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A pathophysiological framework of hippocampal dysfunction in ageing and disease

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Abstract

The hippocampal formation has been implicated in a growing number of disorders, from Alzheimer's disease and cognitive ageing to schizophrenia and depression. How can the hippocampal formation, a complex circuit that spans the temporal lobes, be involved in a range of such phenotypically diverse and mechanistically distinct disorders? Recent neuroimaging findings indicate that these disorders differentially target distinct subregions of the hippocampal circuit. In addition, some disorders are associated with hippocampal hypometabolism, whereas others show evidence of hypermetabolism. Interpreted in the context of the functional and molecular organization of the hippocampal circuit, these observations give rise to a unified pathophysiological framework of hippocampal dysfunction.

Neuroimaging and neuropsychological studies have, among other observations, implicated the hippocampal formation in Alzheimer's disease, temporal lobe epilepsy (TLE), cognitive ageing, post-traumatic stress disorder (PTSD), transient global amnesia, schizophrenia, and depressive and anxiety disorders. Although overlaps exist, these disorders are clearly not phenocopies and, more importantly, are thought to have distinct pathogenic mechanisms. Resolving how the hippocampal formation can be affected by a broad range of disorders is the goal of this Review.

Until recently, most neuroimaging and neuropsychological tests have evaluated the hippocampal formation as a singular structure, but it is in fact a complex circuit made up of functionally and molecularly distinct subregions. Moreover, the complexity of the hippocampal formation extends beyond its internal circuit organization. Most neural information funnels into the circuit through a restricted area, whereas the outflow fans out,

monosynaptically connecting with a broad range of cortical and subcortical sites. In the first section of this Review we will briefly summarize the internal circuitry of the hippocampal formation and describe how hippocampal efferents connect with separate brain networks (FIG. 1).

In the second section of this Review we survey studies that use high-resolution variants of structural and functional MRI (fMRI) that can visualize and assess the integrity of individual hippocampal subregions (BOX 1; FIG. 1). By simultaneously assessing multiple subregions, the hippocampal formation can be interrogated as a circuit, and these imaging approaches are well suited to pinpoint subregions that are differentially affected by, or resistant to, a particular disorder. Over the past few years, these high-resolution imaging methods have been applied to numerous disorders. As will be reviewed, when these studies are examined together, a clear picture emerges in which patterns of regional changes can differentiate disorders that are associated with hippocampal dysfunction.

The concept of regional vulnerability, across the brain and within the hippocampal formation in particular, is not new. For example, the CA1 subfield is known to be the hippocampal subregion differentially vulnerable to vascular disease^{1,2}, whereas the dentate gyrus is known to be differentially vulnerable to the effects of adrenalectomy³. Recent gene-expression studies have established that each hippocampal subregion has a distinct molecular profile⁴⁻⁶, and this ‘molecular anatomy’ provides a partial substrate for regional vulnerability. So, the relatively high expression of certain NMDA receptors⁷ in CA1 helps to explain its vulnerability to excitotoxicity in the context of hypoxia and ischaemia associated with vascular disease^{1,2}, whereas the high levels of mineralocorticoid receptors in the dentate gyrus confer vulnerability to the effects of reductions in the level of circulating corticosteroids⁸. As will be discussed, demonstrating and reinforcing the concept of regional vulnerability is important for providing insights into pathogenic mechanisms as well as for explaining phenotypic variability.

In the third section of the Review we show that functional imaging techniques have identified another, perhaps more surprising, factor by which diseases that affect the hippocampal formation can be segregated. Imaging techniques such as positron emission tomography (PET) and some variants of fMRI can identify the metabolic state of the hippocampal formation. In many diseases the hippocampal formation is found to be hypometabolic. Hypometabolism is expected because these disorders are characterized by hippocampal ‘loss of function’, such as memory deficits. In other disorders, however, there is evidence that the hippocampal formation is in a hypermetabolic state. This observation introduces the interesting idea that, as in TLE, some hippocampus-based disorders may cause ‘gain of function’ symptoms by stimulating hippocampal outflow areas. We will review studies that raise the intriguing possibility that hypermetabolism in the anterior hippocampus might mediate a gain of psychotic and affective symptoms.

When the concepts of regional vulnerability and metabolic state are viewed in light of the functional and molecular organization of the hippocampal circuit, a general pathophysiological framework of hippocampal dysfunction begins to emerge. In the final section of this Review, we will elaborate on this framework, showing its utility in explaining and better characterizing phenotypes that distinguish hippocampal disorders, and demonstrating how it can be used to shed light on pathogenic mechanisms.

Organization of the hippocampal formation

The hippocampal formation spans the posterior-to-anterior extent of the base of the temporal lobes (FIG. 1a). In its transverse axis the hippocampal formation is organized as a mainly unidirectional circuit made up of multiple subregions — the entorhinal cortex, the dentate

gyrus, the CA1 and CA3 subfields, and the subiculum⁹ (FIG. 1b). Notably, in his classic study, Lorente de N6 — who first parsed the hippocampal formation using the Cornu Ammonis nomenclature — also described the CA2 subfield¹⁰. Although there has been renewed interest in CA2, it is not commonly imaged *in vivo*, and will therefore not be featured in this Review.

In the ‘trisynaptic pathway’, layer II of entorhinal cortex connects to the dentate gyrus through the perforant pathway, and the dentate gyrus connects to CA3 through the mossy fibres. CA3 neurons interconnect with other CA3 neurons up and down the hippocampal long axis through ‘auto-associative’ tracts, or with CA1 through the Shaffer collaterals. Finally, CA1 connects to the subiculum. In addition to this pattern of connectivity, layer II of the entorhinal cortex projects to CA3, and layer III of entorhinal cortex can send direct connections to CA1 and the subiculum (FIG. 1b).

The entorhinal cortex serves as the gateway into the hippocampal formation and receives monosynaptic input from numerous regions, including the perirhinal cortex, the parahippocampal cortex, the auditory and olfactory cortices, and the amygdala. The input to the entorhinal cortex is organized in an anterior-to-posterior gradient, and the gradient is largely preserved in the way that the entorhinal cortex connects with the rest of the hippocampus (FIG. 1a). So, for example, input from the amygdala connects with anterior and medial aspects of the entorhinal cortex, which then communicate this input to anterior aspects of the hippocampus. By contrast, visual cortex input is delivered to the hippocampal formation through the perirhinal and parahippocampal cortices, which connect to posterior and lateral aspects of the entorhinal cortex, and from there to posterior aspects of the hippocampus.

The subiculum and, to a lesser extent, CA1 are the hippocampal subregions that provide the dominant out-flow of the hippocampal circuit¹¹. Both CA1 and the subiculum connect to deep layers of the entorhinal cortex (FIG. 1b), which then re-connects — via the parahippocampal gyrus — with neocortical sites that provided the original hippocampal input¹². This hippocampal–neocortical network has historically received the most attention for its role in the consolidation of long-term memories¹³. Anatomical tracing studies have established that the outflow from the subiculum and CA1 can also bypass the entorhinal cortex, monosynaptically connecting with a range of other brain areas, including the amygdala, the medial prefrontal and orbitofrontal cortices, the anterior and posterior cingulate cortex, and the nucleus accumbens^{14–16} (FIG. 1c). These direct hippocampal efferents show a striking topographical organization, such that the more posterior (dorsal in rodents) part projects to the posterior cingulate cortex, whereas more anterior parts show denser projections to medial prefrontal and orbitofrontal cortices and the amygdala.

The observed input and outflow gradients suggest that the hippocampal long axis is functionally organized, an interpretation that agrees with data from gene expression profiling^{4,6}, fMRI studies¹⁷, and electrophysiological studies that have documented long-axis differences in mechanisms of plasticity^{18–20} and place field activity^{21,22}. Building on these observations, a range of studies have established that the posterior extents of the hippocampal long axis are more likely to be involved in memory and cognitive processing, whereas the anterior extents of the hippocampal long axis are more involved in other complex behaviours such as stress, emotion, sensory–motor integration and goal-directed activity^{4,23–26}. Notably, and as will be discussed below, the anterior hippocampus and some of its associated outflow areas are key components of networks that are implicated in schizophrenia and depression.

Hippocampal regional vulnerability

Rather than reviewing all disorders in which the hippocampal formation is implicated, we will focus on a number of major ones, with the purpose of establishing the principle of differential vulnerability. The term ‘differential’ (rather than ‘selective’) is deliberately used: because of connections within the hippocampal circuit, a primary lesion or dysfunction in any subregion can, over time, affect neighbouring subregions. By comparing studies that have mapped patterns of anatomical alterations within the hippocampal circuit, patterns that differentiate disorders can be determined, thereby pinpointing differentially vulnerable and resistant subregions.

