

Supplementary figures and tables

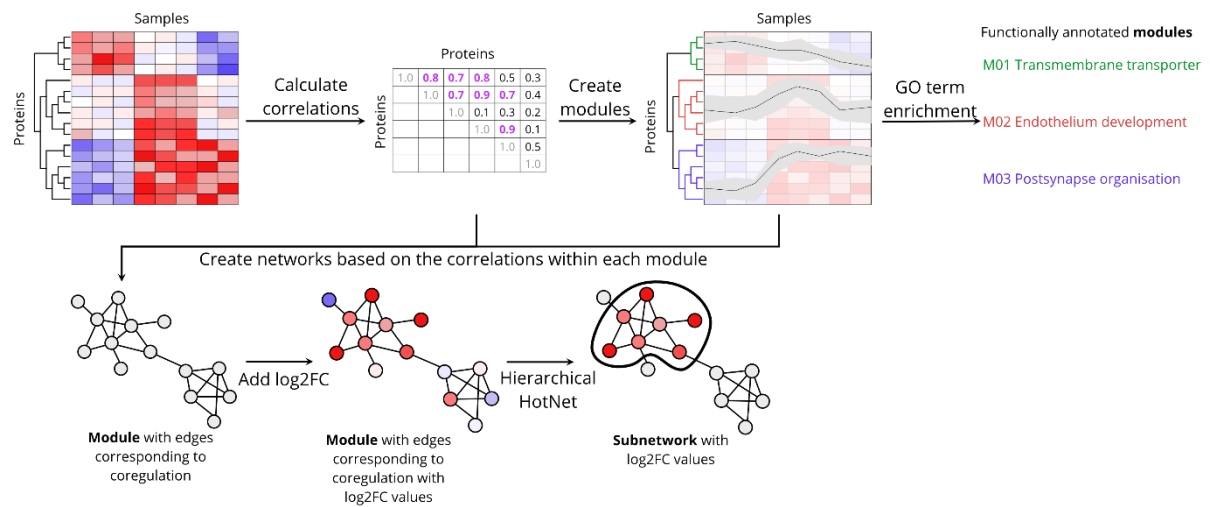


Figure S1 Co-expression analysis and submodule identification workflow.

Proteins were clustered into modules based on expression level correlations. GO term enrichment analysis was used to identify the molecular function most significantly enriched within each module of co-regulated proteins. Within each module, a submodule of most highly and significantly differentially regulated proteins was identified.

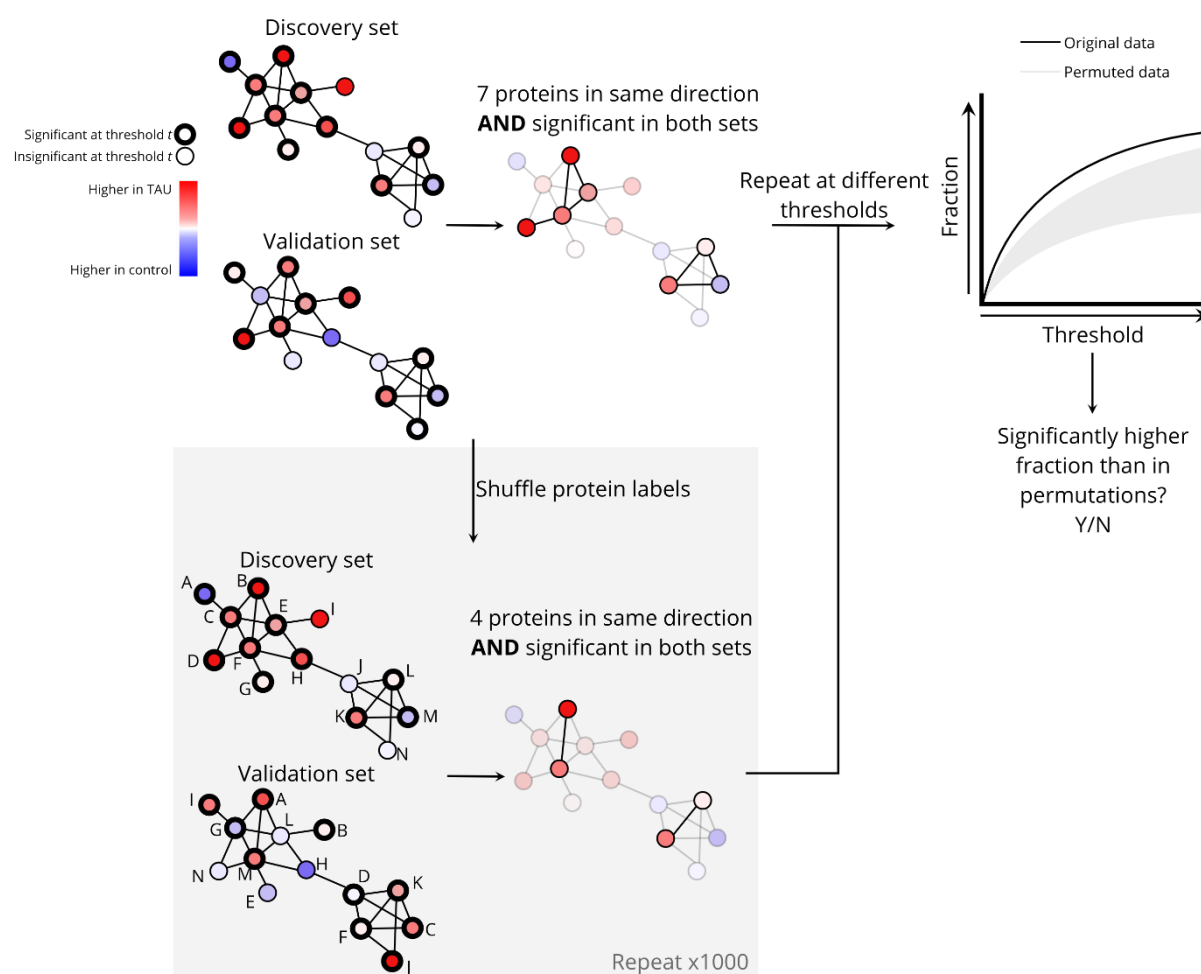


Figure S2 Workflow for module validation

For each module, the log2FC expression values of FTLD-tau versus NHC calculated in the discovery dataset was evaluated in the validation data set. The fraction of proteins that was dysregulated in the same direction in both datasets and that had a p-value lower than a certain threshold in both datasets was calculated. Subsequently, the protein labels in the validation data set were shuffled and the same fraction was calculated. This permutation was repeated 1,000 times. If the fraction of proteins was significantly higher in the comparison with the correct labels than in the comparison with the permuted labels (GESS score (see Methods) taken as the average of the P-values at thresholds 0.1, 0.2, 0.5, 0.8), the module was considered validated.

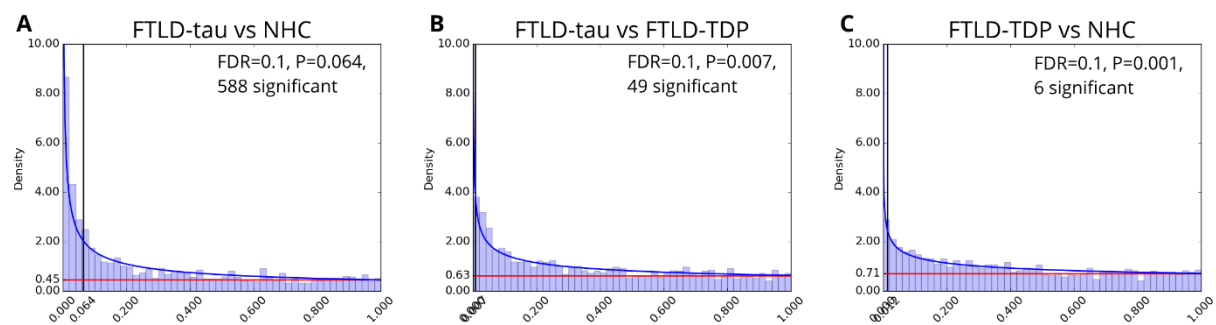


Figure S3 Density distribution of the P-values (x-axis) for pairwise proteome comparisons (A) FTLD-tau versus NHC; (B) FTLD-tau versus FTLD-TDP; (C) FTLD-TDP versus NHC. A beta-uniform mixture (BUM) model was fitted (red and curved black lines) to the distributions to determine the P-value thresholds required for different FDR values (vertical black lines are used to indicate the thresholds at FDR=0.1).

