main

August 11, 2021

1 Comparison with public TWAS associations for SZ

```
[1]: import urllib
import numpy as np
import pandas as pd
from html.parser import HTMLParser
from scipy.stats import fisher_exact
from html.entities import name2codepoint
[2]: class MyHTMLParser(HTMLParser):
    def __init__(self):
        super().__init__()
```

```
class MyHTMLParser(HTMLParser):
    def __init__(self):
        super().__init__()
        self.genes = []

def handle_starttag(self, tag, attrs):
    for attr in attrs:
        if attr[0] == 'href' and attr[1].startswith('/genes/'):
            gene = [attr[1][len('/genes/'):]]
        self.genes += gene
```

1.1 MHC genes

[3]: 136

1.2 Caudate specific of BrainSeq brain regions

```
[4]: fn = "../../libd_twas_comparison/_m/caudate_only_twasList_genes.txt" caudate = pd.read_csv(fn, sep='\t') caudate.shape
```

```
[4]: (445, 47)
```

```
[5]: len(set(caudate.ID) - set(mhc_genes))
 [5]: 356
 [6]: caudate_noMHC = caudate[(caudate["ID"].isin(list(set(caudate.ID) -__
      ⇒set(mhc_genes))))].copy()
      caudate_noMHC.iloc[0:2, 0:5]
 [6]:
                    FILE
                              ID
                                  CHR_TWAS
                                                   P0
                                                                P1
      2 ENSG0000100138
                           SNU13
                                                        41690504.0
                                        22
                                             41673930
      3 ENSG00000204963 PCDHA7
                                         5 140834248 141012344.0
     1.3 TWAS Hub comparison
     1.3.1 Schizophrenia 2014
 [7]: parser = MyHTMLParser()
      html_str = urllib.request.urlopen('http://twas-hub.org/traits/Schizophrenia/').
      →read().decode()
      parser.feed(html_str)
      scz_2014 = np.unique(parser.genes)
 [8]: scz_2014.shape
 [8]: (49,)
     1.3.2 Schizophrenia 2018
 [9]: parser = MyHTMLParser()
      html_str = urllib.request.urlopen('http://twas-hub.org/traits/SCZ_2018/').
      →read().decode()
      parser.feed(html str)
      scz_2018 = np.unique(parser.genes)
     1.3.3 TWAS hub comparison
[10]: twas_hub = np.unique(np.append(scz_2014, scz_2018))
      len(twas_hub)
[10]: 67
[11]: twas_hub
[11]: array(['AC011816.1', 'AC103965.1', 'ACTR5', 'AKT3', 'ALMS1P', 'ANKRD44',
             'AS3MT', 'ATG13', 'BAI1', 'C12orf65', 'C17orf39', 'C2orf47',
             'CACNA2D4', 'CEP170', 'CHRNA5', 'CLCN3', 'CNTN4', 'CPEB1', 'CPNE7',
             'CRELD2', 'CUL3', 'CYSTM1', 'DCP1B', 'ELAC2', 'ERCC8', 'FAM53C',
             'FAM83H', 'FES', 'FLJ10661', 'GIGYF1', 'GMIP', 'HSPD1', 'IK',
```

```
'IMMP2L', 'ITIH4-AS1', 'KIAA0319', 'KLC1', 'LGSN', 'LRRC48',
             'MAD1L1', 'MAP7D1', 'MAPK3', 'MGAT3', 'NAGA', 'NGEF', 'PCCB',
             'PCNX', 'PITPNM2', 'PLEKHO1', 'PPP1R13B', 'PPP1R14B', 'RERE',
             'RP11-981G7.6', 'SDCCAG8', 'SF3B1', 'SLCO4C1', 'SMG6', 'SNAP91',
             'SUGP2', 'THOC7', 'TRIM38', 'TTC14', 'U91328.21', 'VPS29',
             'VPS37A', 'XPNPEP3', 'ZNF318'], dtype='<U12')
[12]: print("There are %d caudate only genes present in the TWAS hub" %
            len(set(caudate.ID) & set(twas_hub)))
      set(caudate.ID) & set(twas_hub)
     There are 8 caudate only genes present in the TWAS hub
[12]: {'AKT3', 'C2orf47', 'ELAC2', 'ERCC8', 'MGAT3', 'PPP1R13B', 'SNAP91', 'VPS29'}
[13]: print("There are %d caudate only genes (no MHC) present in the TWAS hub" %
            len(set(caudate_noMHC.ID) & set(twas_hub)))
      set(caudate.ID) & set(twas_hub)
     There are 8 caudate only genes (no MHC) present in the TWAS hub
[13]: {'AKT3', 'C2orf47', 'ELAC2', 'ERCC8', 'MGAT3', 'PPP1R13B', 'SNAP91', 'VPS29'}
     1.4 Gandal comparison
[14]: gandal = pd.read excel("../h/aat8127_Table_S4.xlsx", sheet_name="SCZ.TWAS")
      gandal_twas = gandal[(gandal['TWAS.Bonferroni'] <= 0.05)].copy()</pre>
      np.sum(gandal.loc[:, 'TWAS.Bonferroni'] < 0.05)</pre>
[14]: 193
[15]: caudate_bonferroni = caudate[(caudate['Bonferroni'] <= 0.05)].copy()</pre>
      np.sum(caudate.Bonferroni <= 0.05)</pre>
[15]: 108
[16]: caudate noMHC bonferroni = caudate noMHC[(caudate noMHC['Bonferroni'] <= 0.05)].
       →copy()
      np.sum(caudate noMHC.Bonferroni <= 0.05)
[16]: 45
[17]: print("There are %d caudate only genes present in the Gandal at Bonferroni < 0.

→05." %

            len(set(caudate_bonferroni.FILE) & set(gandal_twas.GeneID)))
      #list(set(caudate bonferroni.ID) & set(qandal twas.gene name))
      print("There are %d caudate only genes present in the Gandal." %
            len(set(caudate.FILE) & set(gandal_twas.GeneID)))
```

```
