
Research Articles: Behavioral/Cognitive

Mind-wandering in people with hippocampal damage

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43

44 **Abstract**

45 Subjective inner experiences, such as mind-wandering, represent the fundamentals of human
46 cognition. Although the precise function of mind-wandering is still debated, it is increasingly
47 acknowledged to have influence across cognition on processes such as future planning,
48 creative thinking and problem-solving, and even on depressive rumination and other mental
49 health disorders. Recently, there has been important progress in characterizing mind-
50 wandering and identifying the associated neural networks. Two prominent features of mind-
51 wandering are mental time travel and visuo-spatial imagery, which are often linked with the
52 hippocampus. People with selective bilateral hippocampal damage cannot vividly recall
53 events from their past, envision their future or imagine fictitious scenes. This raises the
54 question of whether the hippocampus plays a causal role in mind-wandering and if so, in
55 what way. Leveraging a unique opportunity to shadow people (all males) with bilateral
56 hippocampal damage for several days, we examined, for the first time, what they thought
57 about spontaneously, without direct task demands. We found that they engaged in as much
58 mind-wandering as control participants. However, whereas controls thought about the past,
59 present and future, imagining vivid visual scenes, hippocampal damage resulted in thoughts
60 primarily about the present comprising verbally-mediated semantic knowledge. These
61 findings expose the hippocampus as a key pillar in the neural architecture of mind-wandering
62 and also reveal its impact beyond memory, placing it at the heart of human mental life.

63

64 **Significance statement**

65 Humans tend to mind-wander about 30-50% of their waking time. Two prominent features
66 of this pervasive form of thought are mental time travel and visuo-spatial imagery, which are
67 often associated with the hippocampus. To examine whether the hippocampus plays a causal
68 role in mind-wandering, we examined the frequency and phenomenology of mind-wandering

69 in patients with selective bilateral hippocampal damage. We found that they engaged in as
70 much mind-wandering as controls. However, hippocampal damage changed the form and
71 content of mind-wandering from flexible, episodic, and scene-based to abstract, semanticized,
72 and verbal. These findings expose the hippocampus as a key pillar in the neural architecture
73 of mind-wandering and reveal its impact beyond memory, placing it at the heart of our mental
74 life.

75

76 **Introduction**

77 Even when in the same place and involved in the same activity, at any given moment people
78 can experience the world in different ways. Recently, there have been advances in delineating
79 the various forms of spontaneous inner experiences and their neural correlates (Andrews-
80 Hanna et al., 2014a; Christoff et al., 2016). Self-generated thinking typically refers to the
81 ability to mentally decouple from current perceptual surroundings and generate independent
82 internal thoughts (Smallwood and Schooler, 2015). These thoughts can either be task-related,
83 such as actively thinking about how this manuscript should be structured, or task-unrelated,
84 where there is a spontaneous inner focus, such as suddenly remembering what a nice time I
85 had yesterday with my friends (Seli et al., 2016). These latter thoughts are the focus of the
86 current study, and have been variously described as task-unrelated self-generated thoughts,
87 daydreaming or mind-wandering (Smallwood and Schooler, 2015).

88

89 It has been shown that humans tend to mind-wander about 30-50% of waking time,
90 irrespective of the current activity (Kane et al., 2007; Killingsworth and Gilbert, 2010).
91 Nevertheless, mind-wandering frequency is particularly pronounced during restful periods
92 and low-demanding tasks (Smallwood and Schooler, 2015). The latter is often exploited by
93 experimentalists examining mind-wandering. Although the precise function of mind-

94 wandering is still debated, it is increasingly acknowledged to have influence across cognition
95 on processes such as future planning, creative thinking and problem-solving (Baird et al.,
96 2011; Baird et al., 2012), and even on depressive rumination and other mental health
97 disorders (Ehlers et al., 2004; Andrews-Hanna et al., 2014a). Furthermore, the content of
98 mind-wandering seems wide-ranging, including episodic memory recall (which involves a
99 sense of re-experiencing and is specific in time and place), future planning, mentalizing,
100 simulation of hypothetical scenarios, and involves a variety of emotions and different sensory
101 modalities (Andrews-Hanna et al., 2013; Smallwood et al., 2016). Interestingly, two of the
102 most prominent features of mind-wandering are mentally travelling forwards and backwards
103 in time and visual imagery, which are functions usually associated with the hippocampus
104 (Tulving, 1985, 2002; Hassabis et al., 2007).

105

106 The Default Mode Network (DMN), within which the hippocampus is a node, has been
107 associated with self-generated thoughts such as mind-wandering (Buckner et al., 2008;
108 Andrews-Hanna et al., 2014b). Of particular relevance here, stronger hippocampal
109 connectivity with other regions of the DMN was observed in individuals who experienced
110 more episodic details and greater flexibility in mental time travel during mind-wandering
111 episodes (Karapanagiotidis et al., 2016; Smallwood et al., 2016). Unfortunately, causal
112 evidence for hippocampal involvement in mind-wandering is lacking (Fox et al., 2016).
113 Behavioral studies of patients with lesions are crucial because they permit examination of the
114 causal effects of regional brain damage on the networks established by neuroimaging work.
115 People with hippocampal damage cannot vividly recall events from their past (Lah and
116 Miller, 2008) envision their future (Kurczek et al., 2015) or imagine fictitious scenes
117 (Hassabis et al., 2007). Therefore, whether they experience mind-wandering, and if they do,

118 what form does it take, are important and timely questions which we addressed by examining
119 mind-wandering in patients with selective bilateral hippocampal damage.

120

121 Previous studies have examined the effects of hippocampal damage during demanding tasks,
122 such as autobiographical memory retrieval (Lah and Miller, 2008), designed to challenge the
123 patients' cognitive abilities. In contrast, in order to establish what patients with hippocampal
124 damage think about spontaneously when there is no concurrent task, our focus was on what
125 they do in their mentally "free" time. We initially asked whether or not patients with
126 hippocampal damage were able to mentally decouple from the current perceptual input. If
127 yes, we then had a series of further questions. First, would they engage in mental time travel?
128 Second, what form would their mind-wandering take – spontaneous episodic, detailed
129 thoughts or semantic, abstract thoughts? Lastly, we asked whether they experienced
130 spontaneous visual imagery similar to that typically reported by control participants during
131 mind-wandering (Andrews-Hanna et al., 2013)?

132

133 **Materials and Methods**

134 *Participants*

135 Six patients (all right-handed males, mean age 57.0 years (SD 16.9), age range 27 to 70) with
136 selective bilateral hippocampal lesions and selective episodic memory impairment took part
137 (see Tables 1 and 2 for demographic information and neuropsychological profiles). Of note,
138 these patients were the same high-functioning individuals that took part in our previous
139 studies (McCormick et al., 2016, 2017a). Hippocampal damage (see example in Fig. 1a)
140 resulted in all cases from voltage-gated potassium channel (VGKC)-complex antibody-
141 mediated limbic encephalitis (LE). Two of the patients had bilateral signal hyperintensities in
142 the hippocampi on presentation, but hippocampal atrophy was observed in all patients.

143 Testing took place a median of seven years post-onset of hippocampal damage. In line with
144 previous reports of this patient population (Dalmau and Rosenfeld, 2014; Miller et al., 2017),
145 manual (blinded) segmentation of the hippocampi from high-resolution structural MRI scans
146 confirmed that our patients showed volume loss confined to the left (Patients – HPC:
147 2506mm^3 (mean) ± 394 (standard deviation), control participants – CTL: $3173\text{mm}^3 \pm 339$,
148 $W=4.0$, $p=0.002$) and right (HPC: $2678\text{mm}^3 \pm 528$, CTL: $3286\text{mm}^3 \pm 301$, $W=8.0$, $p=0.01$)
149 hippocampus. To rule out pathological differences between patients and controls elsewhere in
150 the brain, an automated voxel-based-morphometry (VBM; Ashburner, 2009) analysis was
151 carried out on whole brain T1 weighted MRI images and, in line with previous reports on
152 patients of this sort (Wagner et al., 2015; Finke et al., 2017; Miller et al., 2017), did not result
153 in any significant group differences outside of the hippocampus even at a liberal uncorrected
154 p-value of less than 0.001.