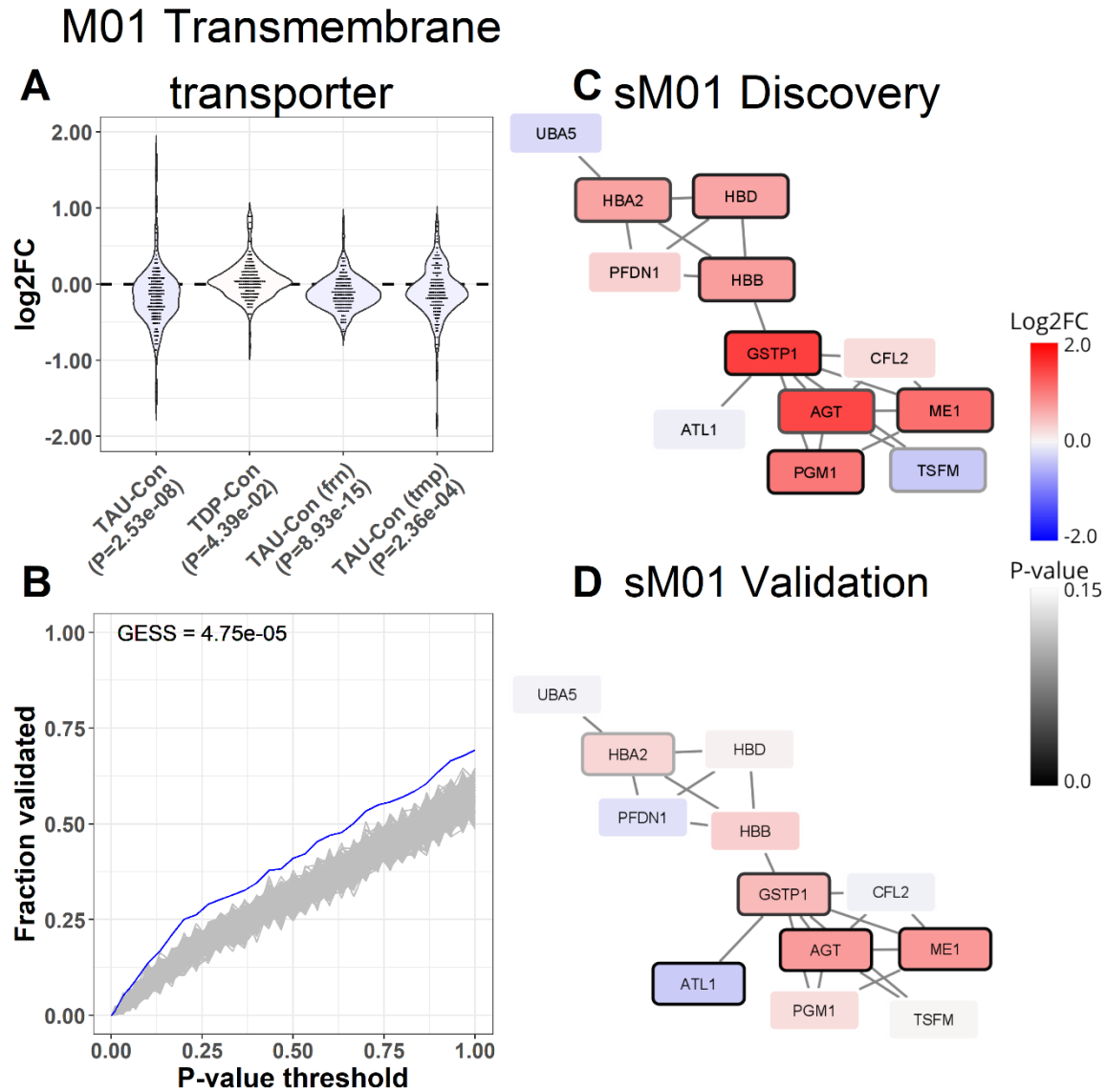
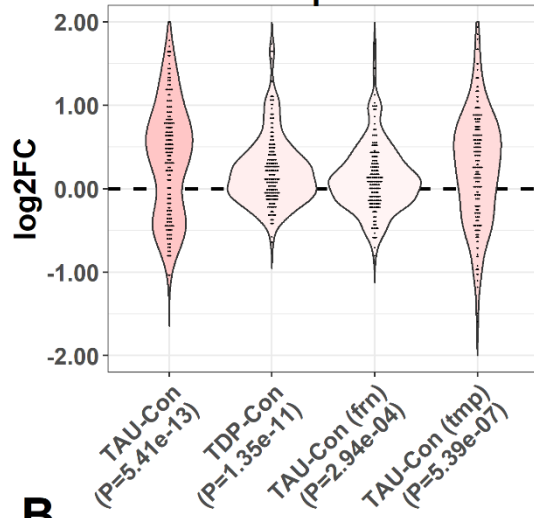


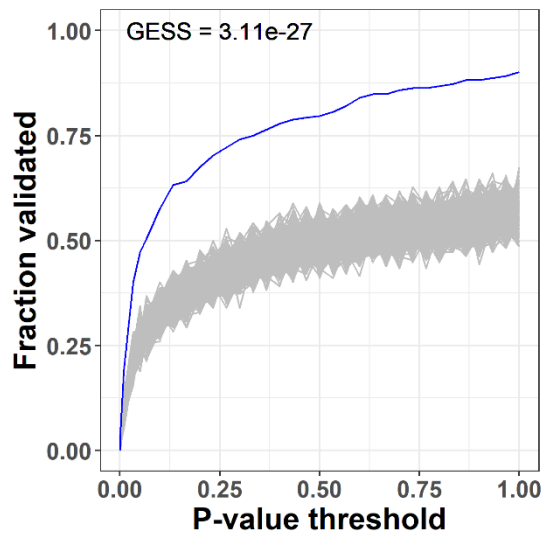
Figure S4-S16 Modules and subnetworks of coregulated proteins that are dysregulated in FTLD-tau and FTLD-TDP for each of the modules not discussed in the main manuscript. **A** Density curves of the log₂ fold-change (log₂FC) values of all the proteins belonging to a module in the discovery data set and the validation data set. The p-value indicates the significance of the median log₂FC difference compared to NHC. **B** Permutation test to validate the modules identified in the discovery data set. A module was validated if the log₂FC measured in the validation data set was in the same direction as in the one measured in the discovery data set, and if the p-value was below the threshold (blue line) in both the discovery and validation data sets. We used a permutation test, where we repeated this procedure with the protein labels in the validation data set randomly reassigned 1,000x (gray lines). The module was considered validated if the average p-value at thresholds 0.1, 0.5, 0.5, and 0.8 was lower than 0.05 (GESS score). Fraction validation: fraction of the proteins in the module that have a log₂FC in the same direction as in the discover data set. **C** Subnetworks of most highly and significantly dysregulated proteins within the corresponding module of co-regulated proteins in the discovery data set. Red nodes indicate a positive log₂FC and blue nodes a negative log₂FC. The darkness of the borders reflects the significance. **D** Subnetworks of most highly and significantly dysregulated proteins in the validation data set. Red nodes indicate a positive log₂FC and blue nodes a negative log₂FC. The darkness of the borders reflects the significance.

