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Evidence for a Large-Scale Brain System Supporting Allostasis and Interoception in Humans

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Supplementary Discussion

Detailed description of subcortical, hippocampal, brainstem, and cerebellar connectivity within the interoceptive system. Here, we briefly justify the inclusion of each non-cortical region in our connectivity analyses, report its observed intrinsic connectivity patterns with the allostatic/interoceptive system seeds (Fig. 4 and Supplementary Figure 6), and compare our results with published tract-tracing studies showing monosynaptic anatomical connectivity among these regions (Table 2). Discrepancies between our results and tract-tracing are only indicated if our fMRI results failed to show connectivity between regions that are monosynaptically connected. fMRI intrinsic connectivity reflects both direct and indirect (multisynaptic) connections^{46,47} and thus our results sometimes show connectivity for regions that are disynaptically connected.

Thalamus. We examined connectivity to two thalamic nuclei: the posterior part of the ventromedial nucleus (VMpo; for a review, see ¹) for interoceptive input specifically and the larger ventral posterior (VP) complex for somatic input more broadly². Our fMRI results revealed that all cortical and amygdalar seeds exhibited connectivity with the VMpo and VP. This is entirely consistent with tract-tracing studies showing direct projections of VMpo or VP to dpIns, dmIns, and dorsal ACC/aMCC (for a review, see ¹). Our other cortical and amygdalar seeds have multisynaptic connectivity to VMpo and VP by way of aMCC.

Hypothalamus. The hypothalamus is a critical region for allostatic regulation of the body; the paraventricular nucleus in the medial zone is particularly responsible for visceromotor control of autonomic, endocrine, and immune function³. Our review of tract tracing studies clearly indicated connectivity from each cortical and amygdalar seed except with dpIns, despite

evidence from tract-tracing, and with lvaIns to the hypothalamus (which was not predicted)⁴⁻¹⁰.

The lack of functional connectivity between lvaIns and the hypothalamus is not surprising because the lateral portion of the ventral anterior insula is part of a sensory integration network in orbitofrontal cortex in monkeys and humans^{4,10} that has little connection to the hypothalamus, except for light connections to the posterior lateral hypothalamus at the level of the mammillary bodies⁴. We were unable to identify the expected connectivity between the dAmy and any part of the hypothalamus.

Cerebellum. The cerebellum is a key structure in sensorimotor regulation because it sends efferent copies (i.e., predictions) to the cortex to help the brain distinguish between the anticipated sensory consequences of self-initiated by the body vs. those that are unexpected^{11,12}. All cortical seeds and the dAmy seed exhibited connectivity with the cerebellum. More specifically, all seeds exhibited connectivity to lobules IV, V, VI and VIIIB, consistent with the cerebellar “somatosensory” network. The default mode portion of the interoceptive system is additionally connected to lobule IX and Crus I, whereas the salience portion of system is additionally connected to lobule VIIIA (for a specific parcellation of cerebellar intrinsic connectivity, see¹³).

Amygdala. The amygdala is a key subcortical region for both interoceptive input (via its lateral nucleus) and visceromotor control (via its central nucleus)¹⁴, and is part of both the salience and the default mode networks^{15,16}. All of our cortical seeds exhibited connectivity to the amygdala seed except for the pACC seed, which had limited connectivity to the left amygdala (only the dorsal section, which contains the central nucleus). This is consistent with results from tract-tracing studies, which show that each of our cortical seeds projects monosynaptically to the dorsal amygdala^{5,17-21}.

Hippocampus. The hippocampus is a key subcortical hub in the default mode network¹⁶ that is strongly connected to the amygdala (for a review, see ²²). Our fMRI results showed that all cortical seeds and the dAmy seed exhibited connectivity to the entire hippocampus except for the aMCC seed, which exhibited connectivity to only the posterior hippocampus, and the dmIns seed, which exhibited connectivity only to the anterior and posterior hippocampus. Our results are consistent with tract-tracing studies indicating direct projections from the amygdala to the hippocampus and indirect projections from many regions of the cortex to the hippocampus; specifically, vaIns, sgACC, pACC, and aMCC all project to the entorhinal cortex, which projects to the hippocampus (for a review, see ²²). Other cortical regions such as the aMCC and dmIns can connect to the hippocampus in three steps: first to a cortical hub (e.g., vaIns), then to and the entorhinal cortex, then to the hippocampus.

Ventral striatum. The striatum is a subcortical region in the basal ganglia comprising the caudate, putamen, and nucleus accumbens, and its ventral portion is important for controlling inhibitory signals to brainstem visceromotor targets^{23,24}. Specifically, cortical regions send excitatory (glutamatergic) signals to the striatum, which enhance inhibition of brainstem visceromotor targets; these connections also have the capacity to release visceromotor targets from tonic inhibition via striatal connections to the pallidum²⁴. All of our cortical and amygdalar seeds exhibited connectivity to the ventral striatum, except for dmIns, which exhibited connectivity to a portion of the ventral striatum. This is consistent with tract-tracing studies showing monosynaptic connections from each of our seeds to the ventral striatum^{7,25-29}.

Periaqueductal gray (PAG). The PAG is a midbrain nucleus important visceromotor control³⁰. It is difficult to precisely localize using 3-T scanning procedures because it encircles the cerebral aqueduct (e.g., ³¹). Nonetheless, the results of our intrinsic connectivity analysis

largely mirror those for the hypothalamus. The tract-tracing literature has identified monosynaptic connectivity to the PAG from all seeds^{60,121} except the dmIns and dpIns³², and the lvaIns³². All cortical visceromotor seeds demonstrated the expected connections with the PAG: aMCC, pACC, and sgACC. We did not observe the expected connectivity with the mvaIns nor with the dAmy. As expected, connectivity with the lvaIns and dpIns was not observed.

Parabrachial nucleus (PBN). The PBN is a nucleus in the pons that relays interoceptive input from the body to the brain¹ and also serves visceromotor functions³³. All cortical seeds and the dAmy seed exhibited connectivity with the PBN. This is in agreement with the tract-tracing literature showing monosynaptic connectivity to the PBN from each of our seeds^{7,26,27,34,35} except for the aMCC³⁶, which must first project to another cortical hub (e.g., vaIns) before projecting to the PBN.

Nucleus of the solitary tract (NTS). The NTS is a key relay nucleus in the medulla that is on a cranial interoceptive pathway from the viscera to the brain^{12,13,22,23} and also contributes to visceromotor control^{1,33,37-39}. We observed NTS connectivity with the dAmy seed and with all cortical seeds except for the sgACC. This is largely consistent with the tract-tracing literature showing monosynaptic projections to the NTS from each of our seeds^{27,35,40,41}. Failure to observe the sgACC connection is perhaps due to the small size of the sgACC and increased noise due to partial-volume effects of the nearby white matter in the corpus callosum.