155
156 Neuropsychologically, the patients displayed an impairment in immediate and delayed recall
157 on the Logical Memory (short stories) test (Wechsler, 1997), and they recollected
158 significantly fewer episodic ('internal'), but not semantic ('external') details on the
159 Autobiographical Interview (Levine et al., 2002), as detailed in Table 2. All other cognitive
160 and emotional aspects of cognition were intact in these patients. In summary, these patients
161 seemed to have a selective difficulty in re-constructing internal events. Importantly, their
162 working memory capacity did not differ from that of controls, suggesting that the differences
163 in mind-wandering episodes we report here are unlikely to be due to an inability to remember
164 the thoughts.

165

166 Twelve healthy control participants also took part (all male, one left-handed, mean age 57.2
167 (16.6) years, age range from 25 to 77). In addition to comparing the two groups, we ensured
168 that each patient was matched closely to two of the control subjects on sex, age, and general

169 cognitive ability (measured by the Matrix Reasoning and Similarities subtests of the
170 Wechsler Abbreviated Scale of Intelligence – WASI; Wechsler, 1999). There were no
171 significant differences between patients and controls on age, general cognitive ability and on
172 neuropsychological tests assessing semantic memory, language, perception, executive
173 functions and mood (see Table 2). All participants gave informed written consent in
174 accordance with the local research ethics committees.

175

176 *Characterization of hippocampal damage*

177 *High resolution T2-weighted structural MRI scans of the medial temporal lobes*

178 Five of the patients and 10 of the control participants underwent structural MR imaging
179 limited to a partial volume focused on the temporal lobes using a 3.0-T whole body MR
180 scanner (Magnetom TIM Trio, Siemens Healthcare, Erlangen, Germany) operated with a
181 radiofrequency (RF) transmit body coil and 32-channel head RF receive coil. These structural
182 images were collected using a single-slab 3D T2-weighted turbo spin echo sequence with
183 variable flip angles (SPACE; Mugler et al., 2000) in combination with parallel imaging, to
184 simultaneously achieve a high image resolution of $\sim 500\mu\text{m}$, high sampling efficiency and
185 short scan time while maintaining a sufficient signal-to-noise ratio (SNR). After excitation of
186 a single axial slab the image was read out with the following parameters: resolution=0.52 x
187 0.52 x 0.5 mm, matrix=384 x 328, partitions=104, partition thickness=0.5 mm, partition
188 oversampling=15.4%, field of view=200 x 171 mm², TE=353 ms, TR=3200 ms, GRAPPA x
189 2 in phase-encoding (PE) direction, bandwidth=434 Hz/pixel, echo spacing=4.98 ms, turbo
190 factor in PE direction=177, echo train duration=881, averages=1.9. For reduction of signal
191 bias due to, for example, spatial variation in coil sensitivity profiles, the images were
192 normalized using a prescan, and a weak intensity filter was applied as implemented by the
193 scanner’s manufacturer. It took 12 minutes to obtain a scan.

194

195 *High resolution T1-weighted structural MRI scans of the whole brain at 3.0 Tesla*

196 In addition, five of the patients and 11 of the control participants underwent a whole brain
197 structural T1weighted sequence at an isotropic resolution of 800 μ m (Callaghan et al., 2015)
198 which was used for the automated VBM analysis (one control participant could not be
199 scanned). These images had a FoV of 256mm head-foot, 224mm anterior-posterior (AP), and
200 166mm right-left (RL). This sequence was a spoiled multi-echo 3D fast low angle shot
201 (FLASH) acquisition with a flip angle of 21⁰ and a repetition time (TR) of 25ms. To
202 accelerate the data acquisition, partially parallel imaging using the GRAPPA algorithm was
203 employed in each phase-encoded direction (AP and RL) with forty reference lines and a
204 speed up factor of two. Gradient echoes were acquired with alternating readout polarity at
205 eight equidistant echo times ranging from 2.34 to 18.44ms in steps of 2.30ms using a readout
206 bandwidth of 488Hz/pixel (Helms and Dechent, 2009). The first six echoes were averaged to
207 increase SNR (Helms and Dechent, 2009) producing a T1-weighted image with an effective
208 echo time of 8.3 ms.

209

210 *High resolution T1-weighted MRI scans of the whole brain at 7.0 Tesla*

211 One patient could not be scanned at our Centre due to recent dental implants. We therefore
212 used a whole brain T1-weighted image acquired previously on a 7.0 Tesla MRI scanner - a
213 three-dimensional whole-brain T1-weighted phase sensitive inversion recovery sequence
214 (Mougin et al., 2015) at an isotropic resolution of 600 μ m, with a tailored inversion pulse for
215 magnetization inversion at ultrahigh field (Hurley et al., 2010), providing inherent bias field
216 correction.

217

218

219

220 *Hippocampal segmentation*

221 To improve the SNR of the anatomical images, two or three T2-weighted high resolution
222 scans were acquired for a participant. Images from each participant were co-registered and
223 denoised following the Rician noise estimation (Coupe et al., 2010). The denoised images
224 were averaged and smoothed with a full-width at half maximum kernel of 2x2x2mm. In each
225 case, left and right hippocampi were manually (blindly) segmented and volumes extracted
226 using the ITK Snap software version 3.4.0 (Yushkevich et al., 2006).

227

228 *VBM analysis*

229 An automated VBM analysis was performed using SPM12 (Statistical Parametric Mapping,
230 Wellcome Centre for Human Neuroimaging, London, UK). The averaged T1-weighted
231 images were segmented into grey and white matter probability maps using the unified
232 segmentation approach (Ashburner and Friston, 2005). Inter-subject registration of the tissue
233 classes was performed using Dartel, a nonlinear diffeomorphic algorithm (Ashburner, 2007).
234 The resulting Dartel template and deformations were used to normalize the tissue probability
235 maps to the stereotactic space defined by the Montreal Neurological Institute (MNI) template.
236 For VBM analysis, the normalization procedure included modulating the grey matter tissue
237 probability maps by the Jacobian determinants of the deformation field and smoothing with
238 an isotropic Gaussian smoothing kernel of 8 mm full width at half maximum (FWHM). The
239 normalized grey matter from controls and the patients with hippocampal damage were
240 contrasted using a two sample t-test and thresholded at $p < 0.001$ uncorrected and a cluster
241 extend of 50 voxels.

242

243

244 *Experimental design and procedure*

245 We had the opportunity to shadow the patients with selective bilateral hippocampal damage
246 over two days during day-time hours, and so we adapted for use a well-established method,
247 descriptive experience sampling (DES), in which participants are asked frequently over an
248 extended period of time to describe what was on their minds just before they were aware of
249 being asked (Hurlburt, 1979; Hurlburt and Heavey, 2001; Hurlburt and Akhter, 2006). DES
250 has the advantage that thought probes can extend over a long period of time and the sampling
251 interval can be more extensive than alternative approaches in which a few thought samples
252 are taken while participants perform low-demanding distractor tasks (Smallwood et al., 2002;
253 Smallwood and Schooler, 2015). Furthermore, using DES, participants are encouraged to
254 describe freely what was on their minds, rather than categorizing thoughts into pre-specified
255 classes.

