain injury, exhibiting symptoms of depression apathy, and reduced sex drive and self esteem, was described by Zappalà *et al.* (2012). MRI scans showed damage to this patient's left uncinate and left cingulate. In another study, Johnson *et al.* (2011) specifically compared the uncinate fasciculus

in 15 children 3 months after traumatic brain injury to 15 healthy control subjects, and discovered that uncinate fasciculus fractional anisotropy measurements were predictive of executive function outcomes, especially emotional control, 1 year later.

Finally, the fifth and most speculative line of evidence suggesting a social role for the uncinate in social–emotional processing comes from case studies of patients suffering from the related disorders of Capgras delusion and reduplicative paramnesia. Capgras delusion is characterized by a patient's insistence, despite all evidence to the contrary, that his or her loved one has been replaced by an impostor (Hirstein and Ramachandran, 1997; Edelstyn *et al.*, 2001). Patients with reduplicative paramnesia, on the other hand, believe a particular person or place has been duplicated and/or relocated to a familiar place (e.g. a hospital room moved to the patient's home) (Politis and Loane, 2012).

Hirstein and Ramachandran (1997) put forth a hypothesis that Capgras delusion (and related delusions) results from severed connections between face processing areas and the limbic regions responsible for emotionally 'colouring' or 'tagging' facial percepts. The logic is that patients with Capgras delusion accurately perceive the face of a familiar loved one but without feeling the expected feelings of warmth, and therefore conclude that it must be someone else's face.

Regions connected with the uncinate fasciculus have been implicated in Capgras delusion and other delusional misidentification syndromes, although lesions are often diffuse and the evidence for any single region's involvement is inconclusive (Bouckoms *et al.*, 1986; Drake, 1987). There is, therefore, reason to believe that the uncinate fasciculus might play a limited role in this disorder. Since uncinate lesions have been reported in the absence of symptoms that characterize Capgras delusion, we can be sure that uncinate fasciculus dissection alone is insufficient to produce Capgras delusion. The ablation of a subset of uncinate fasciculus cell bodies, or the cells on which they terminate, may be necessary. Uncinate fasciculus damage may, however, contribute to Capgras-like symptoms.

The uncinate itself has been implicated in at least two studies. Lee *et al.* (2011) found an uncinate lesion in a patient suffering from reduplicative paramnesia in the absence of other delusions (aside from anosognosia). This patient had an average IQ and normal language and autobiographical memory. Lesions were also observed in several other white matter tracts. Edelstyn *et al.* (2001) reported a patient with abnormal frontal lobe white matter near or in the uncinate who suffered from episodes of Capgras delusion. Retrieval of stored social semantic information (e.g. pairing a person's occupation with their face) was also poor in this patient (see also a case of Capgras delusion following temporal lobectomy, Lipson *et al.*, 2003).

Grey matter regions connected by the uncinate are also implicated in the abovementioned disorders, as well as in social and emotional processing in general. Orbitofrontal and anterior temporal regions are involved in behavioural variant FTD and psychopathy (Hodges, 2001; Yang *et al.*, 2009; Hornberger *et al.*, 2011; Gregory *et al.*, 2012), whereas amygdala dysfunction has been implicated in autism spectrum disorder (Dalton *et al.*, 2005; Nacewicz *et al.*, 2006; Kleinhans *et al.*, 2009). Patients with orbitofrontal lesions often have impaired social functioning, as well as

impaired theory of mind and empathy (Shamay-Tsoory *et al.*, 2003; Mah *et al.*, 2005).

Although none of the reported findings on their own is convincing, as a whole, the literature indicates that the uncinate fasciculus plays an important, albeit underspecified, role in social–emotional processing.