M02 Endothelium

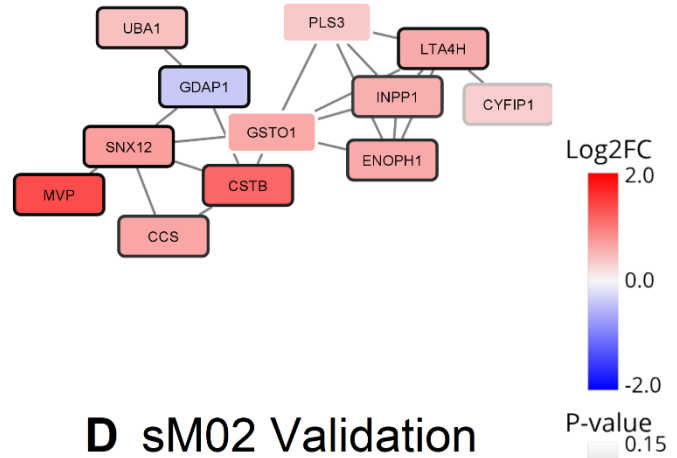
A development



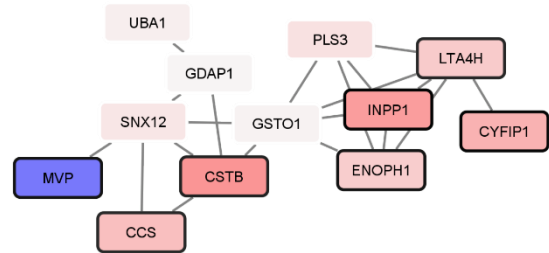
B



C sM02 Discovery

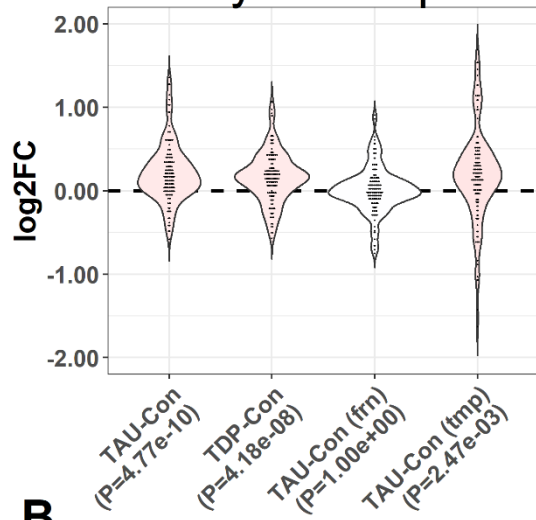


D sM02 Validation

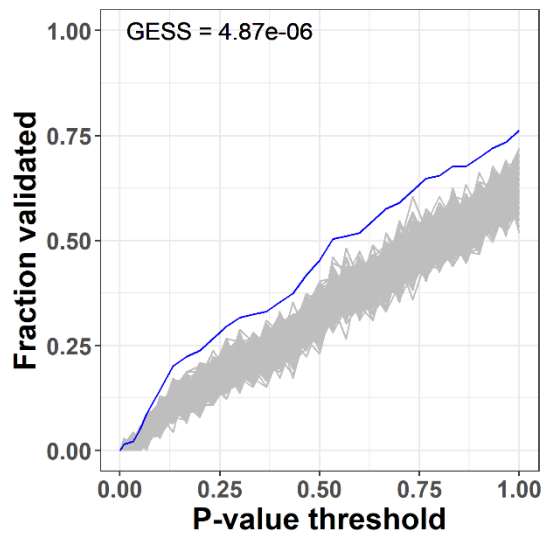


M05 Cell differentiation

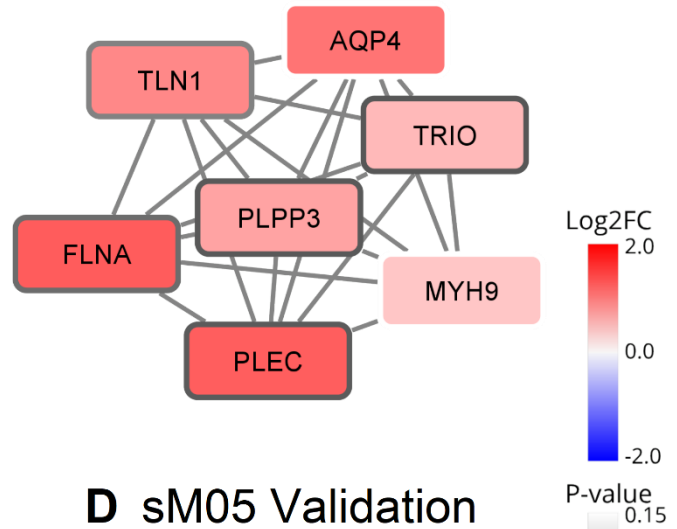
A in kidney development



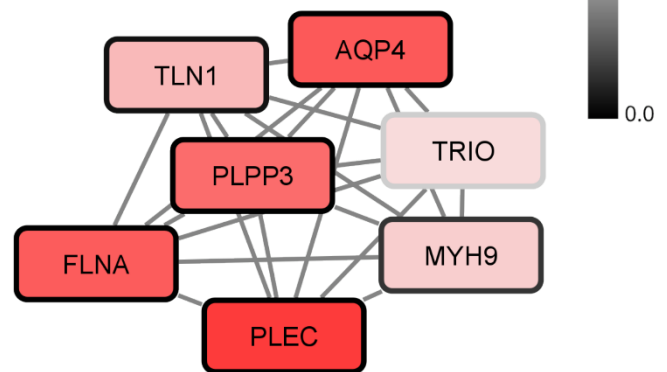
B



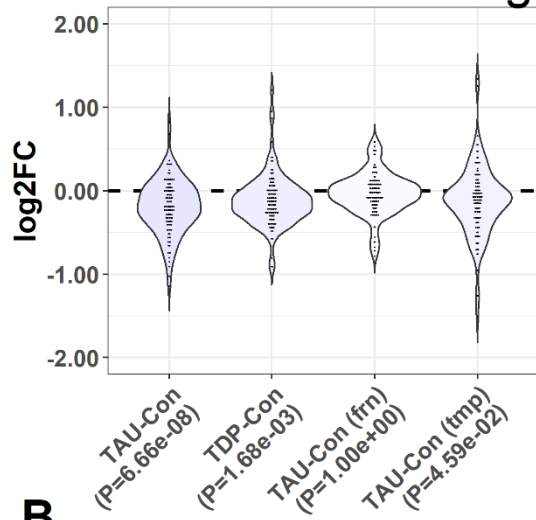
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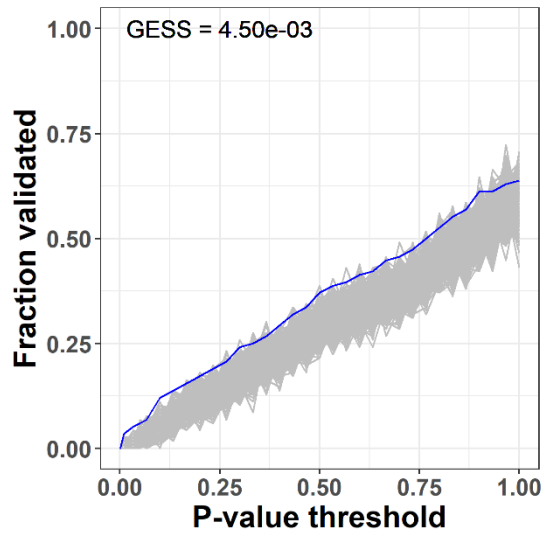
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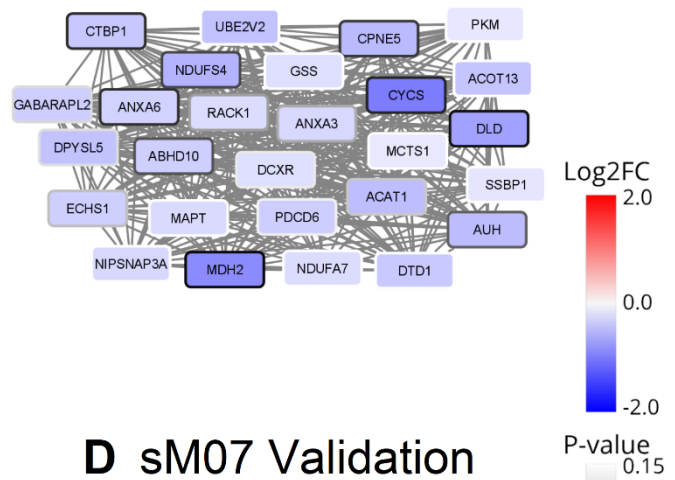
A M07 Filamin binding



