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Regional protein expression in human Alzheimer's brain correlates with disease severity

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1 **Abstract**

2 Alzheimer's disease (AD) is a progressive neurodegenerative disorder that currently affects 36 million
 3 people worldwide with no effective treatment available. Development of AD follows a distinctive
 4 pattern in the brain and is poorly modelled in animals. Therefore, it is vital to widen both the spatial
 5 scope of the study of AD and prioritise the study of human brains. Here we show that functionally
 6 distinct human brain regions show varying and region-specific changes in protein expression. These
 7 changes provide novel insights into the progression of disease, novel AD-related pathways, the
 8 presence of a 'gradient' of protein expression change from less to more affected regions, and the
 9 presence of a 'protective' protein expression profile in the cerebellum. This spatial proteomics
 10 analysis provides a framework which can underpin current research and opens new avenues of interest
 11 to enhance our understanding of molecular pathophysiology of AD, provides new targets for
 12 intervention and broadens the conceptual frameworks for future AD research.

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1 Alzheimer's disease (AD) is a multifactorial neurodegenerative disorder characterized by progressive
2 dementia^{1,2}. Accumulation of A β peptide and microtubule-associated protein tau, which exhibits
3 hyperphosphorylation, and oxidative modifications into so-called 'plaques' and 'tangles' are
4 considered to be central to the pathology of AD³. Other prominent features of AD include early
5 region-specific decline in glucose utilisation and mitochondrial dysfunction and consequently
6 depleted ATP production and increased reactive oxygen species production in neurons⁴.
7 Excitotoxicity in the AD brain arising from altered glutamatergic signalling⁵, and dysregulation in
8 other neurotransmitters has also been documented, including abnormalities of adrenergic, serotonergic
9 and dopaminergic neurotransmission⁶. In response to pathological stimuli associated with AD,
10 inflammatory events mediated through both innate and cell-mediated immune mechanisms are also
11 present³.

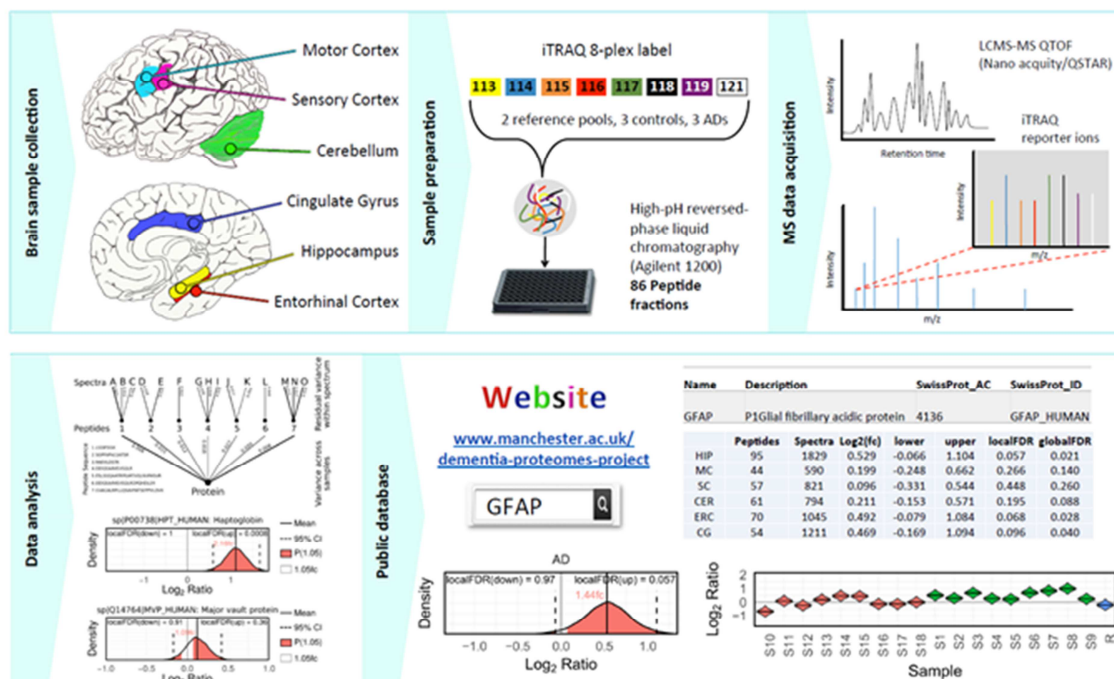
12 Despite an increase in research into the underlying pathology of AD over the last decade, there
13 remains controversy around what underpins this disease process, which in turn affects the pipeline of
14 new disease modifying agents. There remains a lack of detailed mechanistic knowledge about what
15 happens in the human brain in AD. This is exacerbated by the fact that different brain regions develop
16 pathology at different times in the disease process, adding a spatial element to the disease which is not
17 captured by work in cell culture models and is often overlooked in human studies, which tend to focus
18 on single regions. Animal models also fail to capture the full disease process, at either the behavioral
19 or biochemical levels⁷, such that translation of both basic biological findings and/or the activity of
20 potential disease-modifying interventions from animals into humans is relatively unsuccessful. While
21 there have been several studies which have focused on the transcriptome in human AD, there is a
22 wealth of evidence that suggests many protein expression changes in biological systems can occur
23 independently of transcript-level regulation, and that studying the proteome can prove new insights on
24 the regulation of functionally active molecules in a given biological or disease state⁸.

25 Mass spectrometry based proteomics has been recognised as a powerful tool with the potential to
26 uncover detailed changes in protein expression⁹. To date, however, there are few studies of protein
27 expression in AD carried out using human brain tissue, and those that exist typically examine a single
28 AD affected brain region^{10,11}, and use different patient cohorts and analytical methods that makes
29 between-region comparisons difficult. Such studies also frequently use either small numbers of
30 samples (n<4) or cohorts poorly matched for age or tissue *post-mortem* delay^{10,12,13}. This study aims to
31 overcome some of these existing limitations by providing a spatially-resolved analysis of protein
32 expression in six regions of human control and AD-affected brain in well matched, short *post-mortem*
33 delay tissue.

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Results

In this study, we analysed six functionally distinct regions of human *post-mortem* brain; hippocampus (HP), entorhinal cortex (ENT), cingulate gyrus (CG), sensory cortex (SCx), motor cortex (MCx), and cerebellum (CB), by mass spectrometry to gain a more comprehensive understanding of protein expression changes within the AD brain. These regions were selected to represent parts of the brain known to be heavily affected (HP, ENT, CG), lightly affected (SCx, MCx) and relatively ‘spared’ (CB) during the disease process. Donors were well matched for age and *post-mortem* delay times were short, with no significant difference between cases and control. Donor data is provided in Supplementary Table 1. Relative protein expression was determined using an isobaric tagging approach followed by 2-dimensional liquid chromatography and mass spectrometry. Peptide-level data were then analysed using a Bayesian model that infers a posterior probability distribution for the relative levels of each protein between ‘cases’ and ‘controls’ based on the underlying relative peptide levels. To promote sharing and usage of these data, we have developed a searchable web interface that hosts all of our results (www.manchester.ac.uk/dementia-proteomes-project; described in Supplementary Information), which also includes Bayesian probability distributions for each protein across all individual brains examined in this study. The complete workflow is illustrated in Figure 1. The complete processed data for each region (at protein identification FDR <1%) can be found in Supplementary Table 2. Raw mass spectral data can be accessed via PRIDE, with initial search outputs prior to Bayesian modelling available via the Open Science Framework at DOI 10.17605/OSF.IO/6BXJQ (Supplementary methods).



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Figure 1. Proteomics workflow. Selected brain regions were pre-dissected prior to storage at -80°C until analysis. Each region was lysed, and protein assigned to an iTRAQ 8plex. Following digestion and labelling, samples were pooled, peptides fractionated by High-pH reverse phase chromatography and fractions analysed by standard LC-MS/MS methods. Peptides were identified and quantified based on their iTRAQ reporter area; relative protein quantification was inferred from these values using a Bayesian model. All data are deposited in a searchable online database.

Each brain region was analysed in isolation, adding strength to our comparison of protein expression changes across multiple regions, since these were identified and quantified independently. Combining all protein identifications (at 1% false discovery rate) across the six experiments yielded a total of 5,825 unique protein identifications across all regions. In our data, 990 proteins were quantified with only one or two spectra in any single region, and were subsequently omitted from our downstream cross-regional comparison in order to retain the proteins with the most precise quantification – optimisation data suggests that when the same sample is split and processed independently, >99% of proteins are defined as not being significantly different above this threshold (data not shown). However, many of these will be quantified correctly (we have previously validated expression changes based on a single spectrum, e.g. p53 in⁸), and as such these data have been included in Supplementary Table 2 and our online database. We thus quantified a total of 4,835 distinct proteins in at least one brain region, among which 3,302 proteins were common to at least three regions, and 1,899 to all six regions (Fig. 2a). These data allow us to a) define protein changes as a result of AD in any given region of the human brain being studied, and b) identify differences in how distinct brain regions are affected in AD, and by extension protein changes which occur in multiple regions of the AD brain.

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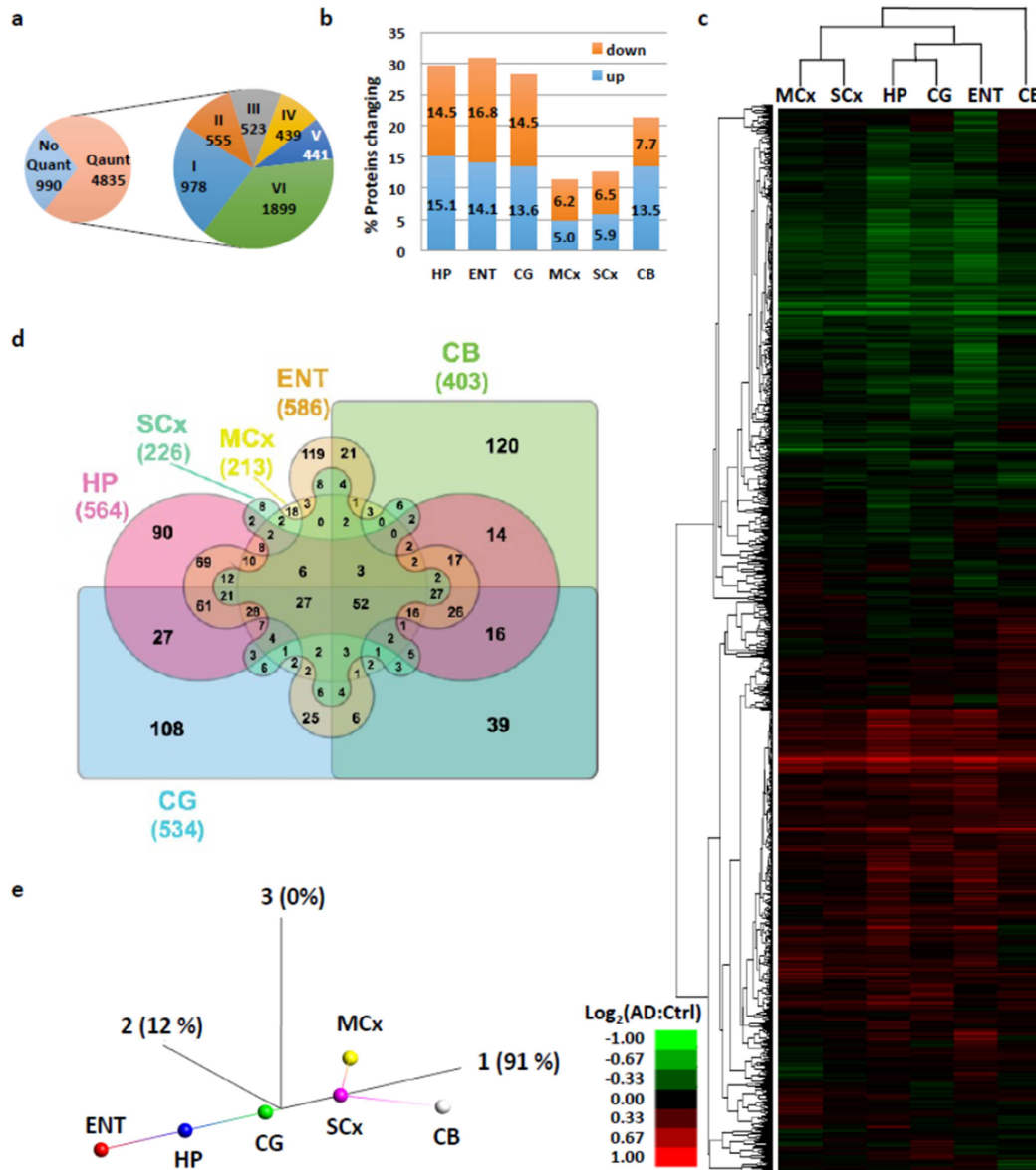


Figure 2. Summary of protein expression data. a) 5,825 proteins were identified, with 990 quantified with only one or two spectra and which were thus omitted from our primary comparative analysis. The remaining 4,835 proteins are classified as to whether they were quantified in six or fewer distinct regions. b) Proportion of identified, quantified proteins showing a change in expression in AD in each of the six regions under study. c) Heat map and dendrogram showing the relationship between protein expression in each region mapped using proteins present in all six regions, with three distinct ‘groups’ based on highly affected (HP, ENT, CG), moderate (MCx, SCx) and spared (CB) clearly visible. d) Edwards-Venn diagram showing the overlap of protein expression changes between brain regions, including only proteins quantified in all regions. e) Isometric mapping (Isomap) representation of protein expression data between brain regions showing a broadly linear relationship from non-affected towards affected regions, with the exception of cerebellum, which shows distinct patterns of protein expression in AD.

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2 Comparison of the total number of proteins whose expression is altered in each region reveals,
3 perhaps unsurprisingly, that the more severely affected areas in AD (HP, ENT, CG) show the largest
4 number of changes in protein expression (~30% of quantified proteins), while less affected regions
5 (MCx, SCx) have fewer changes (11-13%). Strikingly, the CB, which many think to be pathologically
6 ‘unaffected’, shows a substantial number of protein changes (20%; Fig. 2b). This observation
7 accurately recapitulates data from our previous study of the metabolome on these brain samples¹⁴.
8 Unsupervised hierarchical clustering of protein expression changes from all six regions demonstrates
9 that the changes observed in CB are distinct from those seen in the affected HP, CG and ENT (Fig.2c).
10 This is supported by an Edwards-Venn representation of the data which shows that 120/403 (29.8%)
11 of changes in CB are not seen elsewhere (Fig.2d; Supplementary Table 3). While it has long been
12 reported that the CB in AD can contain amyloid plaques¹⁵, it is considered to be relatively ‘spared’ in
13 AD. There is a lack of neurofibrillary tangles in cerebellum¹⁶, and this region does not appear to
14 develop significant neuronal loss, such that this region is often used as a control in imaging studies of
15 the AD brain^{17,18}. However, recent work by Guo *et al.* suggests a distinct pattern of cerebellar atrophy,
16 which spreads from intrinsic connectivity networks within the cerebrum¹⁹, and alterations in
17 cerebellar glucose metabolism have been reported in late stages of the disease^{20,21}. Our data strongly
18 suggest that the CB is heavily affected by AD at the molecular level, at least in late stage disease, and
19 is so to a greater extent than other regions associated with later degeneration such as MCx or SCx,
20 where protein changes were fewer and encompass those seen in the more severely affected regions.
21 That the changes in CB are different from those seen elsewhere in the brain raises the possibility that,
22 rather than being ‘spared’, the CB is affected in a different way to other brain regions and that, given
23 it shows little pathology, these changes may reflect some level of active protection.

24
25 Hereinafter, we refer to HP, ENT and CG as the severely affected, and MCx and SCx as the less
26 affected regions based on the number of significantly altered proteins and pathways observed within
27 this study.

28
29 Unsupervised clustering of brain regions based on their protein expression, by performing a
30 dimensionality reduction on these data using isomeric feature mapping (Isomap), clearly shows this
31 hypothesized ‘evolution’ of the disease from the least affected cortical regions to the most affected,
32 with cerebellum following a distinct pathway from the inception of disease (Figure 2e). This non-
33 linear approach has been shown to be an improvement over the more standard PCA approach for
34 analysis of gene and signalling networks²². These data also further support our previous observation
35 that CB stands out as a single, uniquely affected brain region based on the distinctive patterns of

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changes found here while the other regions line up along the same vector in accordance with disease severity. Previous studies using gene co-expression networks and transcriptomics analysis have demonstrated a pattern where the molecular signatures in less-affected areas of the brain overlap with but are less marked than the grossly affected areas, and have implied that these overlapping changes represent those which occur early in AD-related neurodegeneration²³. Our data at the protein level would support this conclusion - the less affected regions (MCx and SCx) contain very few protein changes which are not seen elsewhere, and a clustering analysis suggests that these regions are simply at an earlier stage down a similar pathway. Therefore, our data shows that by comparing more and less affected brain regions in a multi-regional approach we can observe different stages of the same disease process, enabling identification of early molecular changes, even in patients with late-stage disease.

