Sharing of Genes and Pathways Across Complex Phenotypes: A Multilevel Genome-Wide Analysis

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ABSTRACT Evidence from genome-wide association studies (GWAS) suggest that pleiotropic effects on human complex phenotypes are very common. Recently, an atlas of genetic correlations among complex phenotypes has broadened our understanding of human diseases and traits. Here, we examine genetic overlap, from a gene-centric perspective, among the same 24 phenotypes previously investigated for genetic correlations. After adopting the multilevel pipeline (freely available at <http://grass.cgs.hku.hk/limx/kgg/>), which includes intragenic single nucleotide polymorphisms (SNPs), genes, and gene-sets, to estimate genetic similarities across phenotypes, a large amount of sharing of several biologically related phenotypes was confirmed. In addition, significant genetic overlaps were also found among phenotype pairs that were previously unidentified by SNP-level approaches. All these pairs with new genetic links are supported by earlier epidemiological evidence, although only a few of them have pleiotropic genes in the GWAS Catalog. Hence, our gene and gene-set analyses are able to provide new insights into cross-phenotype connections. The investigation on genetic sharing at three different levels presents a complementary picture of how common DNA sequence variations contribute to disease comorbidities and trait manifestations.

KEYWORDS GWAS; gene-based; complex diseases; genetic sharing; pleiotropy

GENOME-WIDE association studies (GWAS) fall short of explaining most phenotype heritability, and their reported intergenic hits usually cannot link clearly to biological function (Manolio *et al.* 2009; So *et al.* 2011). Nevertheless, the ever-increasing numbers of GWAS datasets actually provide a rich resource for conducting secondary analyses, such as meta-analysis, epistasis analysis, and gene/gene-set analysis (Cantor *et al.* 2010; Califano *et al.* 2012). These analyses are capable of uncovering small-to-moderate genetic effects hidden in previous GWAS by increasing sample size or leveraging external knowledge, for instance, genes or biolog-

ical pathways, protein–protein interaction networks (Li *et al.* 2012a), as well as DNA regulatory machinery elements (Barabasi *et al.* 2011). Since common associated SNPs with moderate effect on complex phenotypes tend to cluster in the same gene, or set of genes, that exert the same key biological function, gene and gene-set analyses tend to increase power (Ramanan *et al.* 2012). Such analyses have been applied successfully to several complex phenotypes, and have led to the discovery of additional disease-gene connections (Eleftherohorinou *et al.* 2009; Chen *et al.* 2010; O'Dushlaine *et al.* 2011; Chen *et al.* 2014).

More recently, pleiotropic effects and cross-phenotype (CP) associations have been identified by joint analysis of GWAS findings or resources from multiple phenotypes (Solovieff *et al.* 2013). Pleiotropy occurs when a gene or genetic variant affects >1 phenotypic trait, and is gradually gaining recognition as a universal property of genetic variants contributing to human phenotypic variation (Solovieff *et al.* 2013; Gratten and Visscher 2016). In an even broader manner, pleiotropy could be explained by single variants or multiple variants in a gene, region, or pathway, or multiple

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Table 1 Summary of associated genic SNPs, genes and gene-sets per phenotype

Phenotypes	Genic SNPs ^a (% of Unique) ^b	Genes (% of Unique)	Gene-sets (% of Unique)
Age at menarche (AM)	683 (75)	155 (54)	0 (NA)
Alzheimer's disease (ALZ)	123 (86)	40 (50)	0 (NA)
Anorexia nervosa (AN)	0 (NA)	1 (0)	1 (100)
Autism spectrum disorder (ASD)	0 (NA)	0 (NA)	0 (NA)
Body mass index (BMI)	654 (65)	171 (46)	7 (71)
Bipolar disorder (BPD)	40 (100)	12 (83)	0 (NA)
Birth length (BL)	1 (0)	6 (33)	1 (100)
Birth weight (BW)	20 (40)	7 (43)	1 (0)
Childhood obesity (CO)	63 (0)	9 (0)	0 (NA)
Coronary heart disease (CHD)	83 (88)	24 (63)	0 (NA)
Crohn's disease (CD)	773 (64)	231 (48)	11 (55)
Major depression disorder (MDD)	0 (NA)	0 (NA)	1 (0)
Ever/never smoked (SMO)	0 (NA)	0 (NA)	0 (NA)
Fasting glucose (FG)	262 (61)	75 (33)	0 (NA)
HDL cholesterol (HDL)	846 (67)	225 (46)	5 (60)
Human height (HH)	10120 (95)	2035 (88)	19 (74)
Infant head circumference (IHC)	0 (NA)	0 (NA)	2 (100)
LDL cholesterol (LDL)	732 (68)	187 (43)	4 (50)
Rheumatoid arthritis (RA)	153 (72)	26 (50)	8 (88)
Schizophrenia (SCZ)	148 (78)	32 (75)	0 (NA)
Type 2 diabetes (T2D)	106 (70)	18 (33)	1 (100)
Triglycerides (TG)	614 (50)	150 (22)	8 (13)
Ulcerative colitis (UC)	506 (63)	251 (63)	20 (70)
Years of education (EDU)	9 (89)	20 (30)	0 (NA)