Laboratory validation of the allostatic/interoceptive system in humans. The following text details our findings of the association between connectivity in the allostatic/interoceptive system in humans and an index of interoception: the concordance between objective and subjective measures of bodily arousal. We measured sympathetic nervous

system arousal using skin conductance responses (SCRs)⁴² while participants viewed each photo for six seconds. We selected SCRs as an index of sympathetic nervous system activity because, although effects from SCRs specifically might not ascend to interoceptive brain systems, the simultaneous non-SCR effects of sympathetic nervous system activity likely are processed through interoceptive pathways. After each image, participants reported their subjective experience of arousal using a validated self-report scale⁴³. Using multi-level regression procedure to account for the nested and repeated-measures design of this experiment (for a review, see ⁴⁴), we found that the number of SCRs in response a picture (either 0, 1, 2, 3, or 4 SCRs to a given picture) predicted the intensity of arousal experiences in response to the same picture across all participants and pictures ($B = 0.21$, $p < 0.001$), consistent with prior research (e.g., ⁴⁵). Furthermore, as predicted, individuals with stronger intrinsic connectivity between primary interoceptive cortex (dpIns) and the aMCC had a stronger correspondence between sympathetic arousal and subjective experience of arousal than did those with weaker connectivity (regression $B = 0.56$; $p < 0.003$; Supplementary Figure 8). We focused on the aMCC because it was an a priori visceromotor seed region (Table 1), a connector hub⁴⁶, and consistently replicated tract-tracing connectivity to non-cortical allostatic nuclei. For completeness, our results focused on the number of SCRs in response to each image and we did not find analogous results using the amplitude of SCRs in response to each image.

Reconciling prior studies that reported a negative correlation between default mode and salience network activity. Many studies have found that the default mode and salience networks have task-related activity that is negatively correlated (i.e., when the BOLD signal in one network goes up, the BOLD signal in the other network goes down). Such findings are often interpreted as evidence that the brain has either an internal focus on an external focus. This

inverse relationship is often the consequence of removing the mean signal change from all voxels in each volume before proceeding with data analysis (called “global-signal regression”; e.g., ⁴⁷). A more reasonable interpretation, however, may be that when one network increases its neural activity compared to some baseline, the other might show a relative decrease in activity from that baseline (which does not mean that network is irrelevant to the task at hand). Alternatively, one network might show a smaller increase than the other (which, when mean signal change is removed, would appear as a negative relationship between the two networks).

Supplementary Tables

Supplementary Table 1. Functional connectivity between interoceptive and visceromotor regions in humans.

Seed	Target	Connectivity (Pearson's r) Mean $\pm$ SD	t -statistic	p -value ^a
dpIns	aMCC	0.11 $\pm$ 0.16	11.21	3.5 $\times$ 10 ⁻²⁴
dpIns	dAmy	0.15 $\pm$ 0.16	16.02	5.2 $\times$ 10 ⁻⁴¹
dpIns	dmIns	0.17 $\pm$ 0.17	17.34	9.9 $\times$ 10 ⁻⁴⁶
dpIns	mvaIns	0.10 $\pm$ 0.16	10.87	4.6 $\times$ 10 ⁻²³
dpIns	lvaIns	0.14 $\pm$ 0.17	13.33	1.8 $\times$ 10 ⁻³¹
dpIns	pACC	0.06 $\pm$ 0.16	5.81	1.8 $\times$ 10 ⁻⁸
dpIns	sgACC	0.07 $\pm$ 0.15	7.74	2.1 $\times$ 10 ⁻¹³
dmIns	aMCC	0.25 $\pm$ 0.17	23.66	1.1 $\times$ 10 ⁻⁶⁷
dmIns	dAmy	0.16 $\pm$ 0.17	15.91	1.3 $\times$ 10 ⁻⁴⁰
dmIns	mvaIns	0.10 $\pm$ 0.16	10.51	7.0 $\times$ 10 ⁻²²
dmIns	lvaIns	0.08 $\pm$ 0.17	7.86	9.5 $\times$ 10 ⁻¹⁴
dmIns	pACC	-0.02 $\pm$ 0.19	-1.42	1.6 $\times$ 10 ⁻¹
dmIns	sgACC	-0.09 $\pm$ 0.17	-8.36	3.4 $\times$ 10 ⁻¹⁵
Note: ^a p-values from one-sample two-tailed t-test to assess whether connectivity is non-zero, $N = 280$ (discovery sample).				

Supplementary Table 2. Correspondence between networks in the interoceptive system and established resting state networks in humans.

Percent of networks 1 and 2 in each established resting state network			
		Default mode network	Salience network
Network 1	Discovery	63%	15%
	Replication	75%	15%
Network 2	Discovery	7%	60%
	Replication	6%	57%
Percent of each established resting state network in networks 1 and 2			
		Default mode network	Salience network
Network 1	Discovery	77%	14%
	Replication	86%	14%
Network 2	Discovery	9%	65%
	Replication	9%	64%
<i>Note:</i> The percent in each cell indicates the fraction of the brain map in the row that overlaps with the brain map in the column. The top part of the table shows that the majority of Network 1 overlaps with the default mode network and the majority of Network 2 overlaps with the salience network. The bottom part of the table shows that the majority of the default mode network overlaps with Network 1 and the majority of the salience network overlaps with Network 2. Calculations of networks 1 and 2 were performed at the threshold shown in Fig. 3 ($p < 10^{-5}$ uncorrected using $N = 280$). The default mode and the salience networks were defined based on default mode and ventral attention networks from Yeo, et al.⁴⁸ at a threshold of $z(r) > 0.05$ where z is the Fisher's z transformation.			

Supplementary Table 3. Anatomical connections from the interoceptive system to visceromotor control regions (hypothalamus or PAG).

	Cortical region	To hypothalamus	To PAG
Default Mode Network Only	From ventromedial prefrontal cortex (BA 12)	Yes	Yes
	From medial orbitofrontal cortex (BA 11, 13)	Yes	No
	From dorsomedial prefrontal cortex (BA 9, 10)	Yes	Yes
	From posterior cingulate cortex (BA 23, 31)	No	BA 23ab but not BA 31
	From ventral precuneus (BA 7)	No	No
	From angular gyrus (BA 39)	No	No
	From middle frontal gyrus (BA 46)	Yes	Yes
Salience Network Only	From dorsal precuneus (BA 5)	No	No
	From medial frontal gyrus (BA 6)	Yes	No
	From fusiform gyrus (BA 37)	No	No
	From middle temporal gyrus (BA 21)	No	Yes
	From precentral gyrus (BA 6)	Yes	Yes
Hubs Connecting Networks Default Mode and Salience Networks	From midcingulate cortex (BA 31)	Yes	Yes
	From medial postcentral gyrus (BA 6)	No	No
	From parahippocampal gyrus (BA 30)	No	No
	From inferior temporal gyrus (BA 36)	No	No
	From cuneus (BA 30)	No	No
	From superior temporal sulcus (BA 13, 22)	No	Yes
	From temporal pole (BA 38)	No	No
	From inferior frontal gyrus (BA 47)	Yes	Yes
<i>Note:</i> Each cell shows the citation number for the paper indicating tract-tracing results in monkey (or sometimes in rat ⁴⁹). Hypothalamus projections obtained from references ^{4,49}; PAG projections obtained from references ^{30,32,50,51}. We only assessed projections from cortical regions to the hypothalamus or PAG (not from hypothalamus or PAG to cortical) because we wanted to assess if these cortical regions support visceromotor control. PAG = periaqueductal gray.			