To probe the differences in AD-related protein expression between brain regions in more mechanistic detail, we performed a pathway enrichment analysis for all differentially expressed proteins for each region. Such analyses enable us to visualise which processes are affected in the AD brain, and also whether two (or more) regions are showing dysregulation in the same pathway even if different subsets of proteins are identified as ‘changing’. These data are summarised in Figs 3a-f (and Supplementary Table 4).

Reflecting the individual protein expression data, HP and CG showed the highest number of biological pathways being affected by AD. The changes in specific molecular pathways were comparable between HP, ENT, and CG. CB, on the other hand, showed altered regulation of a set of molecular pathways with limited overlap with those affected in the other five brain regions, again arguing for the presence of a distinct cellular response to disease in this region.

One of the most consistent features across all brain regions was a significant change in proteins and pathways involved with the innate immune response. In AD, aggregates of A β can trigger both pathogen-associated and initiate immune responses, and a persisting elevation of A β may elicit a chronic reaction of the innate immune system²⁴. In this study, we observed strong evidence for the global activation of the innate immune response, including of the acute-phase response, the complement system (classical and alternative pathways) and the coagulation system, consistent with widespread neuroinflammation, suggesting that this may be a relatively early (prior to atrophy) event in pathogenesis. Previous studies have also implicated complement family proteins as potential AD biomarkers²⁵, and GWAS studies have identified AD risk loci in a number of complement pathway genes²⁶⁻²⁸. It is worthy of note that these studies do not directly inform on the activation state of the

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1 complement pathway, and indeed in our study we see upregulation of SerpinG1, which inhibits
 2 complement C4 cleavage by C1 and MASP2, as well as increased levels of C4, C3 and various
 3 regulators in AD. While it is highly likely that dysregulation of this pathway plays a role in AD, the
 4 precise nature of this role remains to be determined. Overall, HP, ENT and CG showed substantive
 5 evidence for a broader spectrum of changes in immune responses compared to MCx, SCx and CB.
 6 These included specific cellular pathways including granulocyte adhesion and dendritic cell
 7 maturation (Fig. 3a–f, Supplementary Data Table 4 and 5), implying that the innate immune system
 8 becomes activated early, and that the adaptive immune response plays a role later in the disease
 9 process. However the interplay between these two systems is complex and it is yet to be determined if
 10 these changes are a cause, or a consequence of other aspects of AD pathogenesis²⁹.

11

12 This pathway-level analysis also identified signaling pathways involved in apoptosis and cell cycle
 13 regulation as being widely dysregulated in severely affected regions of AD brain, including the
 14 HIPPO, ERK/MAPK, PI3K/AKT, and Wnt/ β -catenin pathways (Fig.3a, b, d), all known to be
 15 critically involved in regulation of apoptosis and the cell cycle. Reduced abundance of proteins
 16 involved in Polo-Like Kinase signaling and G2/M DNA Damage Checkpoint Regulation are likely a
 17 cause of impaired cell cycle regulation, marking these pathways out as potentially key contributors to
 18 neuronal cell death in AD. Strikingly, less affected regions SCx and MCx do not show large changes
 19 in these pathways (Fig.3e, f), reflecting reduced levels of apoptosis seen in these areas. The only
 20 exceptions are the G2/M checkpoint and the Hippo pathway, whose members are significantly
 21 decreased in these regions, suggesting that inactivation of this key developmental pathway, possibly
 22 via the observed upregulation of CD44³⁰, or altered regulation of associated proteins such as the
 23 synaptic scaffolding proteins DLG2, DLG3, and DLG4, all of which are downregulated, is an early
 24 event in AD development. In CB, only granzyme A signaling was identified as an apoptosis-related
 25 pathway, indicative of fewer cell death signals in this region.

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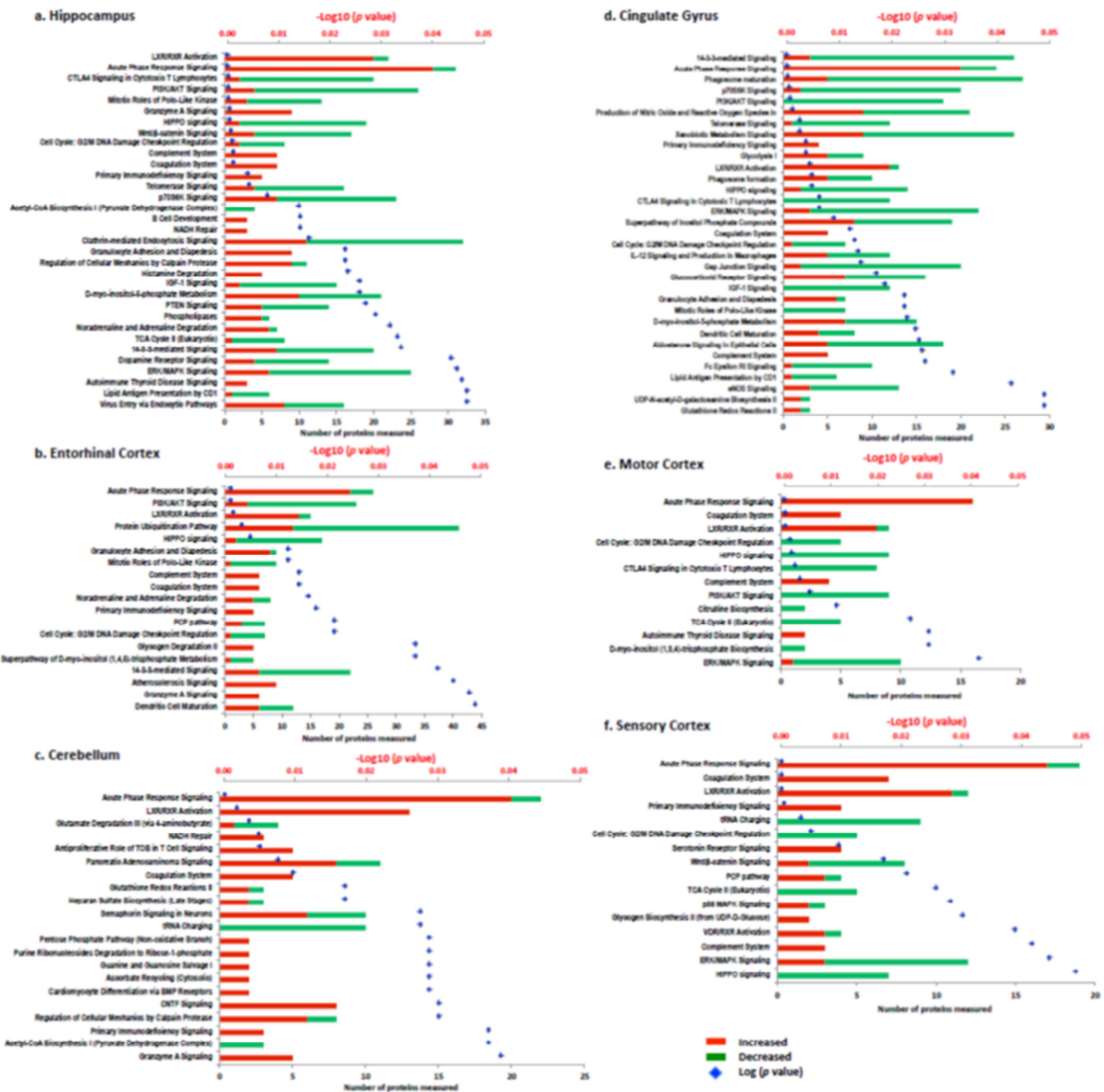


Figure 3. Network analysis summary. Alterations of molecular pathways in human Alzheimer's disease brain across six distinct regions, namely a) Hippocampus, b) Entorhinal cortex, c) Cerebellum, d) Cingulate gyrus, e) Motor cortex, and f) Sensory cortex. In each plot, the numbers of increased and decreased proteins are indicated by the red/green bars, while the blue spots indicate the $\log_{10}(p\text{-value})$ for each pathway.

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We also observed both global and regional metabolic impairments in the AD brain. Defects in brain metabolism and energetics are central to the pathogenesis of AD as evidence by epidemiological, neuropathological, and functional neuroimaging studies³¹. The AD brain characteristically exhibits defective cerebral perfusion³² and glucose uptake³³, which is believed to underlie hypometabolism and cognitive decline³⁴. Alterations in pathways of monosaccharide/glucose metabolism are highly significant in severely affected brain regions and CB (Fig.3a – f, Supplementary Data Table 4), consistent with our previous finding of elevated free glucose levels in AD brain²¹. TCA enzyme abundance was generally decreased in all regions of AD brain, going some way to explaining the previously observed shift from primarily aerobic glycolysis (i.e. glycolysis followed by complete oxidation in mitochondria) to the ketogenic/fatty acid β -oxidation pathway, with impaired mitochondrial bioenergetics³⁵. Severely affected brain regions also showed substantial alterations in signals related to altered regulation of neurotransmitters/hormones (noradrenaline/adrenaline, dopamine, and aldosterone) that were not observed in less affected regions. While this might suggest that altered neurotransmitter biology is a late or downstream process in pathogenesis, it is notable that the enzymes in a key upstream pathway of neurotransmitter production which results in the production of tetrahydrobiopterin (BH4), a precursor of dopamine, noradrenaline and serotonin, is significantly upregulated in all regions studied. Previous work has suggested a decrease in BH4 levels in AD brain³⁶ and the observations at the protein level may reflect a feedback loop where the cell is responding to decreased BH4. The presence of this dysregulation early in disease suggests it is a target which deserves closer attention.

While comparison of affected regions yields a range of interesting and novel observations about the molecular underpinning of AD, the presence of a large number of changes in ‘unaffected’ cerebellum provides a surprising finding, even more so when one observes that these changes are distinct from those manifest elsewhere. To investigate this population of protein changes further, we analysed proteins uniquely affected in CB using both DAVID and STRING. These analyses supported our earlier global pathway analysis in demonstrating that CB additionally showed alteration in Semaphorin and ciliary neurotrophic factor (CNTF) pathway members which play important roles in neuronal survival and neurodevelopment/neuronal regeneration (Fig.3c and Fig.4a, b). SEMA7A, shown here to be upregulated in CB of AD brains, is known to be involved in repair of the glial scar following spinal cord injury and to play a role in the development of multiple sclerosis, but has not previously been linked to the disease process in AD³⁷. CB also showed a significant reduction in levels of both nuclear and mitochondrial aminoacyl-tRNA synthetases. In CB, significantly depleted aminoacyl tRNA synthetases, including those encoded in the mitochondrial genome as well as those from the nuclear genome (Fig.3c and Supplementary Data Table 3), could disrupt translational fidelity, leading to accumulation of misfolded proteins³⁸. However, these proteins are multifunctional.

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For example, Ishimura *et al.* have shown that misregulated tRNA processing can lead to neurodegeneration³⁹, and tRNA synthetases have also been shown to be mediators of inflammation⁴⁰ thus downregulating these proteins may confer some level of protection. This finding could also provide a supportive mechanism for the hypothesis that ribosomal dysfunction is an early event in AD⁴¹. Taken together with its known roles in inflammation and signaling, and in several other neurodegenerative disorders⁴², our data suggest that the role of tRNA synthetases in Alzheimer's disease is worthy of significant further investigation.

One of the most distinct changes observed in this CB-specific analysis was that a much greater number of proteins of electron transport chain (ETC) complex 1 were consistently more reduced in abundance (Fig.4b, c Supplementary Data Table 5) than was found in other areas. Furthermore, CB showed increases in oxidative defense proteins involved in glutathione redox reactions and ascorbate recycling (Fig.3c). These data provide strong additional evidence for a protective mechanism in CB that decreases ROS-production by ETC while simultaneously increasing ROS defenses. Another interesting observation in CB was the activation of a Purine Ribonucleosides Degradation pathway, which could not only contribute substrate to the pentose phosphate pathway, but also participate in guanine/guanosine production in this brain region. Combined with the observed activation of Guanine and Guanosine Salvage I pathway, and an increase in guanosine level in CB as previously reported by our metabolomics analysis¹⁴, these changes may also confer a previously unknown neuroprotective effect in this brain region⁴³.

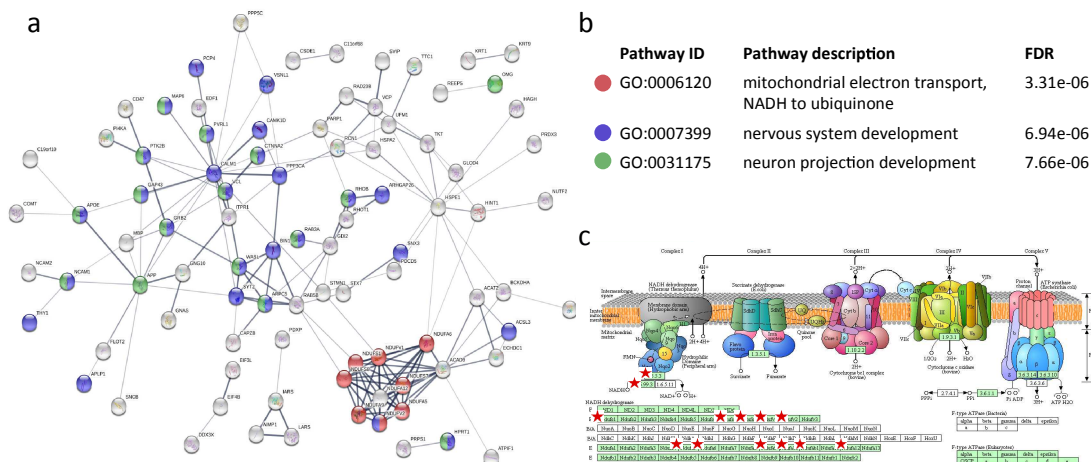


Figure 4. CB-specific biological processes in AD brain. a) 120 proteins that showed CB-specific alterations were enriched for molecular processes in STRING using default setting. Each node represents a protein, and proteins involved in b) significantly enriched pathways were highlighted. c) Dysregulation of the mitochondrial electron transport chain was highlighted by pathway analysis, and

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proteins affected mapped (red star) into the NADH dehydrogenase complex in KEGG oxidative phosphorylation map.

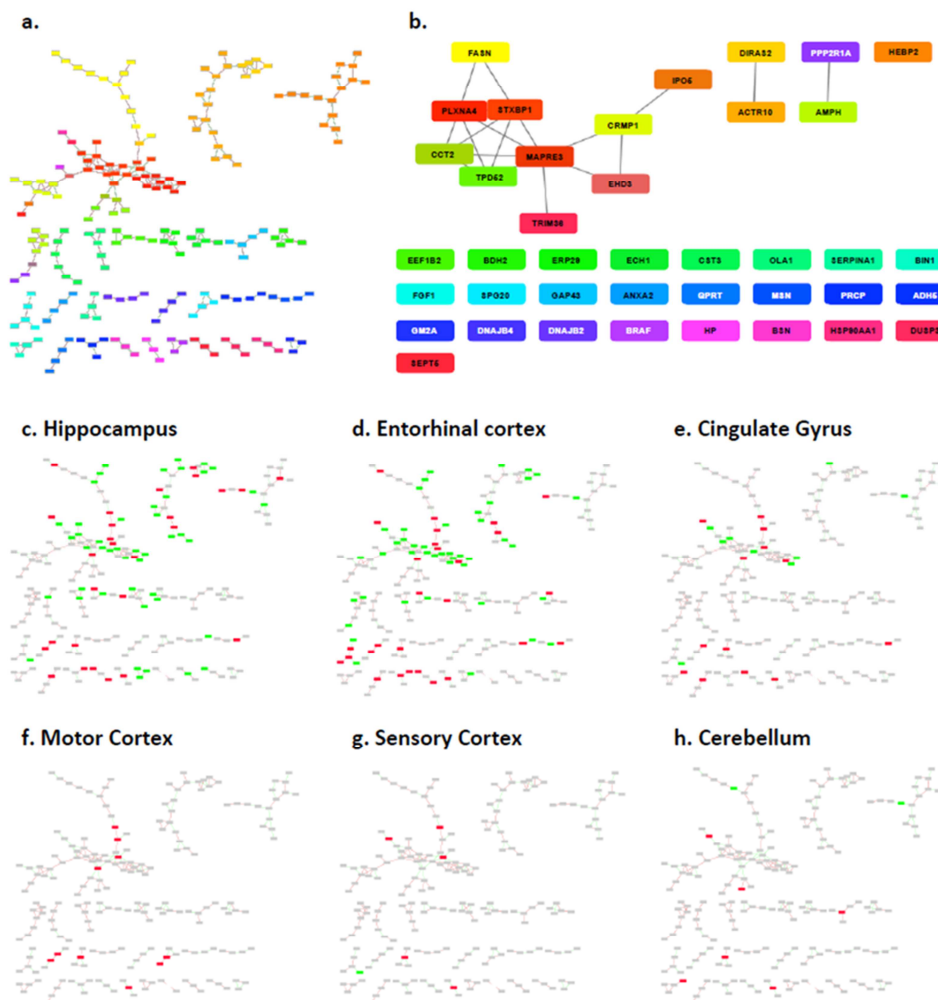
It is well established that CB does not display extensive apoptotic activation seen elsewhere in the brain in Alzheimer's disease, which is unsurprising given its structurally unaffected status. Our findings indicate that the lack of significant neurodegeneration in this region is not merely due to the absence of an apoptotic signal (e.g. Tau tangles) but instead that CB actively induces a unique pattern of upregulated neuronal survival pathways alongside protection against oxidative and inflammatory damage; a protective mechanism of gene/protein expression which limits disease-related degeneration in this region.