^a Only SNPs, genes, or gene-sets with *P*-value significant after Bonferroni correction were counted. Average numbers of genic SNPs, genes, and gene-sets being tested are 583619, 23616, and 2653, respectively.

^b Percent of unique is relative to 24 phenotypes. NA, not available.

associations scattered through the genome. So far, 4.6% of SNPs and 16.9% of genes have been observed to be associated with >1 trait in the GWAS Catalog (Welter *et al.* 2014). Risk score profiling and heritability estimation revealed non-negligible genome-wide sharing across neuropsychiatric diseases (International Schizophrenia Consortium 2009; Cross-Disorder Group of the Psychiatric Genomics Consortium 2013). Recently, an atlas of genetic correlation across human complex diseases and traits has been provided in a study by Bulik-Sullivan *et al.* (2015), which revealed several correlated phenotype pairs that had never been identified before. However, relatively few individual SNPs or genes have been associated with those pairs of genetically correlated phenotypes (Cross-Disorder Group of the Psychiatric Genomics Consortium 2013; Parkes *et al.* 2013). GWAS gene/gene-set analyses may have the potential to identify more susceptibility genes or pathways that contribute to multiple genetically comorbid or correlated phenotypes (Zhernakova *et al.* 2009; Solovieff *et al.* 2013).

Hence, in this study, we utilized our previously developed gene-based approach, GATES (Li *et al.* 2011), and its extension on a pathway level, to identify genetic associations shared among complex phenotypes. In addition, we compared the genetic sharing revealed at gene or gene-set levels to that inferred from individual SNP association and overall genetic correlations so as to determine what additional knowledge could be gained from gene-centric approaches.

Materials and Methods

Real data resources

We included the same 24 phenotypes (full names and abbreviations listed in Table 1; detailed information given in Supplemental Material, Table S1 and Text S3 in File S1) as used in Bulik-Sullivan *et al.* (2015). These phenotypes are all complex traits or diseases for which large-scale GWAS meta-analyses have been done on Caucasian populations.

Raw data (GWAS summary statistics) were downloaded, and then filtered by the script “munge_sumstats.py” provided by linkage disequilibrium score (LDSC) online resource (<https://github.com/bulik/ldsc/wiki>). In brief, variants not covered by the HapMap3 panel, with the estimated imputation “info” score <0.9 if imputed, with mislabeled reference/alternative alleles, with extreme *P*-values (>1, or ≤0), or with small effective sample size (<0.67 times the 90th percentile of sample size) were excluded in following analyses. After this filtering, all 24 datasets contain similar amount of SNPs (~1 million per phenotype; Table S1) as inputs for the following gene-wise analyses.

Gene and gene-set mapping

SNPs passing above filtering were mapped to genes (defined by UCSC human RefGene database) when located within a gene, or within a region of 10 kb up or downstream of the gene, according to their coordinates from human genome build 19 (hg19). In addition, to prevent our gene/gene-set results from being mainly driven by genetic variations in the

extended major histocompatibility complex (MHC) region (Chr6: 27.5 ~ 33.5 Mb), we only considered non-MHC genic SNPs in all our subsequent analyses. After this mapping, on average, 581,619 genic SNPs were retained and mapped to 23,616 human genes (Table S1).