For disorders in which there is death of primary hippocampal neurons, mapping patterns of cell loss provides a histological indication of regional vulnerability. In vascular disease, CA1 shows the most reliable loss of neurons^{27,28}. In Alzheimer's disease, the pattern of cell death and other histological findings suggests that the entorhinal cortex is prominently affected by the disease^{29–31}, that CA1 and the subiculum are also heavily involved, and that the dentate gyrus and CA3 are relatively preserved^{29,30,32}. Although cell death in TLE³³ is extensive — occurring predominantly in the dentate gyrus and CA subfields — two findings differentiate TLE from Alzheimer's disease. First, the subiculum is a subregion with relatively little cell loss in TLE. Second, although entorhinal cortex cell death is observed in both diseases, it is more pronounced in Alzheimer's disease, in which it starts in layer II, whereas in TLE it seems to be confined to layer III neurons³⁴.

High-resolution variants of functional MRI have successfully detected the differential vulnerability of CA1 to vascular disease in living patients². Furthermore, these technologies have been able to detect Alzheimer's disease-associated alterations in entorhinal cortex, CA1 and subiculum, on a background of relatively resistant dentate gyrus and CA3 (REFS 35–37), and have confirmed that the subiculum is relatively unaffected in TLE³⁸. By showing concordance with histologically detected patterns of vulnerability, these observations validate that neuroimaging can accurately map hippocampal pathology *in vivo*.

The other disorders considered in this Review are notable for their relative absence of loss of primary neurons and are therefore often referred to as ‘functional’ disorders. Mapping patterns of differential vulnerability for this class of disorders is particularly challenging — not only because of the absence of cell death but also because the ‘functional’ pathology readily spreads over time. It is for these disorders that high-resolution neuro-imaging has proven most useful (BOX 1). Neuroimaging techniques can map not only patterns of hippocampal dysfunction in living subjects but, as we will highlight, can also map spread over time. We will begin by reviewing disorders that typically affect the hippocampal formation in later-life and then review those that occur more commonly in younger patients.

Pinpointing vulnerable subregions in the hippocampal formation in late life

Although ageing is itself not a disease, cognitive decline that occurs during ageing has deleterious consequences, and because it occurs without prominent cell loss it is considered a prime example of a functional disorder³⁹. A great number of imaging studies have investigated the hippocampal formation in ageing individuals and many findings have been reported. Nevertheless, the effect of ageing on hippocampal function is often confounded by diseases², in particular Alzheimer's disease and vascular disease, which commonly occur in older subjects and can cause hippocampal dysfunction independent of ageing². When attempting to isolate the hippocampal pattern of dysfunction reflective of ageing per se, it is therefore important to exclude the effect of Alzheimer's disease and vascular disease.

The first clues that ageing manifests with a pattern of hippocampal dysfunction that is distinct from Alzheimer's disease or vascular disease came from studies using a high-resolution variant of fMRI that was explicitly developed to map patterns of basal hippocampal metabolism in humans and animal models^{40–42}. The term 'basal metabolism', as historically used in the functional imaging field, refers to the resting-state metabolism in a brain region. This resting-state metabolism can change slowly during ageing or in disease, in contrast with acute changes in metabolism that are evoked by a transient stimulus (BOXES 1,2)). As Alzheimer's disease and vascular disease do not typically occur in non-human primates or rodents, a comparison of fMRI findings across species can isolate a pattern of dysfunction that is differentially linked to ageing. These fMRI variants have been applied to ageing human subjects⁴³ (with or without Alzheimer's disease³⁵ and with or without vascular disease²), ageing rhesus monkeys⁴¹, ageing mice and mice that express disease-causing mutations in the amyloid precursor protein³⁵. Besides confirming the differential link of Alzheimer's disease to the entorhinal cortex and vascular disease to CA1, results from these cross-species studies suggested that ageing itself differentially affects the dentate gyrus^{2,35,41,43}.

Informed by these fMRI findings, recent studies have set out to develop a cognitive task that can assess the functional integrity of the dentate gyrus in humans. As first theorized by Marr⁴⁴, and empirically established in rodent studies^{45–48}, the human dentate gyrus plays a part in 'pattern separation'⁴⁹, a cognitive operation that allows similar stimuli flowing through the hippocampal circuit to be represented with distinct neural codes. Accordingly, memory tasks have been developed that can assess pattern separation ability in humans⁴⁹. Impairments in these tasks have been documented in ageing subjects^{50,51} and have been interpreted as confirming previous fMRI findings that implicated the dentate gyrus in normal ageing^{39,40}. Although the computational process underlying pattern separation occurs within the dentate gyrus itself, impaired performance might also be expected with upstream lesions in the entorhinal cortex, which provides the main dentate gyrus input (FIG. 1b). Indeed, impairment in pattern separation tasks has been suggested in subjects who might be in the early stages of Alzheimer's disease, in which the entorhinal cortex is affected⁵². We will return to this point in the last section, in which we discuss how cognitive tests that engage the entorhinal cortex suggest that there are cognitive phenotypes that differentiate Alzheimer's disease from ageing.

Volumetric techniques using structural MRI were originally used to estimate the volume of the entorhinal cortex and the rest of the hippocampus (including the dentate gyrus, CA3, CA1 and the subiculum) as a single region of interest (ROI). Longitudinal studies have shown that although entorhinal cortex shrinkage can be observed at very old ages, the rest of the hippocampus (not the entorhinal cortex) showed a reliable age-related decline across age groups, tracking the time course of age-related memory decline^{53,54}. More recently, techniques have been developed that can estimate the volumes of the entorhinal cortex as well as the other individual hippocampal subregions, and therefore can pinpoint which subregion is driving the observed effect. Because of the difficulty in visualizing the precise boundaries between the dentate gyrus and CA3, in these studies the two subregions are typically lumped into a single ROI. A study that used these high-resolution techniques to compare patients with Alzheimer's disease with age-matched controls observed Alzheimer's disease-associated shrinkage in the entorhinal cortex, CA1 and subiculum, with sparing of the dentate gyrus or CA3 (REF. 36). When evaluating healthy subjects across the age span, the entorhinal cortex was found to be relatively resistant to ageing, with changes isolated in the dentate gyrus and CA3, and in CA1 (REF. 37). The effect in dentate gyrus and CA3 was more pronounced and better tracked with the temporal profile of age-related memory decline. Moreover, the ageing effect in CA1 was linked to white matter hyperintensities (S. Mueller, unpublished observations), a marker of vascular disease. A more recent volumetric

study also showed age-related effects in CA1, and dentate gyrus and CA3 (REF. 55), with relative preservation of entorhinal cortex volume. Here, shrinkage of dentate gyrus and CA3 correlated with age-related memory decline, whereas CA1 shrinkage was associated with hypertension and was therefore interpreted as reflecting the differential sensitivity of CA1 to vascular disease².

fMRI can also be used to map stimulus-induced changes in the blood oxygen level-dependent (BOLD) response, which is thought to reflect evoked neural activity (BOXES 1,2). Recently, a high-resolution variant of BOLD imaging has been developed and has been applied in ageing individuals and patients with Alzheimer's disease. Normal ageing was characterized by an abnormal stimulus-evoked BOLD response that was restricted to the combined dentate gyrus and CA3 ROI, with normal responses observed in the entorhinal cortex and subiculum⁵¹. By contrast, subjects with presumptive early Alzheimer's disease showed an abnormal evoked BOLD response in both the entorhinal cortex and in the downstream dentate gyrus or CA3 (REF. 52). Finally, a recent study investigated ageing subjects using a novel variant of diffusion tensor imaging (DTI)⁵⁶, an MRI-based technique that may detect white matter and dendritic integrity⁵⁷. The authors interpreted their result as being suggestive of perforant path alterations and, importantly, an age-related decrease in dendritic integrity that was localized selectively to dentate gyrus or CA3, and not observed in entorhinal cortex, CA1 or subiculum⁵⁸.

Taken together, the diverse range of imaging studies show that Alzheimer's disease, vascular disease and ageing all contribute to hippocampal alterations in late life, and over time can affect overlapping subregions. Nevertheless, a direct comparison identifies patterns of alterations within the transverse hippocampal circuit that differentiates the three conditions: the entorhinal cortex is differentially associated with Alzheimer's disease, CA1 with vascular disease, and the ageing process per se seems to differentially target the dentate gyrus (FIG. 2a). Over the hippocampal long axis, posterior CA1 is differentially affected by vascular disease⁵⁹. Although a systematic investigation of age-related dentate gyrus alterations over the hippocampal long axis has not yet been completed, a recent study suggests that age-related pattern separation defects occur in relatively posterior aspects⁵⁸.

Pinpointing vulnerable subregions in the hippocampal formation in adolescence and young adulthood

Although they can occur at any age, schizophrenia and depression are diseases that typically begin in adolescence. Triggered by a traumatic event, PTSD can of course also occur at any age. Nevertheless, reflective of current events, the incidence of PTSD is highest in young adults, and we have therefore included a discussion of PTSD in this section.

