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Disorders of visual perception

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Visual perceptual disorders

Abbreviations

SLF superior longitudinal fasciculus

ILF inferior longitudinal fasciculus

IFOF inferior fronto-occipital fasciculus

LGN lateral geniculate nucleus

Word count 4989

Abstract

Visual perceptual disorders are often presented as a disparate group of neurological deficits with little consideration given to the wide range of visual symptoms found in psychiatric and neurodevelopmental disease. Here we attempt a functional anatomical classification of all disorders linked to visual perception, whatever the clinical context in which they arise, including those disorders that bridge vision, emotion, memory, language and action. Guided by clinical and neuroimaging evidence, visual perceptual disorders are classified by the functional anatomical networks likely to be involved and the class of underlying dysfunction, whether topological (a localised deficit or region of hyperfunction) or hodological (a disconnection or hyperconnection). The wider perspective forces us to consider what visual functions underlie a range of symptoms sidelined by previous classificatory schemes and helps generate novel hypotheses for further research in the area.

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Introduction

If visual perception is the day-to-day experience of ‘seeing’ – the conscious, visual experience of the world we have when we open our eyes - a visual perceptual disorder might be considered a deviation from this experience. Current classificatory schemes focus on disorders traditionally considered neurological (i.e. based on static visual deficits typically related to brain lesions), relating each disorder to a specific brain region or its connections (for examples see ¹ and ²). Visual symptoms such as visual hallucinations, illusions and visual perceptual distortions or symptoms more likely to be found in psychiatric or neurodevelopmental disorders are typically only mentioned in passing or not at all in such schemes. This omission is understandable given that these types of symptom are often transient, and thus difficult to study directly, and may occur without an obvious associated brain lesion. Here we attempt to incorporate both ‘neurological’ and ‘psychiatric’ disorders into a single classificatory scheme using an anatomical network approach. Our aim is to provide an anatomically-structured overview of visual perceptual dysfunction across all clinical contexts that we hope will provoke further research.

A note on terminology

The scope of our classification is set by the way we have interpreted the three constituent terms of ‘visual perceptual disorder’. i) *Visual* The disorders classified here are *visual* in the sense that they are selective for vision without equivalent changes in other sensory modalities. For example, visual hypo-emotionality (a loss of emotional tone for visual stimuli) implies that, in some cases at least, emotions may be preserved for auditory and other sensory stimuli. ii) *Perceptual* If visual perception is the day-to-day experience of ‘seeing’, by this definition, its disorders might include, for example, a blurring of vision through loss of acuity. To limit the scope of the review we have focussed on symptoms attributable to dysfunction within cortico-cortical networks including those that bridge visual perception with memory, attention, emotion and action. We make a distinction between visual *perception* and visual *imagery* as the two differ in terms of their underlying neurobiology (see ³). Visual percepts are seen in the external world and are outside volitional control (one cannot change the percept by force of will) whereas visual imagery is seen in the mind’s eye and is under volitional control. We consider visual hallucinations and illusions as varieties of visual perceptual experience as they are closely related to visual perception in terms of underlying brain activity ³. Phenomena with elements of both perception and imagery are more difficult to

classify. For example, experiences referred to as pseudohallucinations or incomplete hallucinations (see ⁴ for other uses of these terms) are seen in the mind's eye but are outside volitional control. We have highlighted such ambiguous experiences in the classification and, where appropriate, presented both perceptual and imagery viewpoints. iii) *Disorder* We use the term *disorder* for continuity with previous clinical literature, although much of the classification relates to perceptual symptoms that do not imply a specific disease or pathology.

The hodotopic framework

We have classified visual perceptual disorders using the general anatomical framework we set out in 2005 as an update of classical associationist and neo-associationist accounts of higher cognitive function and dysfunction ⁵. Recent advances in functional neuroanatomy, using techniques such as functional magnetic resonance imaging (fMRI) and diffusion tensor tractography (DT-tractography), provide complementary views of higher function and the pathological mechanisms that generate cognitive and behavioural symptoms. One view, termed topological (from *topos* = place), relates to discrete cortical locations. The other, termed hodological (from *hodos* = path), relates to connections between locations. Both views are combined in a generic neuroanatomical model applicable to all higher functions and dysfunctions - the hodotopic framework ⁵. From the perspective of the hodotopic framework, each higher function relates to a specific network of cortical locations and white-matter connections. Topological disorders are localised within a cortical area and result in a loss (deficit) or an increase in function (hyperfunction). Hodological disorders relate to connections between areas, including those in which function in one brain region is altered by changes in another, spatially remote, region. We use von Monakow's term *diaschisis* to refer to such remote effects ⁶; however, the term is not intended here to carry its original implication of transient neurological shock. Connections between areas can be decreased (disconnection) or increased (hyperconnection) in hodological disorders. Deficit and disconnection disorders of visual perception are found in stroke, intracranial compressing masses, neurosurgical procedures, demyelinating disorders, neurodevelopmental conditions (e.g. autism), neurodegenerative disease, schizophrenia and depression. Hyperfunction and hyperconnection disorders of visual perception are found in ophthalmic disease, neurodegenerative disease, Parkinson's disease, psychosis, epilepsy and as normal

phenomena in relation to sleep or in the context of psychedelic drugs, alcohol or medication side-effects.