A proposal and conclusions

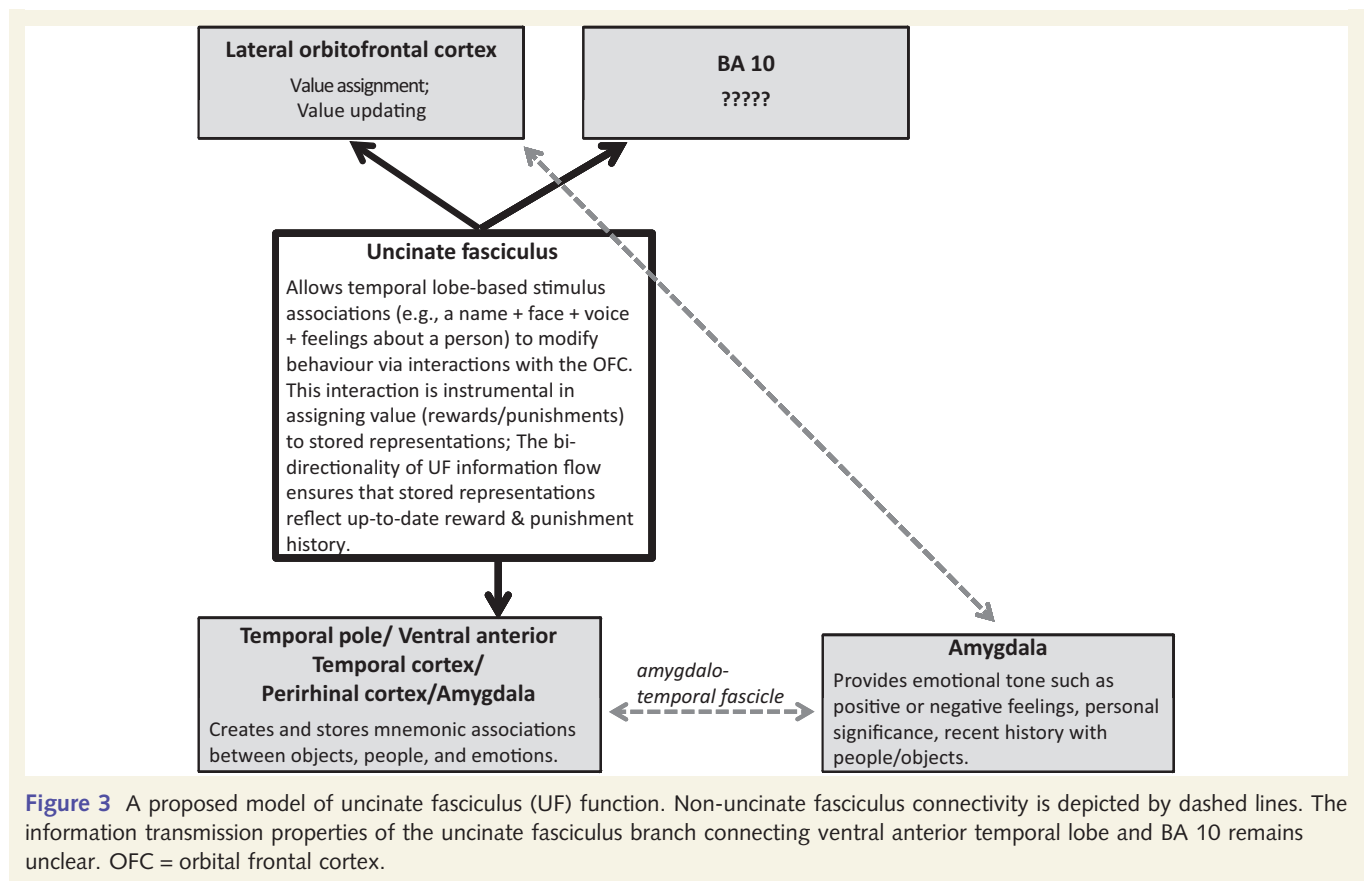
After reviewing the literature on uncinate fasciculus function, two credible accounts of uncinate fasciculus function become apparent. The first is a 'generalist' hypothesis, which suggests that the uncinate fasciculus is involved in various types of memory, language, social–emotional processing, and perhaps other functions that are yet undiscovered. Although it is easy to critique this hypothesis as a cobbled-together patchwork-quilt hypothesis, it is entirely plausible that certain white matter tracts are best conceived as somewhat general routes of information transfer.

The second is a 'specialist' hypothesis, which suggests that the uncinate fasciculus is biased towards a particular type of information transfer. Certain findings stood out in our literature review that we can use to infer a more specific explanation of uncinate fasciculus function (Fig. 3). We note that findings from the monkey literature, which offer the strongest causal power, suggest that the uncinate fasciculus is critically involved in some, but not all, mnemonic functions. Converging evidence from DTI and neuropsychology support this contention. The uncinate fasciculus does not appear to play a general role in episodic memory, however, nor is it critical for forming or retrieving many common types of mnemonic associations.

In the following section we lay out the argument that the uncinate fasciculus does not have general mnemonic, linguistic, or social–emotional functions. Instead, its function lies at the intersection of memory and social–emotional processes, in the explicit service of choice. More specifically, we argue that the uncinate fasciculus allows temporal lobe-based mnemonic associations (e.g. a person's name + face + voice + your feelings about a person) to modify behaviour by interacting with systems in the lateral orbital frontal cortex that are instrumental for making associations between stimuli and rewards, and ultimately, decision making. The bidirectionality of uncinate fasciculus information flow ensures that temporal lobe representations of objects and people reflect up-to-date reward/punishment history.