Supplementary Table 4. Connector hubs within the interoceptive system.

Cluster	Volume (mm ³)	MNI Center of Mass (mm)			Brodmann Area	Anatomic Name	Rich Club Hub	Visceromotor
		X	Y	Z				
1	3712	45	-21	11	Near BA 13, 41	Dorsal posterior insula	No	Yes (e.g., hypothalamus in rat ⁸ , PBN in rat ³⁴)
2	2136	24	17	-17	Near BA 47	Ventral anterior insula Inferior frontal gyrus	Yes	Yes (e.g., hypothalamus ⁴ , PAG ³²)
3	1224	49	28	-4	Near BA 47	Inferior frontal gyrus	No	Yes (e.g., PAG ³²)
4	952	28	-4	-35	Near BA 36	Inferior temporal gyrus	No	Apparently no
5	648	19	-46	3	Near BA 30	Parahippocampal gyrus	Yes	Apparently no
6	536	4	-2	36	BA 24	Anterior midcingulate cortex Posterior midcingulate cortex	Yes	Yes (e.g., hypothalamus ⁴ , PAG ³²)
7	368	53	-41	6	Near BA 22	Superior temporal sulcus	No	Yes (e.g., PAG ⁵¹)
8	320	53	-45	20	BA 13	Superior temporal sulcus	No	Yes (e.g., PAG ³²)
9	304	54	15	-13	BA 22	Superior temporal sulcus	Yes	Yes (e.g., PAG ⁵¹)
10	264	38	7	-39	BA 38	Temporal pole	No	Apparently no
11	256	11	-63	16	Near BA 30	Cuneus Medial occipital	Yes	Apparently no
12	208	7	-23	49	Near BA 31, 6	Midcingulate cortex Postcentral gyrus	Yes	Yes (e.g., hypothalamus ⁴ , PAG ³²)

Note. This table shows clusters from the overlap of two networks displayed in Fig. 3. One of the superior temporal sulcus clusters (7, 8, and 9) is not easily visible in Fig. 3. Clusters are ordered from largest to smallest. “Near” = within 3 mm. To assess whether a cluster was visceromotor, we reviewed the monkey tract-tracing literature for evidence of a monosynaptic anatomical projection to the following subcortical and brainstem visceromotor regions: hypothalamus, periaqueductal gray, parabrachial nucleus, ventral striatum, and nucleus of the solitary tract. We indicated “Yes” if we found evidence of projections to any of those subcortical and brainstem regions and we indicated “No” if we found no evidence of any of those projections using references ^{4,30,32,34,51,52}.

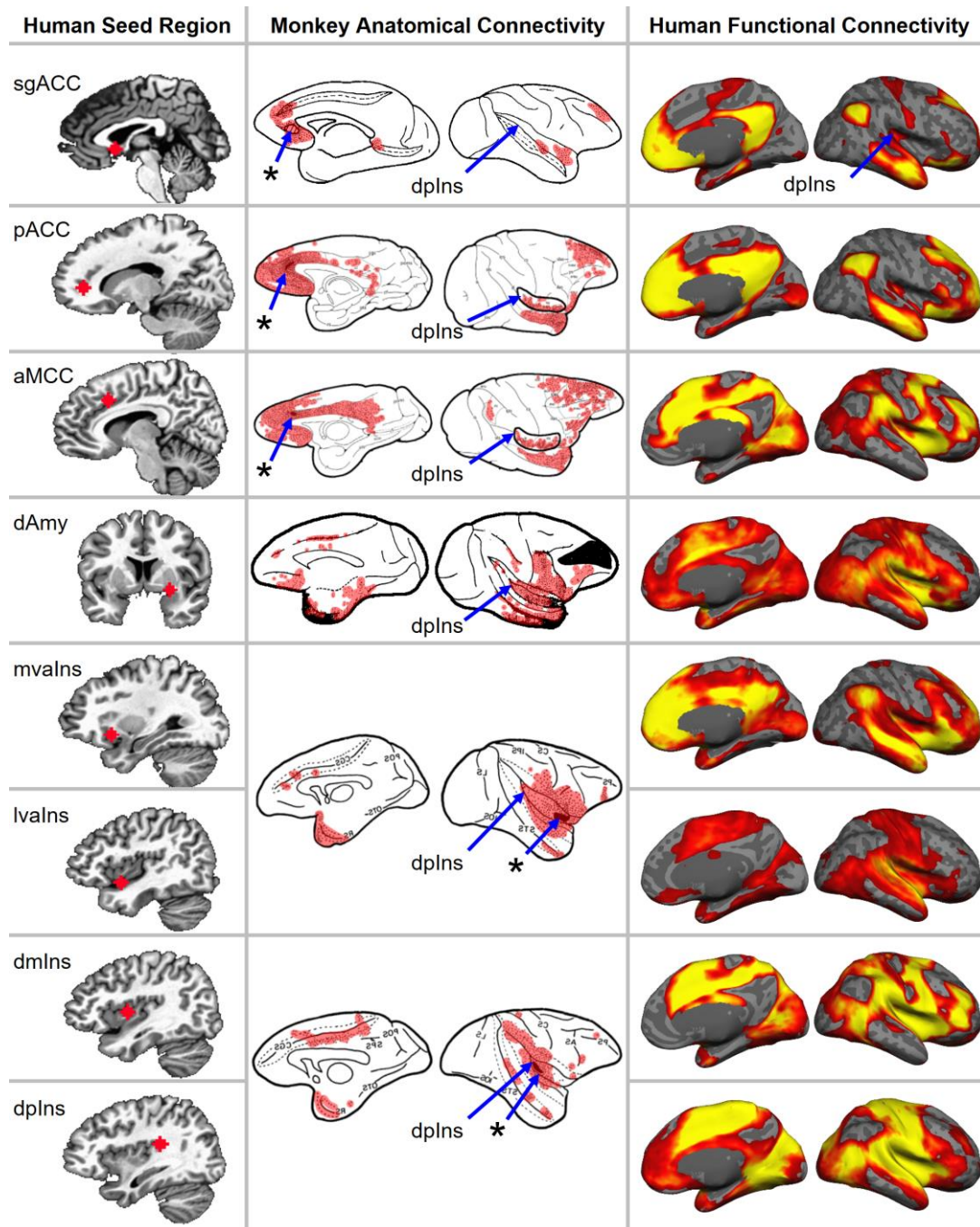
NTS = nucleus of the solitary tract; PAG = periaqueductal gray; PB = parabrachial nucleus; VS = ventral striatum.

Supplementary Table 5. International affective picture system (IAPS) images used in the evocative pictures task.