Various of gene-sets were collected from Molecular Signatures Database version 5.0 (MsigDB; <http://software.broadinstitute.org/gsea/msigdb/index.jsp>), which included 1330 canonical pathways, and 1454 Gene Ontology gene-sets. After restricting gene-set size at 5–300 genes, a total of 2660 sets that cover 10,483 unique human reference genes was retained. To estimate crosstalk effects between different gene-sets, Jaccard distance (JD) was calculated for each pair of gene-sets by formula $A \cap B / A \cup B$, while A and B are the sizes of gene-sets in terms of gene counts. Gene-sets with Jaccard distance < 0.2 are treated as independent sets (Jia *et al.* 2011; Donato *et al.* 2013), which includes in total 979 different gene-sets.

Linkage disequilibrium attenuated rank-sum test (LDRT)

We developed a new gene-set-based association test, LDRT, to combine gene-level *P*-value statistics calculated from our previous gene-based association test, GATES (Li *et al.* 2011), into gene-sets. Under a competitive null hypothesis for a given gene-set (Gui *et al.* 2011), we assume that GATES *P*-values for genes in this gene-set follow the same distribution as *P*-values for all other genes in the genome. A nonparametric Wilcoxon rank-sum test was then used to examine whether the given set of genes was more highly ranked in an ordered list of all genes than would be expected by chance. When multiple genes within a gene set were on the same chromosome, we sorted these genes according to their gene-based *P*-values and checked the LD between the GATES key SNPs of the genes. If two key SNPs from two different genes were in high LD ($r^2 > 0.5$ as default setting), the key SNP in the gene with larger *P*-value were removed. After removing all redundant key SNPs, the gene-based *P*-values were recalculated by GATES. Hence the new gene-based *P*-values were nearly independent of each other, and can be used in a Wilcoxon rank-sum test for gene-set association analysis.

A recent powerful approach, “MAGMA,” that also handles polygenic traits was chosen for comparison (de Leeuw *et al.* 2015) with LDRT (Text S1 and Figure S1 in File S1), using the same SNP-level summary statistics and gene-set database as input.

Intraphenotype genetic association

We identified three levels of genetic associations with biological interest: genic SNPs, genes, and gene-sets. The complete multilevel pipeline was illustrated as shown in Figure 1.

Genic SNP level: Using the 1000 Genome Caucasian (CEU) population as the LD reference, the Genetic type I error calculator (GEC) was used to estimate the effective number (N_1) of genic SNPs across genome (Li *et al.* 2012b), and then to determine the significance cut-off at family wise error rate 0.05 (Bonferroni correction). Genic SNPs with *P*-values

Figure 1 Multilevel pipeline for SNP, gene, and gene-set association and sharing. Genome-wide SNP summary statistics were collected from large-scale GWAS meta-analysis. Step by step: (1) GEC was used to estimate effective number of associated SNPs at genic regions; (2) GATES was then used to perform gene-level association; (3) LDRT was finally used to perform gene-set association for each phenotype. Only genic SNPs, genes, or gene-sets were prioritized if its association *P*-value was significant when family-wise error rate (FWER) was controlled at 0.05. The associated genic SNPs, genes, and gene-sets were intersected for two phenotypes so as to search for genetic sharing.

smaller than the cut-off ($0.05/N_1$; $N_1 = 272,852$) were selected for each phenotype.

Gene level: For a given gene, its SNP-level summary statistics were combined to generate gene-based *P*-value using GATES (Li *et al.* 2011), which adjusts correlations among neighboring markers by considering the LD structure in 1000 Genomes CEU population. Genes with GATES *P*-values significant after Bonferroni correction ($0.05/N_2$; $N_2 =$ number of genes with ≥ 1 SNPs, with mean at 23,616) were retained for each phenotype.

Gene-set level: For each gene-set containing ≥ 5 genes with nonmissing GATES *P*-values, LDRT was adopted to conduct

gene-set-based association for each phenotype, with the same 1000 Genome CEU population to correct for LD between genes in the same gene set. Gene-sets with LDRT P -values below 0.05/ N_3 ($N_3 = 979$ number of independent gene-sets with $JD < 0.2$) were declared significant, which controlled the same family-wise error rate as that for genic SNPs or genes.