256

257 To mitigate any potential difficulties the hippocampal-damaged patients may have had with
258 remembering task instructions over longer time-scales, we made a number of adaptations to
259 the original DES protocol. For example, we changed the type of reminder. The reminder is
260 an important tool as it identifies the precise moment of sampling and happens externally to
261 the participant, meaning that the participant does not have to remember to track their own
262 thoughts (Hurlburt and Stuart, 2014). Usually, DES participants carry a beeper and receive
263 frequent sampling reminders while going about their everyday life (Hurlburt and Akhter,
264 2006). However, we adapted this sampling method to suit an extended experimental setting
265 over two days in which patients and controls experienced the same structured days (three
266 MRI scans, various cognitive tasks, breaks, lunches, etc). In our case, the experimenter
267 provided the external cue for the participant. Equally important as the reminder is the exact
268 time point of the sample. While previous studies have used a random sampling schedule

269 (Hurlburt, 1979; Smallwood and Schooler, 2015), our main goal was to examine the general
270 ability to perceptually decouple and the content of spontaneous thoughts of these rare
271 patients. We therefore tried to maximize our chances of catching perceptually decoupled
272 thoughts. Hence, we probed 20 times over the course of two structured research days (8 hours
273 each) at pre-specified times in restful moments. To keep the experimental context of the
274 sampling time points as closely matched across participants as possible, thoughts for all
275 participants were probed in the same rooms of our Centre, around the same times of day, and
276 in approximately the same experimental situations. This procedure resulted in schedules
277 whereby some samples were separated by several hours (e.g., during which the participant
278 underwent MRI scanning), and other samples which were relatively close in time (e.g., a few
279 minutes). In addition, in order to ascertain that all participants, especially the patients, could
280 remember time spans long enough to report their thoughts, we asked them to describe two
281 experiments unrelated to the current study shortly after completion. All participants were able
282 to provide accurate accounts of those experiments.

283

284 During sampling moments, such as after obtaining consent, and at the beginning of the tea
285 break, the experimenter would allow for a moment of quiet to emerge. That is, the
286 experimenter would fill out some forms or naturally disengage from any conversation. When
287 there was an appropriate time of silence, the experimenter would ask the participant “What
288 were you thinking about just before I asked you?” The participant was encouraged to briefly
289 describe the current thought in one or two sentences. On a prepared note sheet, the
290 participant’s response was written down verbatim. In a follow-up question, the experimenter
291 established whether the thought had been a visual image (if yes, scene or object) or a verbal
292 thought. Then, the experimenter clarified whether that thought had concerned the past,
293 present or future (and if it had been past or future, how far into the past or future). The

294 sampling procedure lasted no longer than approximately one minute to prevent lengthy post-
295 hoc elaboration. Lastly, divergent from other DES reports, we opted not to train our
296 participants before the start of the study. Although the training may have provided useful
297 guidance in monitoring one's own thoughts for the control participants, we felt that patients
298 might not find this as beneficial. Therefore, because the experimenter was present for all
299 samples, none of the participants was required to remember the follow-up questions
300 themselves but were instead cued by the experimenter. Of note, control participants reported
301 equal numbers of decoupled, scene-based thoughts in the first and second half of the samples
302 (first half=5.8 +/-0.9, second half=6.3 +/-1.2, $W=11.0$, $p=0.42$), suggesting that there was no
303 significant training effect. A lack of monitoring was further confirmed by the control
304 participants, because they anticipated the sampling probe for only 3 out of a total of 240
305 sampled thoughts.

306

307 Whereas previous research has examined the frequency and content of mind-wandering
308 episodes in healthy participants for features such as goal-orientation and emotional valence
309 (Andrews-Hanna et al., 2013; Andrews-Hanna et al., 2014a; Christoff et al., 2016), we
310 focused here on examining the effect of hippocampal damage on the frequency, time range,
311 representational content and form of mind-wandering, which are key to understanding
312 hippocampal function.

313

314 In summary, our adapted sampling protocol permitted us to leverage the naturalistic approach
315 of the typical DES reports that sample over an extended period of time, and allowed
316 participants to report their thoughts freely, while equating the daily activities and the
317 sampling moments of patients and controls participants to maximize our chances of catching
318 perceptually decoupled thoughts in an experimentally rigorous manner.

319

320 **Scoring**321 *Perceptually coupled or decoupled thoughts and mind-blanking*

322 An episode was considered mind-wandering when the response indicated that the mind was
323 disengaged from the external world (perceptually decoupled; Smallwood and Schooler,
324 2015). For example, the thought “I see your watch” was considered perceptually coupled,
325 whereas the thought “Time is sometimes slow and sometimes fast” was considered
326 perceptually decoupled. In a few instances, patients and control participants reported
327 thinking about nothing (i.e., mind-blanking; Ward and Wegner, 2013). The frequency of this
328 mind-blanking did not differ between the groups (CTL mean 0.4 +/- 0.88, HPC=1.8 +/-2.8,
329 MWU=23, p=0.19), and we therefore excluded these samples from further analysis.

330

331 *Temporal range*

332 After each sample, we clarified directly with participants whether that thought had concerned
333 the past, present or future, and if past or future, how distant into the past or future. We
334 further sorted participants’ responses from the “present” category based on the observation
335 that patients and control participants reported very different types of thoughts. Consequently,
336 we classified each mind-wandering episode that was labelled by participants as concerning
337 the present moment as either an atemporal scenario or not, in line with the protocol of
338 Jackson et al. (2013). A mind-wandering episode was considered “present”-related if the
339 thought was perceptually decoupled but concerned the now, for example “I’m thinking that
340 you are right-handed” or “I wonder whether I should eat another grape”. On the other hand, a
341 thought was classified as an atemporal scenario if the participant reported a mental event that
342 had no clear temporal direction. For example, a control participant’s thought was, “I noticed
343 this apparatus [EEG box] and I just imagined a picture in my mind in which that box was

344 being used in a horror setting”. By contrast, a patient reported while noticing the same EEG
345 box, “I wonder what this box with all these cables does. But I have no idea”. We display a
346 detailed characterization of the temporal range of mind-wandering episodes in Figure 2. For
347 statistical analysis, thoughts were binned into four main time categories, namely past (any
348 thought related to earlier than the present moment), present (now), future (any thought related
349 to later than the present moment), and atemporal thoughts.

350

351 *Representation type*

352 Thoughts were classified as either semantic or episodic (in line with established methods;
353 (Levine et al., 2002; Andrews-Hanna et al., 2014b) and, in addition, whether they contained
354 self-referential thinking or not (Andrews-Hanna et al., 2014b; Andrews-Hanna et al., 2014a).
355 A thought was classified as semantic if it contained mentalization, or general knowledge
356 about the world or the participant. For example, a semantic, self-referential thought of a
357 patient was: “I am self-pondering. Am I a creative person?” A thought was classified as
358 episodic if it contained specificity of time and place and a feeling of re- or pre-experiencing
359 (Tulving, 1983, 2002). For example, an episodic, self-referential thought of a control
360 participant was: “I am remembering a discussion I had with my friend at King’s Cross
361 concourse a few weeks ago. I can see the scene clearly in front of me.” Of note, we also
362 classified thoughts as episodic that had reference to a specific place and time, even if one or
363 both were fictitious (time was more often fictitious). For example, an atemporal, episodic,
364 non-self-referential thought of a control was: “I’m thinking about my friend. He’s travelling
365 around giving lectures. I imagine an auditorium and see my friend speaking.”

366

367

368

369 *Form of thoughts*

370 We asked participants after each sample whether the thought had been verbal or visual, and if
371 visual, whether it had been a scene or an object. Each thought was sorted into only one of
372 these categories. Some participants reported that some of the visual scenes also contained
373 verbal aspects, however, they regarded the visual scene as being more dominant. Therefore,
374 these thoughts were classified as scenes. This classification was accomplished in agreement
375 with each participant.

376

377 *Interrater reliability*

378 In order to avoid potential rater biases, a second rater, who was blind to group membership,
379 scored all thoughts from the patients and the control participants (except for one control
380 dataset which was used as a training set). Interrater reliability was calculated as the direct
381 correspondence between the two raters. That is, thoughts that were scored identically in a
382 category were given a '1' otherwise they were given a '0'. The reliability was then
383 established as the sum divided by the total amount of rated thoughts. Therefore a value of
384 0.99 indicates that in 99% of samples the raters categorized them identically. The overall
385 agreement between raters ranged between 84 and 99% across the thought categories (i.e.,
386 atemporal: 88%, coupled/decoupled 99%, semantic: 84%, episodic: 85%, and self-referential:
387 87%).