As well as estimating the volume of individual subregions, mapping alterations in the three-dimensional shape of the hippocampus has emerged as another structural MRI technique that can localize anatomical differences in the hippocampal formation. In patients with schizophrenia, this approach has implicated anterior aspects of the hippocampal long axis and, within the hippocampal circuit, changes were found in the anterior CA1 and subiculum^{60,61}. Interestingly, these studies suggest that the left hippocampus is typically more affected than the right. A subsequent high-resolution fMRI study that mapped basal hippocampal metabolism in patients with schizophrenia confirmed this pattern⁶². This study also imaged subjects without psychosis who were at risk of developing the disease and, by following the subjects over time, was able to identify patterns of dysfunction in prodromal stages of the disease. The findings suggested that the disease process might start in anterior CA1 and spread, over time, to the anterior subiculum and other brain regions⁶².

Mapping hippocampal shape has also been used to investigate major depressive illness. As with schizophrenia, the greatest difference between the results from patients with depression and controls was predominantly in anterior aspects of the hippocampal long axis^{63,64}. Within the hippocampal circuit, one study pinpointed shrinkage to the anterior subiculum⁶³ (FIG. 2b). A second study confirmed the finding in the anterior subiculum but also found shrinkage in anterior CA1 (REF. 64). Besides technical issues, a main difference between these two studies is that the latter study included elderly subjects and, as the authors suggest, the involvement of CA1 might in part reflect an interaction with diseases of late life, such as vascular disease.

As with schizophrenia, it would be useful to image subjects who are at risk of developing depression to map the earliest prodromal stages of disease. This has not yet been done in humans, but a recent PET study in rhesus monkeys established that the anterior hippocampus is selectively affected in monkeys with 'anxious temperament'⁶⁵. Notably, as a trait, anxious temperament in children is an established risk factor for developing depression and anxiety disorders in adulthood⁶⁵ (and human studies have suggested a substantial phenotypic and anatomical overlap between anxiety and depressive disorders⁶⁶).

Numerous volumetric MRI studies have documented a smaller hippocampus in PTSD patients (as first described in REF. 67). So far, there is only one high-resolution MRI study that has mapped volumetric differences across hippocampal subregions in PTSD patients and controls⁶⁸. This study reported the greatest volume reduction in a single ROI that included both the dentate gyrus and CA3 (FIG. 2b). In the last section, we discuss how, guided by the functional organization of the hippocampal circuit, neuropsychological testing might be able to disambiguate CA3 from dentate gyrus dysfunction, and how cognitive profiles are expected to vary depending on whether dysfunction occurs in the posterior or anterior aspects of the hippocampal long axis.

Metabolic state and hippocampal dysfunction

Most disorders of the brain are associated with regional changes in basal metabolism. Energy metabolism in neurons requires glucose uptake and oxygen consumption. The metabolic state in any brain structure can be determined *in vivo* by directly measuring glucose uptake or oxygen consumption using PET, or by indirectly measuring correlates of oxygen consumption: cerebral blood flow (CBF) or cerebral blood volume (CBV) with PET, single-photon emission tomography (SPECT) or fMRI.

It is important to note that although the stimulus-evoked BOLD response measured by fMRI is designed to map transient evoked changes in neural activity, studies have established that the BOLD response is not simply coupled to metabolism^{69,70}, and therefore by itself cannot characterize the metabolic state of the brain⁷¹. This limitation applies to normal control subjects, in whom haemodynamic coupling varies across the brain^{72,73}, but is even more problematic when imaging ageing individuals and patients suffering from disease. The BOLD response is in fact influenced by neurovascular factors⁷⁴⁻⁷⁶ that act as confounds when trying to interpret between-group differences in the BOLD response (BOX 2). These neurovascular factors are affected in ageing and disease^{77,78}, and unsurprisingly studies have shown that differences in the BOLD response in ageing and disease do not necessarily reflect changes in activity or metabolism^{77,78}. It is important to emphasize that these limitations can be addressed. For example, it is now possible to calibrate the BOLD response against these neurovascular confounds⁷⁴, thereby deriving a measure of brain metabolism. When changes in the uncalibrated BOLD response are observed in ageing and disease they probably suggest that something is wrong in a particular region, but they do not

necessarily inform on abnormalities in neuronal activity, and they certainly cannot determine metabolic abnormalities (BOX 2).

Considering the volume of evidence for hippocampal atrophy and loss of hippocampus-dependent memory function in ageing and disease, it might be expected that functional imaging studies that assess metabolism should document hippocampal hypometabolism in all conditions. This is certainly the case in Alzheimer's disease, in vascular disease and in ageing, in which evidence for hypometabolism has been localized to the entorhinal cortex^{35,79}, CA1 (REF. 2) and the dentate gyrus, respectively^{41,43} (FIG. 2a). Using global measures of hippocampal metabolism, inconsistent results have been found in PTSD^{80–82}, which perhaps will be clarified in the future with the use of high-resolution technologies.

Notably, however, studies that have imaged hippocampal metabolism in patients with schizophrenia and depression have yielded surprising results. In the case of schizophrenia, a range of functional imaging techniques have clearly established that the hippocampal formation is characterized by an abnormal hypermetabolic state^{83–87}. Most of these studies used techniques that did not possess high spatial resolution, and simply assessed metabolism in the whole hippocampus. In the one published high-resolution fMRI study, anterior CA1 was found to be especially hypermetabolic in people with schizophrenia, together with hypermetabolism in the anterior subiculum and the orbitofrontal cortex⁶². In the early prodromal stages of the disease, evidence for hypermetabolism was found exclusively in anterior CA1 (REF. 62) (FIG. 2b).

In depression, some cross-sectional studies comparing patients with healthy controls have found evidence of hippocampal hypermetabolism^{88–90}. A more consistent observation emerges, however, from a growing number of studies in which patients were imaged before and after receiving pharmacological or behavioural treatments. These studies have found a decrease in metabolism in the global hippocampal formation in association with treatment efficacy (reviewed in REF. 91). Thus, the hippocampal formation in patients with depression seems to be in a relative state of hypermetabolism. In agreement with this conclusion, a PET study showed that an 'anxious temperament' trait in rhesus monkeys⁶⁵ is selectively associated with hypermetabolism in the anterior hippocampus. So far, high-resolution functional imaging techniques have not yet been used in patients with depression, and hence, whether hippocampal hypermetabolism is localized to the subiculum — as might be suggested — remains unknown.

Interestingly, TLE is a disorder in which the hippocampal formation alternates between basal hypometabolism⁹² and transient hypermetabolism (reflective of underlying seizures)⁹³. Loss of memory function in patients with TLE is seen during hippocampal hypometabolic states and exists, of course, in a more profound manner during active seizures. Informatively, during active seizures psychotic and affective symptoms, such as hallucinations and delusions and alterations in mood that often phenocopy schizophrenia and depression, can also occur^{94,95}. As seizure activity begins in, and emanates from, the hippocampal formation, abnormal stimulation of hippocampal outflow areas — those that mediate either psychotic or affective behaviour — has been invoked to explain these gain-of-function symptoms in TLE⁹⁴.

A pathophysiological framework

Based on the wide range of neuroimaging studies reviewed in the previous sections, we propose that any disorder that affects the hippocampal formation does so by differentially targeting specific subregions of the hippocampal circuit, and that this either leads to a decrease or an increase in neuronal metabolism. These two factors — regional vulnerability and metabolic state — provide a framework for how any disorder that affects the

hippocampal formation should be characterized (FIG. 3). In this section we will first discuss how, guided by hippocampal functional anatomy, this framework generates hypotheses on how diseases that affect the hippocampal formation might nevertheless manifest with phenotypic variability. Moreover, guided by the molecular anatomy of the hippocampal formation, we will illustrate how this framework provides the basis for understanding mechanisms of disease.

Using regional vulnerability to understand phenotypic diversity

Hippocampus-dependent tests in humans and other mammals were originally developed to assess the hippocampal formation globally, to contrast its function with that of other brain areas or to implicate the hippocampal formation in patients with cognitive impairments⁹⁶. Recently, however, the cognitive importance of individual hippocampal subregions has come to the fore. When empirical studies are interpreted in the context of computational considerations it is possible to begin charting a ‘functional map’ of the hippocampal formation, whereby specific cognitive or computational operations can be differentially linked to individual subregions (FIG. 3). Again, we emphasize the term differential, because obviously all subregions in the hippocampal circuit participate in multiple operations. Although construction of a functional map has only just begun, once charted it will be possible to develop more nuanced neuropsychological tests to assess the integrity of one subregion over another, thereby improving the ability to phenotype and distinguish disorders. At the same time, applying these tests to disorders for which imaging studies have established patterns of hippocampal vulnerability will help to validate emerging theories about the function of individual subregions. For example, showing that ageing manifests with pattern separation deficits^{50,51} confirms imaging findings (as reviewed above) — which are after all only indirect measures of function — and validates basic studies that have suggested that the dentate gyrus is differentially involved in this cognitive operation.