Classification method

We have allocated each visual perceptual symptom to a visual network and one of four classes of dysfunction (topological versus hodological; hyper- versus hypo-function/connection). For some symptoms, functional imaging, DTI-tractography or lesion evidence is available to help identify the appropriate location in the table (e.g. a localised increase in activity within a specific brain region). However, for many we have been forced to make an educated guess through: i) analogies to related symptoms for which evidence is available and ii) a neurophenomenological approach (see ^{3,7}). This approach is based on the body of evidence suggesting that some, as yet undefined, aspect of activity within a specialised cortical region (resulting in an increase in imaging signal ⁸) contributes directly to visual perceptual experience (e.g. the percept of a face is associated with increased activity within face-specialised cortex ⁹). Where evidence is available, this principle seems also to apply to hallucinatory, illusory and synaesthetic percepts (see ³). Thus the content of visual perceptual experience (whether an object, a face, motion etc.), localises activity to a particular brain network, irrespective of whether the experience is a normal percept, illusion, hallucination etc.

A taxonomy of disordered visual perception

Table 1 and Figure 1 outline a tentative classification of visual perceptual disorders.

Topological disorders of visual perception

Deficits – Prototypical disorders of this nature are characterised by the selective loss of a specific visual perceptual function e.g. achromatopsia ¹⁰ (selective loss of colour vision following lesions of colour-specialised cortex) and akinetopsia ¹¹ (selective loss of motion vision following lesions of lateral occipital motion-specialised cortex; the motion deficit is not absolute with a relative sparing of the perception of slow movement ¹²).

Lesions of V1 result in a scotoma restricted to a single hemifield (a hemianopia if the lesion involves the entire extent of V1 in one hemisphere). Lesions of ventral V2 / V3 are associated with upper quadrantic field defects while those of dorsal V2 / V3 are associated with inferior quadrantic field defects ¹³. Scotomata in the context of migraine aura relate to

regions of cortical spreading depression-like activity in early visual areas (e.g. V1, V2 and V3¹⁴).

Face perception deficits (prosopagnosia) have a variety of subtypes likely to reflect the involvement of different functional modules within the face processing system, although disconnection mechanisms may also be involved¹⁵.

An inability to recognise familiar places (environmental agnosia) is associated with medial temporal lobe lesions¹⁶ and can be considered a topological deficit of cortex specialised for the visual environment (parahippocampal place area¹⁷) or a hodological disconnection of visual and memory representations of places¹⁶.

Size and object distortions are referred to collectively as *metamorphopsias*. Objects appearing larger or smaller (macropsia and micropsia respectively) have been attributed to a dysfunction of object constancy within the temporal lobe (the integration of object distance with retinal visual angle to estimate size), although it is unclear whether this is best conceived as a topological dysfunction (i.e. a deficit or hyper-function of a cortical region specialised for constancy) or a hodological dysfunction (i.e. a disconnection or hyper-connection of relevant specialised modules). The same mechanism may be responsible for objects appearing nearer or further away (pelopsia and teleopsia), although such symptoms can also be interpreted as disorders of depth perception (and are also referred to as enhanced stereopsis⁴). Lesions of lateral occipito-temporal cortex may cause micropsia in one hemifield¹⁸ or bilaterally¹⁹, although as object constancy may be preserved¹⁹, such cases may relate to a deficit of object processing rather than constancy.

Apperceptive agnosia and cortical blindness describe diffuse visual loss affecting many visual domains. They occur in the context of extensive occipital lesions and are likely to involve both topological and hodological dysfunction. Several subtypes of disordered form perception have been described (see²⁰ for review), each likely to depend on the form perception sub-module(s) involved in the lesion. Scieropia refers to darkening of vision and follows diffuse occipital damage (see²¹ for an example: a patient whose vision was dimmed as if seeing everything at twilight). The same symptom occurring without other visual impairments (scierneuropsia) was described in the psychoanalytic literature²² and is likely to be an example of de-realisation (see below).

The Riddoch syndrome is a variant of cortical blindness in which motion-specialised functional networks remain intact. Subjects are able to consciously perceive and discriminate visual motion in their otherwise blind visual field when stimulated with fast motion²³, while

under other stimulus conditions report visual percepts without the ability to discriminate (gnosopsia) or have intact discrimination ability without visual percepts (blindsight or agnosopsia) ²³.

In visuo-spatial neglect, patients have a deficiency of orienting attention to one hemifield (usually left) and (may) fail to perceive stimuli in this region when rival stimuli are present in the non-neglected hemifield (extinction). The disorder has been interpreted as topological (related to the cortical modules involved in visuospatial processing and attention - see ²⁴ for review of regions implicated) but hodological mechanisms have also been suggested ²⁵. Simultanagnosia (an inability to sustain attention across different locations in the visual field) leads to objects appearing fragmented or disappearing from view ²⁶ with shrinkage of the visual attentional field ²⁷. The disorder is likely to relate to parietal lobe specialisations for spatial visual attention, although it is unclear whether the disorder is best conceived as topological or hodological. Rapid fading of vision found with occipito-parietal lesions ²⁸ is likely to be caused by the same mechanism as simultanagnosia.

Selective lesions of the amygdala are associated with perceptual deficits of emotional facial expression ²⁹ with similar deficits described in autism spectrum disorder and psychopathy.

Visual imagery may be selectively impaired (i.e. with intact auditory, olfactory, gustatory etc. imagery – see ³⁰ for review of case reports) with a relative preservation of visual perception. Loss of visual imagery may also be accompanied by a specific loss of visual dreaming (visual anoneria ³¹) or a more general loss of dreams in all sensory modalities (Charcot-Wilbrand syndrome ³²). Such disorders can be interpreted as topological deficits of visual imagery generation or hodological disconnections within visual memory networks.