Given the apparently similarity in protein expression which we seen wining each group (severely affected and less affected), we next attempt to identify key regulators of what appears to be a coordinated alteration in protein expression across the brain in response to AD. We performed a correlation network analysis to identify key nodes which may be responsible for the programme of protein expression observed, using the Cytoscape ModuLand plug-in⁴⁴. The resulting correlation network is shown in Figure 5a. Each cluster is coloured differently according to a distinct meta-node, the key regulators of which can be determined by visualizing higher levels of this hierarchy (Fig. 5b). Using this method, we can identify the most influential genes in this correlation network which we hypothesize to be key regulators of protein expression during the pathogenesis of AD. It is noteworthy that in this correlation matrix we are aiming to correlate what we believe to be two distinct processes – AD pathogenesis (seen in HP, ENT, CG, MCx and SCx) and a protective programme that we observe in CB. By overlaying protein expression data onto this network, we can identify which nodes are associated with which process. This overlay (Fig. 5c-h) clearly demonstrates that the correlation network is mainly constructed from proteins involved in AD pathogenesis in the affected regions – few proteins in the network are changed in CB despite the relatively large number of CB proteins which we observe to be changed in the complete dataset. This is to be expected as CB-specific protein changes have limited correlation to the remainder of the dataset. This network is therefore likely to provide a good representation of the key events in AD pathogenesis, and reveals four proteins with the most overall influence on the correlated expression networks: STXBP1 (syntaxin binding protein 1); CRMP1, (collapsin response-mediator protein 1); ACTR10, (actin-related protein 10 homologue); and AMPH (amphiphysin).

STXBP1 is the regulator with the most influence in this network. It is reportedly upregulated in AD⁴⁵, has been linked to NFTs⁴⁶ and may interact with PS1⁴⁷. It also plays a major role in neurotransmitter

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release. STXBP1 thus provides a potential mechanistic explanation for our observation that pathways of neurotransmitter metabolism including dopamine-, noradrenaline-, and serotonin-related signalling showed significant changes in severely affected regions and SCx, but not in MCx or CB. Another important regulator of the network, CRMP1, is part of the semaphorin signalling pathway which is known to guide axons in developing nervous tissue and participates in shaping of neural circuits⁴⁸. ACTR10 may affect prion susceptibility through its involvement in prion propagation and clearance⁴⁹, and has been identified by large scale computational network analyses as one of a large number of potentially important genes in hippocampal ageing, but our finding is novel in AD⁵⁰. The 4th key network regulator identified here, AMPH, is a candidate AD risk gene that may participate in receptor-mediated endocytosis and hence be involved in APP metabolism/clearance⁵¹. Our finding that these four genes appear to be central to various pathological processes known to be involved in AD development is important, and suggests that further work should be performed to focus on the role of these potentially key mediators of Alzheimer's disease progression.



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Figure 5. Global networks analysis was performed using Cytoscape ModuLand plug-in. a) Correlation network of altered proteins in AD brain, with differently coloured clusters representing different meta-nodes b) key regulators of each meta-node c-h) Overlays of protein expression data from each region and the correlation network.

Since one of the key factors in AD pathogenesis is thought to be the build-up of amyloid consisting of A β peptide generated as a proteolytic product of the Amyloid precursor protein (APP) we examined our data for information about the levels and distribution of these molecules. We found no marked change in APP levels overall but significantly elevated A β peptide levels (Supplementary Figure 2a-b), consistent with previous reports⁵². The extent of the increase in A β between regions does not appear to follow a gradient of ‘affectedness’, albeit there may be a more pronounced increase in hippocampus. There is no way to determine the primary structure of the A β peptide(s) present in each region from these data. Interestingly, while in the AD group almost all samples showed uniformly high levels of A β peptide, there was marked variation in levels in control samples (Supplementary Figure 2c). While the quantification of A β is necessarily from one peptide, these data emanate from between 5 and 12 unique spectra in each sample we consider this observation is likely robust. This variability is therefore likely to be due to inherent variations in the control population. Although all patients in this group were asymptomatic, it is likely that varying degrees of prodromal disease could have been present, given their age. This is most noticeable in our control 115. While initially assigned as a control, a pathological re-examination performed as a result of the findings of this study and our previous metabolomics analyses¹⁴ re-classified this individual as a Braak II pre-clinical AD patient. This patient has the highest level of A β of all of the control samples and interestingly appears to demonstrate some AD-related changes both in their metabolome and in some of the proteins which we observe to be changed in symptomatic disease. This observation supports the idea that increases in A β levels may reflect varying degrees of prodromal disease in these elderly controls. It also demonstrates that studies of the type performed here in earlier stage presymptomatic patients will be critical to further tease out the very earliest events in AD pathogenesis.

In summary, this study provides a map of molecular changes that are present in human *post-mortem* brain tissue in patients with AD and matched controls, providing insights into the brain region specificity of disease at two levels; individual proteins and pathways. We observed global perturbation of protein expression in all six regions of the AD brain which we studied. An association between extent of molecular changes and affectedness was observed for five regions, allowing us to delineate probably ‘early’ and ‘late’ changes in protein expression and revealing previously novel involvement of several pathways and processes. The sixth region, CB, showed an unexpectedly

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distinct pattern of protein changes, suggestive of induction of a protective response. Correlation network analysis identified four candidate genes STXBP1, CRMP1, ACTR10, and AMPH which may underpin significant portions of the protein expression response to AD. Finally, we recognize that these data have significant value to the community and that other researchers will no doubt wish to assess the status of other AD-related changes not discussed here. As such we have provided all results in an accessible format via a freely-available, searchable on-line database, to allow others to probe specific pathways or individual proteins and their expression in regions across the human Alzheimer's disease brain and matched controls.

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2 **Supplementary Information** is available in the online version of this paper

3

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12

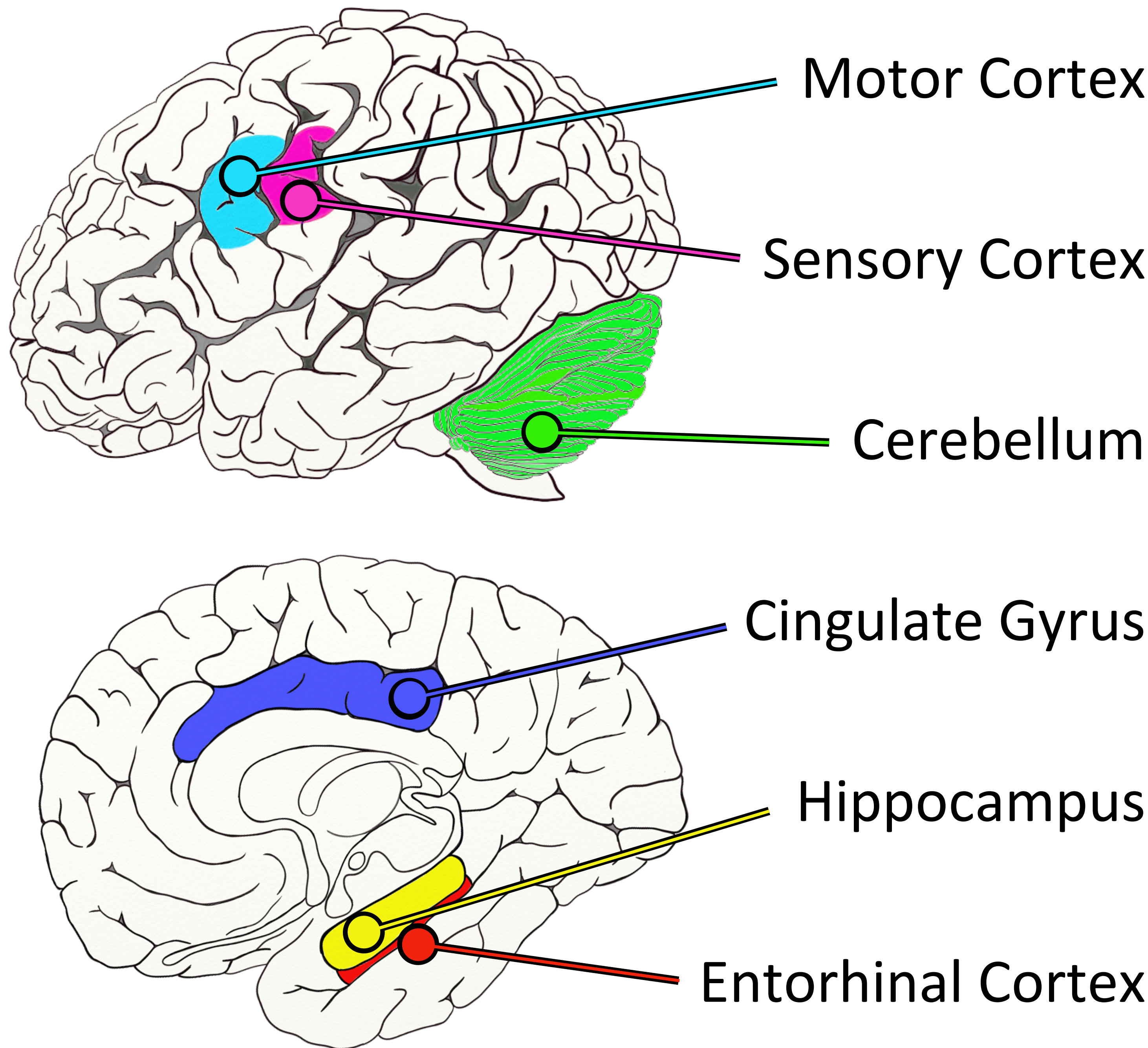
13 **Author Contributions** J.X., S.P., N.R., I.R-G., B.D.H, and H.W. performed the experiments
 14 presented in this manuscript. Initial Bayesian data analysis was developed by A.M.P., A.W.D. and
 15 R.D.U. R.H. and P.B. build the web-based data resource. J.X., A.S. and R.D.U. performed data
 16 interpretation and network analysis. R.L.M.F., G.J.S.C. and R.D.U. supervised the project. All
 17 authors wrote the manuscript.

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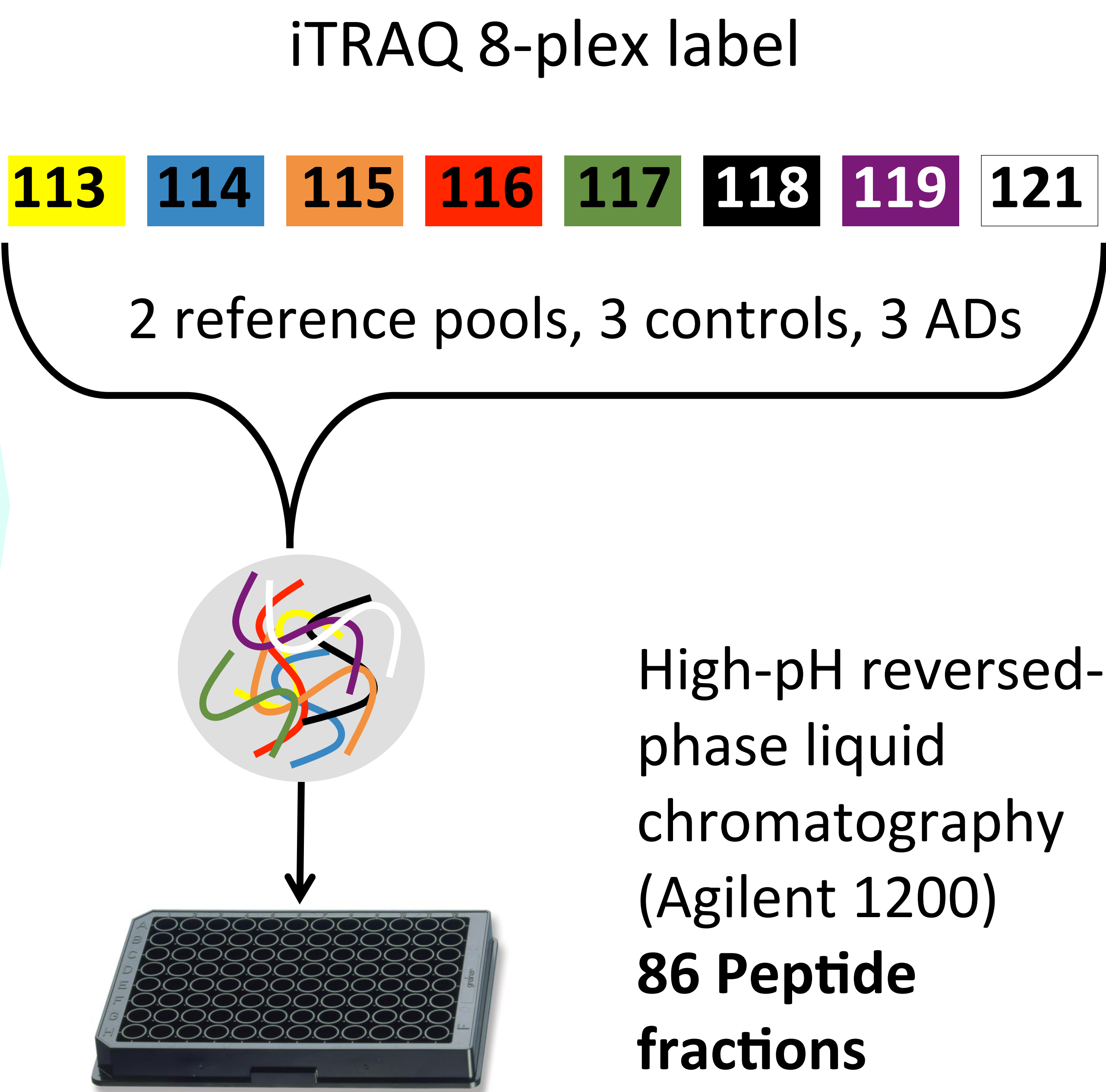
19 **Author information** The authors declare no competing financial interests. Correspondence and
 20 request for materials, methods or data should be addressed to R.U. (r.unwin@manchester.ac.uk).