388

389 *Statistical analyses*

390 Since most of the dependent variables did not meet the assumptions for parametric statistics,
391 non-parametric tests were used for all within- and between-group analyses. Within-group
392 analyses with more than two dependent variables were first conducted using Friedman tests
393 (the non-parametric equivalent of repeated measures ANOVAs) and followed up with two-

394 tailed Wilcoxon Signed Rank tests (the non-parametric equivalent of paired t-tests). Between-
395 group analyses with more than two dependent variables were first conducted using Kruskal-
396 Wallis tests (the non-parametric equivalent of one-way ANOVAs) and followed up with two-
397 tailed Mann-Whitney U tests (the non-parametric equivalent of two-sample t-tests). Analyses
398 with two dependent variables were directly compared using two-tailed Wilcoxon Signed
399 Rank tests (within-group effects) or Mann-Whitney U tests (between-group effects). In all
400 cases, we considered p-values less than 0.05 as statistically significant. For significant results
401 we also report, where appropriate, the effect size (using non-parametric Cohen's d) and we
402 show the data of every participant.

403

404 **Results**

405 *Frequency of mind-wandering*

406 We first examined whether or not patients with hippocampal damage were able to mentally
407 decouple from the current perceptual input (Fig. 1b, c). We found that the percentage of
408 perceptually decoupled thoughts was greater than perceptually coupled thoughts in the
409 controls ($W=78.0$, $p=0.0005$) and patients ($W=21.0$, $p=0.03$); Table 3). Notably, we found no
410 difference between the two groups in the frequency of coupled ($MWU=19.5$, $p=0.12$) or
411 decoupled ($MWU=19.5$, $p=0.12$) thoughts.

412

413 *Temporal range of mind-wandering*

414 Since mental time travel seems to occur frequently during mind-wandering (Smallwood and
415 Schooler, 2015), we next examined whether the patient and control groups spontaneously
416 thought about the past, present or future. After each thought sample, we asked participants
417 whether the thought concerned the present moment, past or future, and if the latter two, how
418 distant was it from the present moment (see Fig. 2 for a detailed visualization of multiple

419 time bins). As described above, we also included an atemporal category in our analyses,
420 comprising thoughts where a participant reported a mental event that had no clear temporal
421 direction.

422

423 Examining the results for control participants in the first instance, we found that there was a
424 significant effect of time category (Friedman statistic=19.99, $df=3$, $p=0.0002$). Post-hoc
425 analyses showed that controls spent more of their mind-wandering time thinking about the
426 past than the present ($W=-62.0$, $p=0.002$) or future ($W=-72$, $p=0.002$). They also spent more
427 time simulating atemporal scenarios than thinking about the present ($W=62$, $p=0.01$). In
428 contrast, there was no overall effect of time category for patients (Friedman statistic=6.86,
429 $df=3$, $p=0.07$).

430

431 Direct comparison between the two groups revealed overall differences (Kruskal-Wallis
432 statistic=31.93, $df=7$, $p<0.0001$). Post-hoc analyses showed that the patients thought less often
433 than controls about past events (MWU=14.0, $p=0.04$, Cohen's $d=1.1$). By contrast, the
434 patients thought more often about the present moment than control participants (MWU=7.0,
435 $p=0.0034$, Cohen's $d=1.7$). There was no difference between the groups in the percentage of
436 future-thinking, which was generally low for both groups (MWU=18.0, $p=0.09$). Lastly, we
437 found that controls more often than the patients imagined atemporal events and hypothetical
438 scenarios that concerned a fictitious reality, which was not attached to any temporal
439 dimension (see the inset of Fig. 2, MWU=3.0, $p=0.0007$, Cohen's $d=2.1$).

440

441 *Representation type*

442 We next investigated what the patients mind-wandered about (Fig. 3; Table 3). Focusing first
443 on the control participants, we found that they reported significantly more episodic than

444 semantic thoughts ($W=78.0$, $p=0.0005$), and more self-related than non-self-related thoughts
445 ($W=78$, $p=0.0005$). The patients with hippocampal damage, on the other hand, experienced
446 more semantic than episodic thoughts ($W=-20.0$, $p=0.04$), and more self-related than non-
447 self-related thoughts ($W=21.0$, $p=0.03$).

448

449 Directly comparing the participant groups revealed that the controls reported more episodic
450 thoughts than the patients ($MWU=0.0$, $p=0.0001$, Cohen's $d=2.6$) and the patients reported
451 more semantic thoughts than the controls ($MWU=0.0$, $p=0.0001$, Cohen's $d=2.6$). As
452 expected, there was no significant difference in the percentage of self-referential
453 ($MWU=35.0$, $p=0.95$) or non-self-referential ($MWU=35.0$, $p=0.95$) thinking between the
454 groups. Together, these results show striking differences in the representational nature of
455 spontaneous inner experiences between control participants and hippocampal-damaged
456 patients.

457

458 *Form of thoughts*

459 Finally, after each sample we asked participants whether the thought had been verbal or
460 visual, and if visual, whether it had been a scene or an object (see Fig. 4, Table 3). For
461 controls, we found overall differences in the frequency of the different forms of thought
462 (Friedman statistic= 21.83 , $df=2$, $p<0.0001$). Post-hoc analyses showed that control
463 participants reported that the majority of their thoughts involved visual scenes, more so than
464 visual objects ($W=-78.0$, $p=0.0005$) or verbal thoughts ($W=-78.0$, $p=0.0005$), but more verbal
465 thoughts than visual objects ($W=58.0$, $p=0.007$). For the patients too there were overall
466 differences in the frequency of the different forms of thought (Friedman statistic= 9.48 , $df=2$,
467 $p=0.005$). In striking contrast to controls, patients thought almost entirely verbally. They

468 reported more verbal thoughts than visual scenes ($W=21$, $p=0.03$) and visual objects ($W=21$,
469 $p=0.03$), with no difference between visual scenes and objects ($W=-3$, $p=0.81$).

470

471 These differences in the experiential form of mind-wandering were confirmed by directly
472 comparing the participant groups (Kruskal-Wallis statistic= 56.33 , $df=7$, $p<0.0001$). Whereas
473 controls reported more visual scenes than patients (MWU= 0.0 , $p=0.0001$, Cohen's $d=2.6$),
474 patients reported more verbal thoughts than controls (MWU= 0.0 , $p=0.0001$, Cohen's $d=2.6$),
475 with no difference between participant groups for visual objects (MWU= 30.0 , $p=0.59$).

476

477 **Discussion**

478 Mind-wandering is pervasive in humans and likely has an important role to play across
479 cognition, influencing processes such as future planning, creative thinking and problem-
480 solving (Baird et al., 2011; Baird et al., 2012; Andrews-Hanna et al., 2013). Here we showed
481 that patients with hippocampal damage were able to perceptually decouple from the external
482 world and experience spontaneous thoughts. Nevertheless, the small, selective lesions of their
483 hippocampi dramatically affected the nature of their mind-wandering. Whereas healthy
484 participants thought about the past, present and future, primarily in terms of episodic, detail-
485 rich visual scenes, the patients mainly experienced verbally-mediated semantic thoughts
486 anchored in the present. Previous studies have examined episodic thought processes in
487 patients with hippocampal damage using explicit tasks, such as the Autobiographical
488 Interview (Levine et al., 2002) or the scene construction task (Hassabis et al., 2007), that
489 were designed to challenge the patients' ability. In contrast, our findings show that even when
490 there is no direct cognitive demand, the thought structure of people with hippocampal
491 damage is strikingly different from healthy controls.

492

493 We first consider whether our results can be explained by a memory deficit that caused the
494 patients to rapidly forget their mind-wandering thoughts before they could be accurately
495 reported. We do not think is the case for a number of reasons. First, the patients had intact
496 working memory and could retain task instructions during neuropsychological tests (Table 2)
497 over longer time-scales than those in the current study. Second, we asked participants to
498 describe two experiments unrelated to the current study shortly after completion, thus
499 mirroring the timescale of reporting their mind-wandering experiences. All participants,
500 including the patients, were able to provide accurate accounts of those experiments. Third, in
501 previously-published studies involving the same patients and control participants using
502 different paradigms, the patients were able to maintain information over time periods that
503 were longer than those required for generating the current mind-wandering samples
504 (McCormick et al., 2016, 2017a). Fourth, our sampling method did not involve any delay or
505 distraction that might have affected the patients, nor did our protocol allow for increased
506 post-hoc elaboration on the part of the control participants. Finally, if patients did not
507 remember what they had been thinking about, the frequency of their mind-wandering would
508 have been lower and they would have reported more mind-blanking, which was not the case.
509 Thus, we are confident the patients were able to accurately report what was on their mind
510 within seconds of the sampling cue.