Hyperfunction –Topological disorders of hyperfunction are related to pathophysiological mechanisms such as local irritation (e.g. in epilepsy or at the margins of a cortical lesion) and de-afferentation (e.g. caused by eye disease [Charles Bonnet Syndrome] or a visual pathway lesion; the mechanism by which loss of visual input results in increased cortical activity is unclear but may relate to changes in thalamocortical circuitry ³³).

Hallucinations linked to V1 and surrounding areas (V2 / V3) are of simple featureless forms, colours and television-like static (visual snow). These phenomena are termed phosphenes or photopsias by some authors although they are perhaps better referred to as simple hallucinations given that they differ from other types of visual hallucination only in the

location of cortex involved ³³. In migraine aura, zigzag lines and patterns (teichopsia ³⁴) relate to the margins of a region of cortical spreading depression-like activity in early visual cortex ¹⁴.

Colour, object and landscape hallucinations are each linked to their respective region of cortical specialisation ^{33, 35, 36} and the same is likely to be the case for motion hallucinations and passage hallucinations (visual hallucinations of persons or animals that pass sideways out of the visual field). Increased vividness of an attribute (e.g. increased vividness for colours: hyperchromatopsia ³⁷) is thought to relate to increased activity within the relevant specialised region (e.g. colour-specialised cortex for hyperchromatopsia). The origin of lilliputian hallucinations (small figures, often in costume) is unclear but has been classified here as related to object specialisations. The reduced size of such hallucinations may relate to visual constancy mechanisms ³⁷.

Text or letter-string hallucinations lacking semantic content are linked to the cortical area specialised for word forms, while those with semantic content (visual command hallucinations and visual verbal hallucinations ⁴) are linked to higher-level representations of visual language ³⁸.

Illusions of distorted faces (prosopometamorphopsia) relate to activity on the lateral convexity of the occipital lobe ³⁹, a region specialised for face features (the occipital face area) and the same region is likely to account for hallucinations of distorted faces. Hallucinations of normal (undistorted) faces relate to activity within ventral occipito-temporal cortex (fusiform face area) ³⁵ and the same region is likely to account for facial intermetamorphosis (a change in the visually perceived identity of a face).

A range of symptoms collectively termed palinopsia ^{37, 40} have been linked to dysfunction within parietal lobe co-ordinate systems ⁴¹. They include: i) polyopia (perceiving multiple copies of the same object often arranged in rows and columns; if the number of copies is large [e.g. > 100] it is referred to as entomopia ⁴²); ii) visual perseveration (an object remaining fixed in retinal co-ordinates with eye movements); iii) delayed palinopsia (an object returning to field of view); iv) illusory visual spread (the spread of a pattern from an object to its surroundings). Parietal lobe regions are also likely to underlie trailing phenomenon (a variety of visual perseveration in which a series of discontinuous stationary images trail behind a moving object) and positive afterimages (afterimages in which objects are seen in their true as opposed to complementary colours).

Patients with Anton's syndrome (visual anosognosia [denial of blindness] in the context of cortical blindness) report visual experiences that are traditionally interpreted as

confabulations (false memories or false reports of visual perceptual experience) or visual imagery experiences ⁴³. These symptoms are likely to be the result of topological hyperfunction within visual memory systems. Flashbulb memories (detail-perfect memories of events that at the time of witnessing evoked a high level of surprise, consequentiality and/or emotional arousal ⁴⁴) and memory hallucinations (fantastic false memories conjured up retrospectively ⁴) may relate to a similar hyperfunction of visual memory systems (and/or visual emotion systems in the case of flashbulb memories). Re-perceptive hallucinations involve the replay of memory fragments (also termed experiential hallucinations ⁴); however, it is unclear whether such experiences are visual imagery or visual percepts ⁴⁵. If visual imagery they are likely to involve hyperfunction of memory systems, if visual percepts they are likely to involve hyperconnection between visual memory networks and occipital cortex.

Autoscopic phenomena describe a range of experiences in which the self is duplicated in external space. In autoscopy, the duplicate self is seen in the external world. In out-of-body experience (OBE) the physical body is seen from the perspective of the duplicated external self. In heautoscopy the hallucinated 'double' is identified as oneself, despite the lack of an exact physical resemblance, and subjective viewpoint may change between the double and physical self in rapid alternation. These phenomena are thought to relate to the disintegration of visual, proprioceptive, tactile and vestibular modalities and have been linked to transient dysfunction in the region of the temporoparietal junction (TPJ) ⁴⁶. Whether this dysfunction is better conceived as topological hyperfunction of one or more of the sensory modalities involved or a hodological connectivity increase between specialised regions is unclear. Visual extracampine hallucinations (hallucinations perceived in impossible locations outside the field of view) could be considered related phenomena in that they involve an alteration in the reconstruction of visual space and body schema. Other disorders likely to be related to hyperfunction or hyperconnection within the TPJ and related regions include: i) the oculogravic illusion (an illusory displacement of objects co-occurring with a change in gravity as in a diving airplane ⁴⁷); ii) the oculogyral illusion (the apparent movement of a stationary light in a dark room during fixation following a period of rotation ⁴⁷); iii) visual hallucinations occurring in association with caloric stimulation ⁴⁸; iv) Zingerle's automatosis in which visual hallucinations occur during transient motor and vestibular dysfunction ³ (e.g. compulsive posturing, torticollis, choreo-athetosis); v) the transient intensification of visual patterns and text, often in association with neuroleptic induced oculogyric crises (paroxysmal perceptual alteration ⁴⁹).