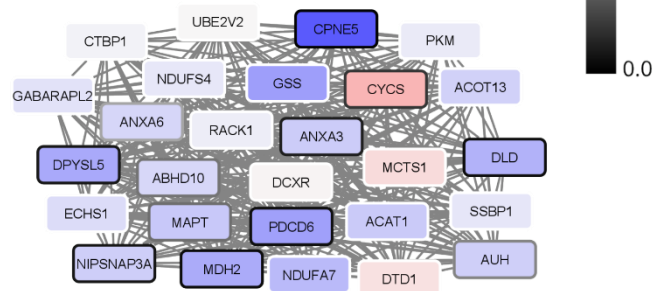
B



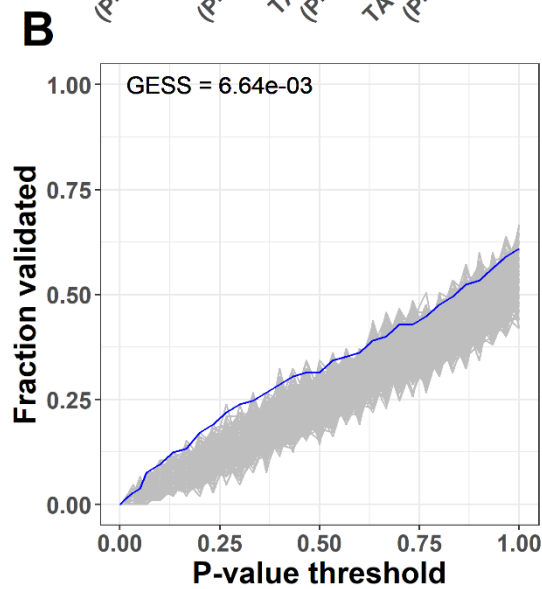
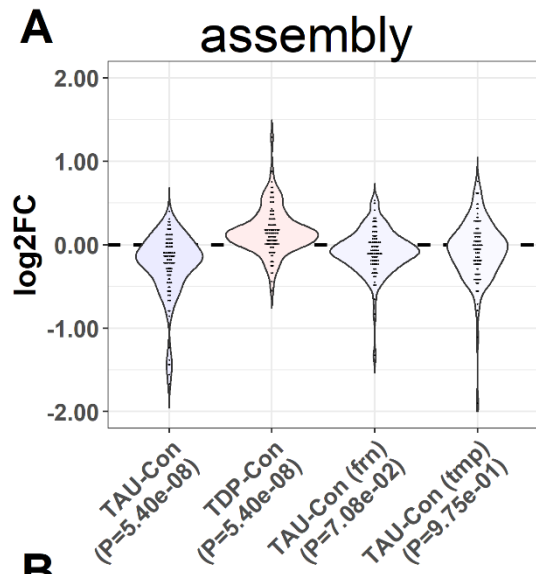
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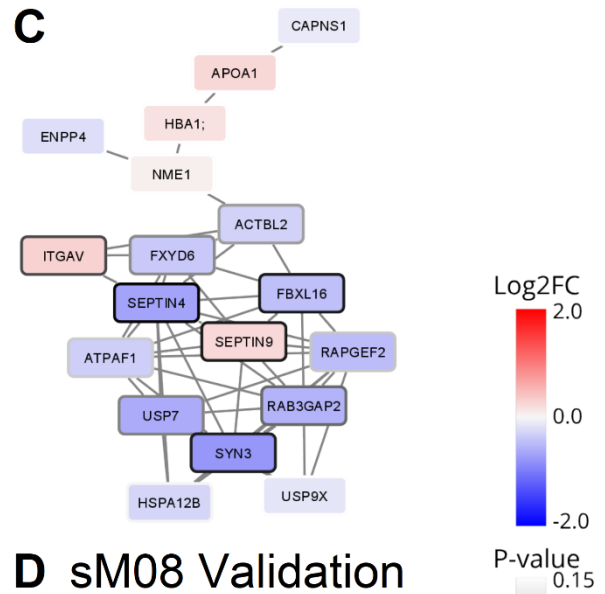
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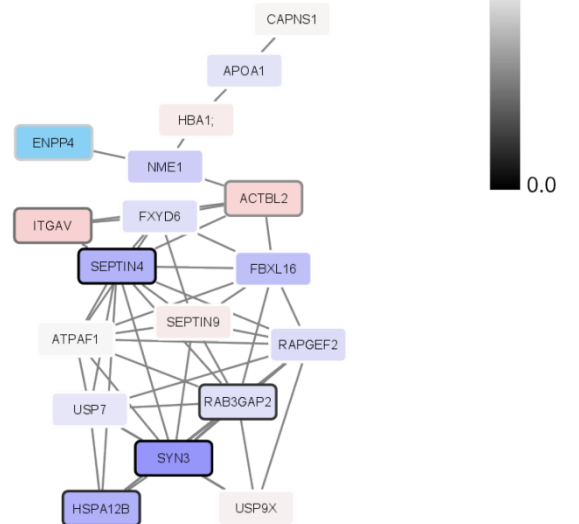
M08 Actin filament assembly