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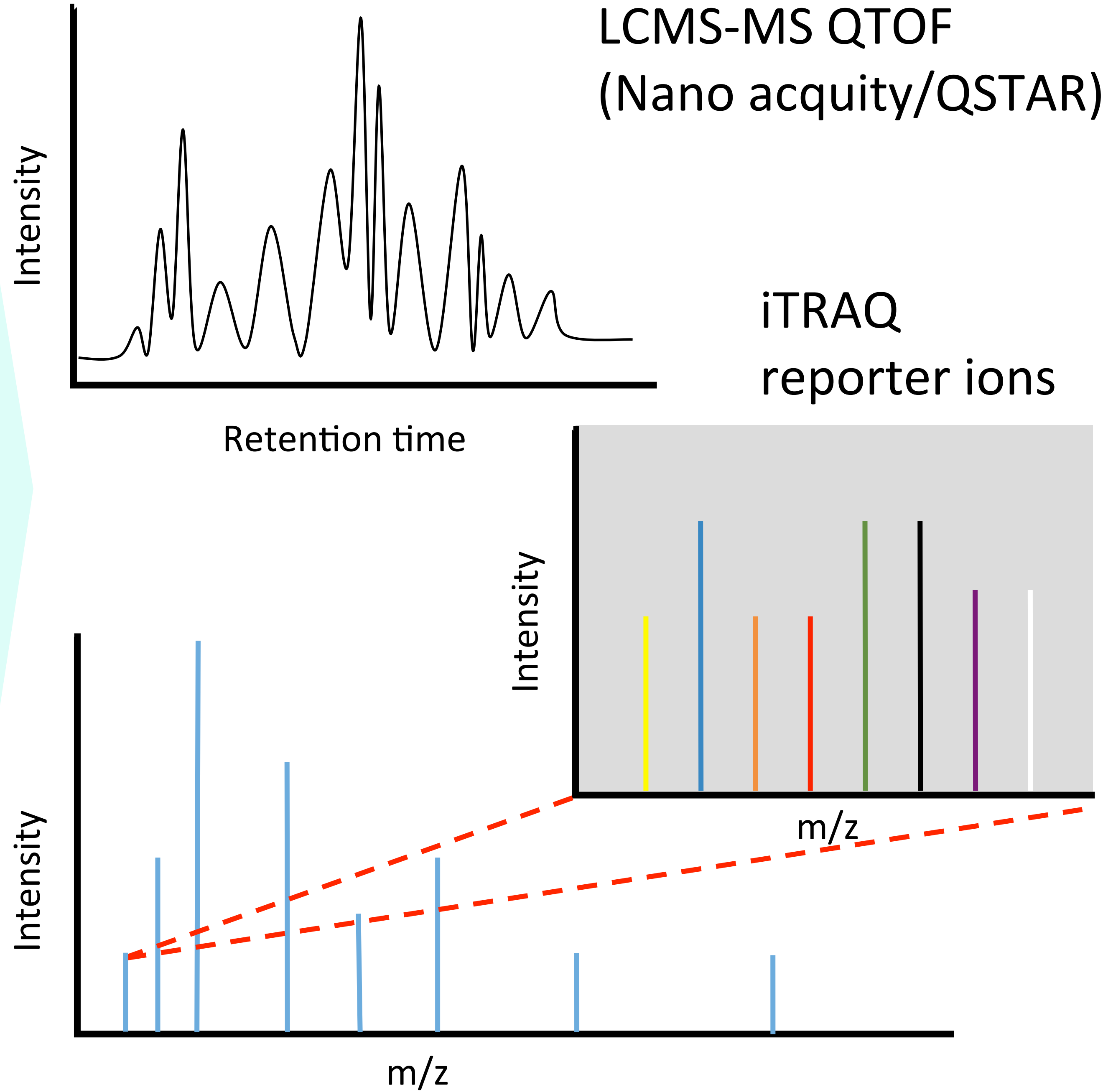
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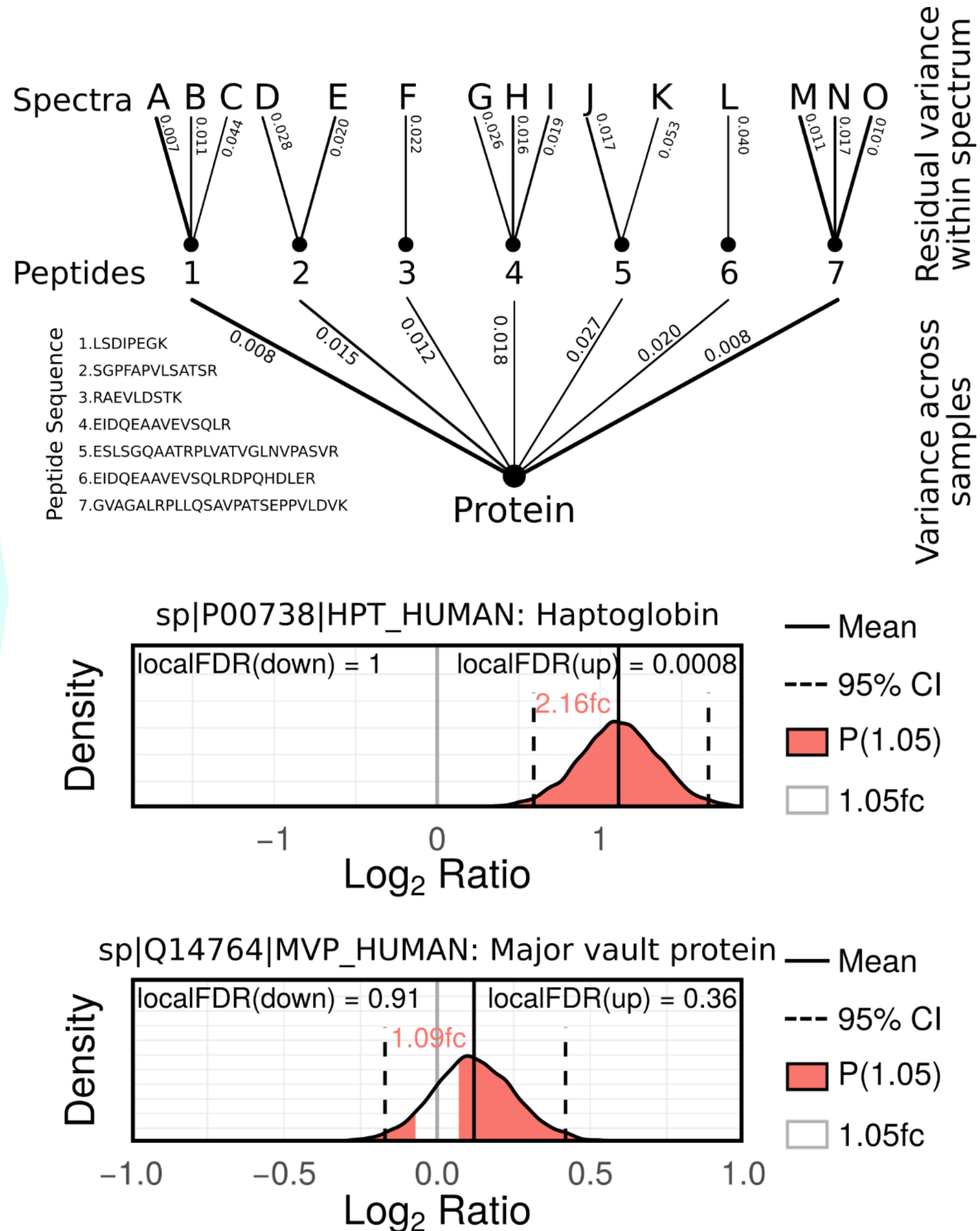
Sample preparation



MS data acquisition



Data analysis

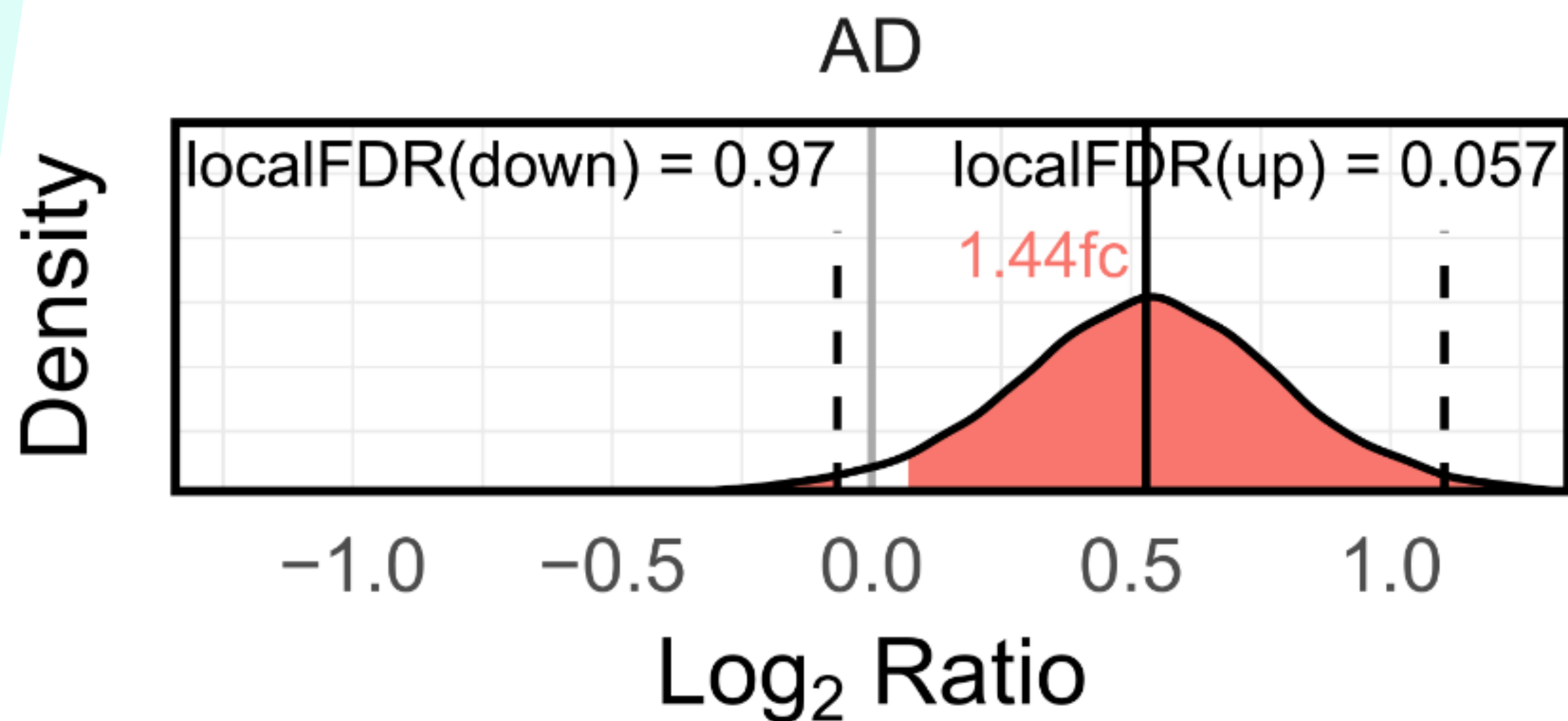


Public database

Website

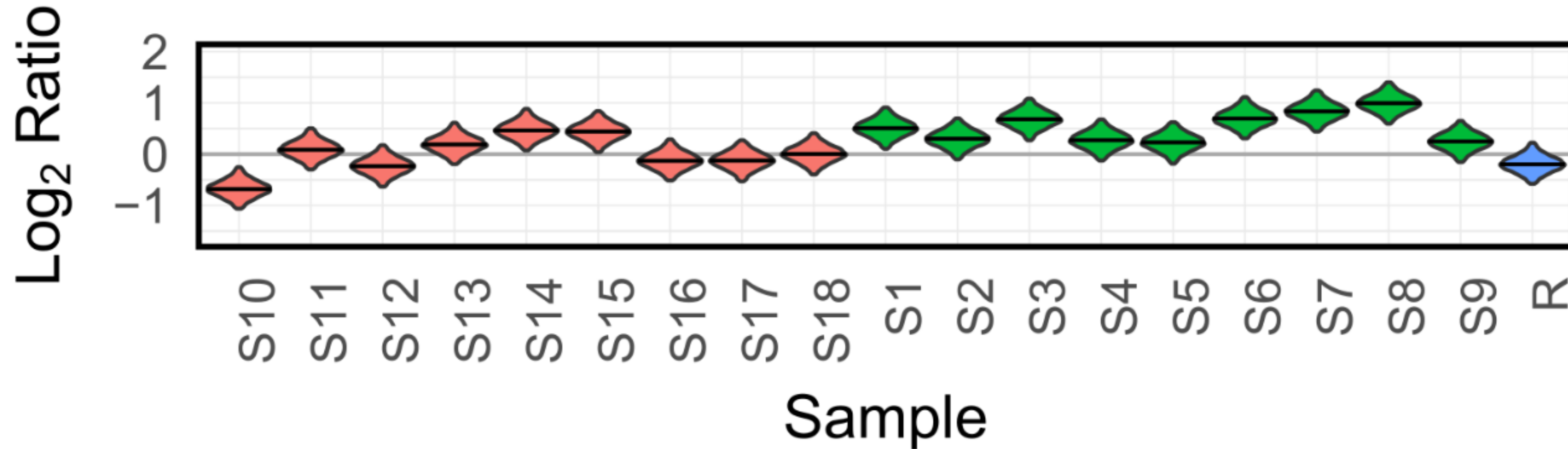
www.manchester.ac.uk/dementia-proteomes-project

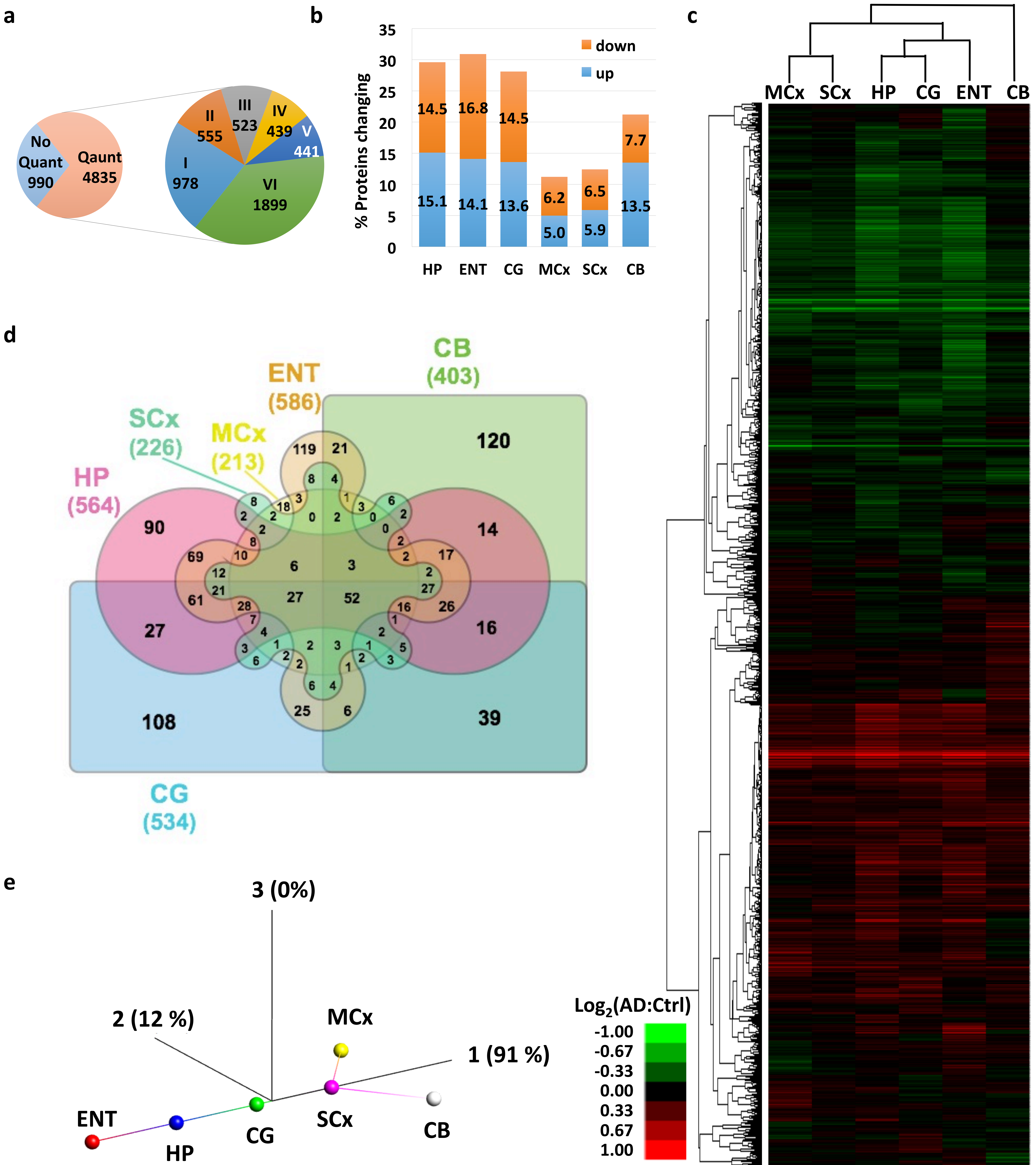
GFAP



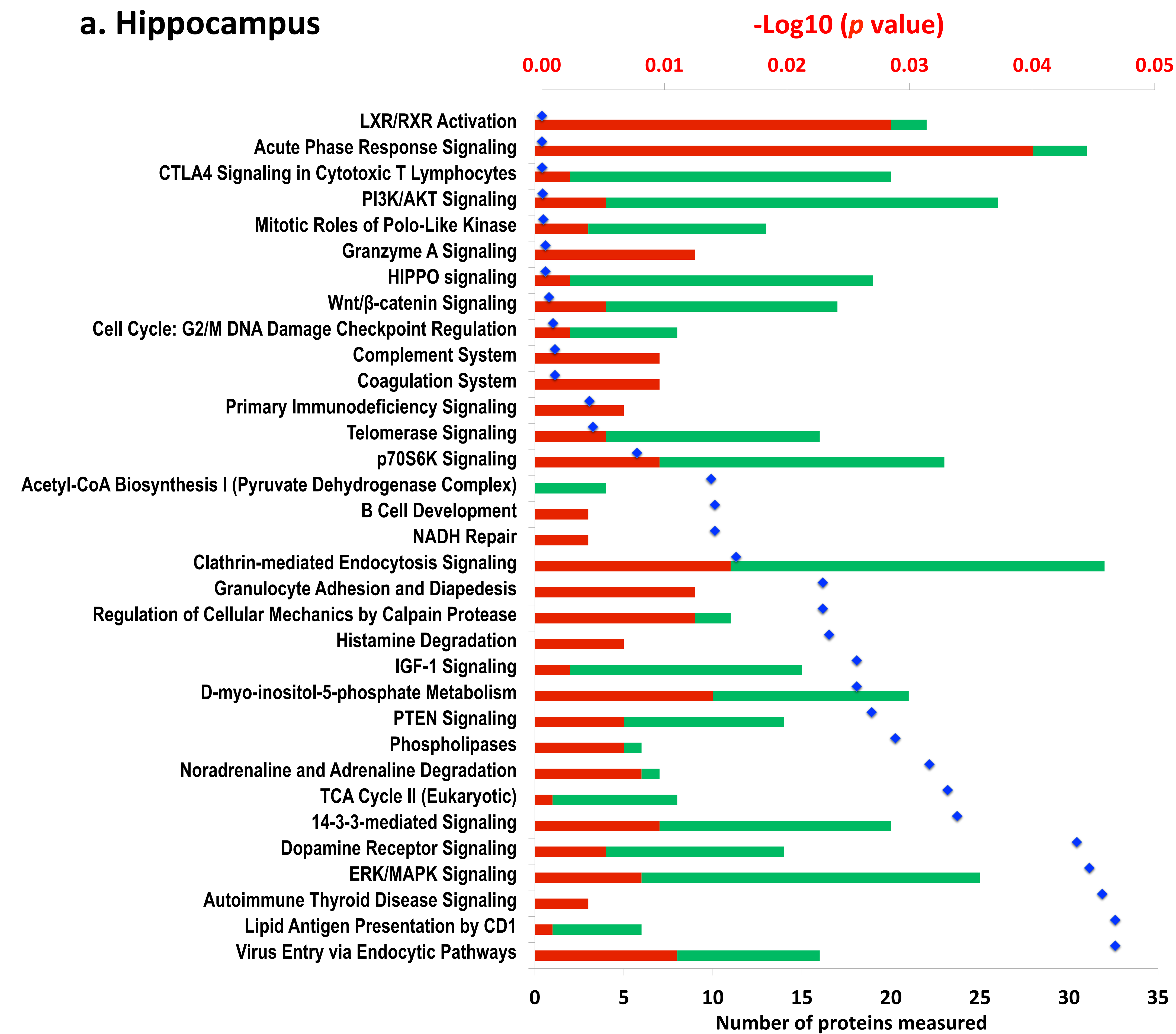
Name	Description	SwissProt_AC	SwissProt_ID
GFAP	P1Glial fibrillary acidic protein	4136	GFAP_HUMAN