Hodological disorders of visual perception

Disconnection – Disconnection of cortical visual networks from sub-cortical inputs as a result of visual pathway lesions or eye disease is associated with simple and complex visual hallucinations and visual illusions^{3, 7} (the Charles Bonnet Syndrome – the change in cortical function secondary to eye disease might be considered a form of diaschisis). Brainstem lesions are associated with peduncular hallucinations (complex hallucinations associated with eye movement disorders and changes in arousal or sleep⁵⁰).

Depth vision (visual space) involves a number of dissociable (see⁵¹) functions, some related to perceptual aspects of depth (e.g. stereopsis [binocular vision] and object constancy), others related to action aspects of depth (e.g. reaching behaviour). One aspect of depth vision (stereopsis) is lost in the central visual field (but not the visual periphery) following splenial section⁵².

In cinematographic vision, moving objects appear as a series of static freeze frames at different spatial locations⁴. The disorder may relate to the disconnection of motion specialised cortex and spatial co-ordinate frames in the parietal lobe.

Associative object agnosia (loss of object meaning with intact object perception) is thought to relate to the disconnection of visual and semantic regions (see²⁰ for review). Specific deficits of colour or object naming (colour anomia, object anomia) are thought to relate to the disconnection of visual and language areas while optic aphasia (an inability to name objects by sight with preserved ability to name by touch or from description) is thought to follow disconnection of left-hemisphere verbal semantic and right-hemisphere visual semantic representations⁵³.

Pure alexia (a deficit of reading words with preserved writing and letter recognition) and left hemialexia (a reading deficit restricted to the left hemifield) have been interpreted as hodological dysfunctions (a disconnection of a left-hemisphere region specialised for word forms from visual inputs) and topological dysfunctions (a deficit of the word form area) (see⁵⁴ for review). In surface dyslexia, the link between word forms and semantics is impaired so that reading occurs through phonology with characteristic errors for irregular words⁵⁵. The disorder might be considered a disconnection of regions specialised for word forms from regions specialised for visual semantics or a topological deficit of visual semantic specialisations.

The action aspect of depth vision (reaching behaviour) is dysfunctional in optic ataxia (impaired reaching with intact motor and visual function) and is thought to reflect an inability to

integrate different co-ordinate frames within the parietal lobe ⁵⁶. The failure of integration might reflect a hodological disconnection of the relevant reference frames or a topological deficit of an area specialised for their integration. A variant (crossed optic ataxia) in which reaching into contra-lateral hemispace is impaired (e.g. reaching with the right hand for an object located on the left side) is thought to relate to the disconnection of the two hemispheres with a combination of splenial and parietal lesions ⁵⁷.

Oculomotor apraxia (a disorder of gaze in which subjects are unable to voluntarily move gaze from one object to another with preserved reflex gaze shifts) can be interpreted as a disconnection of voluntary saccadic systems in the frontal lobe from reflex saccadic systems in the parietal lobe (a network classified here as related to visual attention).

Denial of cortical blindness (Anton's syndrome) might be considered a disconnection of visual cortex from body schema representations in the parietal lobe (classified here as related to visual attention). An inverse Anton's syndrome (denial of seeing) has been interpreted as a disconnection of parietal attentional mechanisms and visual cortex ⁵⁸. Denial of seeing specific objects reported in hypnosis (termed negative hallucination or scotomization ⁴) is likely to relate to the same mechanism.

In reduplicative disorders, a familiar person, place or object is perceived as replaced by a duplicate person, place or object ⁵⁹ (termed Capgras syndrome when involving a person). Reduplicative phenomena are thought to relate to the disconnection of visual and affective or memory regions ⁶⁰. Similar disconnection accounts are given for de-realisation (the feeling that experiences [typically visual] seem strange or unreal), visual hypoemotionality (where visual experience lacks emotional tone) ⁶¹ and visual amnesia (a deficit of registering visual experiences in short-term memory) ⁶².

Visual alloaesthesia involves the transposition, rotation, tilting or inversion of the visual field (also referred to as reversal of vision metamorphopsia ⁶³) and has variants where the visual environment (but not the visual field) is rotated ². The mechanism behind these disturbances is unclear but may relate to a disconnection of vestibular and visual inputs, transcallosal disconnection of visual and parietal inputs or topological lesions affecting regions specialised for the integration of these modalities ². The disorders are associated with occipito-parietal lesions that spare the optic radiations and with brainstem pathology ⁶³. Deficits of visual axis perception (e.g. judgment of true vertical or horizontal) are associated with lesions of the anterior parietal lobe and have been attributed to a disconnection of

vestibular inputs and their cortical representation or topological deficits of vestibular representations in the parietal lobe ⁶⁴.

The inability to mimic meaningless or meaningful actions or postures (visual imitative apraxia) is likely to reflect disconnection of visual and motor representations ⁶⁵. Utilisation behaviour (automatic manual exploration and use of objects placed within the field of view and reach) has been interpreted as a deficit of frontal inhibitory function on parietal-led exploratory behaviour ⁶⁶ and might be considered a form of diaschisis.