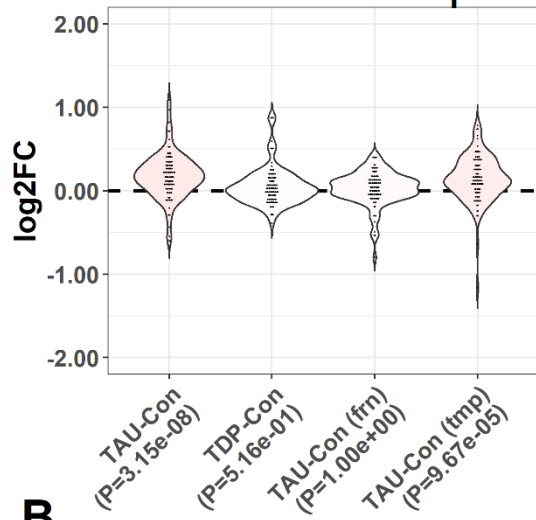
sM08 Discovery



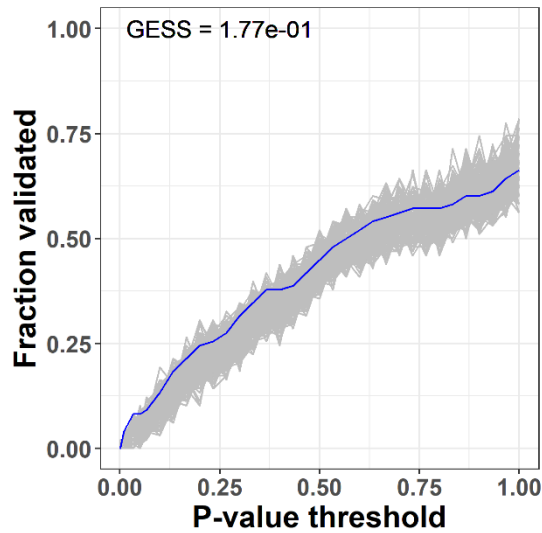
D sM08 Validation



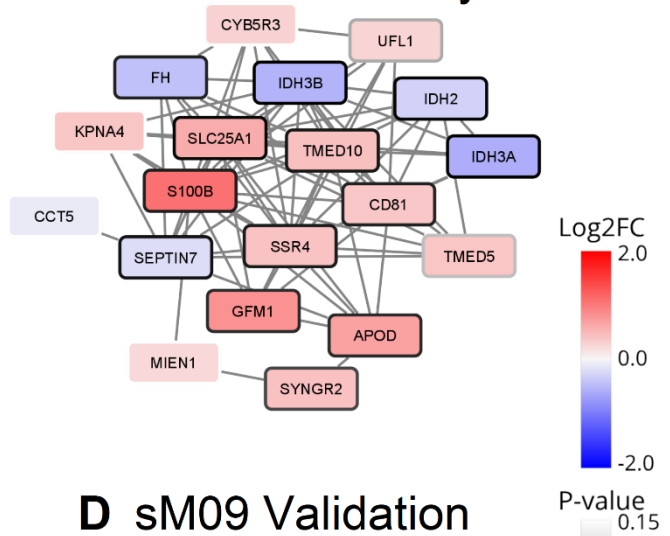
A M09 Viral transcription



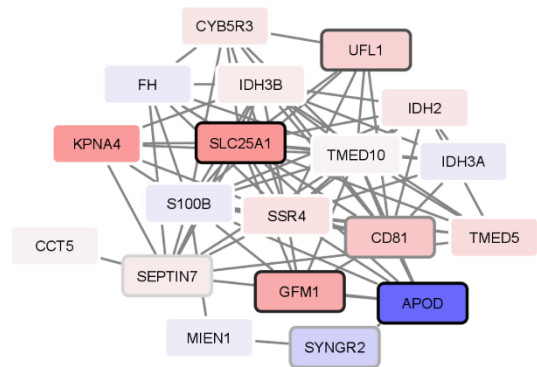
B

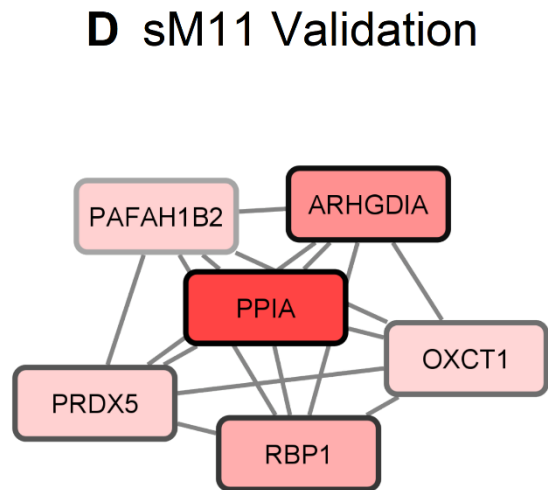
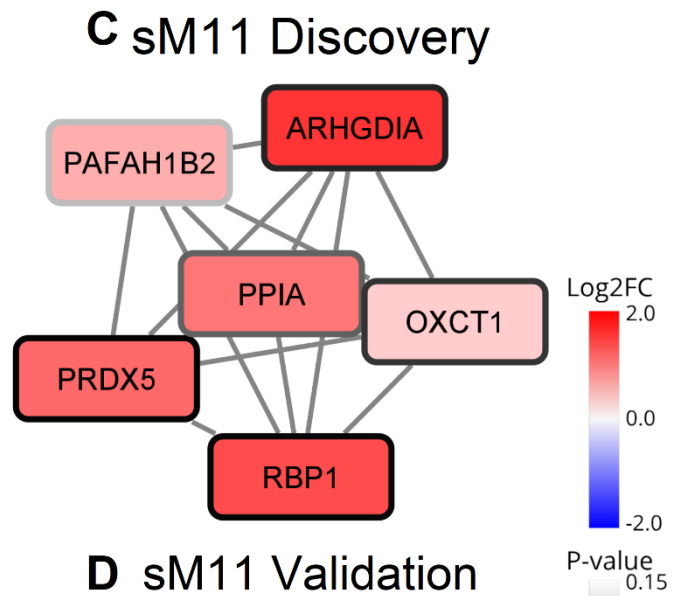
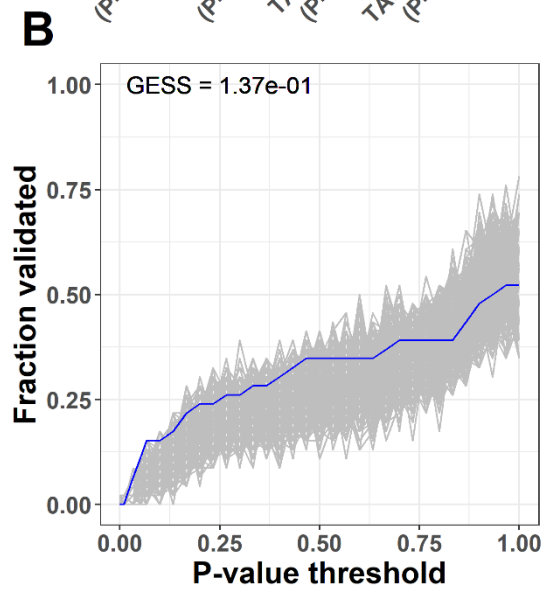
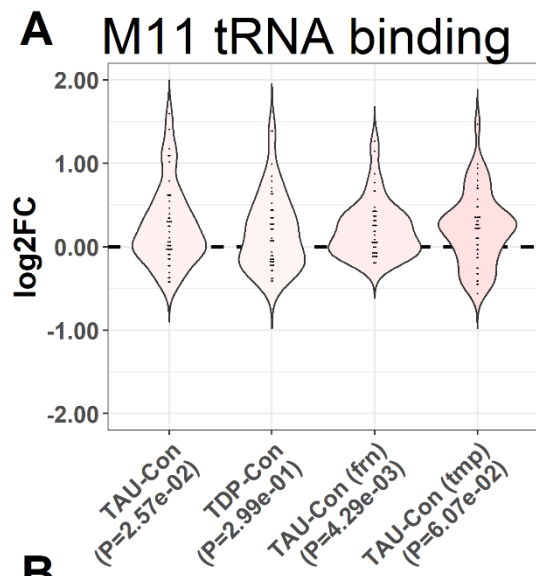


C sM09 Discovery



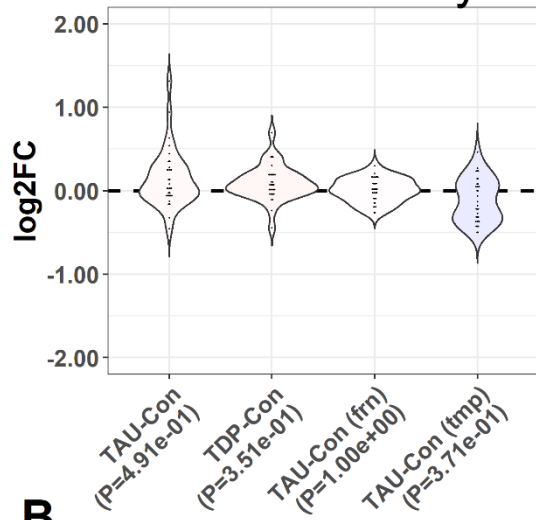
D sM09 Validation



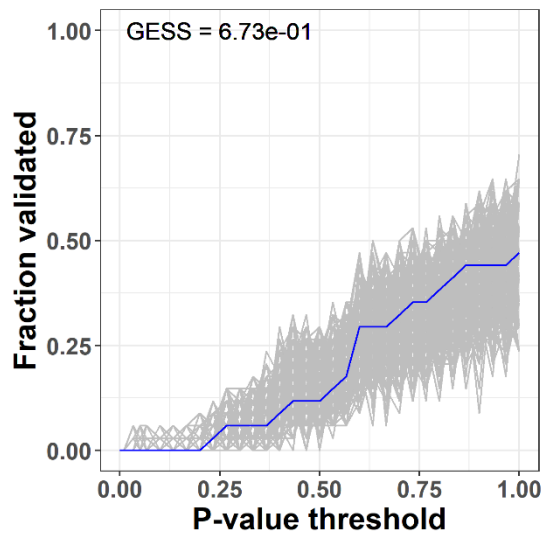


M12 GDP-dissociation

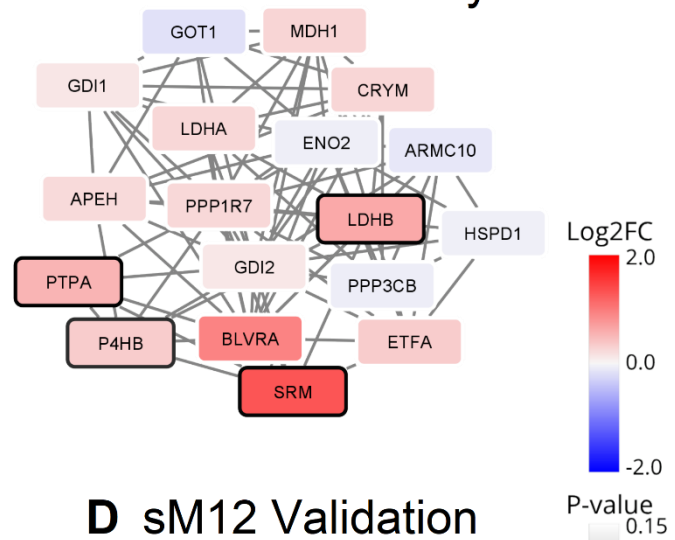
A inhibitor activity



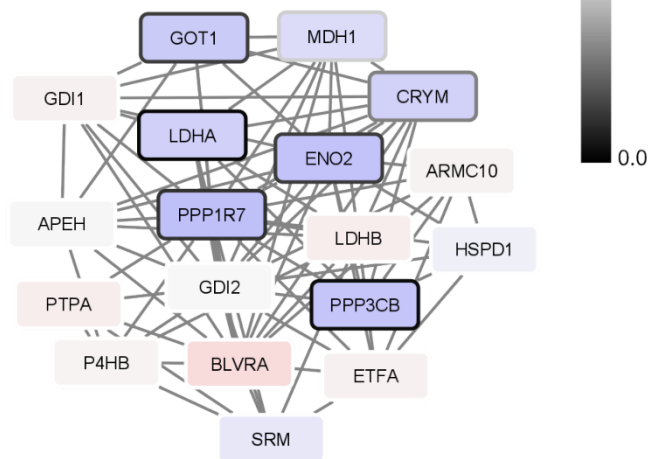
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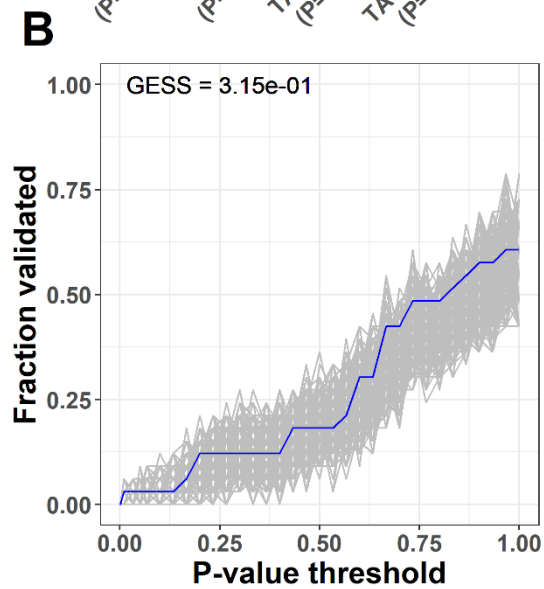
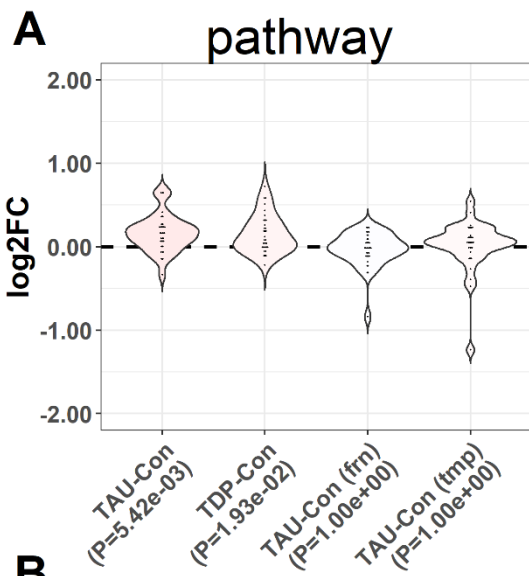
C sM12 Discovery