The protocol of running KGG software for above gene/gene-set association is provided in Text S2 of [File S1](#).

Interphenotype genetic sharing

The genetic similarity between a pair of phenotypes was then estimated by counting the number of overlapping genic SNPs, genes or gene-sets that were significantly associated with each member of the pair in the abovementioned genetic association analysis. We performed hypergeometric tests to calculate P -values of the overlap at each level using R function “1-phyper($k-1, K, N-K, n$)” when $k \geq 1$, or “dhyper($k, K, N-K, n$)” when $k = 0$, with four parameters N, K, n , and k corresponding to total count (N , referring to N_1, N_2 , or N_3 in the section *Gene-set level* above), count of significant items in phenotype 1 (K), count of significant items in phenotype 2 (n), and count of overlapping items (k), respectively. False discovery rate (FDR) was set at 5% to define significance so as to correct for multiple testing in 276 pairs of phenotypes. Specifically, a two-step approach was adopted to evaluate gene-level sharing given the existence of LD dependency between genes (de Leeuw *et al.* 2015). First, the overlapping genes between any pair of phenotypes were tested against hypergeometric distribution, with observed gene counts for each phenotype and their overlap. For phenotype pairs with significant gene-overlap at the first step, their gene counts were recalculated after merging those with genetic distance < 1 cM; accordingly, hypergeometric test and FDR was used to examine the significance of the new counts. Only phenotype pairs with significant overlapping (FDR < 0.05) at both steps were considered with gene-level sharing.

To compare cross-phenotype patterns from above genetic sharing with that from previous genetic correlation on the same 24 phenotypes, we retrieved the pairwise genetic correlation coefficients (denoted as r), and corresponding P -values computed by LDSC from Bulik-Sullivan *et al.* (2015). The same FDR threshold (5%) was used to define nonzero genetic correlation for each pair of phenotypes. In addition, we performed an average linkage hierarchical cluster analysis of those 24 phenotypes using $1-r^2$ as the distance measure. Cross-phenotype genetic sharing at gene/gene-set levels was projected on these clusters to examine how they distributed within or across different phenotype clusters.

GWAS catalog replication

To provide biological interpretations for overall cross-phenotype patterns we observed, the shared SNPs, genes, or gene-sets underlying phenotypes with significant genetic sharing were scrutinized for representative functional genes. To that

end, genes carrying significant shared intragenic SNPs, significant shared genes, and genes with GATES P -value < 0.05 in significant shared gene-sets, were extracted and cross-checked for replication evidence in GWAS Catalog. GWAS findings (in terms of genes) for 24 studied phenotypes were collected from GWAS Catalog in November 2015 (Welter *et al.* 2014). Susceptibility genes for each phenotype were recorded only if the gene (± 10 kb) contains ≥ 1 SNP reported with P -value $< 5 \times 10^{-8}$ in at least one Caucasian population. The intersections of reported gene lists for the corresponding 24 phenotypes were assessed to look up genes with pleiotropic effects, and their overlapping was also evaluated by the same hypergeometric tests as previously stated.

Data availability

GWAS P -values analyzed here can be accessed at the URLs given in [Table S1](#). KGG software that implemented GATES and LDRT is available online at the following URL: <http://grass.cgs.hku.hk/limx/kgg/>. GEC software is available online at URL: <http://grass.cgs.hku.hk/gec/>.

Results

Multilevel survey per phenotype

Total numbers of identified significant genic SNPs, genes, and gene-sets for each phenotype are shown in Table 1. Only ASD and SMO have no genetic risk factors found at all three levels. Human height (HH) is associated with 10,120 SNPs, 2035 genes, and 19 gene-sets, having more associations than any of the other 23 phenotypes. With respect to gene-set association, the results of our LDRT approach overlap significantly with the results of the MAGMA (Table S2 and Table S3 in [File S1](#)). Numbers of identified SNPs, genes, and gene-sets are significantly correlated across phenotypes (Spearman correlation coefficients for SNP/gene, gene/gene-set and SNP/gene-set at 0.96, 0.51, and 0.5, respectively; Bonferroni corrected P -values < 0.05). Apparently, far fewer gene-sets are identified than SNPs or genes, even for those phenotypes with relatively large number of significant SNPs or genes (e.g., SCZ, CHD, ALZ, and FG). As expected, when considering genetic associations (SNP, gene, or gene-set) that are unique to any of the 24 phenotypes, we find a smaller proportion of associated genes than the proportion of associated SNPs (one-sided paired Wilcoxon rank-sum test P -value < 0.05 ; Table 1).