511

512 Previous reports have estimated that humans tend to mind-wander about 30-50% of waking
513 time (Kane et al., 2007; Killingsworth and Gilbert, 2010). Here, we report percentages nearer
514 80-90%. However, we specifically aimed to catch restful periods and so our higher
515 percentage of mind-wandering thoughts suggests that we were successful at probing time
516 points when mind-wandering levels were high.

517

518 Numerous studies have focused on delineating different aspects of inner experiences. For
519 example, self-generated thinking (either intentional or unintentional; Seli et al., 2016)
520 typically refers to the ability to mentally decouple from the current perceptual surroundings
521 and generate independent internal thoughts (Smallwood and Schooler, 2015), which is a
522 dichotomous definition that we employed in the current study. In reality, these self-generated
523 thoughts align on a continuum ranging from closely task-related to totally task-unrelated
524 (Smallwood and Schooler, 2015). What was most important for our research question was
525 whether patients could decouple perceptually from their immediate surroundings in a
526 completely task-free context. We found that they were able to do so and that the frequency of
527 their mind-wandering did not differ from that of the control group. This result is especially
528 noteworthy, given a recent study that found reduced frequency of mind-wandering in patients
529 with ventromedial prefrontal cortex (vmPFC) lesions (Bertossi and Ciaramelli, 2016), a brain
530 region with dense functional and anatomical connections with the hippocampus (Andrews-
531 Hanna et al., 2010; Catani et al., 2012; Catani et al., 2013; McCormick et al., 2017b).
532 Although there were differences in the experimental setup between our study and that
533 involving the vmPFC patients, the difference in mind-wandering frequency observed in these
534 two studies might indicate that the vmPFC is critical for the initiation of endogenous
535 spontaneous thought and the hippocampus for its form and content.

536

537 At first glance, our finding of group differences in the temporal extent of mind-wandering is
538 not surprising given the difficulty patients with hippocampal damage are known to have with
539 recalling recent and remote episodic memories and imagining the future (Rosenbaum et al.,
540 2008; Kurczek et al., 2015). However, these previous results were based on active and
541 cognitively demanding tasks. To the best of our knowledge, this is the first indication that
542 hippocampal-damaged patients experience reduced mental time travel even in their

543 spontaneous thoughts. Of note, we did not replicate previous reports suggesting a near
544 future-thinking bias in the mind-wandering of healthy participants (Stawarczyk et al., 2011;
545 Song and Wang, 2012; Bertossi and Ciaramelli, 2016). The current experimental procedure
546 and the older age of our participants may have influenced these results (Maillet and Schacter,
547 2016). For example, instead of sampling during low-demanding computer tasks or in natural
548 environments that may encourage thoughts about the near future (e.g., “Where am I going
549 after I’m finished here?”), we sampled thoughts across a structured day of stimulating
550 research activities. This may have provided more opportunities to think about the recently-
551 completed cognitive tasks or MRI scans. In addition, many previous studies have not
552 included an atemporal category of thoughts, and it has been argued that thoughts labelled as
553 future-oriented might in some instances be more accurately characterized as atemporal
554 (Jackson et al., 2013). Indeed, in line with our results, it has been reported that healthy older
555 adults experience more atemporal than future-oriented mind-wandering episodes (Jackson et
556 al., 2013).

557

558 Recently, there have been increased efforts to map the complex cognitive processes that
559 support mind-wandering to specific brain regions. While it has been established that the
560 DMN is associated with mind-wandering (Buckner et al., 2008; Andrews-Hanna et al.,
561 2014a; Smallwood and Schooler, 2015), the contributions of specific brain areas within the
562 DMN to mind-wandering remain unclear. Our results provide novel evidence that the
563 hippocampus plays a causal role in episodic mind-wandering. These findings align with
564 recent neuroimaging work that focused on a subsystem of the DMN, of which the
565 hippocampus (and vmPFC) are nodes (Andrews-Hanna et al., 2010), and illustrated that
566 functional and structural connectivity is stronger in individuals who report many detail-rich
567 mental time travel experiences during mind-wandering (Karapanagiotidis et al., 2016;

568 Smallwood et al., 2016). Our results further accord with network analyses in patients with
569 hippocampal damage that showed altered hippocampal-neocortical connectivity patterns
570 (Hayes et al., 2012; McCormick et al., 2014; Henson et al., 2016), which were associated
571 with worse episodic memory capacity (McCormick et al., 2014). Of note, to the best of our
572 knowledge, ours is the first report of a concomitant increase in spontaneous semantic
573 thoughts associated with hippocampal damage. This may help to explain previous findings of
574 increased connectivity between brain areas involved in semantic processing in resting-state
575 fMRI studies involving similar patients (Hayes et al., 2012; McCormick et al., 2014).

576

577 In line with previous studies, our results demonstrate that mind-wandering episodes of
578 control participants typically comprise visual imagery (Andrews-Hanna et al., 2013). We
579 expand on existing studies by showing that visual imagery in task-unrelated mind-wandering
580 of healthy controls primarily consists of spatially coherent visual scenes. In striking contrast,
581 the patients with bilateral hippocampal damage no longer reported visualizing mental scenes,
582 relying instead on a verbal thought structure. A scene construction deficit has been implicated
583 in the impaired autobiographical memory and future thinking of patients with hippocampal
584 damage (Hassabis and Maguire, 2007; Maguire and Mullally, 2013; Clark and Maguire,
585 2016). Our findings support this link between episodic thought and scene imagery.
586 Importantly, this deficit also extends to scene perception tasks (Lee et al., 2005; Aly et al.,
587 2013; McCormick et al., 2017a), suggesting that the lack of mental scenes is not because of
588 faster visual degradation of imagery (Warren et al., 2011), but rather is due to an online scene
589 construction problem. Thus, our results strongly suggest that hippocampal-supported scene
590 construction is also central to the content and form of mind-wandering, and that without it,
591 spontaneous thought seems to be reliant on verbal semantics.

592

593 Although the precise definition of mind-wandering is still debated, our results show that
594 selective bilateral lesions to the hippocampus impair perceptually-decoupled inner thoughts
595 in specific ways, thus informing the nature of mind-wandering and how it is realized at the
596 neural level. That individuals with hippocampal damage experience mind-wandering but very
597 little detail-rich mental imagery are important new insights which indicate the hippocampus
598 is not necessary for the instigation of spontaneous thought per se. Instead, it seems to be
599 crucial for processing the form and content of mind-wandering. Our results also speak to the
600 functions of the hippocampus. By showing it plays a causal role in a phenomenon as
601 ubiquitous as mind-wandering, this exposes the impact of the hippocampus beyond its
602 traditionally-perceived role in memory, placing it at the center of our everyday mental
603 experiences.