	Peptides	Spectra	Log2(fc)	lower	upper	localFDR	globalFDR
HIP	95	1829	0.529	-0.066	1.104	0.057	0.021
MC	44	590	0.199	-0.248	0.662	0.266	0.140
SC	57	821	0.096	-0.331	0.544	0.448	0.260
CER	61	794	0.211	-0.153	0.571	0.195	0.088
ERC	70	1045	0.492	-0.079	1.084	0.068	0.028
CG	54	1211	0.469	-0.169	1.094	0.096	0.040

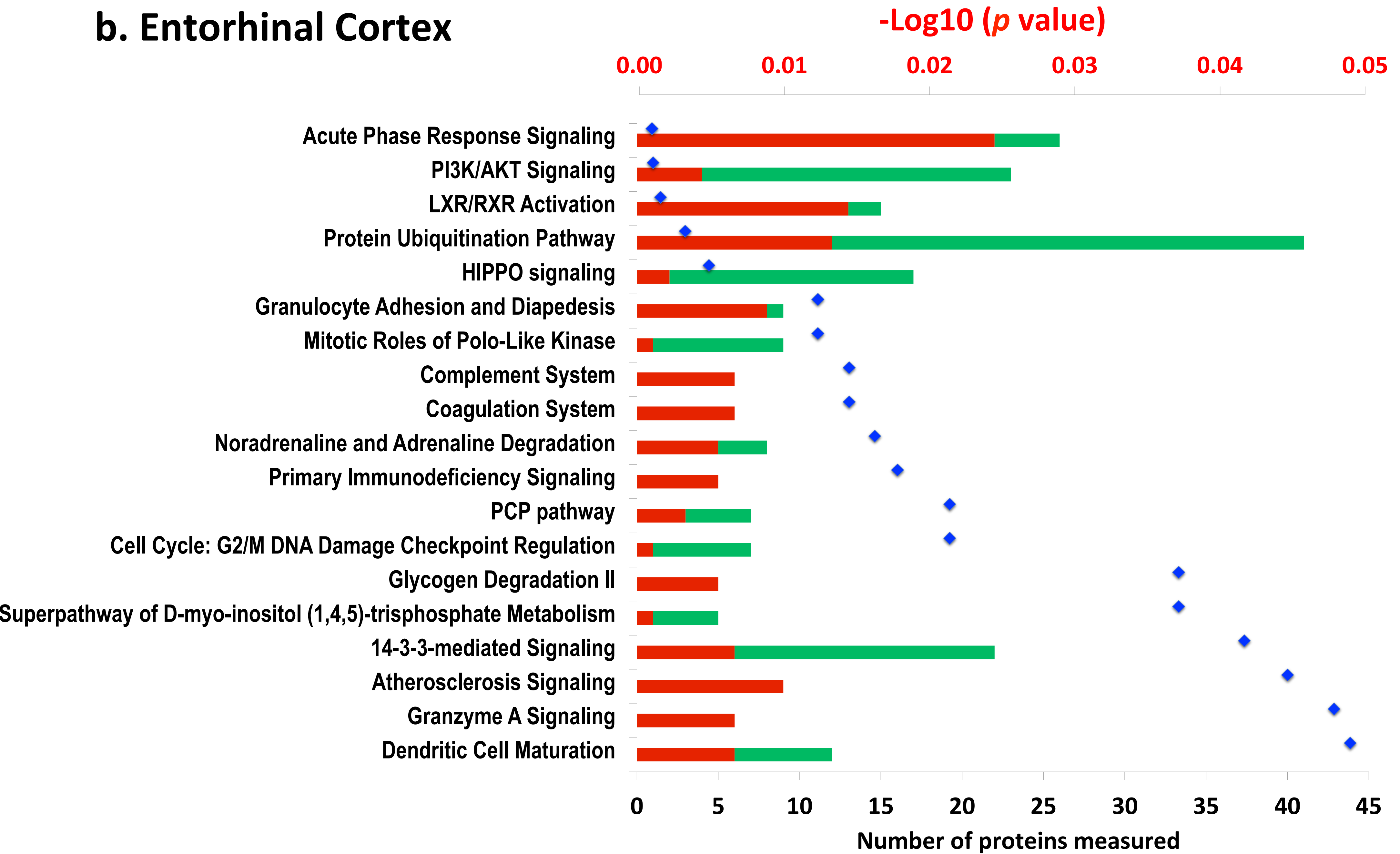




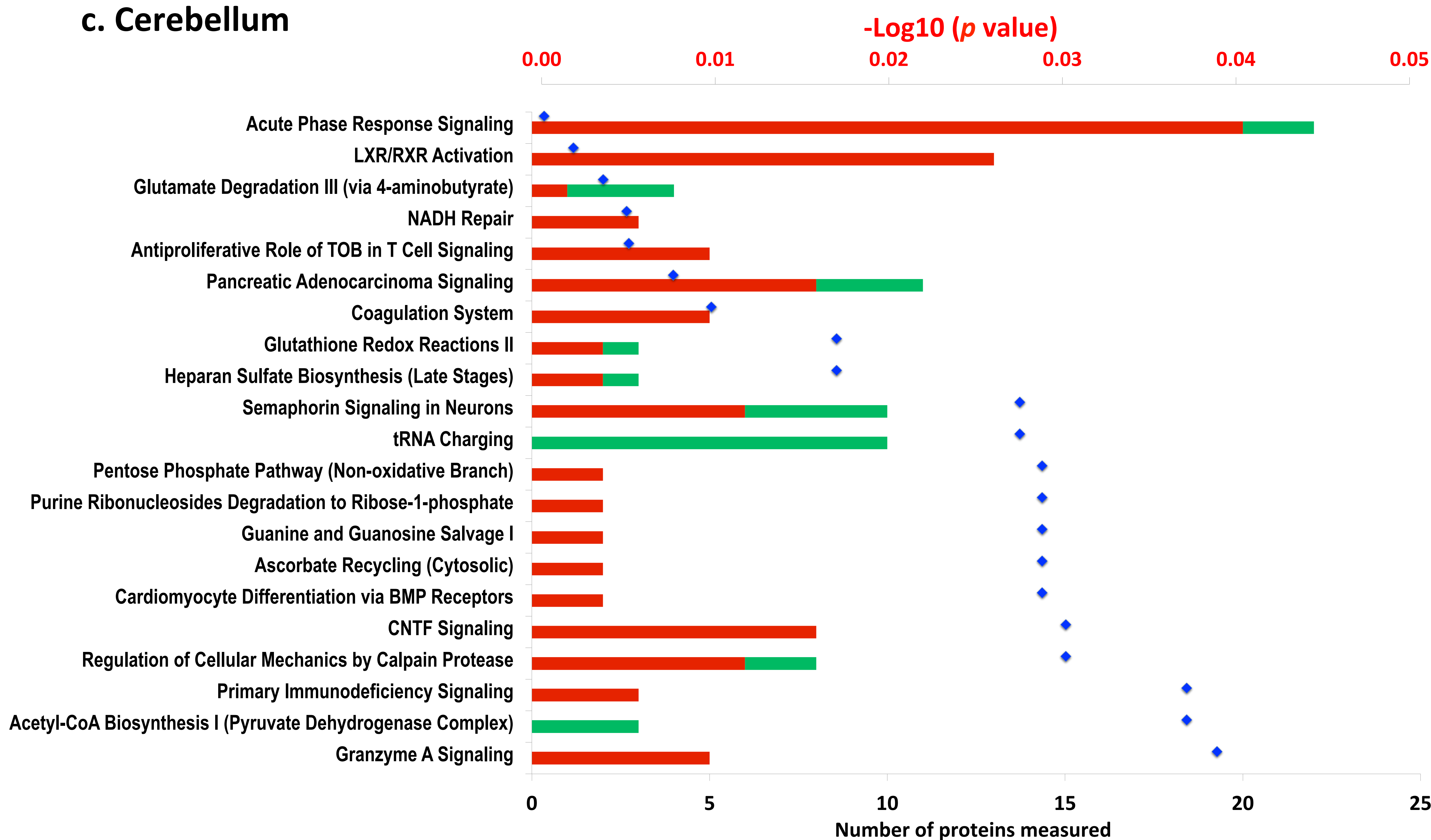