Genetic sharing across phenotypes

Numbers of overlapping genes and gene-sets for each of the 276 pairs of phenotypes among the 24 examined are shown in Table 2, while the overlapping genic SNPs and pairwise genetic correlations are given in Table S4 in [File S1](#). At FDR 0.05 level, we identify 41 and seven phenotype-pairs sharing significantly more than the expected counts of genes or gene-sets by chance (highlighted in Table 2), while six of those pairs share both more than expected genes and gene-sets. In total, 42 different phenotype pairs, comprising

Table 2 Cross-phenotype gene and gene-set sharing

Phenotype	AM	ALZ	AN	ASD	BMI	BPD	BL	BW	CO	CHD	CD	MDD	SMO	FG	HDL	HH	IHC	LDL	RA	SCZ	T2D	TG	UC	EDU
AM	NA	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
ALZ	0	NA	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
AN	0	0	NA	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
ASD	0	0	0	NA	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
BMI	15	1	0	0	NA	0	0	0	0	0	0	0	0	0	0	2	0	0	0	0	0	0	0	0
BPD	0	0	0	0	0	NA	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
BL	0	0	0	0	0	0	NA	0	0	0	0	0	0	0	0	1	0	0	0	0	0	0	0	0
BW	1	0	0	0	0	0	0	NA	0	0	0	0	0	0	0	0	0	0	0	0	0	0	1	0
CO	3	0	0	0	4	0	0	0	NA	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
CHD	0	0	1	0	1	0	0	0	0	NA	0	0	0	0	0	0	0	0	0	0	0	0	0	0
CD	5	0	0	0	3	0	0	0	0	0	NA	0	0	0	0	0	0	0	2	0	0	2	5	0
MDD	0	0	0	0	0	0	0	0	0	0	0	NA	0	0	0	1	0	0	1	0	0	1	0	0
SMO	0	0	0	0	0	0	0	0	0	0	0	0	NA	0	0	0	0	0	0	0	0	0	0	0
FG	0	0	0	0	3	0	0	1	0	0	2	0	0	NA	0	0	0	0	0	0	0	0	0	0
HDL	0	2	0	0	6	0	0	0	0	1	3	0	0	3	NA	0	0	2	0	0	0	2	0	0
HH	29	5	0	0	28	2	2	1	1	3	22	0	0	6	22	NA	0	0	1	0	0	3	0	0
IHC	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	NA	0	0	0	0	0	0	0
LDL	0	3	1	0	5	0	0	0	0	4	4	0	0	3	11	20	0	NA	0	0	0	2	0	0
RA	0	0	0	0	0	0	0	0	0	0	1	0	0	0	0	4	0	0	NA	0	1	1	2	0
SCZ	1	0	0	0	3	0	0	0	0	0	0	0	0	0	0	5	0	0	0	NA	0	0	0	0
T2D	1	1	0	0	2	0	0	1	1	1	1	0	0	2	2	5	0	2	0	0	NA	0	0	0
TG	0	2	0	0	4	0	0	0	0	0	2	0	0	2	16	14	0	10	0	0	2	NA	2	0
UC	8	0	0	0	0	0	0	0	0	1	18	0	0	1	5	24	0	2	0	0	1	0	NA	0
EDU	1	0	0	0	1	0	0	0	0	0	2	0	0	0	0	0	0	0	0	0	0	0	1	NA

Upper triangular matrix: counts of gene-sets shared for each two phenotypes; lower triangular matrix: counts of genes shared for each two phenotypes after two-step approach. Cells with significant counts after FDR correction (0.05) are in boldface.