604

605 **References**

- 606 Aly M, Ranganath C, Yonelinas AP (2013) Detecting changes in scenes: The hippocampus is
607 critical for strength-based perception. *Neuron* 78:1127-1137.
- 608 Andrews-Hanna JR, Smallwood J, Spreng RN (2014a) The default network and self-
609 generated thought: component processes, dynamic control, and clinical relevance.
610 *Ann NY Acad Sci* 1316:29-52.
- 611 Andrews-Hanna JR, Saxe R, Yarkoni T (2014b) Contributions of episodic retrieval and
612 mentalizing to autobiographical thought: Evidence from functional neuroimaging,
613 resting-state connectivity, and fMRI meta-analyses. *Neuroimage* 91:324-335.
- 614 Andrews-Hanna JR, Reidler JS, Sepulcre J, Poulin R, Buckner RL (2010) Functional-
615 anatomic fractionation of the brain's default network. *Neuron* 65:550-562.
- 616 Andrews-Hanna JR, Kaiser RH, Turner AE, Reineberg AE, Godinez D, Dimidjian S, Banich
617 MT (2013) A penny for your thoughts: dimensions of self-generated thought content
618 and relationships with individual differences in emotional wellbeing. *Front Psychol*
619 4:900.
- 620 Ashburner J (2007) A fast diffeomorphic image registration algorithm. *Neuroimage* 38:95-
621 113.
- 622 Ashburner J (2009) Computational anatomy with the SPM software. *Magn Reson Imaging*
623 27:1163-1174.
- 624 Ashburner J, Friston KJ (2005) Unified segmentation. *Neuroimage* 26:839-851.
- 625 Baird B, Smallwood J, Schooler JW (2011) Back to the future: autobiographical planning and
626 the functionality of mind-wandering. *Conscious Cogn* 20:1604-1611.
- 627 Baird B, Smallwood J, Mrazek MD, Kam JW, Franklin MS, Schooler JW (2012) Inspired by
628 distraction: mind wandering facilitates creative incubation. *Psychol Sci* 23:1117-1122.

- 629 Bertossi E, Ciaramelli E (2016) Ventromedial prefrontal damage reduces mind-wandering
630 and biases its temporal focus. *Social Cognitive and Affective Neurosci* 11:1783-1791.
- 631 Buckner RL, Andrews-Hanna JR, Schacter DL (2008) The brain's default network: anatomy,
632 function, and relevance to disease. *Ann NY Acad Sci* 1124:1-38.
- 633 Burgess P, Shallice T (1997) *The Hayling and Brixton Tests*. Test manual. Bury St Edmunds,
634 UK: Thames Valley Test Company.
- 635 Callaghan MF, Josephs O, Herbst M, Zaitsev M, Todd N, Weiskopf N (2015) An evaluation
636 of prospective motion correction (PMC) for high resolution quantitative MRI. *Front*
637 *Neurosci* 9:97.
- 638 Catani M, Dell'acqua F, Thiebaut de Schotten M (2013) A revised limbic system model for
639 memory, emotion and behaviour. *Neurosci Biobehav Rev* 37:1724-1737.
- 640 Catani M, Dell'acqua F, Vergani F, Malik F, Hodge H, Roy P, Valabregue R, Thiebaut de
641 Schotten M (2012) Short frontal lobe connections of the human brain. *Cortex* 48:273-
642 291.
- 643 Christoff K, Irving ZC, Fox KC, Spreng RN, Andrews-Hanna JR (2016) Mind-wandering as
644 spontaneous thought: a dynamic framework. *Nat Rev Neurosci*.
- 645 Clark IA, Maguire EA (2016) Remembering preservation in hippocampal amnesia. *Annu Rev*
646 *Psychol* 67:51-82.
- 647 Coupe P, Manjon JV, Gedamu E, Arnold D, Robles M, Collins DL (2010) Robust Rician
648 noise estimation for MR images. *Med Image Anal* 14:483-493.
- 649 Dalmau J, Rosenfeld MR (2014) Autoimmune encephalitis update. *Neuro Oncol* 16:771-778.
- 650 Delis DC, Kaplan E, Kramer JH (2001) *Delis Kaplan Executive Function System (D-KEFS)*.
651 San Antonio, TX: The Psychological Corporation.
- 652 Ehlers A, Hackmann A, Michael T (2004) Intrusive re-experiencing in post-traumatic stress
653 disorder: phenomenology, theory, and therapy. *Memory* 12:403-415.
- 654 Finke C, Pruss H, Heine J, Reuter S, Kopp UA, Wegner F, Then Bergh F, Koch S, Jansen O,
655 Munte T, Deuschl G, Ruprecht K, Stocker W, Wandinger KP, Paul F, Bartsch T
656 (2017) Evaluation of cognitive deficits and structural hippocampal damage in
657 encephalitis with Leucine-rich, glioma-inactivated 1 antibodies. *JAMA Neurol* 74:50-
658 59.
- 659 Fox KC, Andrews-Hanna JR, Christoff K (2016) The neurobiology of self-generated thought
660 from cells to systems: Integrating evidence from lesion studies, human intracranial
661 electrophysiology, neurochemistry, and neuroendocrinology. *Neuroscience* 335:134-
662 150.
- 663 Gabrovska V, Laws K, McKenna P (1996) Visual form perception in schizophrenia: further
664 evidence for a disorder of semantic memory. *Eur Psychiatry* 11:278.
- 665 Hassabis D, Maguire EA (2007) Deconstructing episodic memory with construction. *Trends*
666 *Cogn Sci* 11:299-306.
- 667 Hassabis D, Kumaran D, Vann SD, Maguire EA (2007) Patients with hippocampal amnesia
668 cannot imagine new experiences. *Proc Natl Acad Sci USA* 104:1726-1731.
- 669 Hayes SM, Salat DH, Verfaellie M (2012) Default network connectivity in medial temporal
670 lobe amnesia. *J Neurosci* 32:14622-14629.
- 671 Henson RN, Greve A, Cooper E, Gregori M, Simons JS, Geerligs L, Erzinclioglu S, Kapur N,
672 Browne G (2016) The effects of hippocampal lesions on MRI measures of structural
673 and functional connectivity. *Hippocampus* 26:1447-1463.
- 674 Hurlburt RT (1979) Random sampling of cognitions and behavior *J Res Pers* 13:103-111.
- 675 Hurlburt RT, Heavey CL (2001) Telling what we know: describing inner experience. *Trends*
676 *Cogn Sci* 5:400-403.
- 677 Hurlburt RT, Akhter SA (2006) The descriptive experience sampling method. *Phen Cogn Sci*
678 5:271-301.

- 679 Hurlburt RT, Stuart SAJ (2014) Grounding the science of inner experience in the
680 apprehension of phenomena. A review of *Consciousness and the Self: New Essays*.
681 *American Journal of Psychology* 127:253-260.
- 682 Hurley AC, Al-Radaideh A, Bai L, Aickelin U, Coxon R, Glover P, Gowland PA (2010)
683 Tailored RF pulse for magnetization inversion at ultrahigh field. *Magn Reson Med*
684 63:51-58.
- 685 Jackson JD, Weinstein Y, Balota DA (2013) Can mind-wandering be timeless? Atemporal
686 focus and aging in mind-wandering paradigms. *Frontiers in psychology* 4:742.
- 687 Kane MJ, Brown LH, McVay JC, Silvia PJ, I. M-G, Kwapil TR (2007) For whom the mind
688 wanders, and when. *Psychol Sci* 18:614-621.
- 689 Karapanagiotidis T, Bernhardt BC, Jefferies E, Smallwood J (2016) Tracking thoughts:
690 Exploring the neural architecture of mental time travel during mind-wandering.
691 *Neuroimage* 147:272-281.
- 692 Killingsworth MA, Gilbert DT (2010) A wandering mind is an unhappy mind. *Science*
693 330:932.
- 694 Kurczek J, Wechsler E, Ahuja S, Jensen U, Cohen NJ, Tranel D, Duff M (2015) Differential
695 contributions of hippocampus and medial prefrontal cortex to self-projection and self-
696 referential processing. *Neuropsychologia* 73:116-126.
- 697 Lah S, Miller L (2008) Effects of temporal lobe lesions on retrograde memory: a critical
698 review. *Neuropsychol Rev* 18:24-52.
- 699 Lee AC, Bussey TJ, Murray EA, Saksida LM, Epstein RA, Kapur N, Hodges JR, Graham KS
700 (2005) Perceptual deficits in amnesia: challenging the medial temporal lobe
701 'mnemonic' view. *Neuropsychologia* 43:1-11.
- 702 Levine B, Svoboda E, Hay JF, Winocur G, Moscovitch M (2002) Aging and autobiographical
703 memory: dissociating episodic from semantic retrieval. *Psychol Aging* 17:677-689.
- 704 Maguire EA, Mullally SL (2013) The hippocampus: a manifesto for change. *Journal of Exp*
705 *Psychol Gen* 142:1180-1189.
- 706 Maillet D, Schacter DL (2016) From mind wandering to involuntary retrieval: Age-related
707 differences in spontaneous cognitive processes. *Neuropsychologia* 80:142-156.
- 708 McCormick C, Rosenthal CR, Miller TD, Maguire EA (2016) Hippocampal damage
709 increases deontological responses during moral decision making. *J Neurosci*
710 36:12157-12167.
- 711 McCormick C, Rosenthal CR, Miller TD, Maguire EA (2017a) Deciding what is possible and
712 impossible following hippocampal damage in humans. *Hippocampus* 27:303-314.
- 713 McCormick C, Ciaramelli E, De Luca F, Maguire EA. 2017b. Comparing and contrasting the
714 cognitive effects of hippocampal and ventromedial prefrontal cortex damage: a review
715 of human lesion studies. *Neuroscience* (doi: 10.1016/j.neuroscience.2017.07.066).
- 716 McCormick C, Protzner AB, Barnett AJ, Cohn M, Valiante TA, McAndrews MP (2014)
717 Linking DMN connectivity to episodic memory capacity: What can we learn from
718 patients with medial temporal lobe damage? *NeuroImage Clinical* 5:188-196.
- 719 McKenna P, Warrington EK (1980) Testing for nominal dysphasia. *J Neurol Neurosurg*
720 *Psychiatry* 43:781-788.
- 721 Miller TD, Chong TTJ, Aimola Davies AM, Ng TWC, Johnson MR, Irani SR, Vincent A,
722 Husain M, Jacob S, Maddison P, Kennard C, Gowland PA, Rosenthal CR (2017)
723 Focal CA3 hippocampal subfield atrophy following LGI1 VGKC-complex antibody
724 limbic encephalitis. *Brain*.
- 725 Mougins O, Abdel-Fahim R, Dineen R, Pitiot A, Evangelou N, Gowland P (2015) Imaging
726 gray matter with concomitant null point imaging from the phase sensitive inversion
727 recovery sequence. *Magn Reson Med* 76:1512-1516.