a. Hippocampus



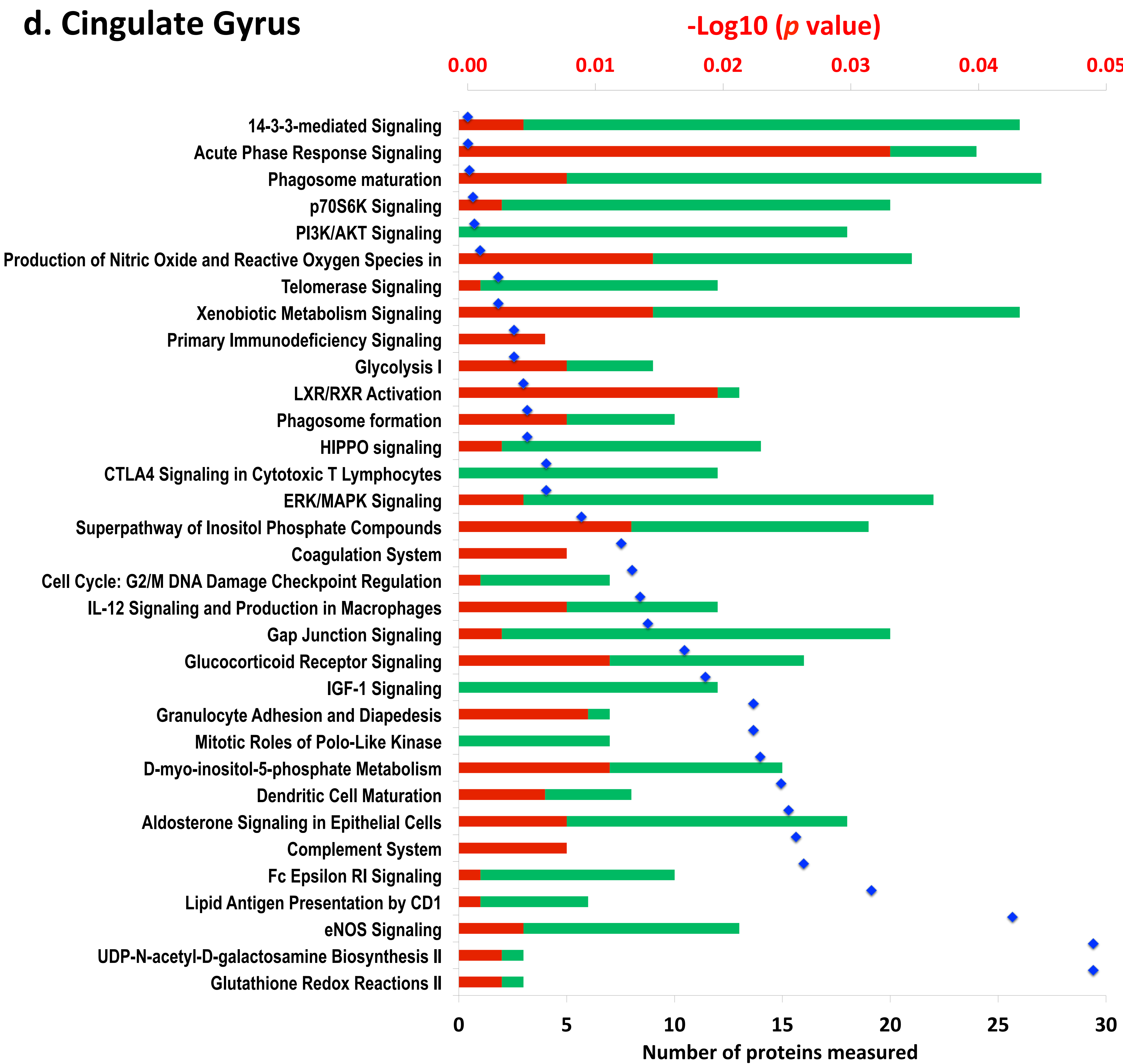
b. Entorhinal Cortex



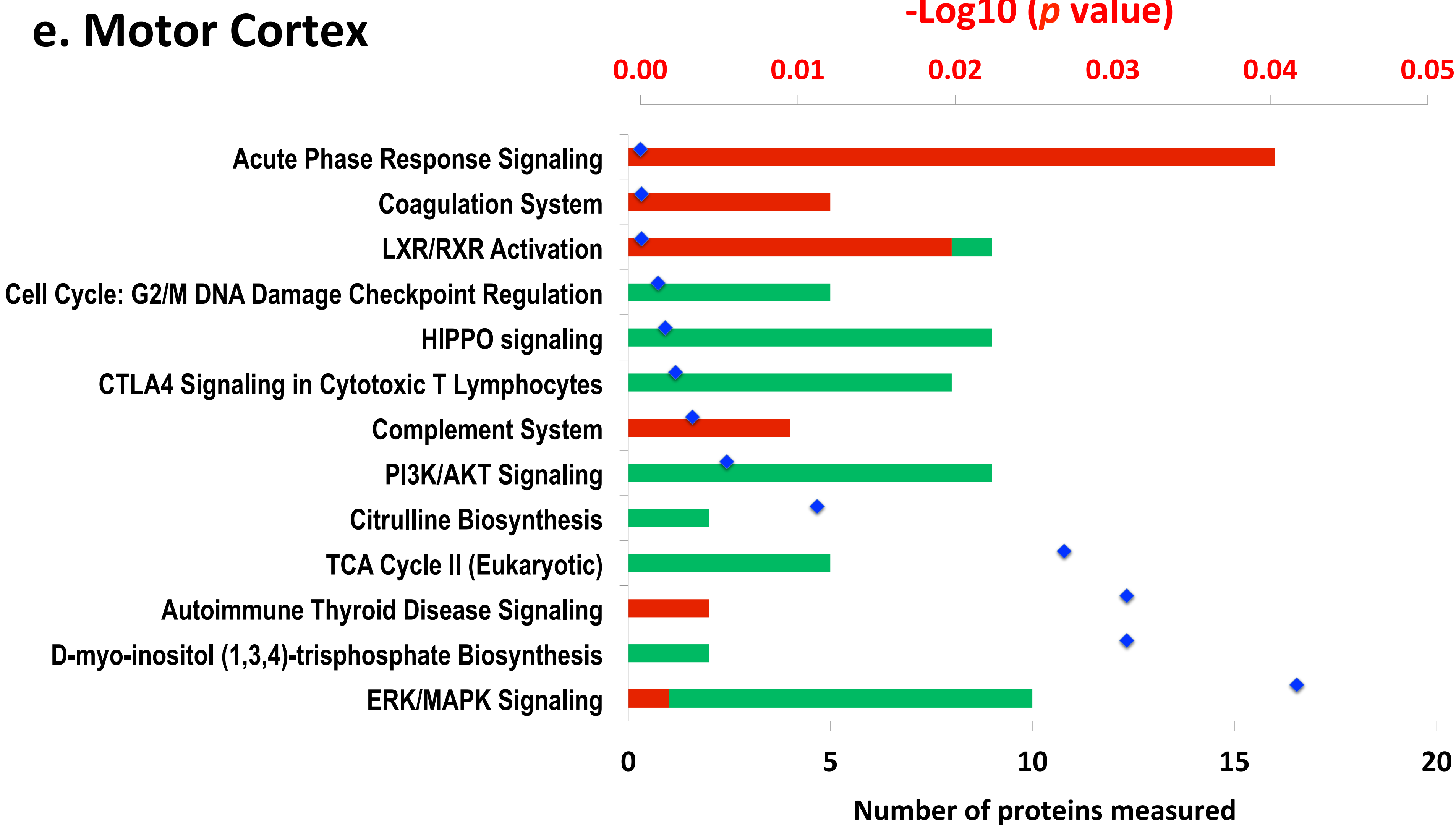
c. Cerebellum



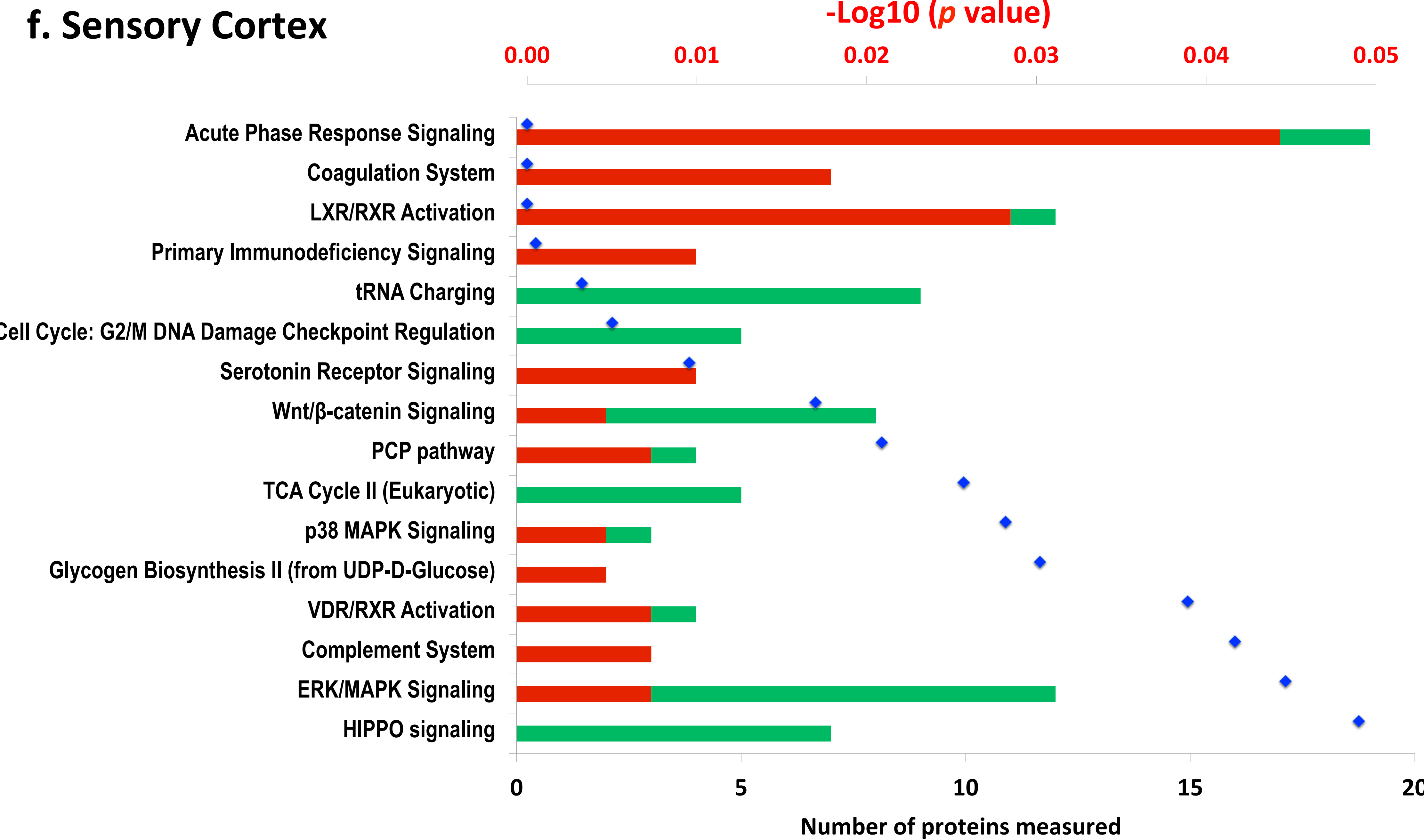
d. Cingulate Gyrus



e. Motor Cortex

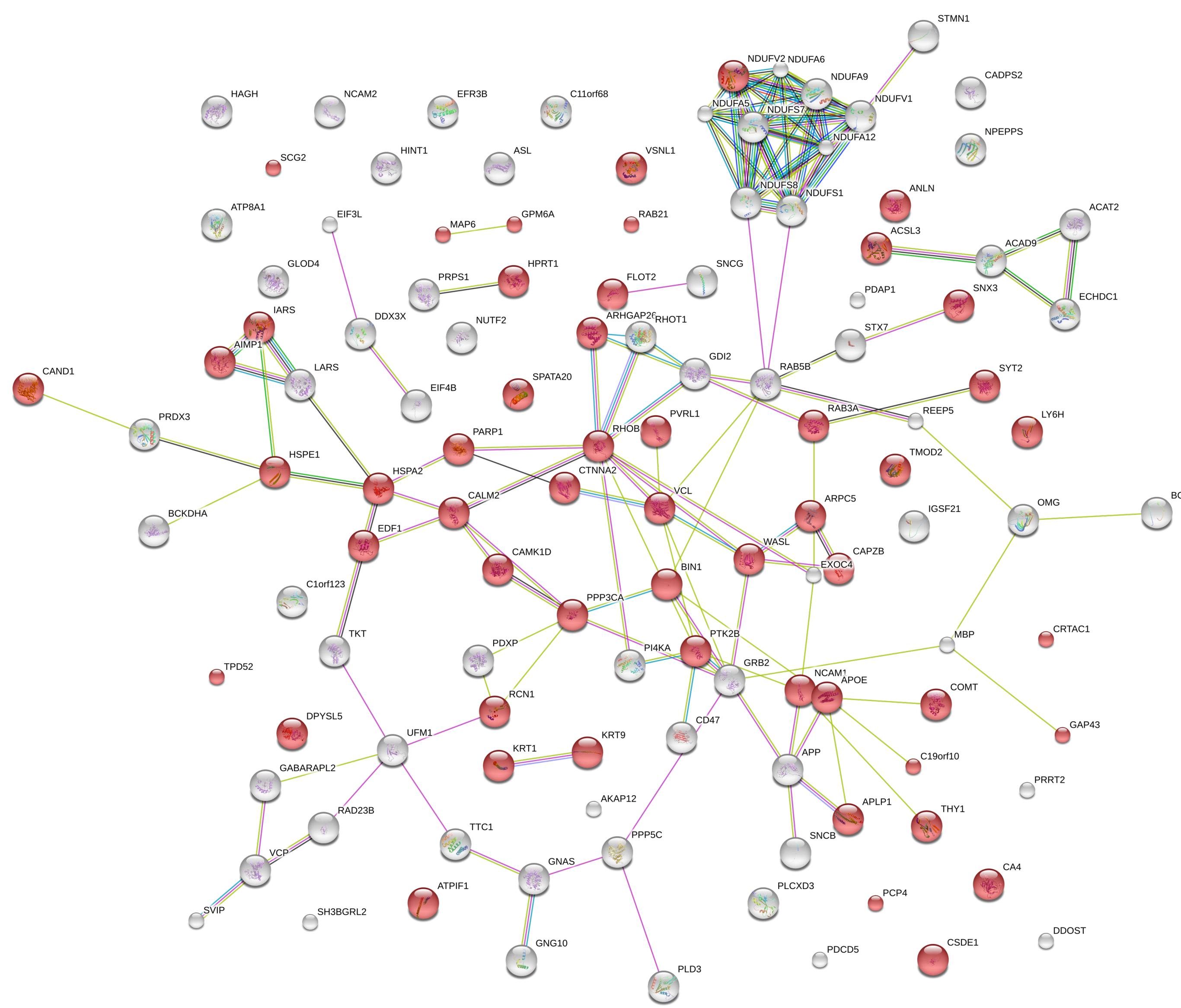


f. Sensory Cortex

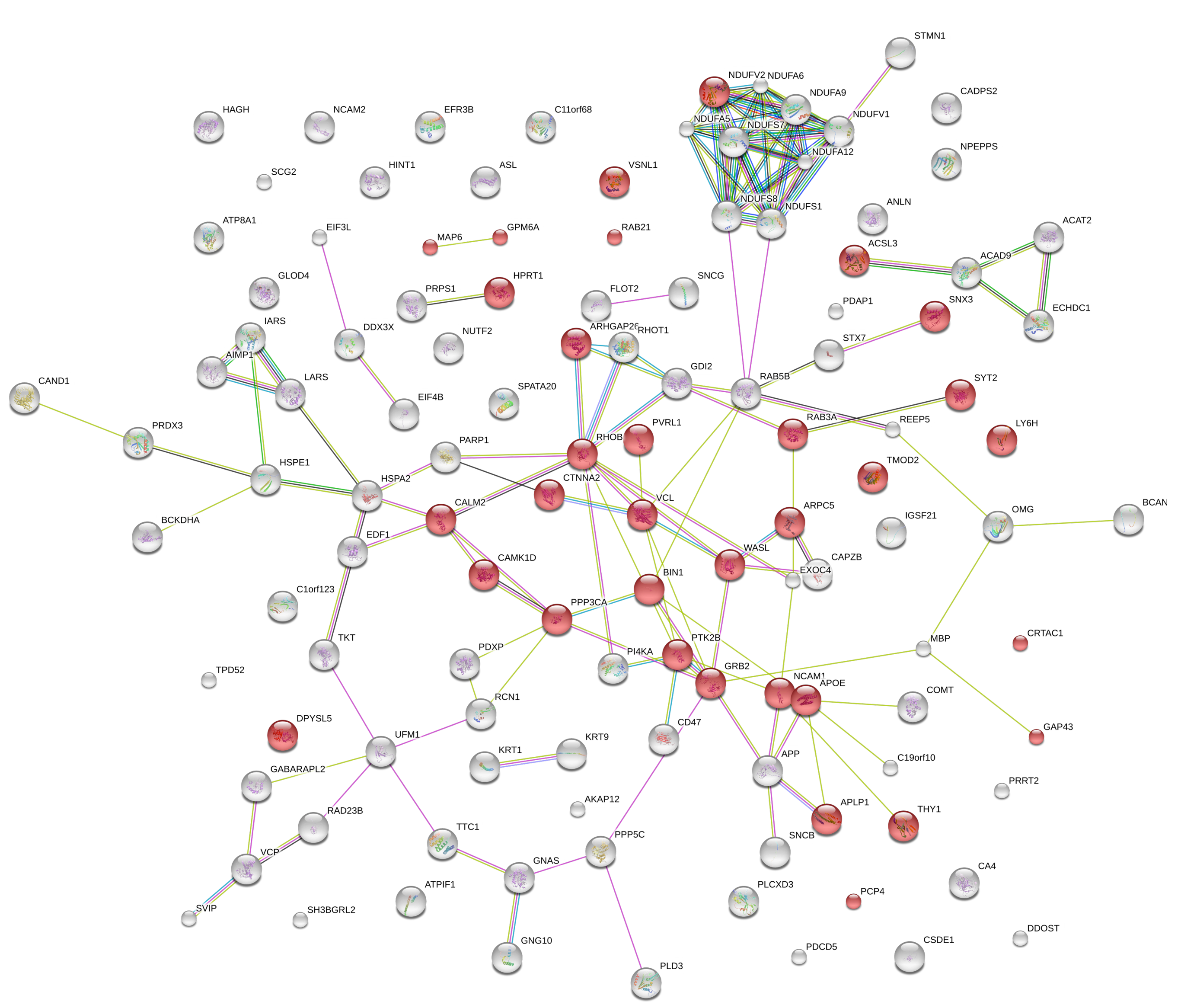


Increased
Decreased
 $\log(p \text{ value})$