As mentioned, because the entorhinal cortex is upstream to the dentate gyrus, cell death in this subregion is also expected to cause pattern separation deficits and defects in cognitive operations that are linked to other downstream subregions. In fact, recent studies have suggested that pattern separation deficits might occur in the early stages of Alzheimer's disease (characterized by entorhinal cortex abnormalities)⁵². From a clinical perspective, this introduces the important point that hippocampal subregions really serve two functions: they perform a distinct operation as neural information flows through the circuit, but they also act as conduits that deliver information onto downstream subregions. In this regard, as the entorhinal cortex is the most upstream subregion in the hippocampal circuit, it is important to identify cognitive operations that differentially engage this subregion.

Rodent studies suggest that some cells in the entorhinal cortex, particularly those in its medial aspect, show unique grid-cell properties⁹⁷. A recent human fMRI study described a cognitive task based on predictions extracted from grid-cell properties, and showed that this task activated the entorhinal cortex⁹⁸. The task, however, was found to also activate numerous other brain regions outside the hippocampal formation. A convergence of other findings suggests that the entorhinal cortex is selectively linked to a cognitive operation performed during brief delay periods of hippocampus-dependent tasks. This link was first suggested by unit recordings in monkeys and rats^{99,100}. Here, sustained activity was observed in the entorhinal cortex during the delay period of matching- or non-matching-to-sample paradigms. In slice preparations, the entorhinal cortex has been found to have electrophysiologically unique ‘persistent-firing’ neurons that show intrinsically-driven, sustained activity lasting minutes¹⁰¹. Although they have not yet been confirmed *in vivo*, persistent-firing neurons have been incorporated into computational models of hippocampal function in which the entorhinal cortex is proposed to act as a ‘memory buffer’ during brief

delays¹⁰², and importantly, these neurons have been linked to the emergence of grid cells in the entorhinal cortex¹⁰³.

These electrophysiological findings have been used to interpret observations in a growing number of human fMRI studies. One study observed sustained activation in a large area that included the entorhinal cortex during a delay in a matching-to-sample task, and sustained activity in this area predicted performance on delayed recognition for visual stimuli^{104,105}. A separate study, using higher spatial resolution, showed sustained activity that was unambiguously localized to the entorhinal cortex during a delay period in a matching-to-sample task of faces¹⁰⁶. Although these fMRI studies were interpreted in the context of previous electrophysiological findings, it is notable that stimulus-induced activity was also observed in multiple downstream hippocampal subregions. Two fMRI studies have documented a selective link with the entorhinal cortex. One study found sustained activity in the entorhinal cortex during a memory encoding phase, and the level of sustained activity predicted performance during delayed cue-recall¹⁰⁷. Another study used fMRI to map basal metabolism in multiple hippocampal subregions in subjects who received an extensive neuropsychological battery of tests outside the scanner. Basal metabolism in the entorhinal cortex was found to differentially correlate with performance on a derived measure of delayed retention¹⁰⁸.

When these electrophysiological, computational and fMRI studies are viewed as a whole, it is possible to conclude that the entorhinal cortex has an important role in ‘delayed retention’ during brief delays (FIG. 3). Although many tasks, such as delayed matching-to-sample, activate the entorhinal cortex, performance on these tasks does not isolate this cognitive operation. In clinical neuropsychological tests, delayed retention is typically isolated by normalizing the amount of information that is recalled after a delay by the amount of information that is recalled immediately. Defects in this derived measure of delayed retention is one of the hallmark cognitive features of Alzheimer's disease¹⁰⁹, confirming its link to the entorhinal cortex. Moreover, defects in delayed retention have also been observed in the preclinical stage of the disease, decades before the onset of dementia¹¹⁰. Most importantly, delayed retention is one of the few hippocampus-dependent cognitive abilities that is often found to be relatively resistant to ageing compared to Alzheimer's disease^{109,111–113}. These observations confirm neuroimaging findings that suggest that the entorhinal cortex is relatively resistant to the ageing process, and shows how our pathophysiological framework can predict phenotypic differences between Alzheimer's disease and ageing.

In addition to distinguishing the entorhinal cortex from other hippocampal subregions, it is also important to be able to use cognitive tests to assess the function of other hippocampal subregions. As mentioned, in the case of PTSD neuroimaging studies have so far been unable to anatomically dissociate the dentate gyrus from CA3. Therefore, in addition to pattern separation tasks, which engage the dentate gyrus, it would be useful to have neuropsychological tests that differentially assess the functional integrity of CA3. In fact, the CA3 subregion has been linked to a cognitive operation termed pattern completion — the ability to reactivate a complete hippocampus-dependent memory when presented with only a fragment of the memory (FIG. 3). As first described by Lorente de Nó¹⁰, CA3 is unique in that it contains an ‘association tract’ that connects CA3 neurons up and down the hippocampal long axis (FIG. 1) and, as computationally suggested by Marr⁴⁴, this auto-association system is an anatomical feature that is ideally suited for pattern completion. A mouse fMRI study¹¹⁴ and *in vivo* recording studies^{115,116} have provided empirical confirmation of the importance of CA3 for pattern completion. Interestingly, studies suggest that performance on cognitive tasks that are sensitive to pattern completion are relatively

preserved in ageing⁵⁰, supporting the idea that compared with the dentate gyrus, CA3 is relatively preserved.

Based on the functional organization of the hippocampal long axis, reviewed above, we propose that the anterior dentate gyrus and CA3 are involved in pattern separation and pattern completion, respectively, of emotionally charged stimuli, whereas the posterior subregions are predicted to be engaged by stimuli that are emotionally neutral. Although future studies are required to test these specific predictions, BOLD fMRI studies support this long-axis organization¹¹⁷, showing, for example, that emotionally charged faces activate the anterior hippocampus²⁶, whereas activation of the posterior hippocampus is more typically found when emotionally neutral faces are used¹⁷. Indeed, the mnemonic defects that are observed in PTSD generally involve memories of emotionally charged stimuli¹¹⁸.

Reflecting the dual input that the CA1 subregion receives from layer III entorhinal cortex and from CA3 (FIG. 1b), CA1 neurons are well positioned to play a part in integrating or comparing information from these converging sources of input. Computational and electrophysiological studies have confirmed that CA1 has a role as an integrator or comparator of converging entorhinal cortex and CA3 input^{119–123}. This role has been invoked to explain a human fMRI study that showed that posterior CA1 is differentially activated when subjects encode allocentric spatial information¹²⁴. A recent study has observed selective lesions to CA1 in patients with transient global amnesia, and showed that this CA1-selective lesion was associated with defects in learning in a virtual water maze¹²⁵.

Finally, we come to the last subregion in the hippocampal circuit — the subiculum. A growing number of human fMRI studies have observed that the subiculum shows a differential BOLD response during the retrieval (compared to the encoding) of new hippocampus-dependent memories^{106,126–129}, probably reflecting the fact that the subiculum receives input from CA1 and the entorhinal cortex¹³⁰.

As highlighted, a definitive functional map of the hippocampal formation has not yet been completed. Nevertheless, the studies that are reviewed above establish the principle that such a map exists (FIG. 3), and illustrate the general approach by which this map may be charted with greater certainty. Informed by computational and experimental observations, human neuroimaging — in particular, high-resolution BOLD¹³¹ — will be needed to link specific cognitive operations to distinct subregions, and are likely to show how these operations vary over the hippocampal long axis. For clinical purposes, such a functional map will hopefully guide the development of a new wave of standardized neuropsychological tests (FIG. 3) designed to assess the function of multiple subregions in a single battery. As emphasized, because the hippocampal subregions are interconnected to form a circuit, a ‘lesion’ in one subregion might cause secondary dysfunction in other subregions, and thus impaired performance might be observed in multiple tests. However, by administering an extensive battery, it should be possible to identify the test in which the greatest impairment occurs, thereby pinpointing the anatomical source of hippocampal dysfunction.

Using metabolic state to understand phenotypic diversity

Patients who have extensive bilateral resections of the medial temporal lobes (such as the famous patient H.M.), including of the hippocampal formation, develop profound cognitive defects but are notably free of changes in personality or affect¹³². As a hippocampectomy can be considered an extreme version of hippocampal loss of function, this confined cognitive phenotype supports the idea that disorders that manifest with hypometabolism are characterized primarily by cognitive symptoms.

Of course, as shown in TLE, hypermetabolism within the hippocampal formation will also interfere with its mnemonic function, although by its very nature hypermetabolism might not be linked to specific cognitive operations. More interestingly, when hippocampal hypermetabolism is observed in TLE, it can (in addition to a loss of memory function) be associated with a 'gain' in psychotic and affective symptoms, thus phenocopying the symptoms of schizophrenia or depression⁹⁵.