- 728 Osterrieth PA (1944) The test of copying a complex figure: A contribution to the study of
729 perception and memory. *Arch Psychol* 30:286-356.
- 730 Rosenbaum RS, Moscovitch M, Foster JK, Schnyer DM, Gao F, Kovacevic N, Verfaellie M,
731 Black SE, Levine B (2008) Patterns of autobiographical memory loss in medial-
732 temporal lobe amnesic patients. *J Cogn Neurosci* 20:1490-1506.
- 733 Seli P, Risko EF, Smilek D, Schacter DL (2016) Mind-wandering with and without intention.
734 *Trends Cogn Sci* 20:605-617.
- 735 Smallwood J, Schooler JW (2015) The science of mind wandering: empirically navigating
736 the stream of consciousness. *Annu Rev Psychol* 66:487-518.
- 737 Smallwood J, Obonsawin M, Reid H (2002) The effects of block duration and task demands
738 on the experience of task unrelated thought. *Imagin Cogn Pers* 22:13-31.
- 739 Smallwood J, Karapanagiotidis T, Ruby F, Medea B, de Caso I, Konishi M, Wang HT,
740 Hallam G, Margulies DS, Jefferies E (2016) Representing representation: Integration
741 between the temporal lobe and the posterior cingulate influences the content and form
742 of spontaneous thought. *PLoS One* 11:e0152272.
- 743 Song X, Wang X (2012) Mind wandering in Chinese daily lives - an experience sampling
744 study. *PLoS One* 7:1-9.
- 745 Stawarczyk D, Majerus S, Maj M, Van der Linden M, D'Argembeau A (2011) Mind-
746 wandering: phenomenology and function as assessed with a novel experience
747 sampling method. *Acta Psychol (Amst)* 136:370-381.
- 748 Tulving E (1983) *Elements of Episodic Memory*. New York: Oxford University Press.
- 749 Tulving E (1985) Memory and consciousness. *Can J Exp Psychol* 26:1-12.
- 750 Tulving E (2002) Episodic memory: from mind to brain. *Annu Rev Psychol* 53:1-25.
- 751 Wagner J, Weber B, Elger CE (2015) Early and chronic gray matter volume changes in
752 limbic encephalitis revealed by voxel-based morphometry. *Epilepsia* 56:754-761.
- 753 Ward AF, Wegner DM (2013) Mind-blanking: when the mind goes away. *Frontiers in*
754 *Psychology* 4:650.
- 755 Warren DE, Duff MC, Tranel D, Cohen NJ (2011) Observing degradation of visual
756 representations over short intervals when medial temporal lobe is damaged. *J Cogn*
757 *Neurosci* 23:3862-3873.
- 758 Warrington EK (1984) *Recogniton Memory Test: Manual*. Berkshire, UK.: NFER-Nelson.
- 759 Warrington EK (2010) The Graded Naming Test: A Restandardisation. *Neuropsychol Rehab*
760 7:143-146.
- 761 Warrington EK, James M (1991) A new test of object decision: 2D silhouettes featuring a
762 minimal view. *Cortex* 27:370-383.
- 763 Wechsler D (1997) *Wechsler Memory Scales - Third Edition*. San Antonio, TX: The
764 Psychological Corporation.
- 765 Wechsler D (1999) *Wechsler Abbreviated Scale of Intelligence*. New York, NY: The
766 Psychological Corporation: Harcourt Brace & Company.
- 767 Yushkevich PA, Piven J, Hazlett HC, Smith RG, Ho S, Gee JC, Gerig G (2006) User-guided
768 3D active contour segmentation of anatomical structures: significantly improved
769 efficiency and reliability. *Neuroimage* 31:1116-1128.
- 770 Zigmond AS, Snaith RP (1983) The hospital anxiety and depression scale. *Acta Psychiatrica*
771 *Scand* 67:361-370.
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774 **Figure legends**

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Figure 1. Hippocampal damage and the frequency of mind-wandering. (a) A T2-weighted structural MR image of an example patient with selective bilateral hippocampal damage and an age, gender and IQ-matched healthy control participant. Images are displayed in native space corresponding approximately to the position of $y=-10$ in the MNI coordinate system. (b) Examples of mind-wandering experiences from controls (CTL) and patients with hippocampal damage (HPC). (c) The average percentage of perceptually coupled and decoupled spontaneous thoughts (minus ‘blank’ thoughts) during quiet restful moments for individual patients with hippocampal damage (red symbols) and healthy control participants (blue circles). Both groups reported a high level of mind-wandering experiences, with no differences between patients and control participants.

Figure 2. The temporal range of mind-wandering. Mean percentages of mind-wandering thoughts of patients with hippocampal damage (HPC, red circles with a dot) and controls (CTL, blue circles) for the past, present and future. For display purposes, thoughts are classified into time bins according to past (including earlier today), the present (now) and future (including later today); m=months, y=years. Control participants reported more thoughts related to the past than patients. In contrast, patients reported more thoughts related to the present than controls. The inset graph shows the percentage of thoughts during which patients (red symbols) and controls (blue circles) engaged in the imagining of atemporal scenarios.

Figure 3. Semantic and episodic thinking during mind-wandering. Percentages of mind-wandering samples classified as semantic, episodic, self-referential or non-self-referential for patients with hippocampal damage (HPC, red symbols) and controls (CTL, blue circles). The patients had predominantly semantic thoughts, whereas the thoughts of the control participants were mainly episodic.

Figure 4. Cumulative percentages of visual and verbal mind-wandering thoughts. The average percentage of verbal thoughts is depicted per group (HPC=hippocampal-damaged patients, CTL=controls) as an orange bar; the individual data points are illustrated with orange symbols. The average cumulative percentage of thoughts containing visual objects is depicted as a grey bar above the average percentage of the verbal thoughts. The individual data points of thoughts containing visual objects (grey symbols) are illustrated as cumulative percentages above the orange data points (i.e., the patient represented as a square symbol reported around 70% verbal and around 25% visual object thoughts). Lastly, the average cumulative percentage of thoughts containing visual scenes is depicted as a green bar on top of the grey bar (green symbols all adding up to 100%). Whereas patients with hippocampal damage reported thinking in words for the majority of samples, healthy control participants’ thoughts were predominantly in the form of visual scenes.