np.array(list(set(caudate.FILE) & set(gandal_twas.GeneID)))
     There are 33 caudate only genes present in the Gandal at Bonferroni < 0.05.
     There are 42 caudate only genes present in the Gandal.
[17]: array(['ENSG00000085788', 'ENSG00000110492', 'ENSG00000249839',
             'ENSG00000216901', 'ENSG00000030110', 'ENSG00000198315',
             'ENSG00000262074', 'ENSG00000006744', 'ENSG00000231389',
             'ENSG00000175662', 'ENSG00000219392', 'ENSG00000204963',
             'ENSG00000197238', 'ENSG00000161896', 'ENSG00000124613',
             'ENSG00000185829', 'ENSG00000261353', 'ENSG00000158406',
             'ENSG00000111237', 'ENSG00000189298', 'ENSG00000259404',
             'ENSG00000124610', 'ENSG00000162972', 'ENSG00000168237',
             'ENSG00000100162', 'ENSG00000177096', 'ENSG00000228223',
             'ENSG00000197279', 'ENSG00000186522', 'ENSG00000163938',
             'ENSG00000204252', 'ENSG00000186470', 'ENSG00000088808',
             'ENSG00000272009', 'ENSG00000174939', 'ENSG00000168405',
             'ENSG00000117020', 'ENSG00000219891', 'ENSG00000226314',
             'ENSG00000065609', 'ENSG00000187987', 'ENSG00000205702'],
            dtype='<U15')
[18]: print("There are %d caudate only genes (no MHC) present in the Gandal at ⊔
       →Bonferroni < 0.05." %
            len(set(caudate_noMHC_bonferroni.FILE) & set(gandal_twas.GeneID)))
      #list(set(caudate bonferroni.ID) & set(qandal twas.gene name))
      print("There are %d caudate only genes (no MHC) present in the Gandal." %
            len(set(caudate_noMHC.FILE) & set(gandal_twas.GeneID)))
      np.array(list(set(caudate_noMHC.FILE) & set(gandal_twas.GeneID)))
     There are 16 caudate only genes (no MHC) present in the Gandal at Bonferroni <
     0.05.
     There are 22 caudate only genes (no MHC) present in the Gandal.
[18]: array(['ENSG00000085788', 'ENSG00000110492', 'ENSG00000249839',
             'ENSG00000262074', 'ENSG00000006744', 'ENSG00000175662',
             'ENSG00000204963', 'ENSG00000185829', 'ENSG00000111237',
             'ENSG00000162972', 'ENSG00000259404', 'ENSG00000168237',
             'ENSG00000100162', 'ENSG00000177096', 'ENSG00000186522',
             'ENSG00000163938', 'ENSG00000088808', 'ENSG00000174939',
             'ENSG00000168405', 'ENSG00000117020', 'ENSG00000065609',
             'ENSG00000205702'], dtype='<U15')
     MHC is present within Gandal analysis
[19]: gandal[(gandal["TWAS.Bonferroni"] < 0.05)].shape
[19]: (193, 31)
```

```
[20]: ## Not MHC
      gandal[~(gandal['gene_name'].isin(list(set(mhc_genes)))) &
             (gandal["TWAS.Bonferroni"] < 0.05)].shape
[20]: (174, 31)
[21]: print("There are {} MHC genes within Gandal significant TWAS (Bonferroni < 0.
       →05)!"\
            .format(len(gandal[(gandal['gene_name'].isin(list(set(mhc_genes)))) &
             (gandal["TWAS.Bonferroni"] < 0.05)].gene_name)))</pre>
      gandal[(gandal['gene_name'].isin(list(set(mhc_genes)))) &
             (gandal["TWAS.Bonferroni"] < 0.05)].gene_name
     There are 19 MHC genes within Gandal significant TWAS (Bonferroni < 0.05)!
[21]: 2
              HIST1H4J
              HIST1H3C
      3
      5
             ZSCAN12P1
              HIST1H1A
               ZSCAN23
      10
      15
                BTN3A2
      22
               ZKSCAN3
      29
               ZKSCAN8
      34
                 HCG11
                ZNF165
      38
      39
              HIST1H4H
      59
              ZNF192P1
      70
                 IP6K3
      71
                BTN2A2
      74
               HLA-DOA
      94
              HIST1H3A
      124
                ZNF391
      146
              HLA-DPA1
      174
                  BAK1
      Name: gene_name, dtype: object
     1.4.1 Calculated enrichment with Gandal
[22]: caudate.shape
[22]: (445, 47)
[23]: dft = caudate.loc[:, ['FILE', 'ID', 'Bonferroni']]\
                   .merge(gandal.loc[:, ['GeneID', 'gene_name', 'TWAS.Bonferroni']],
                          left_on='FILE', right_on='GeneID',
                          suffixes=["_Benjamin", "_Gandal"])
      dft.shape
```

```
[23]: (312, 6)
[24]: table =
               [[np.sum((dft['Bonferroni']<0.05) & ((dft['TWAS.Bonferroni']<.05))),
                np.sum((dft['Bonferroni']<0.05) & ((dft['TWAS.Bonferroni']>=.05)))],
                [np.sum((dft['Bonferroni']>=0.05) & ((dft['TWAS.Bonferroni']<.05))),</pre>
                np.sum((dft['Bonferroni']>=0.05) & ((dft['TWAS.Bonferroni']>=.05)))]]
     print(table)
     fisher_exact(table)
     [[33, 31], [9, 239]]
[24]: (28.268817204301076, 6.91682701358258e-19)
     1.4.2 Extract and save overlapping genes
     Bonferroni < 0.05