This raises the intriguing possibility that the hippocampal hypermetabolism that is observed in schizophrenia and depression may, by stimulating specific hippocampal outflow areas, actually drive the psychotic and affective symptoms that are characteristic of these diseases (FIG. 4). If this is true, then selectively deactivating the hippocampal formation pharmacologically or by other means¹³³ should ameliorate symptoms. As far as we know, this has never been formally tested as most experimental manipulations do not exclusively target the hippocampal formation. Vagal nerve stimulation reduces hippocampal metabolism and has proven to be an effective treatment for depression, but it reduces metabolism in other brain areas as well¹³⁴. It is also possible that hippocampal hypermetabolism in schizophrenia and depression might, in the long term, entrain hypermetabolism in hippocampal outflow areas by altering their metabolic set-point. In this scenario, hypermetabolism in these areas becomes hippocampus-independent over time, and hippocampus-selective interventions would therefore be effective only if administered during the earliest stages of the disease.

Currently, only correlative observations and animal studies provide evidence for the idea that the hippocampal formation acts as an anatomical driver of gain-of-function symptoms in schizophrenia and depression (FIG. 4). In the case of schizophrenia (FIG. 4a), one study found that hippocampal hypermetabolism occurs first, and that orbitofrontal hypermetabolism is observed only later in the disease course⁶². Furthermore, CA1 hypermetabolism was found to be associated with measures of delusions and hallucinations. As the hippocampus monosynaptically connects with the orbitofrontal cortex¹⁴, it is possible that hippocampal hypermetabolism acts to drive this later extra-hippocampal effect. The hippocampus also monosynaptically connects with the nucleus accumbens, and animal studies have established a mechanism by which hippocampal hypermetabolism links to increased dopamine release in the ventral striatum. Specifically, increased activity in the ventral hippocampus (equivalent to the anterior hippocampus in humans) stimulates the nucleus accumbens through monosynaptic connections, leading to increased striatal dopamine release¹³⁵ (FIG. 4a). Not only is increased striatal dopamine release observed in schizophrenia^{136,137} but elevations in dopamine are a neurochemical cause of psychotic symptoms¹³⁷.

In depression, hypermetabolism occurs, in addition to the hippocampal formation, in multiple regions of the frontal lobe — in particular, the subgenual anterior cingulate and in subregions of the amygdala, caudate, occipital lobe, cerebellum and thalamus⁹¹. A brain network linked to depression has been suggested based on the connectivity among some of these areas¹³⁸. An important feature of this network is the connection between the hippocampus and the subgenual anterior cingulate (FIG. 4b), a connection that was first suggested by tracer studies in non-human primates¹⁴ and confirmed more recently by *in vivo* imaging in humans¹³⁹. This connection is particularly interesting because selectively reducing hypermetabolism in the subgenual anterior cingulate with deep brain stimulation ameliorates affective symptoms in some patients with depression and anxiety disorders⁶⁶.

Future studies are needed to establish whether there is a causal link between the hippocampal hypermetabolism that is observed in schizophrenia and depression, and their respective symptoms. If this link is established, the pathophysiological framework generates

a number of predictions. Although there are some overlaps between the symptoms of schizophrenia and depression, we predict that differences in symptoms might be explained by differential regional hippocampal hypermetabolism. As reviewed, there is already some evidence that these disorders target different hippocampal subregions, with schizophrenia differentially targeting CA1 and depression targeting the subiculum. We predict that the hippocampal long axis will also show this regional vulnerability. As well as explaining phenotypic differences between schizophrenia and major depressive illness, the framework might also explain phenotypic variability within the two diagnostic categories. Specifically, it might explain why some patients with depression have accompanying psychotic symptoms, and why there is variation in the degree of affective symptomatology of patients diagnosed with schizophrenia and related psychotic disorders.

Using the framework to understand pathogenic mechanisms

Clearly, for nearly all of the disorders reviewed here, the hippocampal formation is not the only brain area that is affected. However, by mapping patterns of vulnerability within the hippocampal formation and determining whether vulnerable subregions are hypo- or hypermetabolic, a basic assumption of the framework is that this information can be exploited to clarify pathogenesis. Showing that two diseases target different subregions would provide compelling evidence that they are mediated by separate processes. For example, there has been an ongoing debate about whether Alzheimer's disease and ageing overlap mechanistically. The observed anatomical dissociation — with Alzheimer's disease differentially targeting the entorhinal cortex and ageing differentially targeting the dentate gyrus with relative sparing of the entorhinal cortex — suggests that Alzheimer's disease and ageing have distinct aetiologies.

More importantly, if a pattern of vulnerability is established, it can be used experimentally to isolate cellular and molecular mechanisms of disease. For example, using gene-expression profiling, a study relied on the differential pattern of hippocampal dysfunction in Alzheimer's disease to isolate deficiencies in molecules related to membrane sorting and trafficking in those subregions that are more affected in Alzheimer's disease¹⁴⁰. Studies have confirmed that these intracellular pathways contribute to the pathogenesis of the disease¹⁴¹, leading to a general cell biological hypothesis of late-onset Alzheimer's disease¹⁴².

These studies in Alzheimer's disease show the framework's utility. As long as post-mortem samples can be accessed for a given disease, molecular correlates of regional vulnerability and resistance can be identified. Even if post-mortem samples are unavailable, pinpointing a hippocampal subregion that is differentially vulnerable to a given disorder can provide clues regarding pathogenesis. For example, the molecular map of the hippocampal formation shows differential expression of glucocorticoid receptors in the CA3 (REFS 143,144), which may link this subregion to PTSD. Indeed, previous rodent studies have found that both acute and chronic stress differentially target the CA3 subregion^{143–149}. Interestingly, recent studies in rodents suggest that the effects of stress-induced glucocorticoid release is greatest in the anterior (or ventral) hippocampus¹⁵⁰. Similarly, when the dentate gyrus is implicated in a disorder, a potential link of the disease to adult neurogenesis — a feature that is unique to this subregion^{151,152} — must be considered. One straightforward possibility is that a reduction in neurogenesis underlies hippocampal dysfunction. Indeed, age-related reductions in neurogenesis have been observed in rodents¹⁵³, and recent studies have shown that blocking neurogenesis in mice causes defects in pattern separation⁴⁸, whereas increasing neurogenesis enhances pattern separation¹⁵⁴. Alternatively, neurogenesis during development — which persists longest in the dentate gyrus compared to other subregions — can 'imprint' the expression of molecules that mediate transcriptional regulation. This form of gene-expression imprinting has been documented¹⁵⁵, and some studies suggest that the

dentate gyrus is differentially engaged in histone modification and other epigenetic mechanisms important for the regulation of transcription.

Interestingly, a range of high-resolution fMRI studies in humans, non-human primates and rodents have shown that high levels of blood glucose causes metabolic defects selectively in the dentate gyrus². An increase in post-prandial blood glucose (owing to age-related insulin resistance) is observed in ageing humans and other mammals^{156,157}. Thus, an age-related increase in blood glucose might account for why dentate gyrus dysfunction is observed with ageing across species (as reviewed above). These observations suggest that manipulations that improve glucose homeostasis might reduce or prevent age-related memory decline. Indeed, physical exercise is one manipulation that improves glucose homeostasis¹⁵⁸, differentially increases dentate gyrus basal metabolism¹⁵⁹ and brain-derived neurotrophic factor (BDNF) expression¹⁶⁰, and ameliorates age-related atrophy measured in the whole hippocampus¹⁶¹.

The occurrence of hippocampal hypermetabolism offers a separate set of clues about underlying mechanisms. For example, studies have suggested that in schizophrenia there might be an abnormal elevation in synaptic glutamate levels^{162,163}, and therefore tonic stimulation of AMPA receptors might act as a source of hyper-metabolism. The loss of CA3 GABAergic interneurons in patients with schizophrenia¹⁶⁴ could cause hyperactivity in conjunction with increased synaptic glutamate release and, based on hippocampal circuit properties, this effect has been proposed to occur most prominently in the CA1 subregion¹⁶⁴. Of course, many molecules regulate synaptic glutamate and some, like glutaminase, have been genetically^{165,166} and functionally¹⁶⁷ implicated in schizophrenia. However, future studies are required to establish whether molecules that regulate glutamate metabolism are differentially altered in CA1 of patients with schizophrenia.