820 **Table 1. Summary of demographic information.**

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Group	N	HD	Age	Chronicity	LHPC vol*	RHPC vol*
HPC group	6 (M)	6 (R)	57.0 (16.9)	6.8 (2.1)	2506 (394)	2678 (528)
CTL group	12 (M)	11 (R)	57.2 (16.6)	n.a.	3173 (339)	3286 (301)
p-value			0.97	n.a.	0.002	0.01

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823 For both groups, means are displayed with standard deviations below the corresponding mean in
824 parentheses. HPC=hippocampal-damaged patients; CTL=healthy control participants; M=Male;
825 HD=Handedness; n.a.=not applicable; R=Right; L=Left; vol=volume in mm³. *One control
826 participant could not be scanned, therefore hippocampal volumes are based on all six patients and 11
827 control participants. Age and chronicity are described in years. p-value=p-value of between-group
828 non-parametric Mann-Whitney U tests with significant differences depicted in bold.

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831 **Table 2. Summary of neuropsychological information.**

	Controls		HPC Patients		P-Value
	M	SD	M	SD	
General Cognition					
WASI Matrix Reasoning	13.8	1.5	13.2	2.2	0.51
WASI Similarities	11.8	2.6	12.8	1.8	0.54
Episodic memory					
Autobiographical Interview int*	51.3	13.6	31.7	6.7	0.01
Autobiographical Interview ext*	5.9	2.2	6.1	3.8	0.67
WMS Logical Memory (immediate recall, units)	12.6	3.2	8.7	2.4	0.03
WMS Logical Memory (immediate recall, thematic)	13.8	3.0	9.2	2.6	0.01
WMS Wordlist (immediate recall)	13.3	3.2	10.2	3.9	0.14
Rey-Osterrieth Complex Figure (copy /36)	35.5	1.4	33.4	4.2	0.19
Rey-Osterrieth Complex Figure (immediate recall /36)	23.8	7.4	19.0	5.9	0.12
WMS Logical Memory (delayed recall, units)	13.2	3.7	7.8	4.0	0.01
WMS Logical Memory (delayed recall, thematic)	13.5	3.2	7.0	4.8	0.01
WMS Wordlist (delayed recognition)	11.7	1.4	9.3	4.5	0.47
Rey-Osterrieth Complex Figure (delayed recall, /36)	23.8	7.8	18.1	6.4	0.06
Warrington Recognition Memory test for Words	12.0	2.3	12.3	2.4	0.99
Warrington Recognition Memory test for Faces	11.3	3.0	9.2	4.3	0.14
Semantic memory					
Warrington Graded Naming Test	13.7	2.4	12.5	3.0	0.34
Attention/Working memory					
WMS Digit Span (forward)	13.3	3.2	12.0	2.5	0.76
Executive Functions					
DKEFS Letter Fluency (FAS)	14.3	3.2	12.7	3.7	0.32
DKEFS Category Fluency	13.9	4.4	12.5	5.2	0.51
DKEFS Category Switch Test	12.7	2.9	12.3	3.5	0.59
DKEFS Stroop Word-Colour Interference Test	12.4	2.2	13.3	2.2	0.41
Hayling Sentence Completion Test (coherent)	6.1	1.0	5.8	0.4	0.73
Hayling Sentence Completion Test (incoherent)	5.8	0.8	5.8	0.4	0.95
Hayling Sentence Completion Test (errors)	6.8	1.1	6.5	1.9	0.93

Hayling Sentence Completion Test (total)	18.7	1.6	18.2	2.1	0.75
DKEFS Trails Test (visual scanning)	12.0	1.3	11.2	1.0	0.17
DKEFS Trails Test (number sequencing)	11.8	2.5	10.2	2.3	0.12
DKEFS Trails Test (letter sequencing)	12.5	1.5	11.0	2.4	0.21
DKEFS Trails Test (letter-number sequencing)	12.8	1.0	10.7	2.2	0.04
DKEFS Trails Test (motor speed)	11.8	1.1	10.2	4.5	0.99
Visual perception					
VOSP Dot Counting (/10)	10.0	0.0	9.7	0.8	0.33
VOSP Position Discrimination (/20)	20.0	0.0	19.7	0.8	0.33
VOSP Cube Analysis (/10)	9.6	0.8	9.7	0.8	0.99
VOSP Overall (/40)	39.6	0.8	39.0	2.4	0.99
Mood					
HADS Anxiety	4.3	2.9	4.3	3.3	0.83
HADS Depression	2.3	2.7	2.5	2.3	0.71

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For both groups, means (M) and standard deviations (SD) are displayed. HPC=hippocampal-damaged patients; p-value of between-group non-parametric Mann-Whitney U tests with significant differences are depicted in bold; WASI=Wechsler Abbreviated Scale of Intelligence (Wechsler, 1999); Scaled scores of the WASI Matrix Reasoning subtest and WASI Similarities subtest. Autobiographical Interview (Levine et al., 2002): int=average number of internal (episodic) details over five memories, ext=average external (semantic) details over five memories. ⁴Of note, autobiographical memory performance of the patients was compared to a separate control group (5 males, 1 female, mean age 55.2+/-18 years, range 22-69, all right-handed). Scaled scores of the WMS=Wechsler Memory Scale III (Wechsler, 1997), LM=logical memory immediate and delayed units and thematic scores, wordlist immediate total recall and delayed recognition. Rey=Rey-Osterrieth complex figure copy, immediate, and delayed recall (Osterrieth, 1944). Scaled scores of the Warrington Recognition Memory Test for Words and Faces (Warrington, 1984). Scaled scores of the Warrington Graded Naming Test (McKenna and Warrington, 1980; Warrington, 2010). Scaled scores of the DKEFS=Delis-Kaplan Executive Function System, letter fluency (FAS), category fluency (animals/boys names), category switch test (fruit/furniture), Stroop word-colour interference test, trails tests, including visual scanning, number sequencing, letter sequencing, number-letter switching and motor speed tests (Delis et al., 2001). Scaled scores of the Hayling Sentence Completion Test (Burgess and Shallice, 1997). VOSP=Visual Object and Space Perception Battery dot counting, cube analysis, and position discrimination subtests (Warrington and James, 1991; Gabrovska et al., 1996), HADS=Hospital Anxiety and Depression Scale (Zigmond and Snaith, 1983).

854 **Table 3. Summary of mind-wandering data.**
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	CTL		HPC		p
Mind-wandering					
<i>Perceptually coupled</i>	6.8	(7.5)	13.4	(13.6)	n.s.
<i>Perceptually decoupled</i>	93.1	(7.5)	86.6	(13.6)	n.s.
Temporal range					
<i>Past</i>	36.9	(12.7)	21.2	(16.4)	*
<i>Present</i>	18.9	(8.3)	32.9	(7.4)	**
<i>Future</i>	15.6	(9.9)	24.6	(9.2)	n.s.
<i>Atemporal</i>	31.8	(8.7)	17.8	(5.5)	****
Representational type					
<i>Episodic</i>	72.6	(12.4)	24.8	(14.3)	****
<i>Semantic</i>	27.4	(12.4)	75.2	(14.3)	****
<i>Self-referential</i>	75.2	(10.1)	75.6	(8.4)	n.s.
<i>Non-self-referential</i>	27.8	(10.1)	24.4	(8.5)	n.s.
Form of thought					
<i>Scenes</i>	63.5	(5.4)	12.3	(11.7)	****
<i>Objects</i>	9.8	(6.9)	9.1	(9.9)	n.s.
<i>Words</i>	26.8	(8.4)	78.7	(16.2)	****

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 857 For both groups, means (percentages) are displayed with standard deviations next to them in
 858 parentheses. CTL=healthy control participants; HPC=hippocampal-damaged patients; p=p-value for
 859 between-group non-parametric Mann-Whitney U tests. *=p<0.05, **=p<0.01, ****=p<0.001;
 860 n.s.=not significantly different.
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