Hyperconnection – Of the four classes of dysfunction, hodological hyperconnection is the most speculative; however, there is indirect evidence that connectivity increases are linked to perceptual symptoms in the auditory domain ⁶⁷. Increased connectivity might lead to perceptual symptoms through, for example, enhanced feed-forward or feed-back processing or resonant activation through ephaptic (non-synaptic) transmission. The prototypical disorder in this class is synaesthesia. In the clinical domain, synaesthesia has traditionally referred to a percept in one sense modality evoking a parallel hallucinatory percept in a different sense modality, e.g. percepts of taste evoking visual percepts of colour (coloured taste synaesthesia). More recently, work in the cognitive sciences has lifted the requirement of synaesthetic experience to cross modalities through the study of, for example, coloured grapheme synaesthesia (visual percepts of text evoking visual percepts of colours). The induced synaesthetic experience may be perceived in the external world (projector synaesthesia) or inside the mind's eye (associator synaesthesia) ⁶⁸. Other parallel sensory experiences recognised clinically include reflex hallucinations (a sensory stimulus in one modality evokes a hallucination in another e.g. seeing a butterfly whenever the thumb is touched) and functional hallucinations (hallucinations are triggered in the same modality as the inducing stimulus e.g. hearing voices in parallel with the sound of running water). Both clinical symptoms would be classed as synaesthetic by contemporary cognitive science. We would argue the content of the parallel experience in such disorders is defined by the cortical specialisation of the area with increased connectivity. Thus, for example, synaesthesia, reflex or functional hallucinations that involve colour relate to the connectivity of colour-specialised cortex, while experiences involving spatial location (i.e. number forms - a variety of synaesthesia in which numbers and other ordinal lists [e.g. days of the week] are located at specific spatial locations ⁶⁹) relate to the connectivity of spatial representations in the parietal lobe.

Pareidolia refers to images of objects or faces seen in random visual patterns (e.g. cloud formations or tree bark). The experience is an illusion in which the inducing visual stimulus is typically of simpler form than the illusory percept and can be considered the result of hyperconnection between visual areas.

The Frégoli syndrome, in which unfamiliar people are perceived as familiar (typically as a person in disguise with malevolent intent), can be conceived as hyperconnection within visual emotional or memory networks. In PTSD flashbacks, patients relive stressful events, typically with multiple sensory modalities involved and a strong emotional component. The flashbacks might be considered the result of hodological hyper-connection between visual and emotional systems or topological hyper-function of visual memory systems.

The Perky effect originally referred to an ambiguity between visual imagery and visual perception such that normal subjects were unable to detect the presence of a real visual stimulus when asked to form a visual image of it. More recently the term has been used to describe the reduction of visual perceptual sensitivity during visual imagery⁷⁰. The effect is likely to relate to connectivity between visual memory networks and occipital cortex. Eidetic imagery (sometimes referred to as photographic memory)⁷¹ is likely to relate to increased connectivity between these regions.

Visual phobias and visual objects appearing beautiful and comforting (kalopsia) or ugly and menacing (kakopsia)⁷² might be considered the result of hodological hyperconnection between visual emotional networks and occipital cortex or topological hyperfunction of visual specialisations within the amygdala.

Dazzle (painful brightness of visual stimuli) is thought to be a visual variant of a thalamic pain syndrome⁷³ and can be interpreted as hodological hyperconnection of the thalamus and visual cortex. The same mechanism may account for the intense photophobia of children with cortical visual impairment and the paradoxical worsening of their visual acuity (as measured by the evoked response) for lighter (higher luminance) compared to darker (lower luminance) visual stimuli⁷⁴.

Limitations

The classificatory scheme has a number of limitations and is perhaps better viewed as a structured review of symptoms from which to generate and test further experimental hypotheses than a definitive pathophysiological taxonomy or clinical *aide memoire*. (i) It does not easily accommodate disorders involving combinations of networks or combinations of

topological and hodological dysfunction (the latter marked with an asterisk in the Table). Typical lesions have both hodological and topological effects so that pure disorders of one or other class may be largely theoretical. (ii) The classification deconstructs traditional clinical syndromes by placing their component parts in different locations, losing associations between symptoms related to the anatomical proximity of the cortical regions underlying them. For example, in migraine aura, the same region of cortical spreading depression may result in a hyper-functional symptom at its margins (teichopsia) and a deficit in its centre (a scotoma) but each is placed in a different location in the table. Similarly the Balint syndrome (simultanagnosia, optic ataxia and oculomotor apraxia in association with bi-parietal lesions⁷⁵) is spread across the table. (iii) Hyper- and hypo- function are convenient concepts to map on the statistical signal increases and decreases of imaging and tractography data. However, the neurophysiological meaning of such changes is unspecified. An increase in activity as measured by fMRI may have a number of neurophysiological causes (e.g. a change in excitatory input or inhibitory processing, a shift from burst to tonic mode firing [see³³], neuromodulatory inputs etc.). Similarly DT-tractography dissects *virtual* connections between brain regions and different causes of micro-structural white-matter change may lead to the same change in connectivity. By focussing on increases or decreases in imaging signal, the classification is likely to miss key pathophysiological distinctions. (iv) The classification does not easily accommodate interactions between pre-disposing factors and changes occurring at the time of a symptom. The Charles Bonnet Syndrome, for example, appears both as a hodological disconnection (because of the predisposing factor of loss of visual inputs from the eye) and as topological hyperfunction (because of the localised increases in fMRI signal during hallucinations).