[25]: overlapping_twas = np.append(np.
      →array(caudate_bonferroni[(caudate_bonferroni['FILE'].
      →isin(list(set(caudate_bonferroni.FILE) &
      ⇒set(gandal_twas.GeneID))))].ID),
                                  np.array(list(set(caudate bonferroni.ID) &
      →set(twas hub))))
     len(overlapping_twas)
[25]: 37
[26]: caudate_bonferroni[~(caudate_bonferroni['ID'].isin(overlapping_twas))].
      →to_csv("caudate_only_twasList_genes_bonferroni.txt",
                                                             sep='\t', index=False)
     caudate_bonferroni[~(caudate_bonferroni['ID'].isin(overlapping_twas))].shape
[26]: (75, 47)
[27]: drop_caudate = caudate_bonferroni[~(caudate_bonferroni['ID'].
      →isin(overlapping_twas))].copy()
     drop_caudate[(drop_caudate['P'] > 5e-8)].shape
[27]: (6, 47)
[28]: drop_caudate[(drop_caudate['P'] > 5e-8)].sort_values('FDR')\
                                             .loc[:, ['ID', 'our_snp_id', _
      [28]:
                       ID
                                                               FDR
                                                                               Ρ
                                                                                 \
                                    our_snp_id CHR_TWAS
     192
                            chr15:43782086:A:G
                                                      15 0.000062 2.750000e-06
                     STRC
     132
                  ANKRD45
                            chr1:173743105:T:C
                                                       1 0.000129 5.050000e-07
                                                      12 0.000156 7.380000e-07
     358 ENSG00000269938 chr12:123996254:A:G
```

```
220
                   ZNF852
                             chr3:44034110:G:A
                                                      3 0.000220 1.870000e-06
     259
                                                     13 0.000241 2.780000e-07
          ENSG00000283361 chr13:114134675:T:C
     71
                   SPHKAP
                            chr2:228452776:C:T
                                                      2 0.000298 7.070000e-08
                     FILE
     192 ENSG00000242866
     132 ENSG00000183831
     358 ENSG00000269938
     220 ENSG00000178917
     259 ENSG00000283361
     71
          ENSG00000153820
     FDR < 0.05
[29]: overlapping_twas = np.append(np.array(caudate[(caudate['FILE'].
      →isin(list(set(caudate.FILE) &
                                                                           Ш
      ⇒set(gandal_twas.GeneID))))].ID),
                                 np.array(list(set(caudate.ID) & set(twas_hub))))
     len(overlapping_twas)
[29]: 50
[30]: caudate[~(caudate['ID'].isin(overlapping twas))].
      sep='\t', index=False)
     caudate[~(caudate['ID'].isin(overlapping_twas))].shape
[30]: (401, 47)
[31]: drop_caudate = caudate[~(caudate['ID'].isin(overlapping_twas))].copy()
     drop_caudate[(drop_caudate['P'] > 5e-8)].shape
[31]: (250, 47)
[32]: drop_caudate[(drop_caudate['P'] > 5e-8)].sort_values('FDR')\
                                            .loc[:, ['ID', 'our_snp_id', _
      → 'CHR_TWAS', 'FDR', 'P', 'FILE']].head(25)
[32]:
                       ID
                                   our_snp_id CHR_TWAS
                                                             FDR.
                                                                                \
     192
                     STRC
                            chr15:43782086:A:G
                                                     15 0.000062 2.750000e-06
     132
                            chr1:173743105:T:C
                                                      1 0.000129 5.050000e-07
                  ANKRD45
          ENSG00000269938 chr12:123996254:A:G
     358
                                                     12 0.000156 7.380000e-07
     220
                             chr3:44034110:G:A
                                                      3 0.000220 1.870000e-06
                   ZNF852
                                                     13 0.000241 2.780000e-07
     259
          ENSG00000283361 chr13:114134675:T:C
     71
                           chr2:228452776:C:T
                                                      2 0.000298 7.070000e-08
                   SPHKAP
     333 ENSG00000279726
                                                      5 0.000384 8.660000e-07
                            chr5:140841554:G:A
     52
                 SYNDTG11.
                            chr14:74416653:A:C
                                                     14 0.000454 3.090000e-05
```