In the case of depression, mechanistic insight is provided by the fact that, when effective, anti-depressants suppress hippocampal hypermetabolism⁹¹. Serotonin reuptake inhibitors (SSRIs) can, by increasing synaptic serotonin levels and thereby stimulating specific serotonin receptors, suppress hippocampal synaptic activity, including in the subiculum¹⁶⁸, and has recently been shown to suppress an fMRI correlate of hippocampal metabolism¹⁶⁹. Moreover, *in vivo* radioligand and postmortem studies have suggested that in depression there might be a deficiency in hippocampal serotonin receptors¹⁷⁰, which could contribute to hypermetabolism. An alternative mechanism by which SSRIs might suppress hippocampal hypermetabolism is suggested by an intriguing finding showing that a reduction in dentate gyrus neurogenesis causes an increase in spontaneous synchronized activity¹⁷¹. Newly born neurons in the dentate gyrus seem, therefore, to have an inhibitory effect on the hippocampal circuit. Although recent studies suggest that increasing neurogenesis per se does not have antidepressant effects¹⁵⁴, studies nevertheless show that the effect of SSRIs are mediated in part by enhancing neurogenesis¹⁷². Finally, very low doses of the NMDA receptor antagonist ketamine can be rapidly effective in depression, and a recent study showed that, among its many effects, low-dose ketamine reduces spontaneous hippocampal activity¹⁷³.

Conclusions

The proposed pathophysiological framework of hippocampal dysfunction explains, on the one hand, how alterations in single brain structure can lead to phenotypic diversity, and on the other hand, how hippocampal dysfunction can be caused by a range of mechanistically distinct disorders. As its core feature, the framework is guided by the awareness that the hippocampal formation is not a singular structure; the hippocampal formation is a complex

circuit that is functionally and molecularly segregated over its longitudinal and transverse axes.

The functional anatomy of the hippocampal circuit can explain how diseases manifest non-overlapping symptoms by differentially targeting specific sites within the circuit. Moreover, the framework informs recent theories about the computational operations performed by individual hippocampal subregions. It supports the view that the function historically assigned to the hippocampal circuit should be broadened, from an exclusive focus on memory to include affect, personality and other complex behaviours. As reviewed, the molecular anatomy of the hippocampal circuit and the unique cellular features of individual subregions provide a general principle for why the hippocampal formation is vulnerable to a broad range of mechanistically distinct disorders.

As we have emphasized, many specific questions about phenotypic features that distinguish each disorder, and the manner in which hippocampal dysfunction is causally linked to these phenotypes, remain undetermined. Moreover, although in some cases clues have emerged, the full complement of pathogenic mechanisms underlying hippocampal dysfunction have not yet been isolated. Just as advances in functional and structural MRI have generated findings that form the basis of the framework, newer technological innovation promises to address these outstanding issues. With higher field strength scanners, it will be possible, for example, to visualize divisions within the entorhinal cortex¹⁷⁴ and other hippocampal subregions, and magnetic resonance spectroscopy (MRS) could be used to pinpoint putative glutamate abnormalities¹⁷⁵ in CA1 or neurogenesis defects in the dentate gyrus¹⁷⁶. Higher resolution and better signal-to-noise ratios will accelerate the development of ‘molecular imaging’, with the promise of visualizing receptors and other molecular abnormalities that are thought to be pathogenic in specific diseases. Additionally, a new crop of technologies that are designed to map patterns of connectivity — including DTI, resting-state functional connectivity MRI¹⁷⁷ and manganese-enhanced MRI¹⁷⁸ — are already available. These technologies are well suited to further clarify patterns of connectivity within the hippocampal circuit, to map how the hippocampal long axis differentially interconnects with other brain areas, and to understand how disease and ageing affect the hippocampal circuit and the larger brain networks to which the hippocampal circuit belongs. Lastly, all of these imaging technologies, both old and new, can be used to longitudinally follow large number of subjects, tracking how ageing and disease spreads over the hippocampal circuit.

By providing a basis for better understanding hippocampal dysfunction in ageing and disease, the framework proposed in this Review will help to guide future studies, with the ultimate goal of identifying pathogenic mechanisms and of developing effective interventions for some of the most common disorders of the brain.

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Box 1 | fMRI and the hippocampal circuit

A long history of positron emission tomography (PET) and other functional imaging studies have established that cerebral blood flow (CBF) and cerebral blood volume (CBV) are correlates of brain function and metabolism. Informed by these observations, the first human functional MRI (fMRI) study used a technique that measured CBV¹⁷⁹, and shortly thereafter, studies established that fMRI can also measure CBF¹⁸⁰ and blood oxygen level-dependent (BOLD)¹⁸¹ responses to map brain function (see BOX 2 for a detailed discussion of the BOLD response).

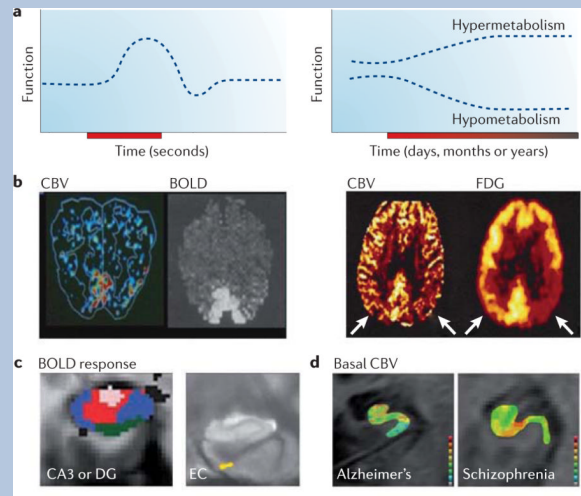
When imaging brain function, it is important to consider that, neurophysiologically, function can change acutely and transiently, such as when changes in neural activity are evoked by an external and transient stimulus (see the figure; part **a**, left panel; part **b**, left panel; and part **c**). Changes can also occur slowly and chronically, affecting a region's basal metabolic state (see the figure; part **a**, right panel; Part **b**, right panel; and part **d**). Examples of chronic stimuli that change basal metabolism include the dendritic remodelling that underlies long-term memory, neuronal dysfunction that occurs during ageing and in many diseases, and the effects of therapeutic interventions. fMRI measures of CBV, CBF or BOLD responses can map acute changes in brain function (see the figure, part **b**, left panel, in which visual stimulation is shown to activate visual cortex using either CBV or BOLD measurements). As fMRI measures of CBV and CBF are quantitative, they are better suited to mapping chronic changes in basal metabolism that are associated with ageing and disease (see the figure, part **b**, right panel, in which CBV defects (as detected with fMRI) match fluorodeoxyglucose (FDG) uptake defects detected with PET) (see BOX 2 for a discussion of the limitations of BOLD in mapping metabolism in ageing and disease).

Different approaches have been used to assess group data in fMRI studies to generate functional maps of the hippocampal formation using CBV, CBF or BOLD. The challenge in evaluating the hippocampal circuit is inherent in the fact that the subregions are very small structures, a few millimetres in dimension. Accordingly, most fMRI studies that evaluate the hippocampal circuit try to generate maps with higher than usual spatial resolution. Because of image acquisition protocols, CBF and BOLD maps are typically generated with an in-plane resolution that nears 1 millimetre. CBV is currently the only fMRI measure that can generate functional maps with sub-millimetre resolution¹⁸².

Another challenge is the large degree of subject-to-subject variability in the anatomy of hippocampal subregions, particularly in diseased or ageing populations that often have hippocampal atrophy. Different analytic approaches are now commonly used to overcome this challenge¹³¹. One is to evaluate functional maps on a subject-by-subject basis. By drawing regions of interest (ROIs) within hippocampal subregions, group data analysis is performed on functional information that is acquired from each subject (see the figure, part **d**, which shows individual CBV maps of a patient with Alzheimer's disease³⁵ and of a patient with schizophrenia⁶². Warmer colours indicate greater CBV).

With the development of sophisticated co-registration techniques, it is now possible to co-register the hippocampal formation across subjects and to perform voxel-based statistical analyses (reviewed in REF. 131). These approaches have been used to show a BOLD response in the dentate gyrus (DG) or CA3 area using a pattern separation task⁴⁹ (see the figure, part **c**; pink colour in the left image) and a BOLD response in the entorhinal cortex (EC) during a brief delay in a match-to-sample task¹⁰⁶ (see the figure, part **c**; yellow colour in the right image). By using cortical unfolding techniques, it is also possible to perform voxel-based analyses on 'flat maps' of the hippocampal formation¹²⁶. Figure part **c**, left image is reproduced, with permission, from REF. 49 ©

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Box 2 | The complexity of interpreting the BOLD response in ageing and disease

The blood oxygen level-dependent (BOLD) response is a sensitive indicator of where neural activity has acutely changed in response to a transient stimulus. The interpretation of altered responses in disease populations, in ageing, or even in healthy controls on medication⁷⁵, is intrinsically ambiguous. BOLD is sensitive to changes in deoxyhaemoglobin levels¹⁸³, which are reduced during acute activation because cerebral blood flow (CBF) increases much more than the cerebral metabolic rate of oxygen (CMRO₂) consumption. The BOLD response, however, turns out to be strongly modulated by two additional confounding factors⁷⁴ that act independently of neural activity: baseline levels of deoxyhaemoglobin and the precise ratio of the fractional changes in CBF and CMRO₂.