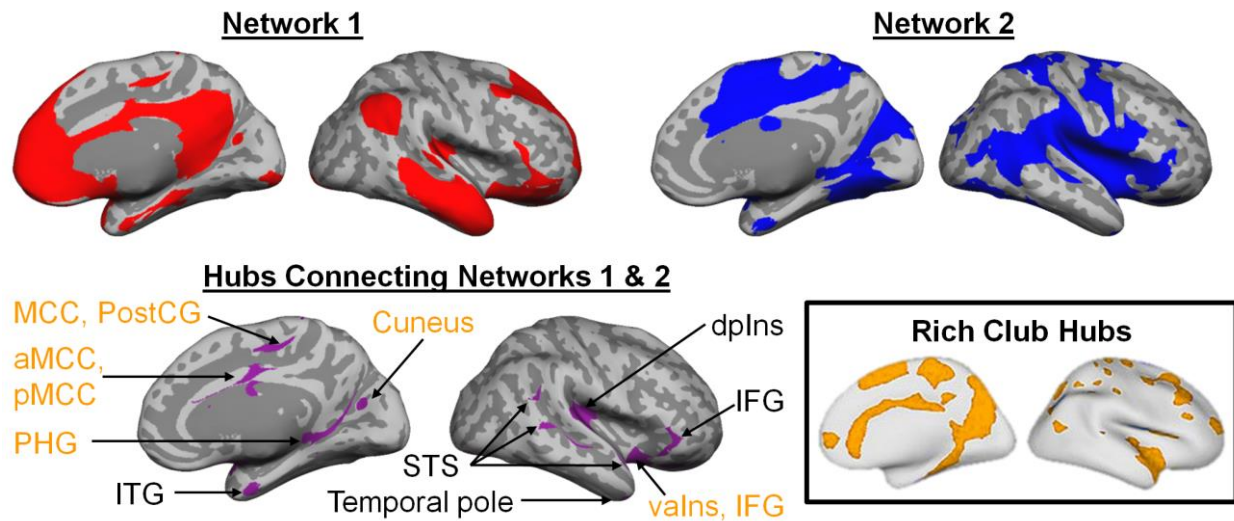
Unpleasant High arousal	Unpleasant Low arousal	Neutral Low arousal	Pleasant High arousal	Pleasant Low arousal
3001	2039	1122	4660	1441
3010	2101	2038	4664	1601
3053	2104	2214	4668	1604
3080	2221	2377	4698	1610
3102	2271	2381	5621	1620
3110	2525	2385	5629	2299
3120	2722	2411	5950	2304
3266	4233	2485	8030	2370
3530	5130	2487	8080	2388
6313	6010	2495	8163	2530
6520	7011	2514	8179	2540
9163	7060	2518	8180	2560
9412	7078	2575	8186	2598
9413	7234	2870	8370	5891
9635	7290	5395	8492	7325
		5510		
		5531		
		5731		
		6150		
		7000		
		7003		
		7004		
		7021		
		7056		
		7077		
		7080		
		7207		
		7503		
		7550		
		7700		

Note. Pictures are ordered by IAPS number. The experimental picture order was block-randomized (see Methods: Procedure). Normative ratings for IAPS pictures have been reported elsewhere⁵³.