The mnemonic functions of the uncinate fasciculus are necessarily limited to the types of information, and the type of processing functions, supported by cortical anatomy associated with this tract. Likewise, the social–emotional information that is transmitted by the uncinate should be viewed in light of known functions of the cortical anatomy associated with the uncinate fasciculus. The cortical anatomy of the uncinate fasciculus is described in the following points:

- (i) Uncinate fasciculus cell bodies are found in the anterior temporal lobe. Cells in the ventral anterior temporal lobe have rapid association formation capacities. Sakai and Miyashita (1991) reported that single neurons in the ventral anterior



temporal lobe of monkeys that initially responded to only one abstract pattern would later respond to a second abstract pattern that had been associated through training with the first. Recordings from these cells show that they can represent an associative pairing between faces and abstract patterns, acquired through training (Eifuku *et al.*, 2010). Tsukiura *et al.* (2010) showed that successful encoding of person-related semantics with their names was associated with blood oxygen level-dependent activity in the left anterior temporal lobe. Patients with damage to this region have difficulty forming new associations between names and pictures of objects (Sharon *et al.*, 2011). It has been proposed that the rapid association formation capacity of this region underlies some aspects of semantic memory (Eifuku *et al.*, 2010), most likely the association between a visual stimulus such as a face and more abstract forms of semantic knowledge, such as an occupation. The association formation capabilities of the anterior temporal lobe may be reliant on the hippocampus during the initial encoding phase (Alvarez and Squire, 1994; Frankland and Bontempi, 2005; Nieuwenhuis *et al.*, 2011). These structures are strongly interconnected by tracts through the entorhinal cortex and subiculum (Chabardès *et al.*, 2002);

(ii) The temporal pole of the anterior temporal lobe has bi-directional interactions with the basomedial nucleus and the border of the basolateral nucleus of the amygdala through a small white matter tract called the 'amygdalo-temporal fascicle' (Klingler and Gloor, 1960; Ghashghaei and Barbas, 2002). The amygdala has a known role in emotion generation,

social attention, and salience tagging (Adolphs, 2010). The connections between the anterior temporal lobe and the amygdala may serve to give the representations stored in the anterior temporal lobe emotional tone, such as positive or negative feelings, and personal importance and significance (Olson *et al.*, 2013). One plausible role of the uncinate fasciculus is to transmit salience-laden stimulus representations stored in anterior-medial aspects of the temporal lobe to the lateral orbital frontal cortex so that decisions can be made based on the emotional tone or incentive value of the stimuli;

(iii) Most uncinate fasciculus fibres terminate in lateral orbital frontal cortex rather than midline orbital frontal cortex. The lateral orbital frontal cortex's function is variously related to the evaluation of losses related to ongoing behaviour (Kringelbach and Rolls, 2004), or in assigning rewards and punishments to behaviours (Noonan *et al.*, 2010). Of late, more evidence has accrued for the second idea especially in the context of associative learning. For instance, Noonan *et al.* (2010) showed that when lateral orbital frontal cortex was lesioned, monkeys failed to make the correct associations between behaviours and rewards, and the history of what was rewarded in the past was lost (Noonan *et al.*, 2010). Another group found that lesions to this region in non-human primates disrupted the rapid updating of object value (Rudebeck and Murray, 2011).

Our hypothesis predicts that bilateral damage of the uncinate fasciculus should cause problems with the manner in which

memory is used to alter behaviour—approaching or avoiding stimuli based on reward and punishment history. It should also cause problems in the expression of certain memories since the link between associative memory and certain decision processes would be disrupted. It is possible that some types of learning, such as devaluation, would be perturbed by uncinate fasciculus damage, as feedback from lateral orbital frontal cortex to the temporal lobe would be perturbed. These types of learning and memory problems could present clinically as gambling addiction, or poor decision making, for instance. Problems associated with uncinate fasciculus damage should extend beyond memory to include social–emotional problems due to people and objects being stripped of personal value and lacking in higher-level motivational value, potentially explaining aspects of Capgras delusion and some of the symptoms associated with autism spectrum disorder. In autism for instance, failure to form emotional attachments and friendships and failure to learn from praise and other social rewards could potentially be explained by uncinate fasciculus dysfunction. Behavioural variant FTD is characterized by disruptions in affiliative social behaviour and social disinhibition (Mummery *et al.*, 2000; Hodges, 2001), both of which can be conceptualized as disturbances in social reward processing and social learning. Findings reported earlier in psychopaths with reduced fractional anisotropy values in the right uncinate (Craig *et al.*, 2009; Motzkin *et al.*, 2011; Sundram *et al.*, 2012) may reflect difficulties in learning from punishment.