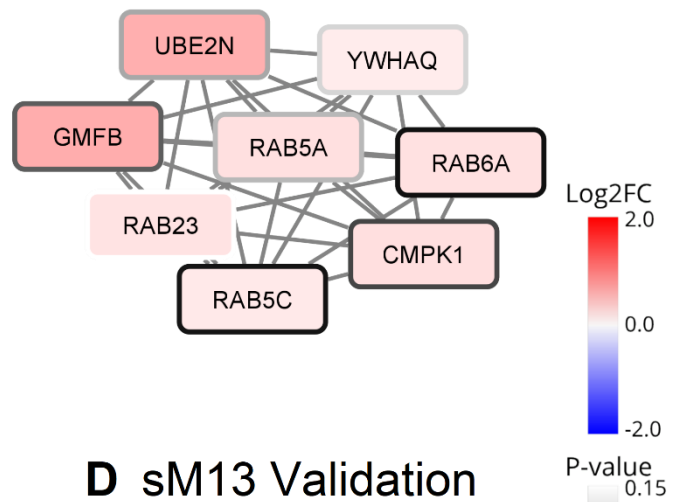
D sM12 Validation



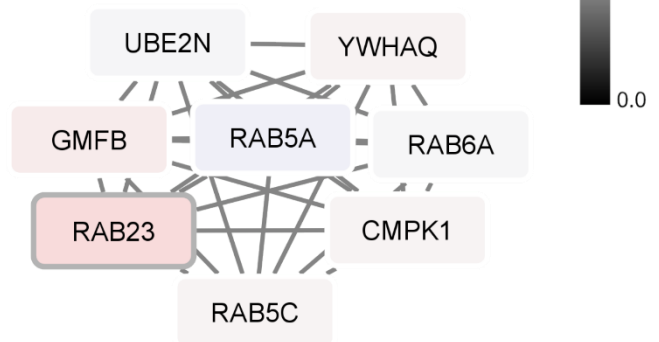
M13 Apoptotic signaling



C sM13 Discovery

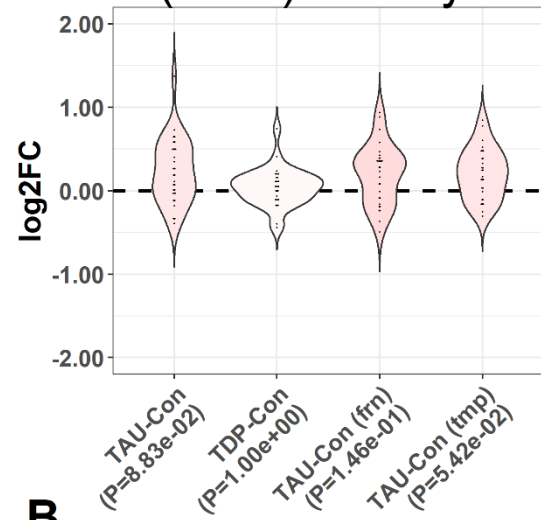


D sM13 Validation

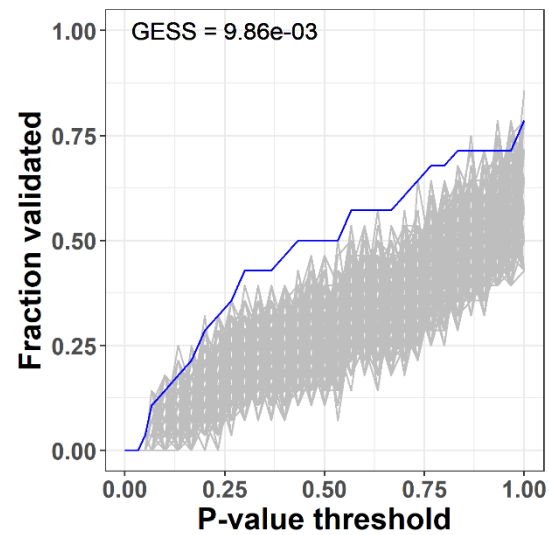


14 Aldehyde dehydrogenase

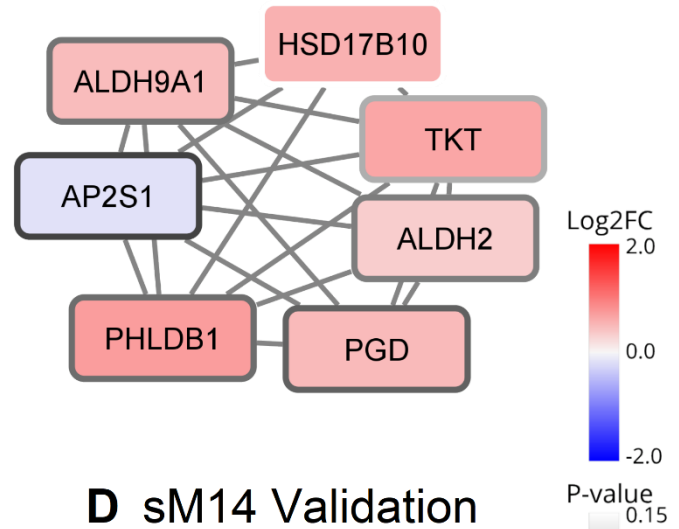
A (NAD) activity



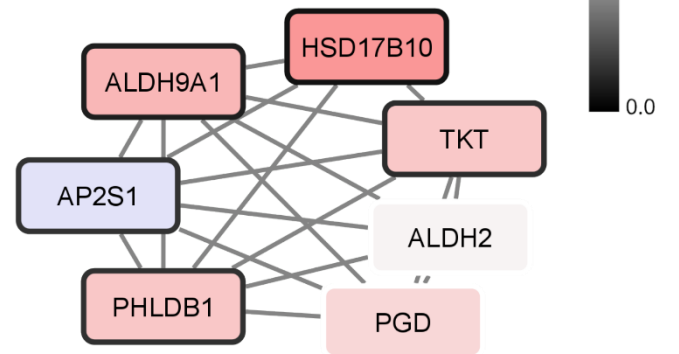
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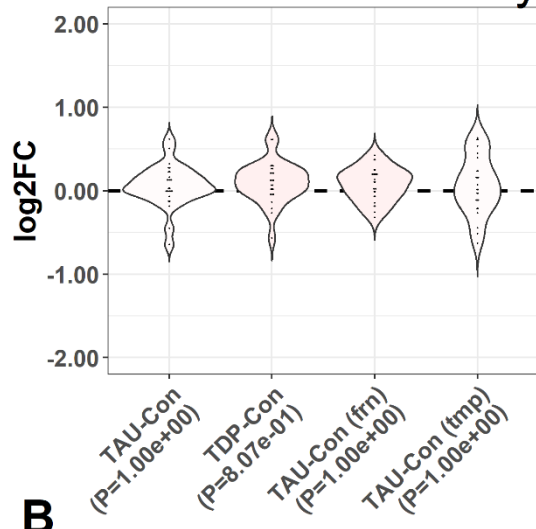
C sM14 Discovery