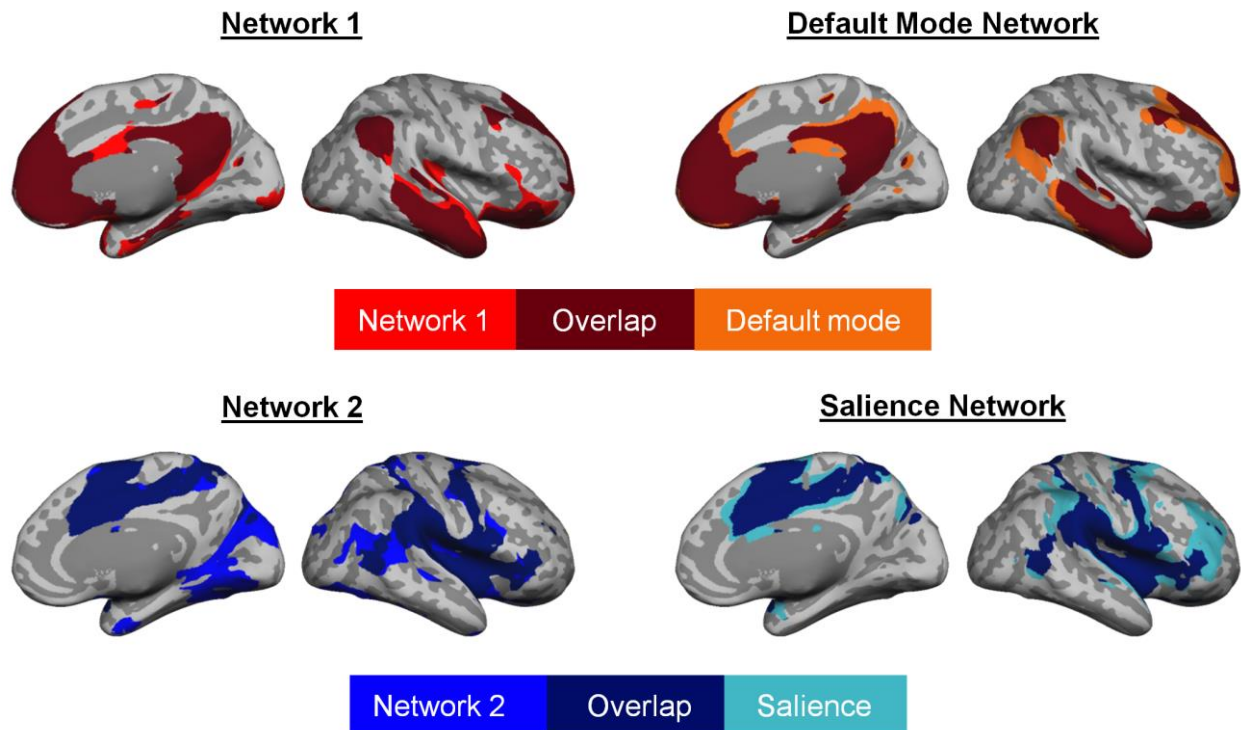
Supplementary Figures



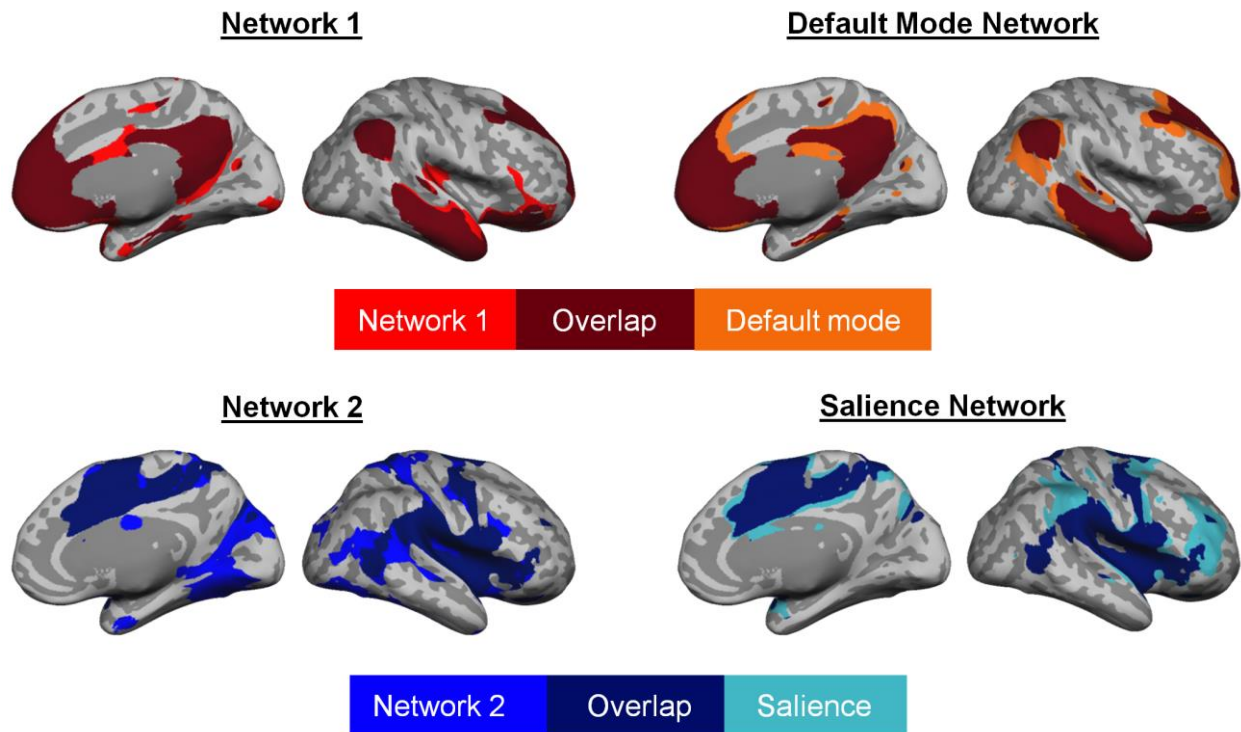
Supplementary Figure 1. Replication of the interoceptive system connecting the cortical and amygdalar visceromotor regions and primary interoceptive regions shown in Fig. 2 using the replication sample ($N = 270$). The color map ranges from $p = 10^{-5}$ in red to $p = 10^{-40}$ in yellow, uncorrected and on a log scale.



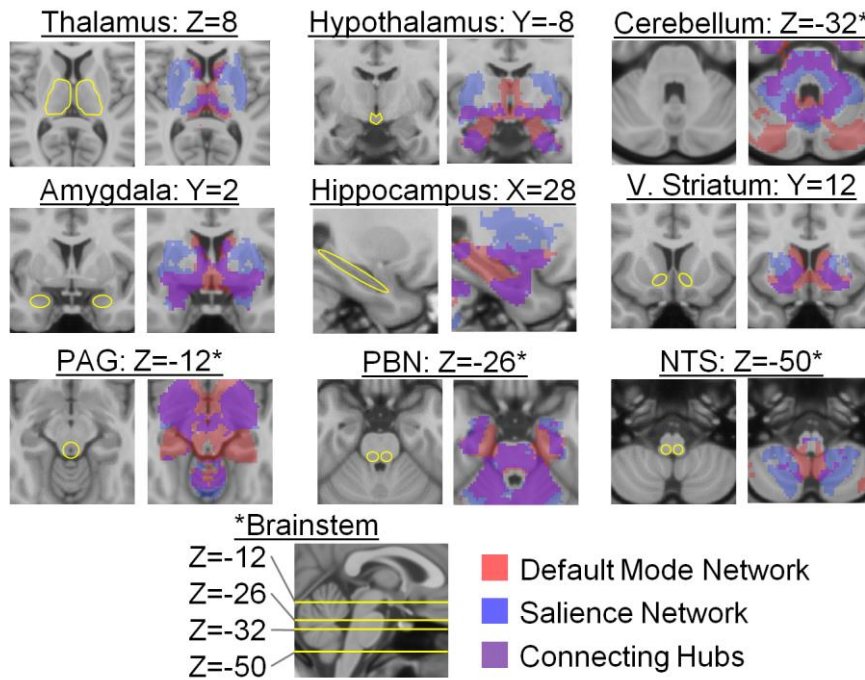
Supplementary Figure 2. Replication of the interoceptive system shown in Fig. 3 using the replication sample ($N = 270$). Intrinsic connectivity maps binarized at $p < 10^{-5}$ uncorrected. Rich club hubs figure adapted with permission from van den Heuvel & Sporns (2013)⁵⁴.



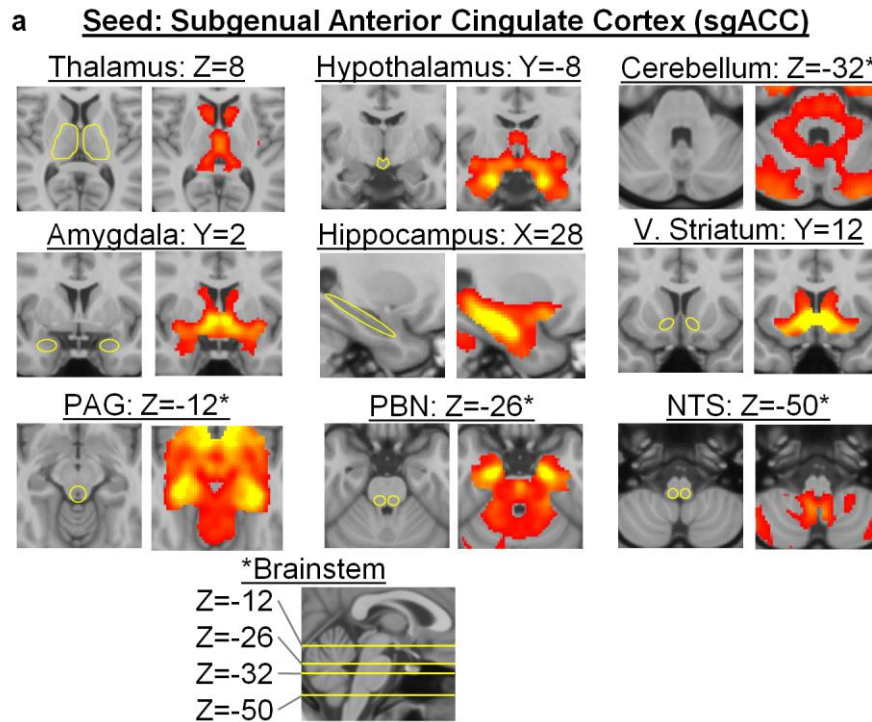
Supplementary Figure 3. Correspondence between allostatic/interoceptive system networks (binarized at $p < 10^{-5}$ uncorrected) and established default mode and salience networks in the discovery sample ($N = 280$). The masks of the established networks were computed based on Yeo et al. (2011)⁴⁸ using a traditional volume-based approach^{55,56} on a sample of 150 subjects (see Methods for description of subjects). The correspondence between the interoceptive system networks and established networks was replicated in a second sample of $N = 270$ (Supplementary Figure 4).



Supplementary Figure 4. Replication of correspondence between interoceptive system networks (binarized at $p < 10^{-5}$ uncorrected) and established default mode and salience networks shown in Supplementary Figure 3 using the replication sample ($N = 270$). The masks of the established networks were computed based on Yeo et al.⁴⁸ using a traditional volume-based approach^{55,56} on a sample of 150 subjects (see Methods for description of subjects).