Clinical implications

Despite its limitations, the classificatory scheme and its underlying assumptions are of relevance to current clinical practice in a number of areas. First it suggests that, in the same way that seizure semiology is used in the context of epilepsy, the contents of hallucinations and illusions can be used as localising signs. By indicating increased activity in a specific functionally specialised cortical area, different types of visual hallucinations (e.g. a face, colour or geometrical pattern) point to specific cortical locations, although not the location of the pathology causing the activity increase. Second, the classification highlights the fact that many symptoms held traditionally as distinct, appear closely related, if not indistinguishable,

when considered from the perspective of their underlying mechanism. For example, synaesthesia, functional hallucinations and reflex hallucinations are all likely to relate to the same underlying pathophysiology. Third it predicts disorders which are, as yet, undescribed, or at least not recognised as distinct clinical entities. For example, disorders classified here as hyperfunction of spatial co-ordinate frames within the parietal lobe (polyopia, visual perseveration, palinopsia etc.) do not have obvious deficit or disconnection counterparts. It is conceivable that their natural counterparts are, in fact, spatial neglect, simultanagnosia, oculomotor apraxia, classified here as related to visual attention, as all are linked to the parietal lobe and dorsal stream. Finally, the classification highlights a number of areas which pose a challenge to visual neuroscience. For example, lilliputian hallucinations are not easily accommodated within current models of object perception, while polyopia and extracampine hallucinations are not easily accommodated within current models of spatial vision. In pointing to deficiencies in such models, visual perceptual disorders advance our understanding of normal visual function.

Conclusions

The last decade has seen advances in white-matter and functional imaging (including EEG imaging) that allow new classes of clinically relevant questions to be answered (see ⁷⁶). We envisage that such imaging techniques, applied to single case and group studies, will be used to test and expand the classification presented here in the future (see^{5, 76}). For now, we hope that our broad overview of visual perceptual disorders will provide a fresh perspective from which to challenge and refine the neuroscience of visual perception and visual perceptual dysfunction.

Figure legend

Figure 1 The anatomy of visual networks. The upper left panel shows the major associative tracts with connection to the occipital lobe in the human brain. SLF = Superior longitudinal fasciculus; ILF = Interior longitudinal fasciculus; IFOF = Inferior longitudinal fasciculus. The Brodmann areas corresponding to the cortical projections of each fibre tract are given in the lower left panel. The right panels show Brodmann areas colour coded on lateral (upper) and medial (lower) views of the brain. Images modified from ⁷⁷. The primary visual cortex (V1, area 17) is located on the polar and medial surface of the occipital lobe. V2 is adjacent to V1 and is divided on the medial occipital surface into a dorsal portion and a ventral portion. V3 is adjacent to V2 and is similarly divided. The occipital lobe, lateral and ventral occipitotemporal cortex, limbic cortex and the parietal lobe contain regions specialised for different visual perceptual attributes including: familiar faces, colours, extended landscapes, objects, text and visual textures (ventral occipito-temporal cortex) and visual motion, body parts, face features, intermediate-level object processing and eye and mouth movements (posterior-lateral cortex) (see ⁷). The predominant pathway from the eye to V1 relays through the lateral geniculate nucleus (LGN), although a clinically important pathway passes directly from the LGN or pulvinar to motion specialised cortex on the lateral occipital surface ²³. The detailed anatomy of short association fibres within and between occipital regions and from the occipital lobe to the pulvinar has yet to be characterised in man. Chains of U-shaped fibres and parallel long association pathways passing to the parietal lobe form a cortical network (dorsal stream) ⁷⁸ thought to underlie spatial aspects of vision while equivalent pathways to the temporal lobe form a cortical network (ventral stream) thought to underlie object vision, colours, faces and visual language. The inferior longitudinal fasciculus (ILF) carries reciprocal connections between the occipital and medial / lateral anterior temporal lobes related to visual emotion and visual memory ⁷⁹. The inferior fronto-occipital fasciculus (IFOF) connects territories in the occipital lobe and ventral frontal lobe ⁷⁸. Connections between the visual parietal lobe and frontal cortex are carried in the superior longitudinal fasciculus (SLF) while those between occipital and limbic cortex are carried in the cingulum and ILF. The SLF and IFOF are likely to relate to functions such as visual attention, visual working memory, visual praxis, visual-vestibular function and eye movement control. Cortico-cortical visual connections between hemispheres pass predominantly through the splenium of the corpus callosum.