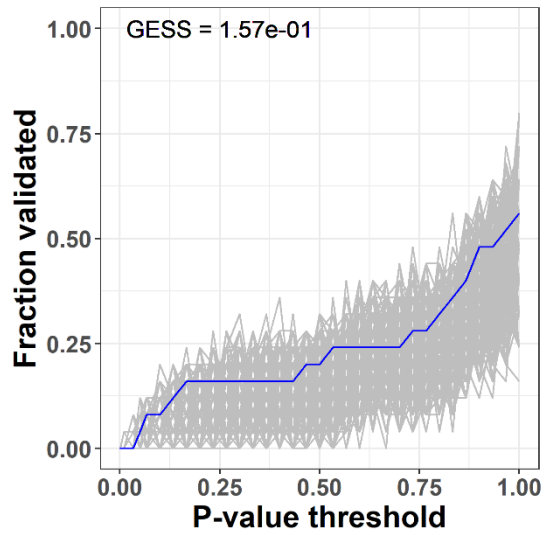
D sM14 Validation



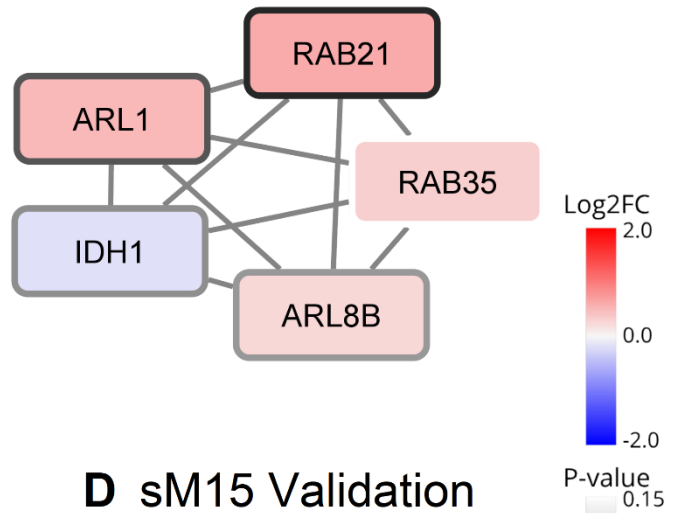
A M15 GTPase activity



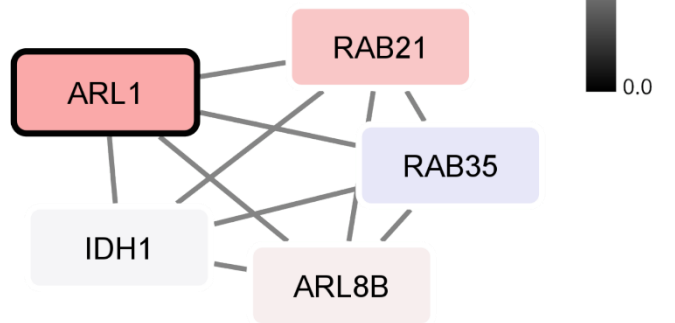
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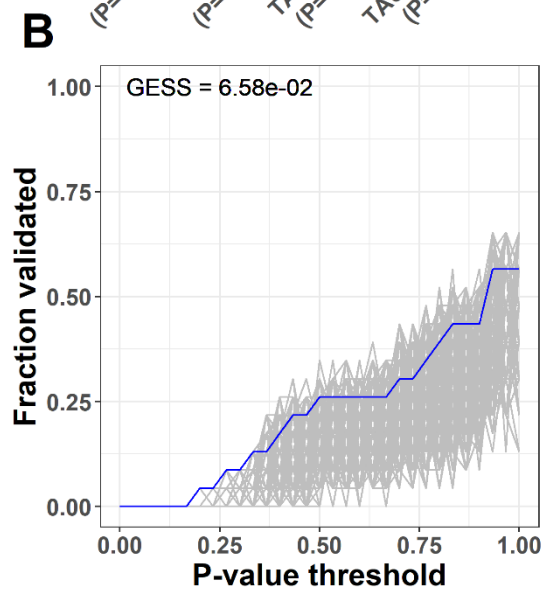
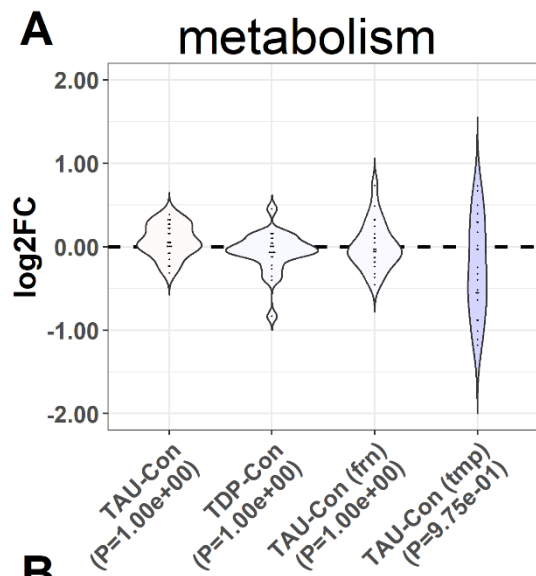
C sM15 Discovery



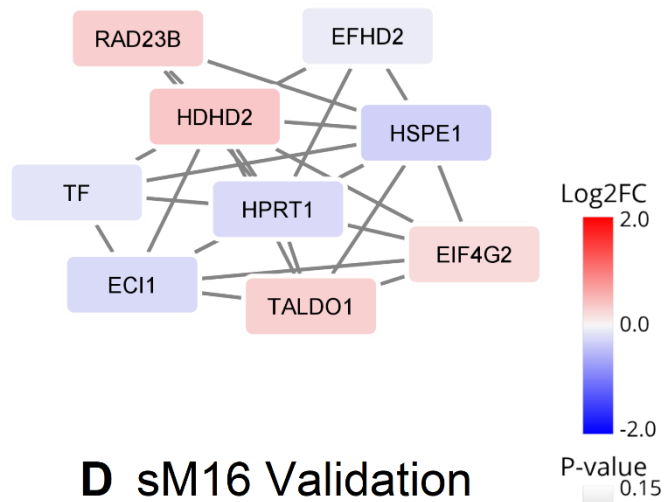
D sM15 Validation



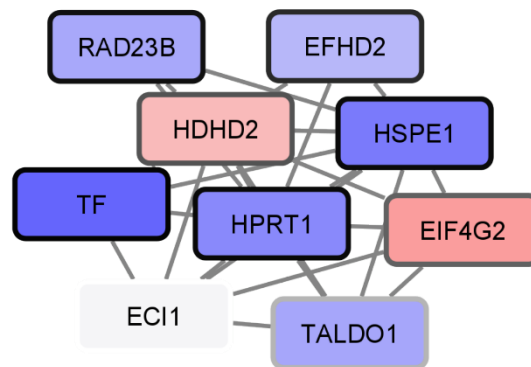
M16 Purine metabolism



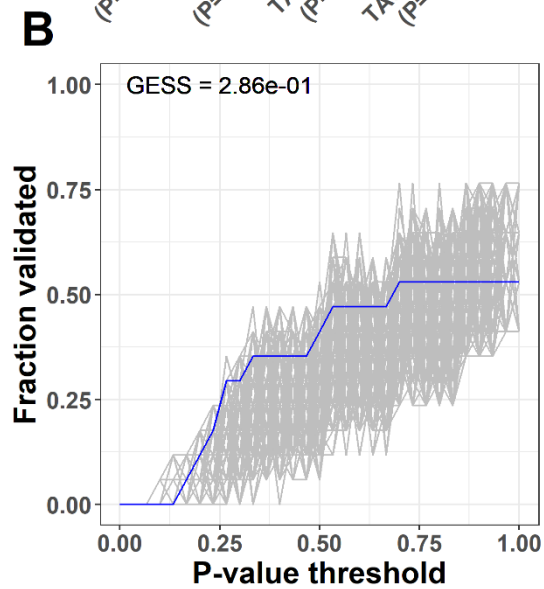
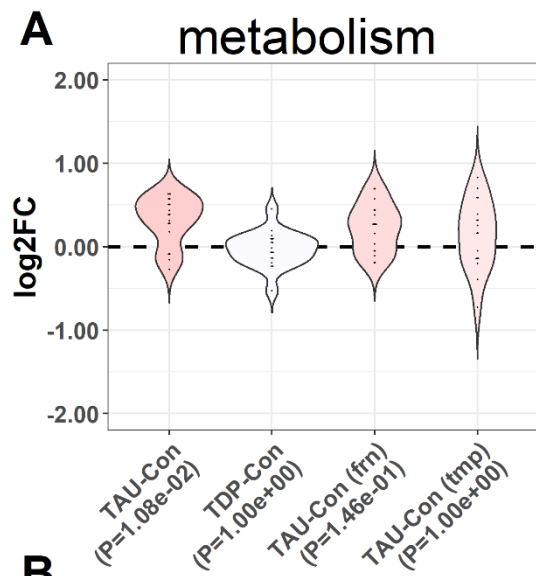
C sM16 Discovery