301	NTN5	chr19:48746940:T:C	19	0.000477	1.100000e-06
106	TBC1D5	chr3:16834975:C:T	3	0.000628	5.770000e-08
213	PCDHAC1	chr5:140841554:G:A	5	0.000643	8.660000e-07
425	CCDC92	chr12:123431550:A:G	12	0.000703	7.100000e-07
200	GST01	chr10:104732662:T:C	10	0.000712	2.010000e-05
380	NLRP1	chr17:5267575:T:C	17	0.000735	1.300000e-05
138	CEBPZOS	chr2:37017150:C:T	2	0.000815	3.140000e-06
91	IFT57	chr3:108058204:C:A	3	0.000961	2.010000e-05
153	TRPS1	chr8:115455975:A:G	8	0.000972	4.280000e-07
289	PDIA3	chr15:43782086:A:G	15	0.000974	2.750000e-06
250	RPF2	chr6:111221185:A:G	6	0.001017	8.300000e-07
90	FAM134A	chr2:219196879:T:C	2	0.001115	6.300000e-06
198	DND1	chr5:140841554:G:A	5	0.001170	8.660000e-07
409	RN7SKP101	chr15:47038124:C:T	15	0.001186	2.670000e-06
241	ZBTB37	chr1:173743105:T:C	1	0.001201	5.050000e-07
48	NPIPB3	chr16:21680256:A:G	16	0.001547	1.790000e-07
12	ADAM10	chr15:58749813:T:C	15	0.001766	1.610000e-06