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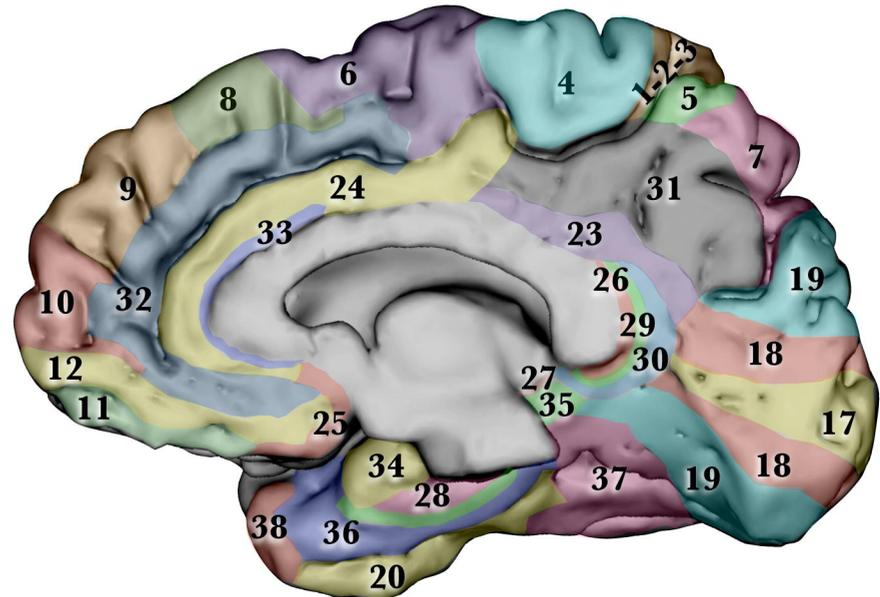
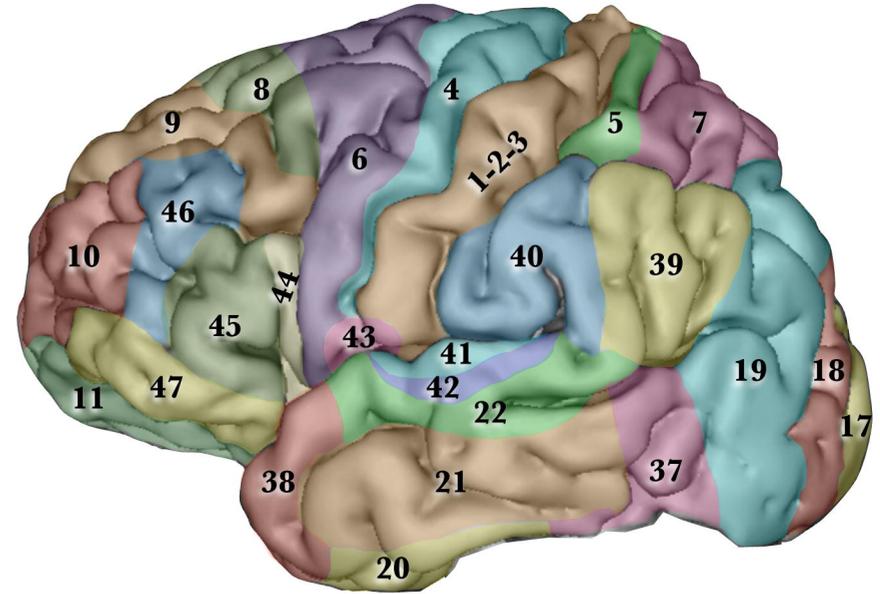
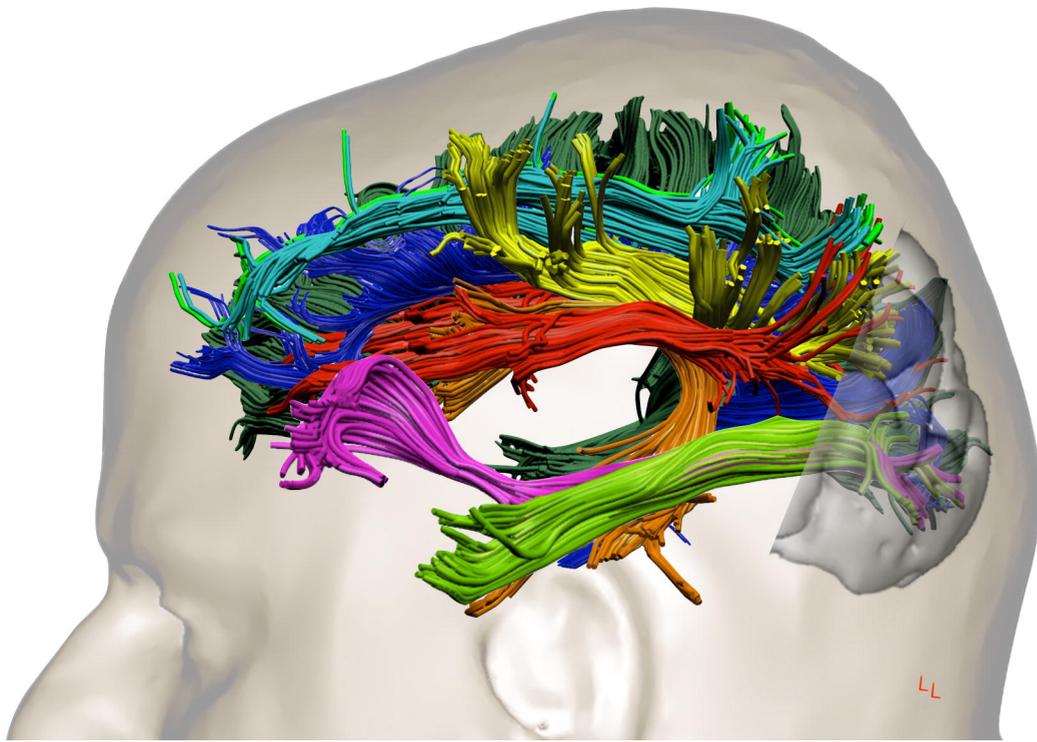
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		Topological		Hodological			
		Cortex	Hyperfunction	Deficit	White matter	Hypoconnection	Hyperconnection
Subcortical / Brainstem			Phosphenes	Brainstem / subcortical visual deficits	Subcortical pathways	Charles Bonnet syndrome Peduncular hallucinosis	Dazzle
V1 / V2 / V3	BA17/18		Phosphenes/ Simple hallucinations Visual snow Teichopsia	Cortical blindness* Scotoma Apperceptive agnosia* Scieropia* Riddoch syndrome	U-shaped	Cortical blindness* Apperceptive agnosia* Scieropia*	
<i>Depth (stereopsis)</i>					Splenium	Astereopsis (split brain)	
Ventral Stream	BA18/19		Colour hallucination/illusion/ hyperchromatopsia	Achromatopsia	ILF/Arcuate U-shaped	Colour anomia*	Coloured hearing, coloured music, coloured grapheme synaesthesia
<i>Colour</i>							
<i>Face gestalt</i>	BA37		Face hallucination/illusion Facial intermetamorphosis*	Prosopagnosia*	ILF/IFOF	Prosopagnosia*	Pareidolia for faces
<i>Face feature</i>	BA19		Prosopometamorphopsia	Prosopagnosia*	ILF/IFOF	Prosopagnosia*	
<i>Object</i>	BA18/19/37		Object hallucination/illusion Lilliputian hallucinations	Object agnosia* /Micropsia	ILF/IFOF/ Arcuate	Object agnosia* Object anomia Optic aphasia	Pareidolia for objects
<i>Places</i>	Parahippocampal gyrus		Landscape hallucinations	Environmental agnosia*			
<i>Constancy</i>			Micropsia / macropsia* Pelopsia/telopsia*	Micropsia / Macropsia* Pelopsia/telopsia*		Micropsia / macropsia* Pelopsia/telopsia*	Micropsia / macropsia* Pelopsia/telopsia*
<i>Text</i>	BA37		Visual text hallucination	Alexia*	ILF/Splenium/ U-shaped	Alexia* Left hemialexia (split brain)	Coloured grapheme synaesthesia
<i>Text Semantics</i>	BA20/21/38		Visual command hallucination	Surface dyslexia*	ILF/IFOF	Surface dyslexia*	
Dorsal Stream	BA19/37		Motion hallucination/illusion Passage hallucination	Akinetopsia	U-shaped	Cinematographic vision	
<i>Motion vision</i>							
<i>Spatial co-ordinate frames</i>	BA7/31		Polyopia /Entomopia Visual perseveration Trailing phenomenon Delayed palinopsia Illusory visual spread Positive afterimages		U-shaped		Number forms
<i>Depth (reaching)</i>	BA7/19/39			Optic ataxia*	U-shaped	Optic ataxia*	