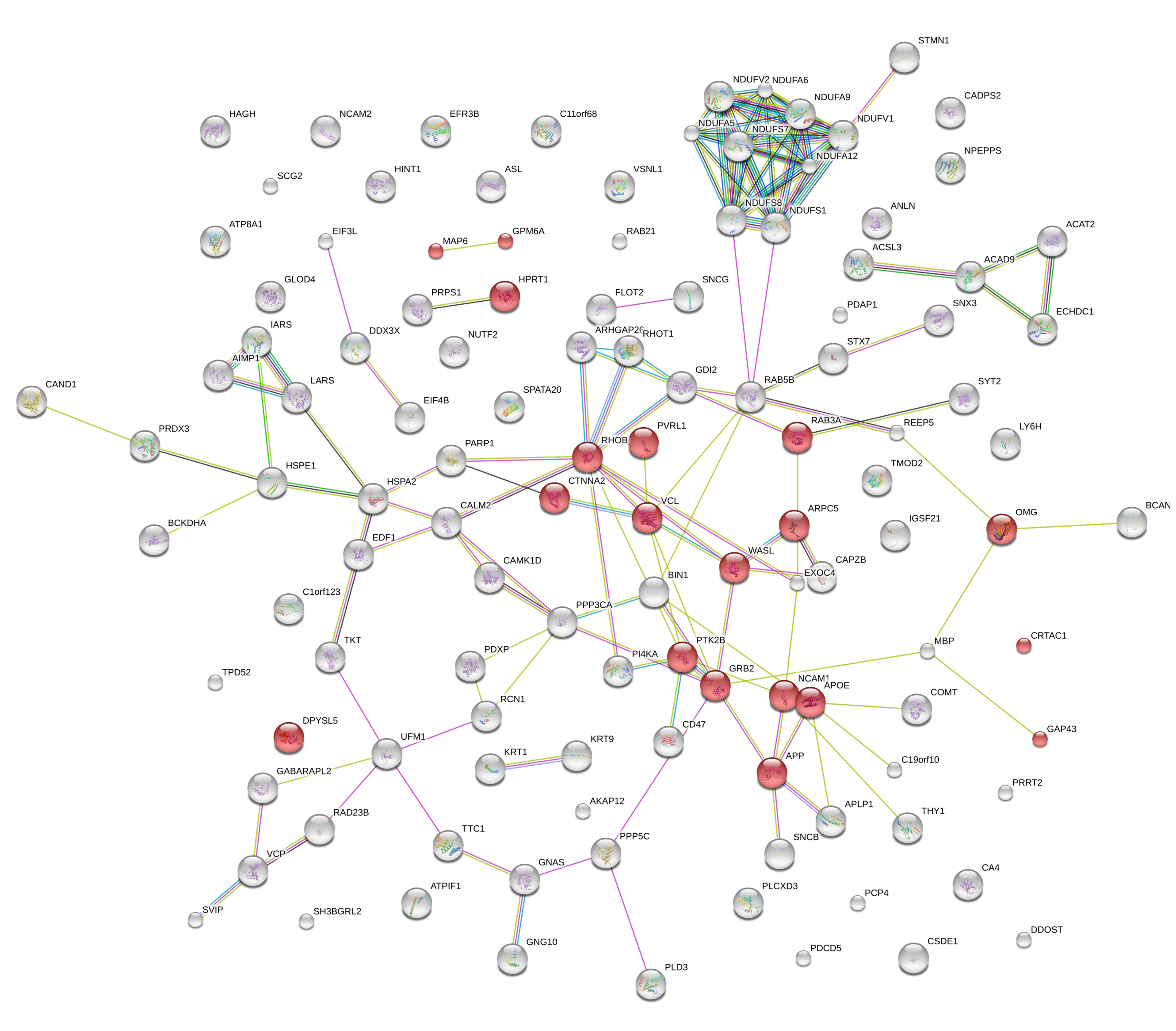
a. Developmental process



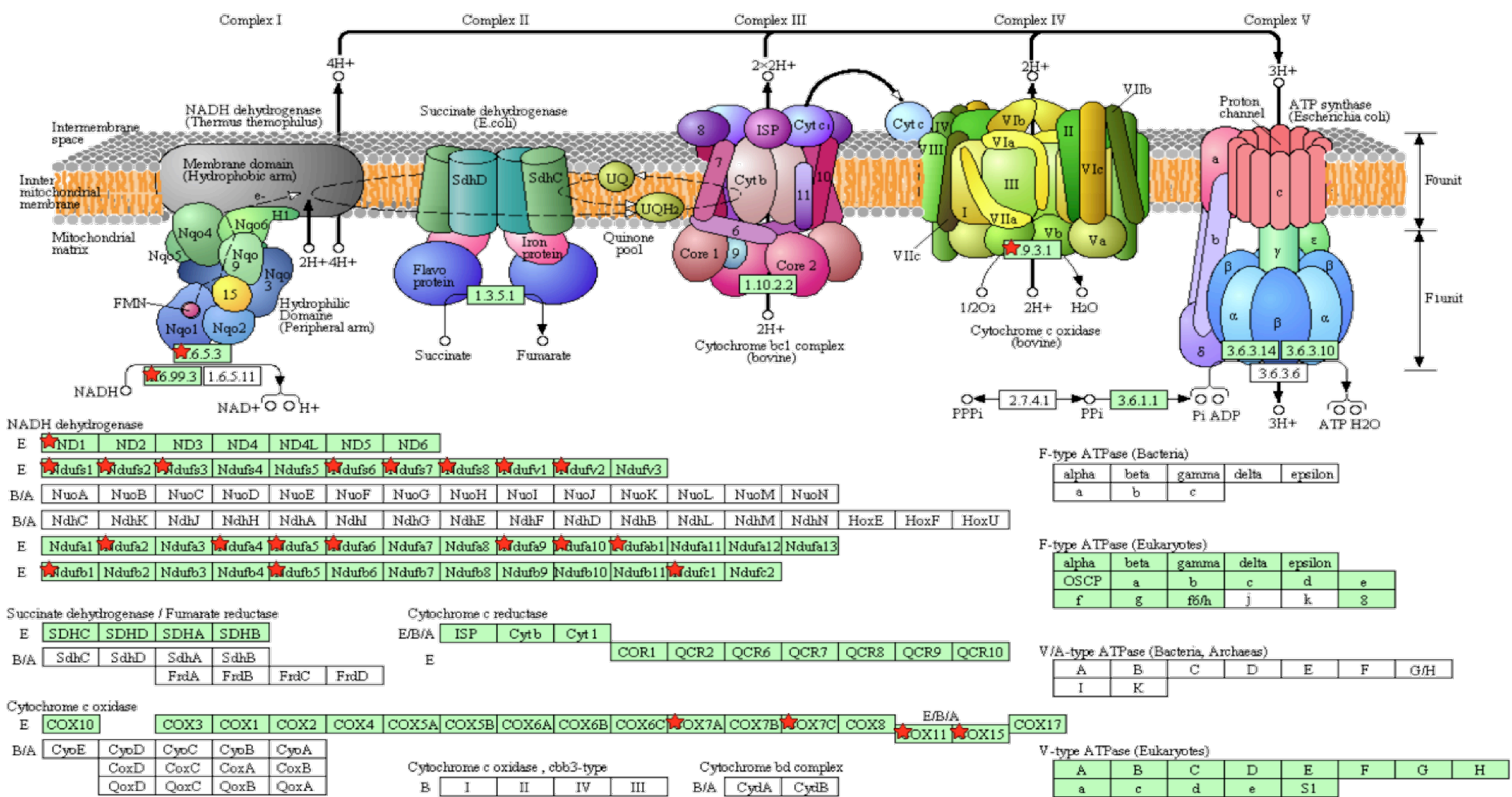
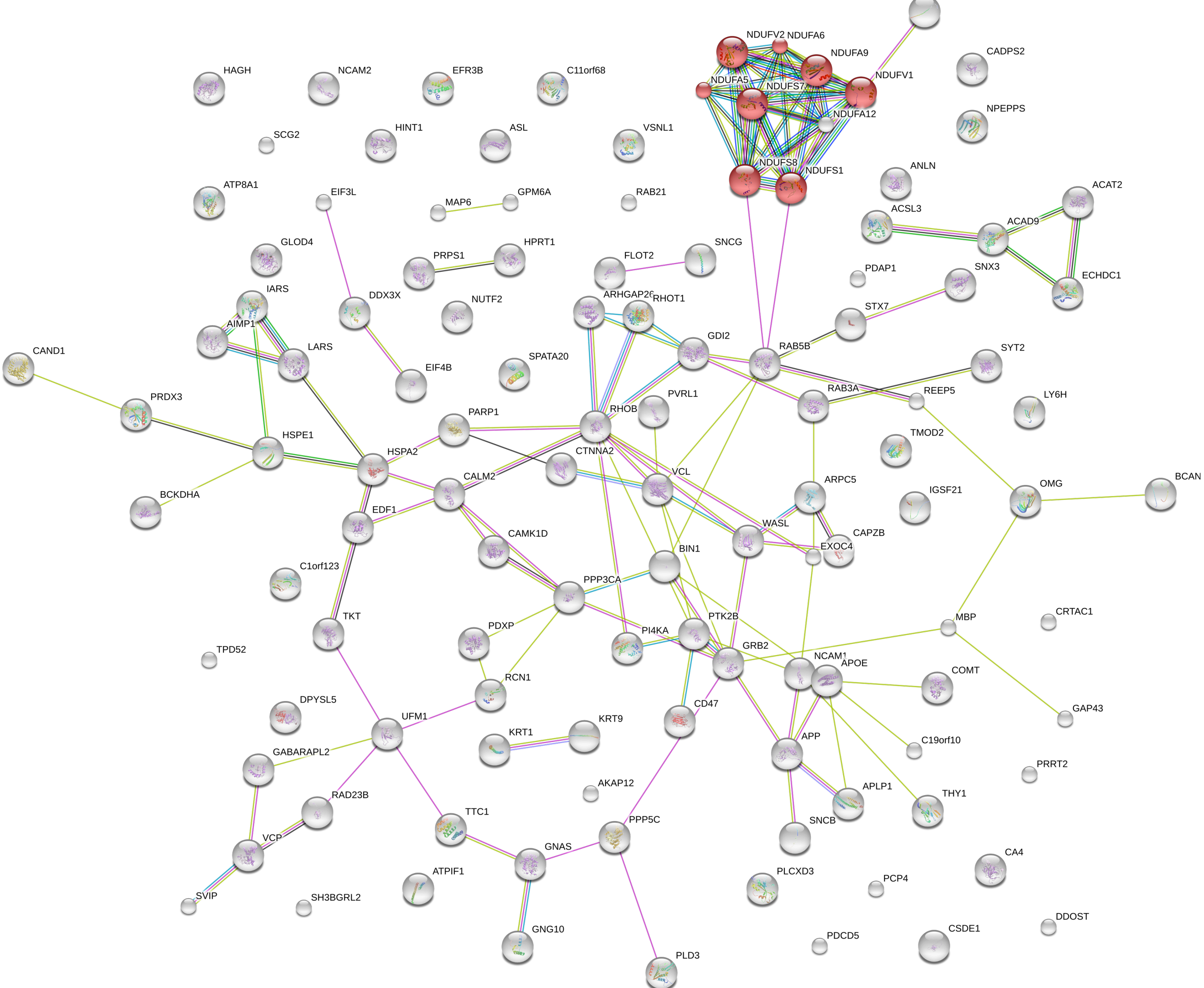
b. Nervous system development



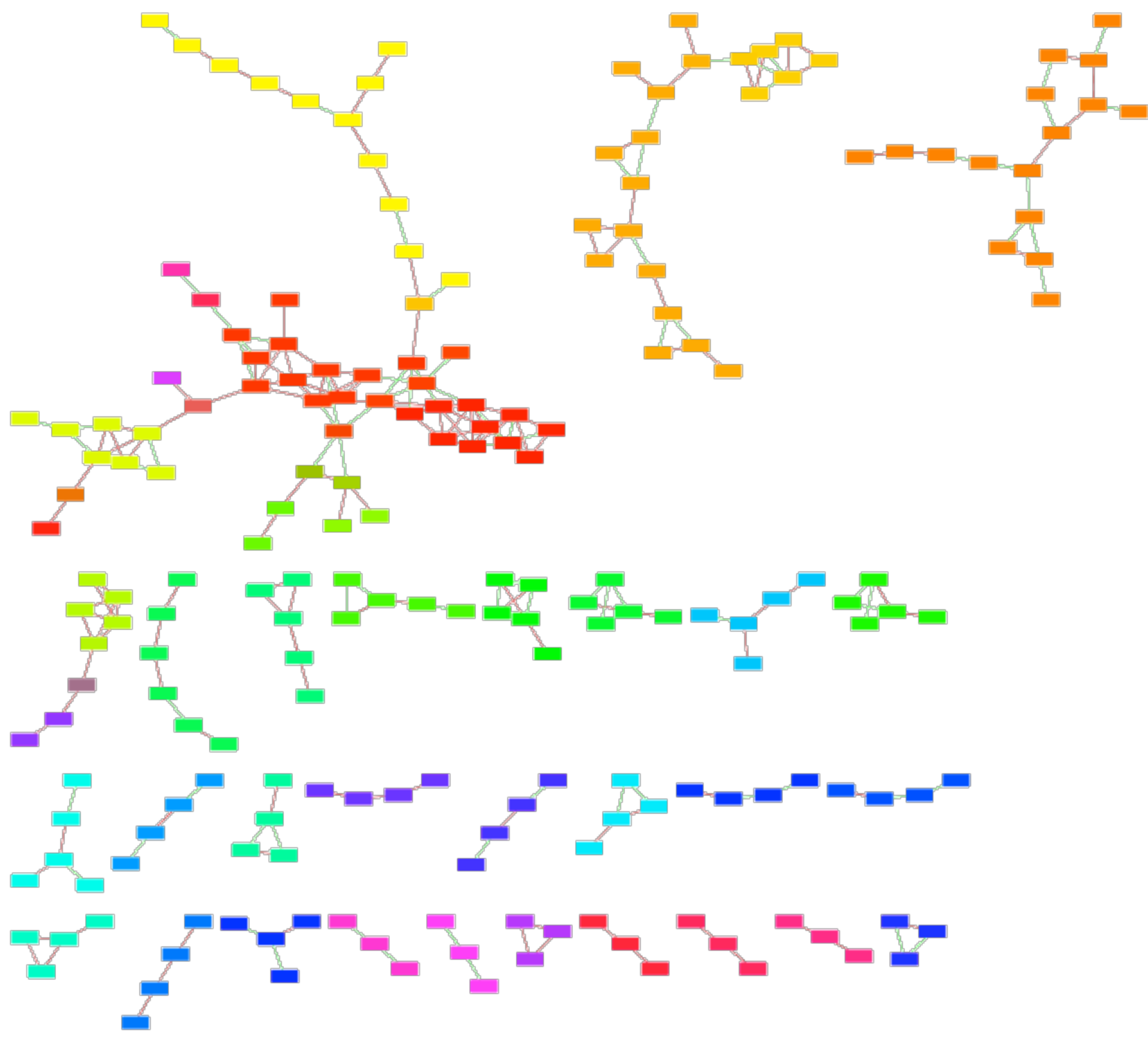
c. Neuron projection development



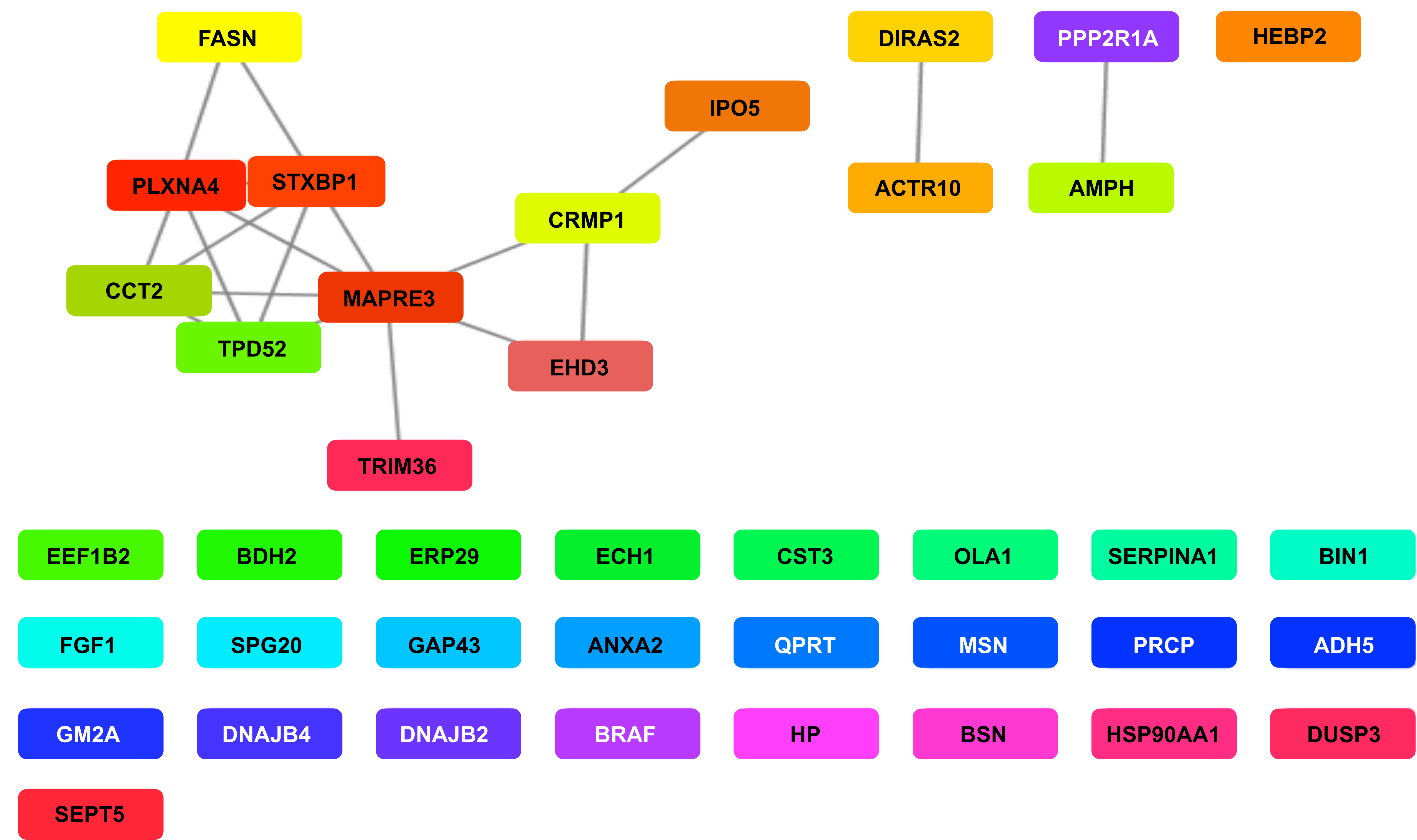
d. Mitochondrial electron transport, NADH to ubiquinone



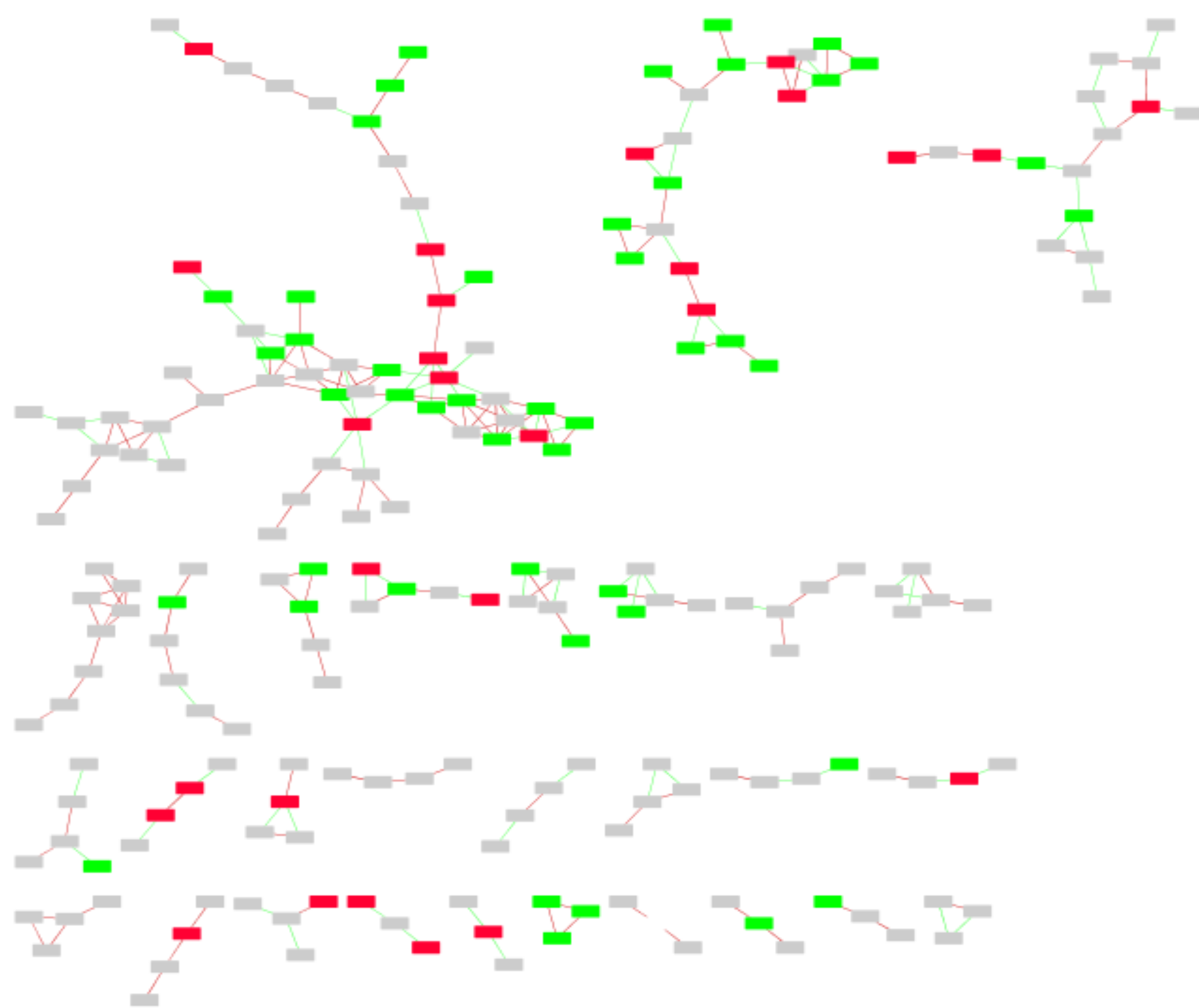
a.