FILE

192 ENSG00000242866 132 ENSG00000183831 358 ENSG00000269938 220 ENSG00000178917 259 ENSG00000283361 71 ENSG00000153820 333 ENSG00000279726 52 ENSG00000183379 301 ENSG00000142233 106 ENSG00000131374 213 ENSG00000248383 425 ENSG00000119242 200 ENSG00000148834 380 ENSG00000091592 138 ENSG00000218739 91 ENSG00000114446 153 ENSG00000104447 289 ENSG00000167004 250 ENSG00000197498 90 ENSG00000144567 198 ENSG00000256453 409 ENSG00000223308 241 ENSG00000185278 48 ENSG00000169246

ENSG00000137845

12

1.5 TWAS tissue summary

```
[33]: brainseq = pd.read_csv("../../libd_twas_comparison/_m/TWAS_gene_tissue_summary.
       ⇔csv")
      brainseq.shape
[33]: (10387, 11)
[34]: brainseq.head(2)
[34]:
                            Symbol
                                    Caudate_TWAS.Z Caudate_FDR Caudate_GWAS.SNP
                  Geneid
      0 ENSG0000000457
                             SCYL3
                                           1.090068
                                                        0.597981
                                                                             Other
      1 ENSG00000000460 C1orf112
                                          -0.372763
                                                        0.892471
                                                                             Other
         DLPFC_TWAS.Z DLPFC_FDR DLPFC_GWAS.SNP HIPPO_TWAS.Z HIPPO_FDR \
      0
                  {\tt NaN}
                             NaN
                                             {\tt NaN}
                                                           {\tt NaN}
                                                                       {\tt NaN}
                             NaN
                                                                       {\tt NaN}
      1
                  NaN
                                             NaN
                                                           {\tt NaN}
        HIPPO_GWAS.SNP
      0
                   NaN
      1
                   NaN
[35]: bb = brainseq.merge(pd.DataFrame({'Symbol': twas_hub, 'inTWAS_HUB': 1}), __
       .merge(pd.DataFrame({'Geneid': gandal_twas.GeneID, 'inGandal': 1}),
                          on="Geneid", how='left')
      bb.to_csv('TWAS_gene_tissue_summary.csv', index=False)
```