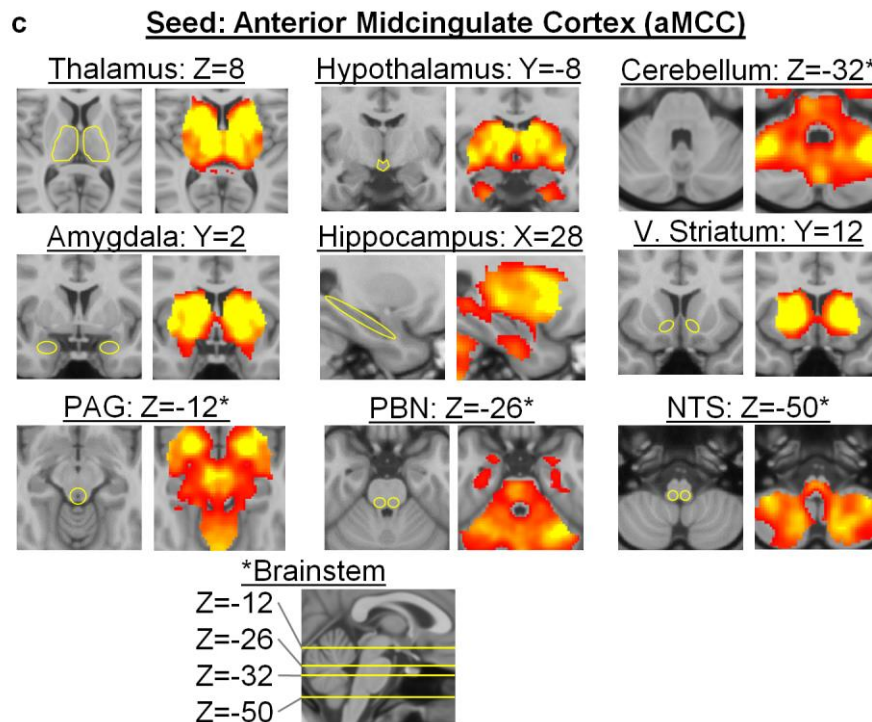
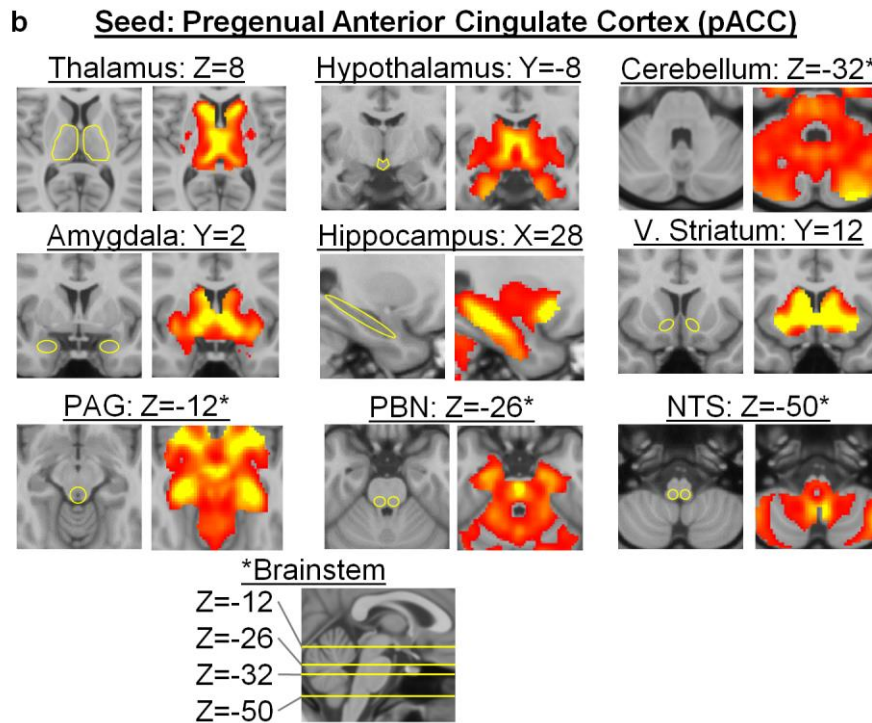


Supplementary Figure 5. Replication of subcortical connectivity of the two intrinsic networks within the interoceptive system in Fig. 4 using the replication sample ($N = 270$; $p < 10^{-5}$ uncorrected, one-sample two-tailed t-test). PAG = periaqueductal gray; PBN = parabrachial nucleus; V. Striatum = ventral striatum; NTS = nucleus of the solitary tract.

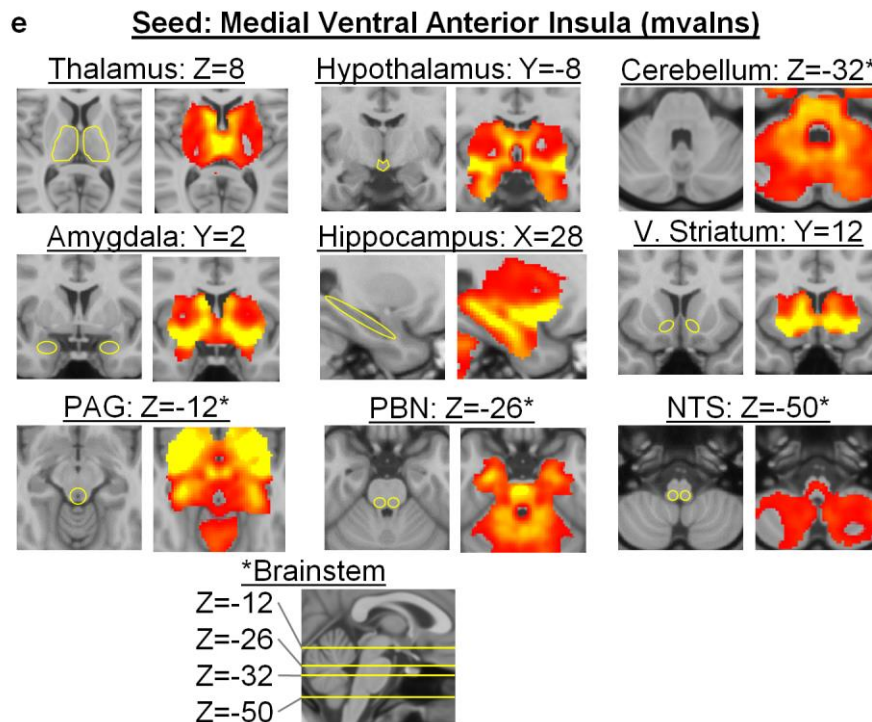
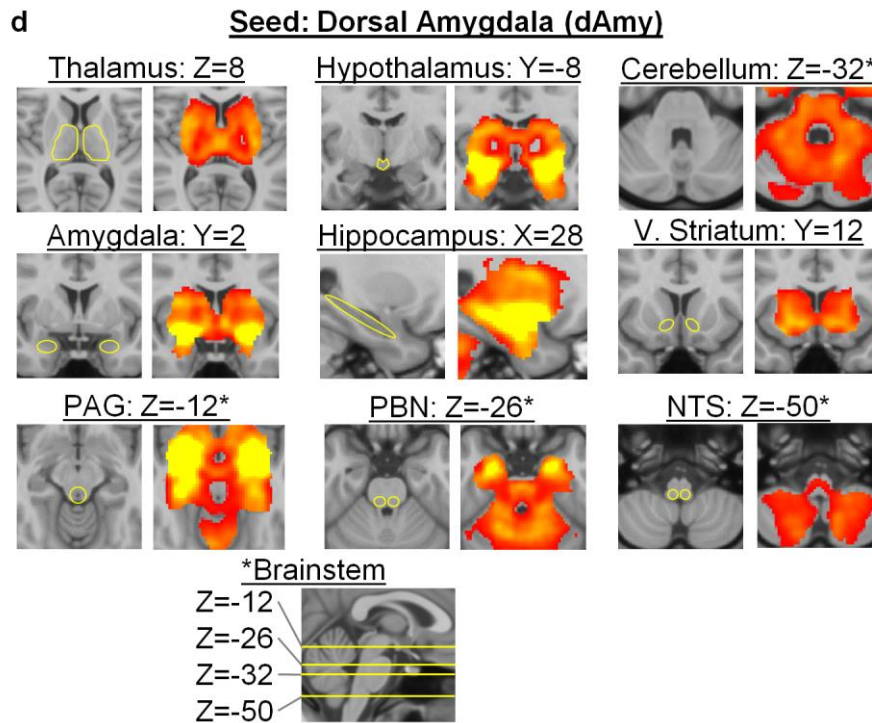