Although mathematical modelling of the BOLD response is complex¹⁸⁴, the dependence of the BOLD response ($\Delta S/S_0$) on CBF changes and on these two confounding factors¹⁸⁵ can be approximated as:

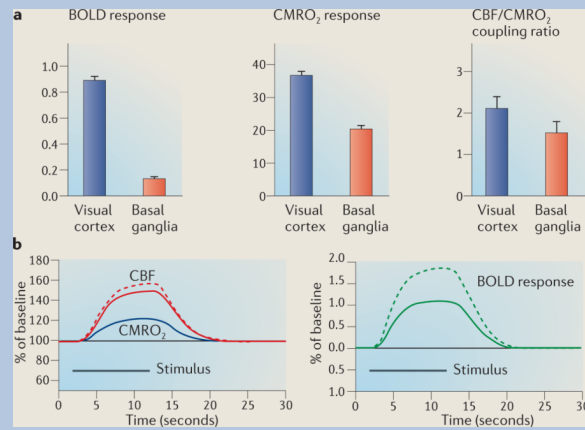
$$\frac{\Delta S}{S_0} = \frac{\Delta CBF}{CBF(base)} \cdot A \cdot \left(1 - \frac{b}{n}\right)$$

The first term, $\Delta CBF/CBF(base)$, represents the CBF change. The second term, A , is proportional to baseline deoxyhaemoglobin, which is affected by cerebral blood volume (CBV) and oxygen extraction fraction. The third term, n , is the CBF/CMRO₂ coupling ratio, defined as the per cent change in CBF divided by the percent change in CMRO₂ (the numerical constant b has a value of about 1.2).

If both baseline deoxyhaemoglobin (A) and the CBF/CMRO₂ coupling ratio (n) were invariant, interpreting differences in the BOLD response would be straightforward. Both of these factors, however, are highly variable across different brain regions and in ageing and disease. Therefore, when differences in the BOLD response are observed they do not necessarily reflect differences in neural or metabolic activity. So, for example, subjects at risk of Alzheimer's disease were found to have a reduced BOLD response in the hippocampal formation evoked by a memory task compared to controls⁷⁸. However, this effect was found to be related to differences in baseline CBF, which probably affects baseline deoxyhaemoglobin and confounds the interpretation of the BOLD response. Similarly, another study showed that age-related changes in baseline deoxyhaemoglobin levels can cause misleading differences in the BOLD response when comparing young and older subjects⁷².

High variability in the CBF/CMRO₂ coupling ratio further contributes to the difficulty in interpreting the BOLD response. For example, a visual-evoked BOLD response in visual cortex was sevenfold larger than a motor-evoked BOLD response in the basal ganglia⁷² (see the figure, part **a**, left panel). However, only a twofold difference was observed in the metabolic response between the two regions (see the figure, part **a**, middle panel). The dramatic mismatch between the BOLD and metabolic responses was due to a modest difference in the CBF/CMRO₂ coupling ratio (see the figure, part **a**, right panel). Using the standard BOLD model it is possible to show how, if a disease subtly increases the CBF/CMRO₂ coupling ratio (see the figure, part **b**; shown by solid and dashed red curves), the same evoked metabolic response (see the figure, part **b**; shown by the blue curve) can lead to dramatic differences in the BOLD response (see the figure, part **b**; shown by the green curves). In principle, the opposite can also occur if the disease causes a decrease in coupling.

The confound factors A and n can interact with each other, further adding to the complexity of interpreting the BOLD response. Indeed, this interaction can lead to scenarios in which no differences in the BOLD response are observed, despite underlying metabolic differences. This possibility was recently demonstrated in a study that examined the effects of caffeine intake on the BOLD response¹⁸⁶, and which showed that caffeine had countervailing effects on A and n . Notably, in the basal state (that is, without acutely altering metabolism with a transient stimulus) the relationship between CBF, CBV and $CMRO_2$ is typically invariant and tightly coupled^{69,187}, and so fMRI measures of basal CBF or CBV often provide an accurate map of metabolic state. Data in the figure, part **a**, from REF. 72.



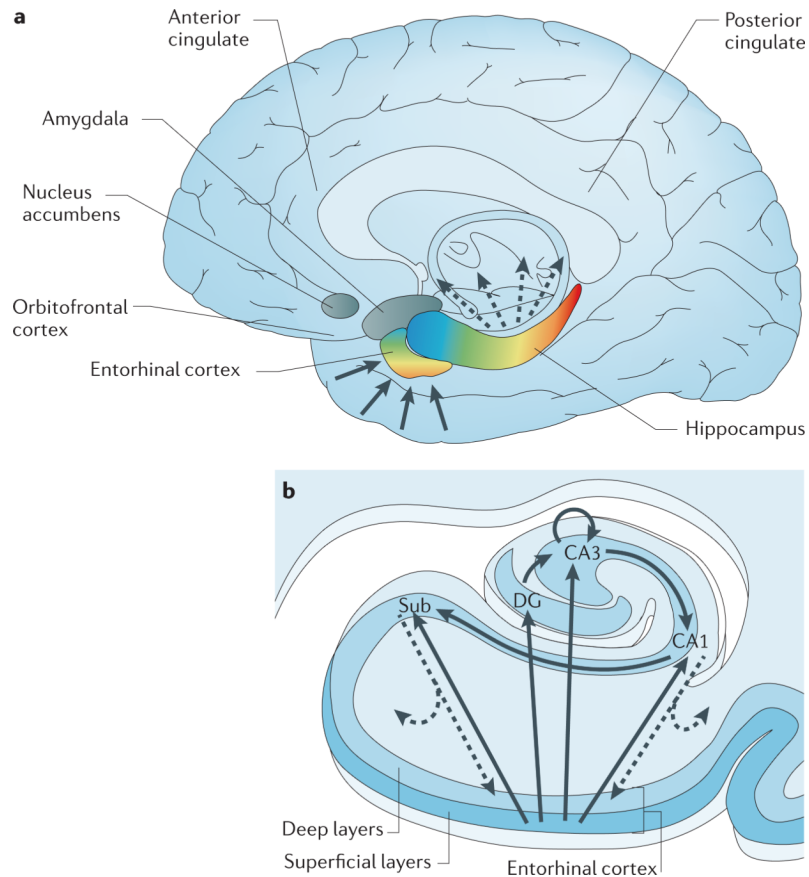


Figure 1. The organization of the hippocampal formation

a | The hippocampal formation, which is made up of the entorhinal cortex and hippocampus (shown in colour) extends over the anterior-to-posterior axis of the brain. The colour gradients reflect the topological input–output relations between the hippocampal formation and other brain areas, as well as its internal functional and molecular organization. Input to the hippocampus is shown by solid arrows, hippocampal outflow is shown by dashed arrows. Cortical and subcortical information funnels onto superficial layers of the entorhinal cortex, and this input is organized in an anterior–medial to posterior–lateral gradient (shown by the colour gradient in the entorhinal cortex). This anatomical gradient is largely preserved as the entorhinal cortex conveys this information to the hippocampus (shown by the corresponding colour code in the hippocampus). The long-axis gradient is preserved in the output pattern of the hippocampus. As well as reconnecting with the entorhinal cortex, the hippocampus monosynaptically connects with — from anterior to posterior — the orbitofrontal cortex, anterior cingulate, amygdala, nucleus accumbens and posterior cingulate. **b** | In the hippocampal transverse axis, superficial layers of the entorhinal cortex connect with the dentate gyrus (DG), CA3, CA1 and the subiculum (Sub). The trisynaptic circuit connects the dentate gyrus to CA3, to CA1 and to the subiculum. Through auto-association fibres, CA3 neurons interconnect with other CA3 neurons throughout the long axis. The CA1 and primarily the subiculum provide the main hippocampal outflow (shown by dashed arrows), back to the deep layers of the entorhinal cortex and also to a range of cortical and subcortical sites (as shown in part **a**).