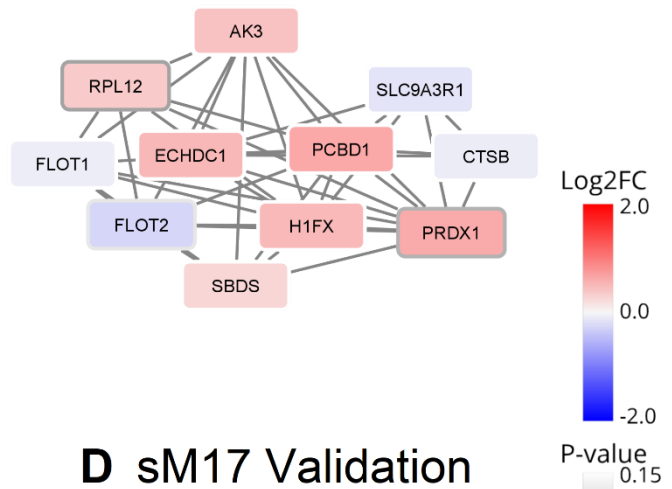
D sM16 Validation



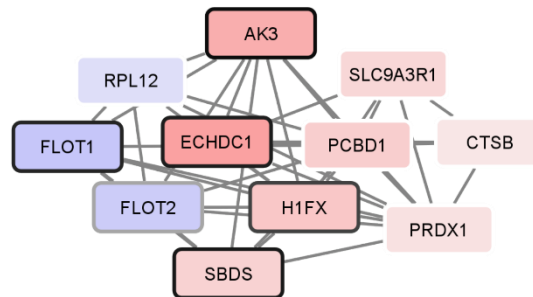
M17 Aromatic aa metabolism



C sM17 Discovery



D sM17 Validation



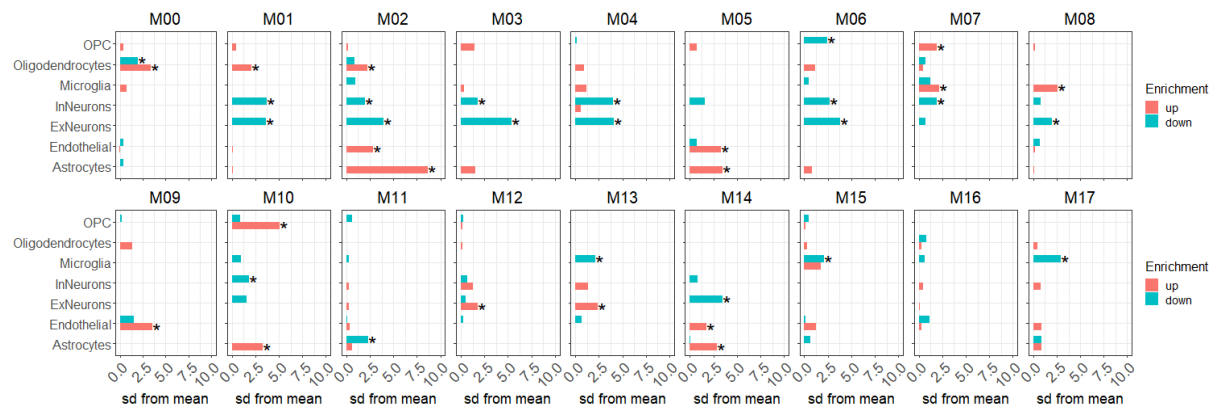


Figure S17 Cell type enrichment in all modules. Cell type enrichment in each of the modules was determined based on the log2FC values of specific CNS cell type protein markers. Included cell types are astrocytes, endothelial cells, excitatory neurons, inhibitory neurons, microglia, oligodendrocytes and oligodendrocyte precursor cells. Enrichment up (red bar) means that the markers for that cell type had high log2FC values in the FTLD-tau samples compared to the control group, enrichment down (blue bar) means that the markers for that cell type had low log2FC values in the FTLD-tau samples compared to the control group.

Table S1 Discovery and validation cohorts. mfg, medial frontal gyrus; temp, temporal lobe; NHC, neurologically healthy control; AD, autosomal dominant; S, sporadic; Fa, familial; NA, not applicable; f, female; m, male; PMI, postmortem interval in hours and minutes. Age is in years.

COHORT	ANATOMICAL REGIONS	PATHOLOGY (TDP SUBTYPE)	GENOTYPE (MUTATION)	AGE	SEX	PMI
DISCOVERY (N=26)	mfg	Tau	AD (MAPT)	68	m	05:25
	mfg	Tau	AD (MAPT)	54	F	05:40
	mfg	Tau	AD (MAPT)	64	F	05:10
	mfg	Tau	AD (MAPT)	46	M	05:35
	mfg	Tau	AD (MAPT)	61	F	06:00
	mfg	Tau	AD (MAPT)	56	m	04:05
	mfg	TDP43 (B)	AD (C9ORF)	63	m	04:45
	mfg	TDP43 (ND)	AD (C9ORF)	58	m	17:30
	mfg	TDP43 (C)	AD (C9ORF)	64	m	06:20
	mfg	TDP43 (B)	AD (C9ORF)	52	f	08:15
	mfg	TDP43 (ND)	AD (C9ORF)	60	f	06:20
	mfg	TDP43 (A)	AD (C9ORF)	70	f	04:40
	mfg	TDP43 (A)	AD (C9ORF)	75	f	05:15
	mfg	TDP43 (B)	AD (C9ORF)	75	m	06:25
	mfg	TDP43 (A)	AD (GRN)	69	f	06:50
	mfg	TDP43 (A)	AD (GRN)	59	m	04:45
	mfg	TDP43 (A)	S	57	f	07:20
	mfg	TDP43 (C)	S	68	f	06:35
	mfg	TDP43 (C)	S	69	M	06:45
	mfg	TDP43 (C)	Fa (none identified)	67	M	05:45
	mfg	TDP43 (C)	Fa (none identified)	69	m	05:20
	mfg	NHC	NA	82	M	12:55
	mfg	NHC	NA	75	F	07:45
	mfg	NHC	NA	55	M	07:15
	mfg	NHC	NA	57	F	07:40
	mfg	NHC	NA	60	f	08:10
VALIDATION (N=24)	mfg and temp	Tau	AD (MAPT)	66	M	05:00
	mfg and temp	Tau	AD (MAPT)	51	M	04:25
	mfg and temp	Tau	AD (MAPT)	60	M	05:23
	mfg and temp	Tau	AD (MAPT)	64	F	07:30
	mfg and temp	Tau	AD (MAPT)	60	M	04:40
	mfg and temp	Tau	AD (MAPT)	67	F	04:10
	mfg and temp	Tau	AD (MAPT)	75	F	07:25
	mfg and temp	Tau	AD (MAPT)	66	F	06:40
	mfg and temp	Tau	AD (MAPT)	54	F	07:30
	mfg and temp	Tau	AD (MAPT)	49	M	05:10
	mfg and temp	Tau	AD (MAPT)	52	M	11:30
	temp	Tau	AD (MAPT)	64	M	06:25
	temp	Tau	AD (MAPT)	46	M	05:35
	mfg and temp	NHC	NA	91	M	08:00
	mfg and temp	NHC	NA	84	F	05:36
	mfg and temp	NHC	NA	83	F	06:03
	mfg and temp	NHC	NA	60	F	07:30
	mfg and temp	NHC	NA	70	M	03:35
	mfg and temp	NHC	NA	83	M	05:45
	mfg and temp	NHC	NA	76	F	07:15
	mfg and temp	NHC	NA	89	F	05:40
	mfg	NHC	NA	82	M	05:50
	mfg	NHC	NA	87	F	07:00
	mfg	NHC	NA	75	F	05:25

Table S2 Module summary. Module identity (number), module size and largest submodule size (number of proteins in the module/submodule), and validation in the FTLT-tau medial frontal gyrus cortex or temporal cortex data set of the validation cohort. Module function indicates the Gene ontology (GO) term enrichment analysis biological function most significantly associated with module

#	Module size	Subnetwork size	Module validated in the frontal cortex validation data set	Module validated in the temporal cortex validation data set	Module function
M00	26	5	No	No	--- (not a module)
M01	272	12	Yes	Yes	Transmembrane transporter activity
M02	233	12	Yes	Yes	Chromatin regulation
M03	212	13	Yes	Yes	Postsynapse organisation
M04	159	26	Yes	Yes	Oxidative phosphorylation
M05	148	7	No	Yes	Protein-containing complex binding
M06	129	37	Yes	Yes	Regulation of vesicle-mediated transport
M07	124	27	No	Yes	Filamin binding
M08	121	18	No	No	Positive regulation of actin filament bundle assembly
M09	103	19	No	Yes	Co-translational protein targeting to membrane
M10	56	25	Yes	Yes	Phenylalanine-tRNA ligase activity
M11	49	6	Yes	No	tRNA binding
M12	34	18	No	No	Carbohydrate metabolic process
M13	34	8	No	No	Apoptotic signalling pathway
M14	30	7	Yes	Yes	Aldehyde dehydrogenase (NAD) activity
M15	28	5	No	No	GTPase activity
M16	25	9	No	No	Superoxide metabolic process
M17	18	11	No	No	Oxidoreductase activity