Supplementary Figure 6. Subcortical connectivity of each of the eight cortical and amygdalar seeds used to delineate the interoceptive system using the discovery sample ($N = 280$). The color scale ranges from $p = 0.05$ in red to $p = 10^{-50}$ in yellow, uncorrected and on a log scale. (a) Seed at subgenual anterior cingulate cortex (sgACC). (b) Seed at pregenual anterior cingulate cortex (pACC). (c) Seed at anterior midcingulate cortex (amMCC). (d) Seed at dorsal amygdala (dAmy). (e) Seed at medial ventral anterior insula (mvaIns). (f) Seed at lateral ventral anterior insula (lvaIns). (g) Seed at dorsal mid insula (dmIns). (h) Seed at dorsal posterior insula (dpIns). PAG = periaqueductal gray; PBN = parabrachial nucleus; V. Striatum = ventral striatum; NTS = nucleus of the solitary tract.

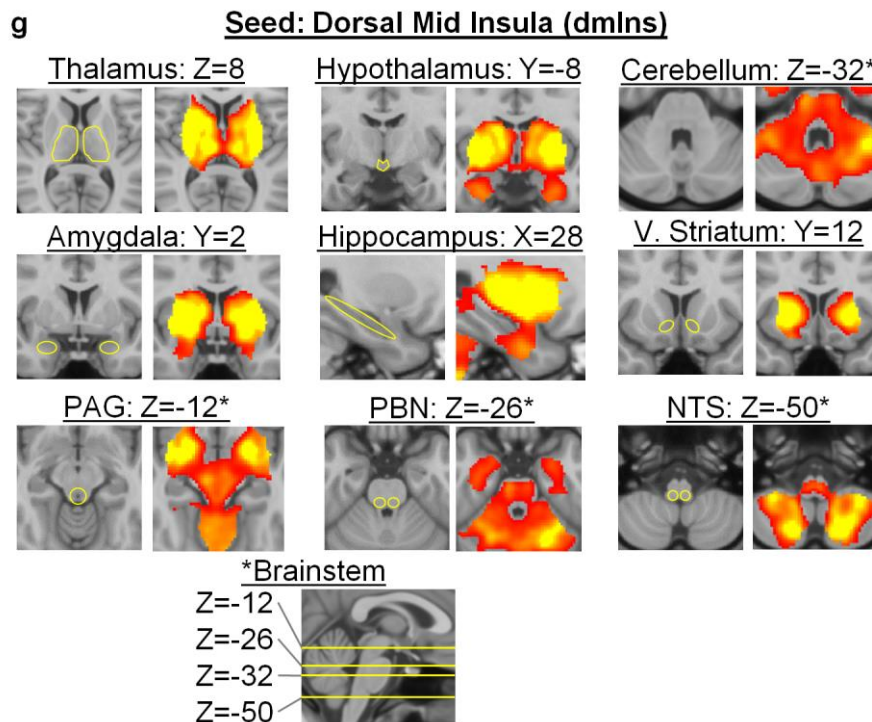
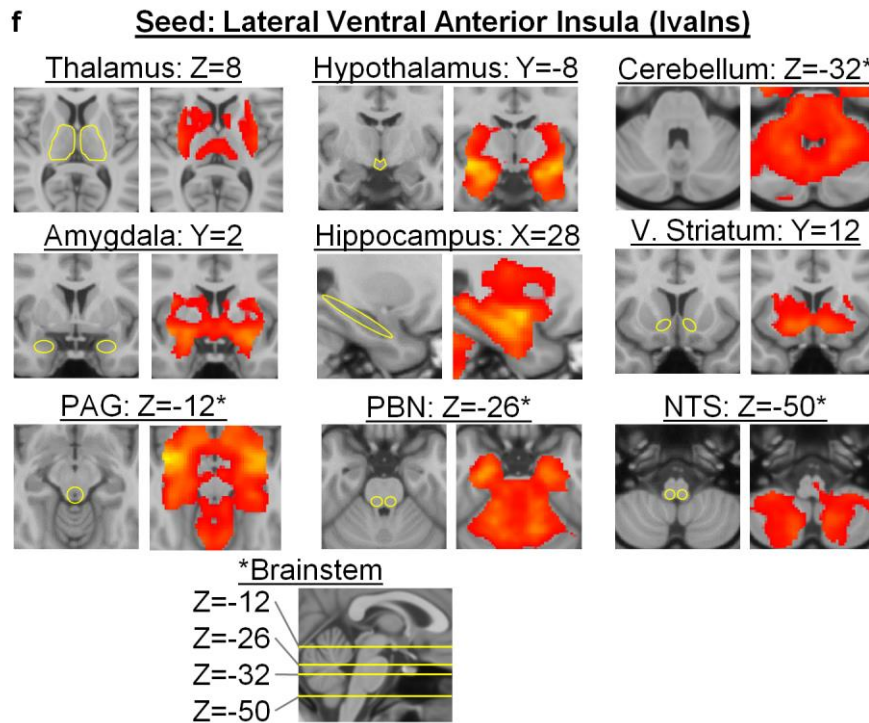
Supplementary Figure 6 continued