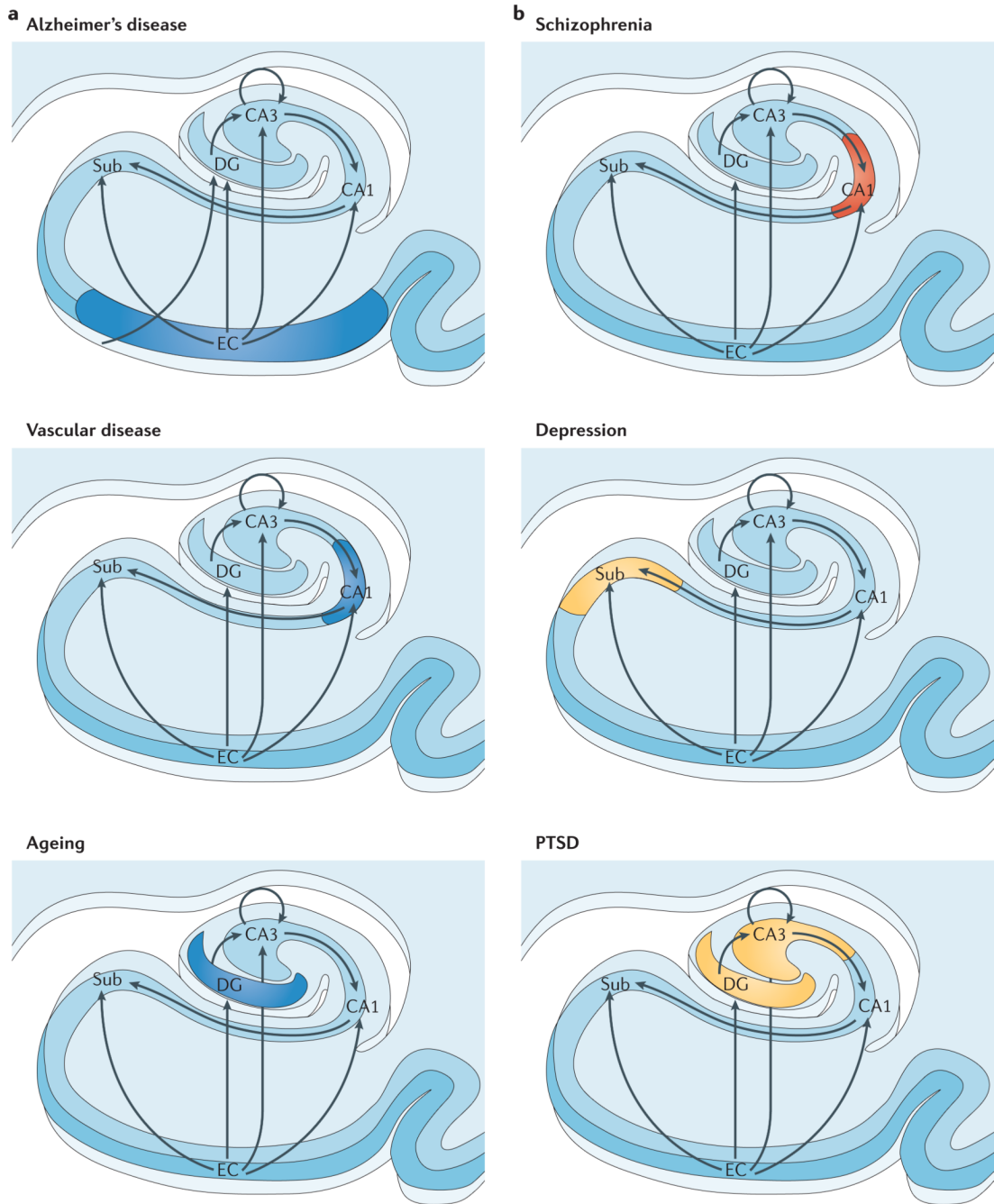


Figure 2. Regional vulnerability and metabolic state differentiate disorders that affect the hippocampal formation

Although multiple hippocampal subregions can be affected in disorders, by comparing patterns of alterations that are observed by functional and structural MRI it is possible to isolate individual subregions differentially affected by each disorder. Furthermore, functional imaging techniques that are sensitive to metabolic state have suggested that some hippocampus-based disorders are characterized by hypometabolism (shown in blue), whereas others are abnormally hypermetabolic (shown in red). **a** | Alzheimer's disease, vascular disease and ageing all contribute to hippocampal alterations in late life. A direct comparison suggests that the entorhinal cortex (EC) is differentially associated with

Alzheimer's disease and the CA1 with vascular disease, whereas the ageing process per se seems to differentially target the dentate gyrus (DG). Hypometabolism has been localized to the EC in Alzheimer's disease, to CA1 in vascular disease and the DG in ageing. **b** | Schizophrenia and depression typically begin during adolescence, and post-traumatic stress disorder (PTSD) currently has its highest incidence in young adulthood. A direct comparison suggests that CA1 is differentially associated with schizophrenia and that the subiculum (Sub) may be differentially associated with depression. In the case of PTSD, both CA3 and the DG have been implicated. Evidence for global hippocampal hypermetabolism exists for both schizophrenia and depression. In schizophrenia, this hypermetabolism may be driven by hypermetabolism in CA1, whereas subicular hypermetabolism in depression is a hypothesis that remains untested (shown in yellow). In PTSD the metabolic state of the hippocampal formation is as yet undetermined (shown in yellow).

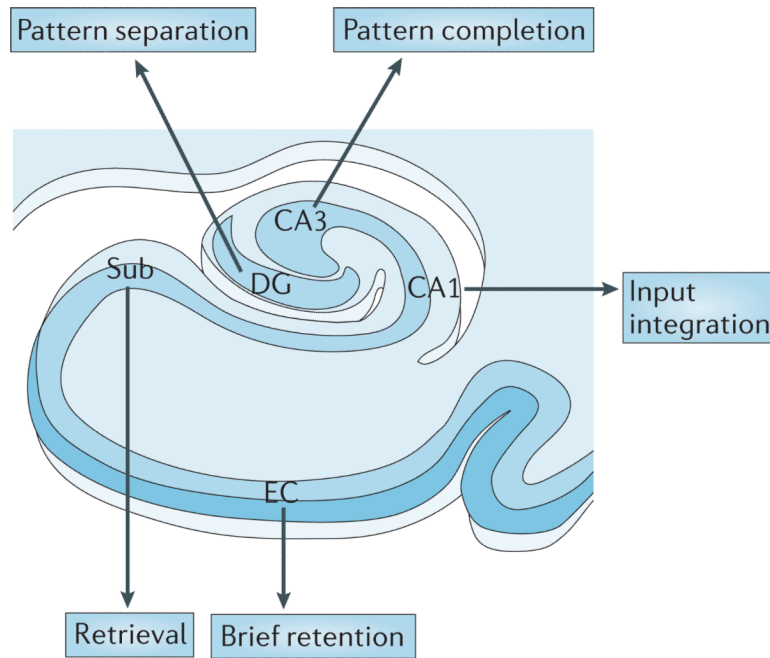


Figure 3. A proposed ‘functional map’ of the hippocampal circuit

Although multiple subregions are engaged by different tasks, as neural information flows through the hippocampal circuit, each hippocampal subregion is thought to perform a distinct cognitive or computational operation. It is possible to begin to chart a ‘functional map’ of the hippocampal circuit, linking distinct operations to individual hippocampal subregions. Here, CA1 is important in integration of inputs, the dentate gyrus (DG) has a role in pattern separation, CA3 has a role in pattern completion, the subiculum (Sub) is involved in memory retrieval and the entorhinal cortex (EC) is involved in brief retention in hippocampus-dependent memory tasks.

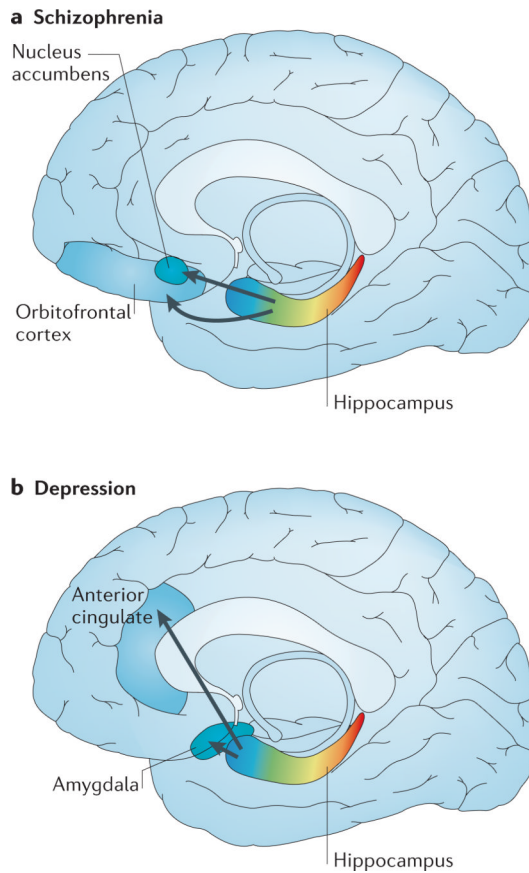


Figure 4. Proposed hippocampus-based networks in schizophrenia and depression

a | In schizophrenia, hypermetabolism in the anterior hippocampus occurs early in the disease process, and through monosynaptic connections this can be linked to orbitofrontal hypermetabolism that emerges later in the disease course. In animal models, hyperactivity in the anterior hippocampus stimulates the nucleus accumbens, leading to increased striatal dopamine release. **b** | Studies in patients with depression suggest that hypermetabolism in the anterior hippocampus occurs early in the disease process, and via monosynaptic connections this may be linked to hypermetabolism observed in the amygdala and the anterior cingulate.