				Splenium	Crossed optic ataxia (split brain)	
Visual attention	BA6/7/8/39/40		Visuo-spatial neglect* Extinction* Visual fading* Simultanagnosia* Inverse Anton's* Negative hallucination*	SLF/ Cingulum/ U-shaped	Visuo-spatial neglect* Extinction* Visual fading* Simultanagnosia* Inverse Anton's* Negative hallucination*	
	BA6/7		Oculomotor apraxia*	SLF	Oculomotor apraxia*	
<i>Awareness of body schema</i>				U-shaped	Anton's syndrome (anosognosia)	
Visual memory Imagery	Hippocampus Parahippocampal gyrus BA11/47/45	Anton's syndrome (confabulation) PTSD flashbacks* Flashbulb memories Memory hallucination Re-perceptive hallucination*	Visual imagery deficits* Visual anoneria*	ILF/IFOF	Visual imagery deficits* Visual anoneria* Visual amnesia	Perky effect PTSD flashbacks* Eidetic imagery Re-perceptive hallucination*
<i>Faces</i>				ILF/IFOF	Capgras syndrome	Frégoli's phenomenon
<i>Places</i>				ILF/IFOF	Reduplication of places Environmental agnosia*	
Visual emotion	Amygdala BA11/47/45	Phobic disorders* PTSD flashbacks* Flashbulb memories	Fear recognition deficits	ILF/IFOF	Visual hypo-emotionality De-realisation	Phobic disorders* PTSD flashbacks* Kakopsia/Kalopsia
<i>Faces</i>				ILF/IFOF	Capgras syndrome	Frégoli's phenomenon
<i>Places</i>				ILF/IFOF	Reduplication of places	
Visual vestibular	BA22/39/40	Autoscopy/Heautoscopy/OBE* Extracampine hallucinations* Oculogravic illusion* Oculogyral illusion* Caloric visual hallucinations* Zingerle's automatosis* PPA*	Visual alloaesthesia / Inverted vision*	U-shaped/ Arcuate	Visual alloaesthesia / Inverted vision*	Autoscopy/Heautoscopy/OBE* Extracampine hallucinations* Oculogravic illusion* Oculogyral illusion* Caloric visual hallucinations* Zingerle's automatosis* PPA*
<i>Visual axis</i>	BA40/39/insula		Axis impairment* Environmental tilt*	U-shaped/ Arcuate	Axis impairment* Environmental tilt*	
Visual motor Limb Praxis				SLF	Utilisation behaviour	
				SLF/Arcuate	Visual imitative apraxia	

* disorders likely to relate to both hodological and topological dysfunction. BA = Brodmann Area; PPA = Paroxysmal perceptual alteration; PTSD = Post-traumatic stress disorder



- | | | | |
|---|---|---|----------------------------|
|  | Arcuate (BA42,21, 22,37,6,44) |  | SLF II (BA6,39,40) |
|  | SLF I (BA6,7,8,9) |  | SLF III (BA39,40, 6,44) |
|  | Corpus callosum |  | IFOF (BA18,19, 37, 11, 47) |
|  | ILF (ventral BA18,19,20,21,37, 38, amygdala, hippocampus) | | |
|  | Cingulum (BA6, 7,8,24,26,31,32,33,34) | | |