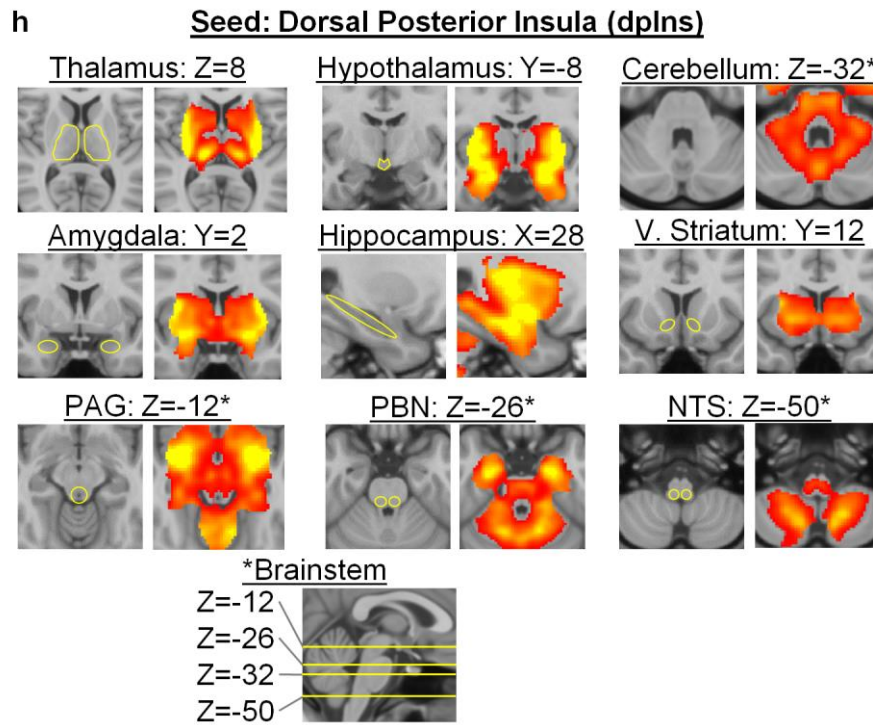
Supplementary Figure 6 continued

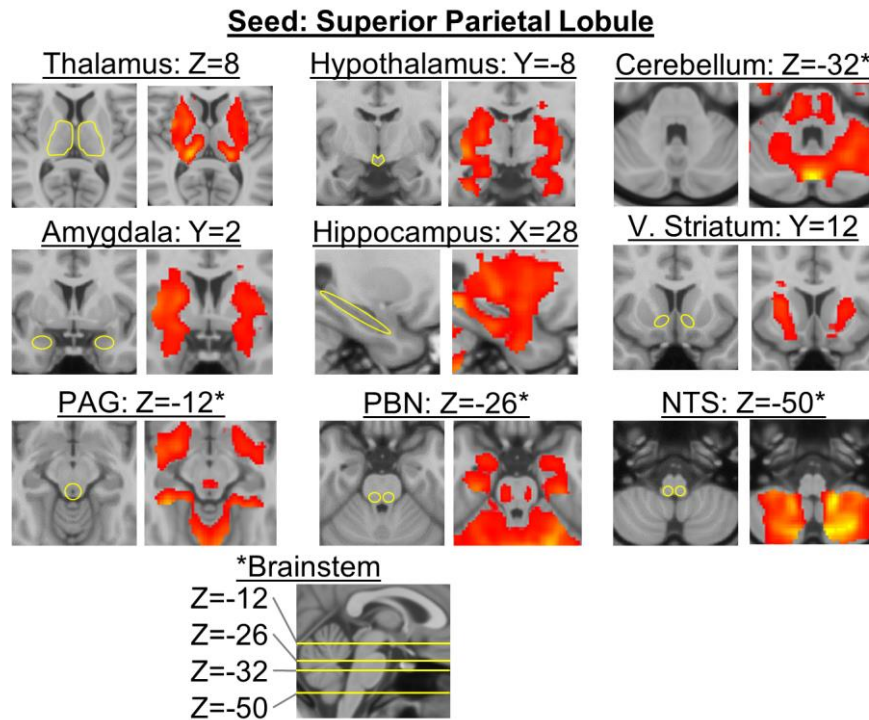


Supplementary Figure 6 continued

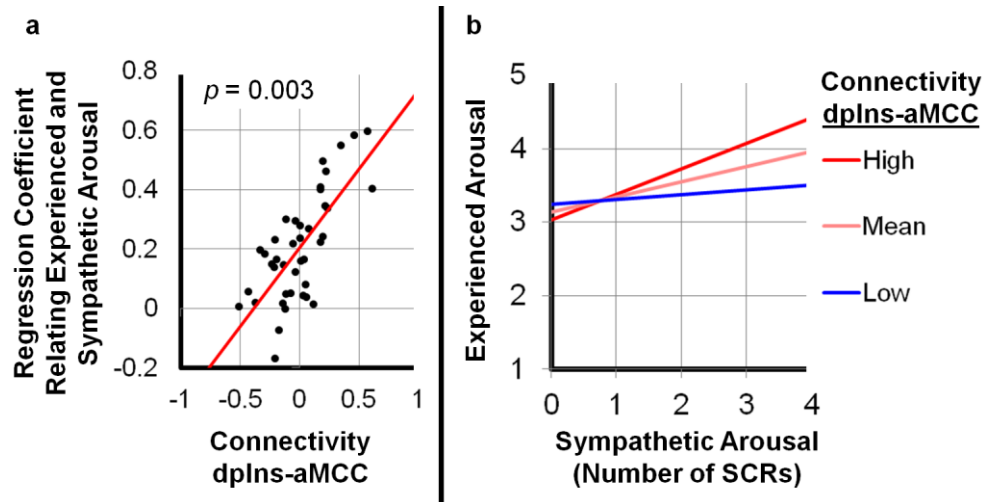


Supplementary Figure 6 continued





Supplementary Figure 7. Subcortical connectivity of the superior parietal lobule does not involve non-cortical targets important for allostasis (hypothalamus, PAG, PBN, NTS), suggesting a degree of specificity in the functional connections between cortical regions in the allostatic/interoceptive system and subcortical regions that support allostasis. The seed coordinates are MNI X, Y, Z = 46, 12, 28. These results are from the discovery sample ($N = 280$). Identical conclusions were obtained using results for the replication sample ($N = 270$). The color scale ranges from $p = 0.05$ in red to $p = 10^{-50}$ in yellow, uncorrected and on a log scale. PAG = periaqueductal gray; PBN = parabrachial nucleus; V. Striatum = ventral striatum; NTS = nucleus of the solitary tract.



Supplementary Figure 8. The strength of intrinsic connectivity within the allostatic/interoceptive system predicts the degree of interoception during allostatic fluctuations. (a) Each data point corresponds to one participant ($N = 41$). Interoceptive ability is represented on the y-axis as the unstandardized linear regression coefficient B for the relationship between sympathetic nervous system arousal (i.e., the number of skin conductance responses) and the intensity of experienced arousal (rated on a 1 to 5 scale) while participants viewed evocative pictures. The strength of intrinsic connectivity between a primary interoceptive region (dorsal posterior insula, or dpIns) and a key visceromotor control region (the anterior portion of the midcingulate cortex, or aMCC) is represented on the x-axis as the Fisher's z transformation of the Pearson's correlation coefficient of the two BOLD time courses (the red line shows the regression line with coefficient $B = 0.56$; $p = 0.003$; multi-level regression analysis). (b) The relationship between experienced arousal and the number of skin conductance responses (as an index of sympathetic nervous system arousal) is moderated by intrinsic connectivity strength. "High" or "Low" corresponds to one SD above or below the mean connectivity across participants, respectively (mean 0.1, SD 0.23). dpIns = dorsal posterior insula; aMCC = anterior midcingulate cortex; SCRs = skin conductance responses.

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