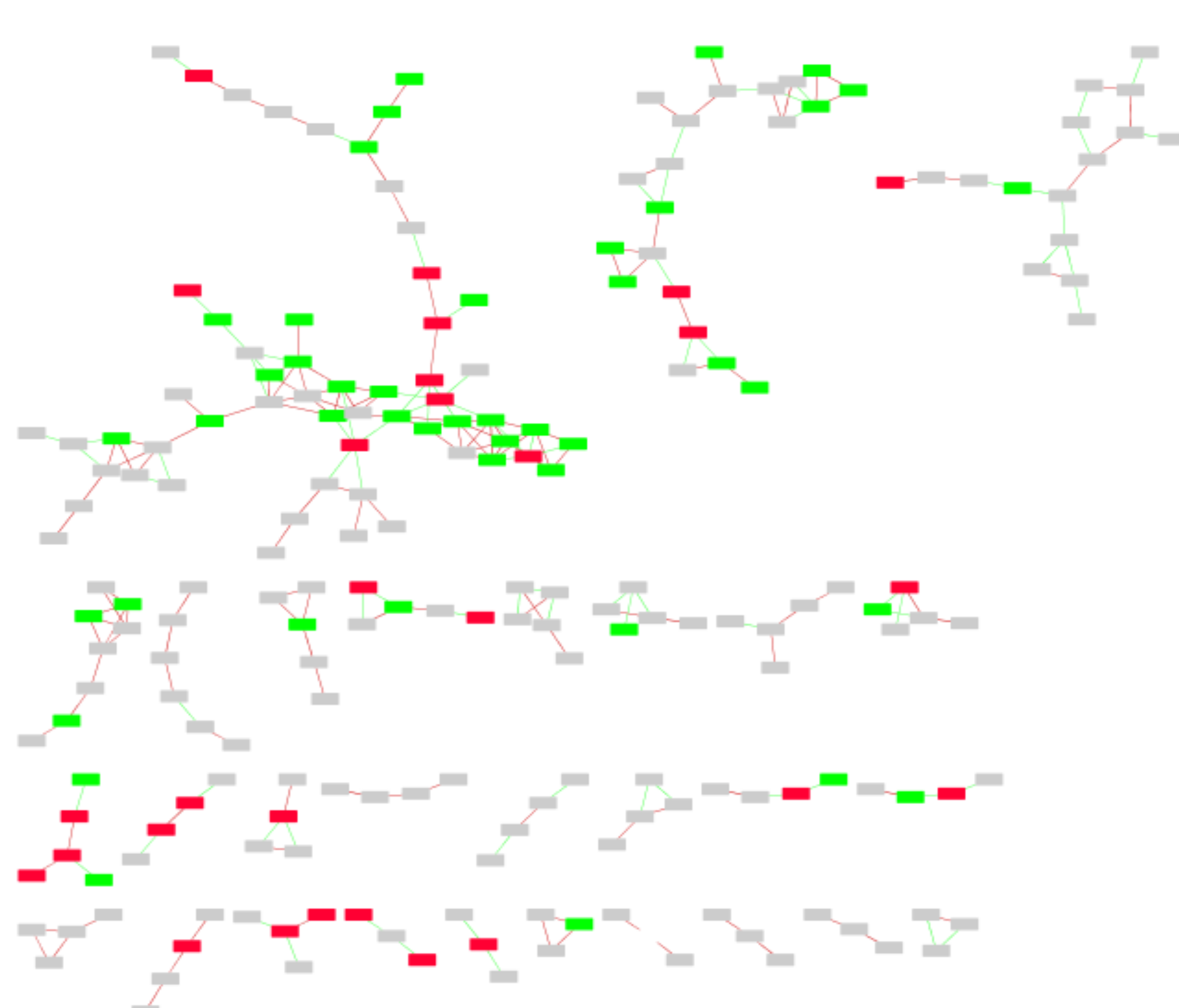
b.



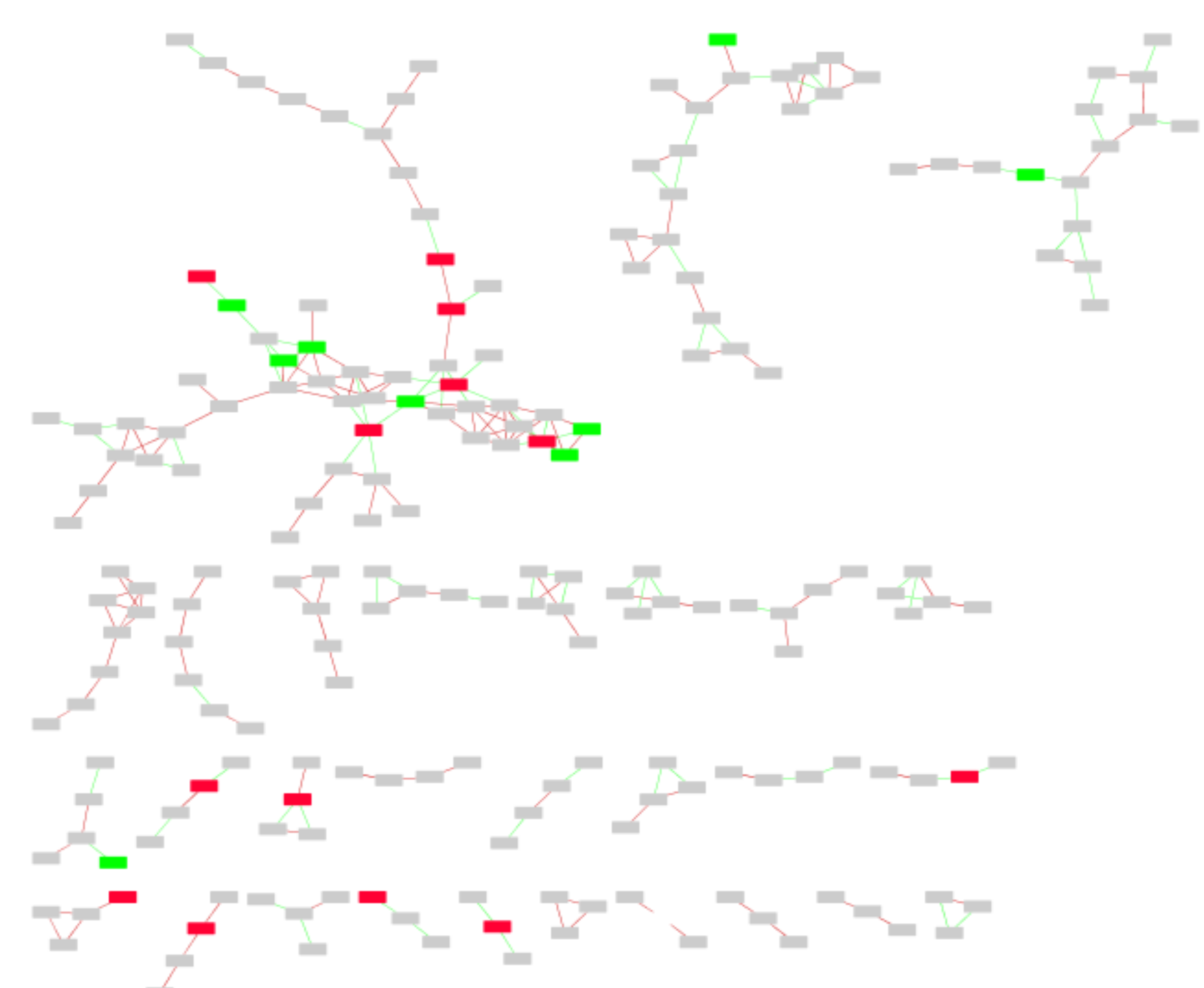
c. Hippocampus



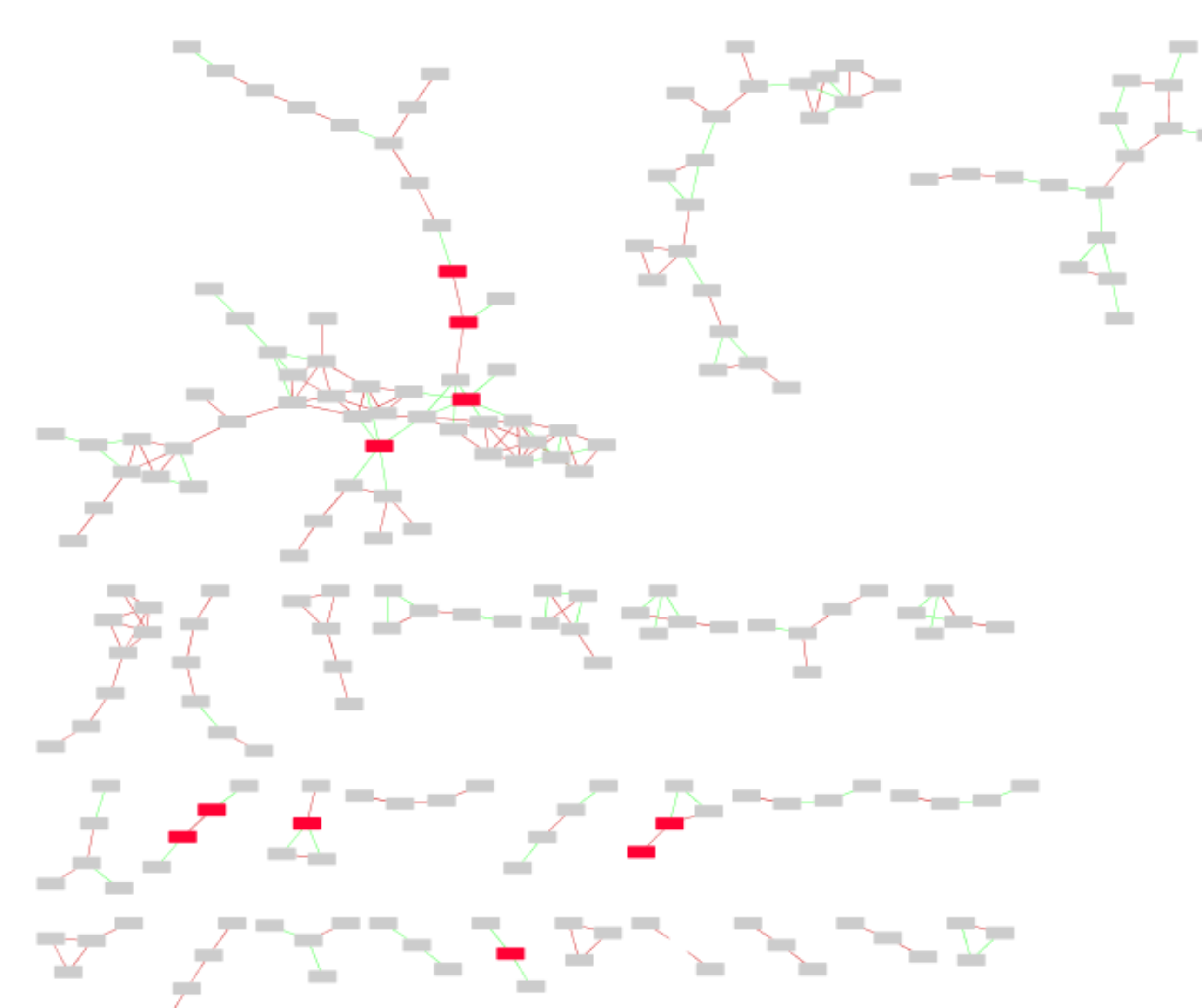
d. Entorhinal cortex



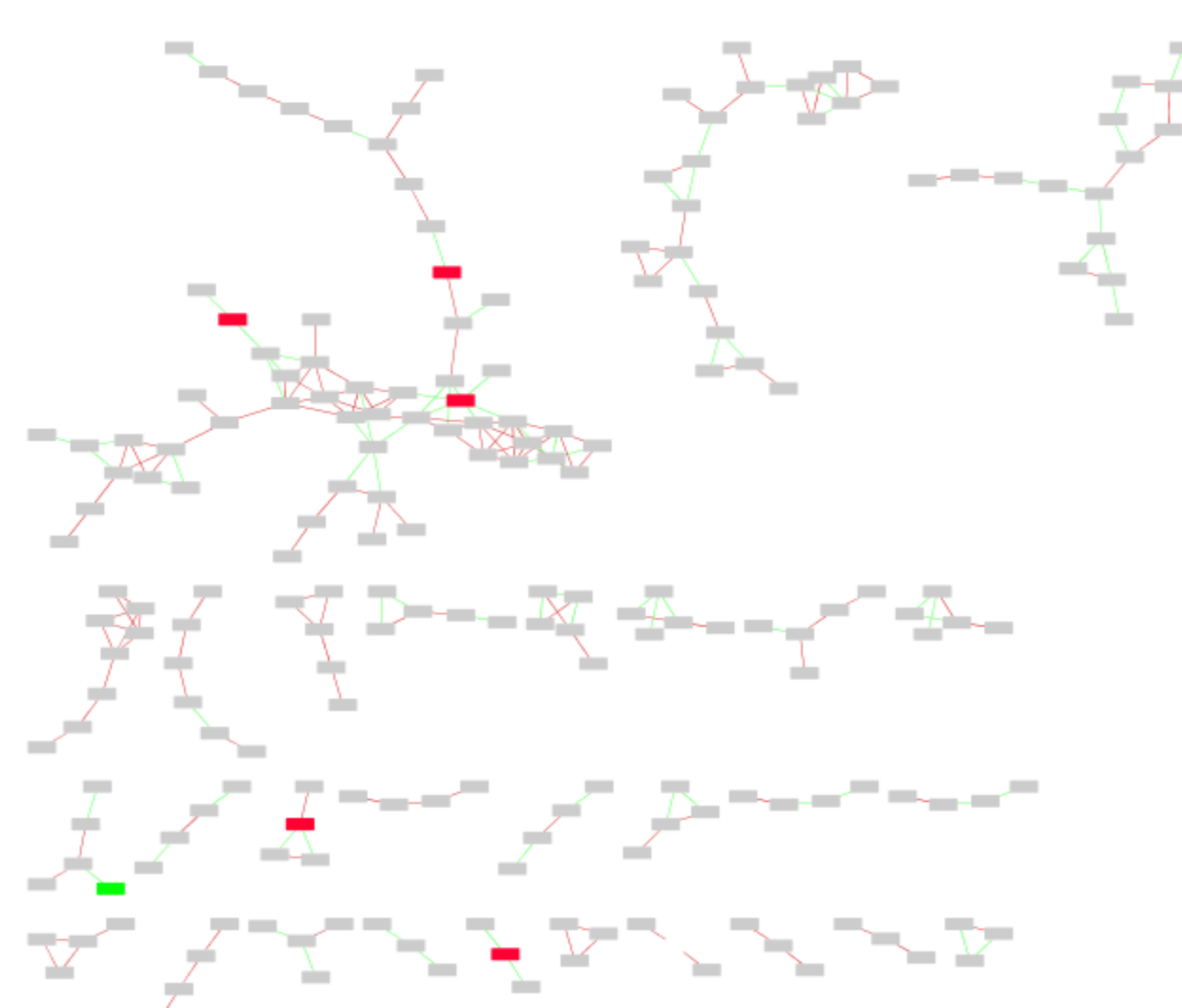
e. Cingulate Gyrus



f. Motor Cortex



g. Sensory Cortex



h. Cerebellum