Finally, Papagno (2011) reported that humans with unilateral uncinate fasciculus ablations had a highly specific semantic retrieval deficit in naming famous individuals. At first glance, these findings contradict our proposed model, namely that the unique connectivity of the uncinate fasciculus allows it to play a particular role in affective tagging of memory during storage and retrieval. However, further consideration is required. Salient, emotional information plays a significant role in person identification. Upon seeing a face that you recognize, you automatically activate a host of often salient and emotional, semantic associations beyond the mere name or identity of the person. An image of Marilyn Monroe, for example, evokes not only a name but also semantic and emotional associations, such as her performance in the film *Gentlemen Prefer Blondes*, her extreme version of female sexuality, her relationships with several famous men such as John F. Kennedy and Joe DiMaggio, and her tragic death by drug overdose. These emotionally laden associations support and strengthen your knowledge of Marilyn Monroe's identity, even if all you intend to do is access her name. Thus, the findings that person naming is associated with the uncinate fasciculus provides another layer of evidence for the model presented here.

Additional evidence for our proposal is clearly needed and there are many unanswered questions (Box 1). For instance, if our hypothesis is true, the question arises as to why there are no reports of patients with epilepsy who undergo incidental disconnection of the uncinate fasciculus who become psychopathic or exhibited abnormalities in forming emotional associations? As noted above, there is surprisingly little research on the uncinate fasciculus in epilepsy resection patients. Historically, research in these patients has focused on episodic memory and there are few studies of social or emotional processes. Even if such studies did exist, it

Box 1 Unanswered questions in the literature

- (1) There is an extensive literature documenting deficits in encoding and retrieving memories about individuals—their names, feelings of familiarity, or biographical knowledge—after anterior temporal lobe damage from resection or lesions (Grabowski *et al.*, 2001; reviewed in Olson *et al.*, 2013). One question that arises from this review is whether these deficits are best attributed to grey matter loss or white matter disconnection, specifically to the uncinate?
- (2) We report a large number of studies that have associated uncinate fasciculus perturbation with various psychiatric disorders. With the exception of psychopathy, most disorders do not appear to be uniquely or even particularly associated with differences in the uncinate fasciculus. Our hypothesis predicts that uncinate perturbation should cause a particular type of social–emotional problem due to the inability to acquire reward or punishment-based associations for people and objects. Within the realm of psychiatric disorders, are there any symptoms or treatment outcomes that can be directly linked to this proposed function?
- (3) Our hypothesis is vague regarding the effects of laterality on uncinate fasciculus function, reflecting the state of the literature. Although psychopathy and antisocial personality disorder are associated with differences in the right uncinate fasciculus, studies of other disorders report differences in the left or bilateral uncinate fasciculus. How do the left and right uncinate fasciculus differ?
- (4) Although we propose a transmission route between ventral anterior temporal lobe and the lateral orbital frontal cortex through the uncinate fasciculus, we neglect to discuss information transmission to BA 10 through the uncinate fasciculus. What are the details of information transmission to this region?

would be difficult to attribute any social/emotional dysfunction to uncinate fasciculus damage given that the amygdala is destroyed in most instances. In future work, it will be important to examine predictions of our model in resection patients with or without verified uncinate fasciculus disconnection.

The dearth of relevant findings forced us to be vague about several issues, such as laterality effects. We acknowledge that our proposal is speculative, given that supporting evidence is circumstantial (e.g. the uncinate fasciculus's connections to social and emotional grey matter regions), correlational (e.g. DTI), or based on lesion research, which is often imprecise. Most of the psychiatric and neurological findings were focused on correlating the presence or absence of a psychiatric diagnosis, such as schizophrenia, with differences in white matter tracts with little emphasis on linking white matter changes to specific clinical symptoms. In other cases, the uncinate fasciculus was not clearly delineated from nearby white matter tracts such as the cingulum bundle. Although each individual piece of evidence is weak, the summation of findings indicates that the uncinate fasciculus

transmits both social–emotional and mnemonic information, but of a particular type. The particulars of this function are only beginning to be understood. We urge future investigators to use larger and more homogenous clinical populations and to correlate specific symptoms, rather than presence or absence of disease state, with changes in white matter. Also, it would be wise to extend the important findings on memory functions in non-human primates to humans using DTI. To our knowledge, only two groups have achieved this (Metzler-Braddeley *et al.*, 2011; Thomas *et al.*, 2012).

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