18 different phenotypes, are found to have significant higher amount of overlapping genes or gene-sets. In order to compare these findings to SNP-based evidence, the patterns of cross-phenotype sharing are tabulated, as shown in Table 3. Accordingly, four cross-phenotype patterns (Y-Y-Y, Y-Y-N, N-Y-N, and N-N-Y; “Y” or “N” for share or not share at the genic SNP, gene, or gene-set level, respectively) are observed when considering all three levels of sharing. Gene/gene-set analyses provide significant genetic sharing for 12 out of 42 pairs that are not detected by genic SNP analysis (Table 3). When comparing with pairwise genetic correlation from Bulik-Sullivan *et al.* (2015), our cross-phenotype analyses identify fewer pairs with close genetic relationship (42 vs. 74; one-sided Fisher exact test P -value <0.01); however, it is more likely to identify significant gene or gene-set sharing among those phenotypes with nonzero genetic correlation than those with nearzero genetic correlation (21/74 vs. 21/202, one-sided Fisher exact test P -value <0.001).

On the basis of reported genetic correlations in the study of Bulik-Sullivan *et al.* (2015), three big phenotype clusters were formed (Figure 2A). The clusters are largely consistent with the EBI ontology classification included in Table S1: with cluster 1 containing anthropometric traits, cluster 2 containing cardiometabolic traits or diseases, and cluster 3 containing autoimmune or psychiatric diseases. To further demonstrate how multiple phenotypes (≥ 3) related to each other through genetic sharing, a network was constructed to connect those phenotypes from the same or different clusters, with edges representing significant molecular sharing

at different levels (Figure 2B). Overall, genetic sharing for phenotypes within the same cluster is significantly higher than sharing across clusters (Fisher exact test P -value <0.001). Interestingly, traits or diseases within the same cluster (e.g., CD-UC and HDL-LDL-TG) tend to share genetic hits at all three levels; however, those from different clusters (e.g., CD-TG and HH-SCZ) are more prone to have sharing at advanced level only (genes or gene-sets). Moreover, phenotype pairs prioritized by both our pipeline and LDSC genetic correlation are mainly from within-category phenotypes (15 out of 21 pairs), while those pairs found only by our pipeline are more likely from cross-category phenotypes (17 out of 21 pairs). Noticeably, among all 42 phenotype connections, 25 are comprised by those seven cardiometabolic traits (HDL, LDL, TG, FG, T2D, CO, and CHD) and/or five anthropometrical measurements (BMI, HH, IHC, BL, and BW) that have apparent epidemiological correlation (Sorensen *et al.* 1999) or clinical comorbidity. Their closer relationships have also been detected by cross-phenotype genetic correlations, GWAS Catalog overlapping analyses, and several previous lines of evidence showing their genetic interconnections (Global Lipids Genetics *et al.* 2013; Locke *et al.* 2015).

Annotation for cross-phenotype patterns

All 12 phenotype pairs with significantly shared genes or gene-sets but not genic SNPs were further annotated by epidemiology literature, LDSC genetic correlation, and GWAS Catalog (Table 4). All these novel pairs we found have epidemiological evidence supporting their relationship (Bregenzner *et al.*

Table 3 Comparison of multilevel genetic sharing and correlation

Genic SNPs	Genes	Gene-sets	LDSC genetic correlation	Counts	Pairs
Y	Y	Y	Nonzero	5	BMI-HH, CD-UC, HDL-TG, LDL-TG, HH-TG
Y	Y	Y	Nearzero	1	HDL-LDL
Y	Y	N	Nonzero	13	AM-BMI, AM-CO, AM-HH, BMI-CO, BMI-FG, BMI-HDL, BMI-T2D, BW-T2D, CO-T2D, CHD-LDL, FG-HDL, FG-T2D, HH-LDL
Y	Y	N	Nearzero	11	ALZ-LDL, ALZ-TG, BMI-LDL, CD-HDL, CD-HH, CD-EDU, FG-HH, FG-LDL, FG-TG, HDL-HH, HH-RA
N	Y	N	Nonzero	3	BMI-SCZ, BMI-TG, T2D-TG
N	Y	N	Nearzero	8	AM-CD, AM-UC, ALZ-HDL, AN-CHD, AN-LDL, CD-FG, CD-LDL, HH-SCZ
N	N	Y	Nearzero	1	CD-TG

Y, for significant sharing; N, for nonsignificant sharing.

2006; Zammit *et al.* 2007; Locke *et al.* 2015). Among them, three pairs had significant genetic correlations in Bulik-Sullivan *et al.* (2015), and two other pairs had enriched overlapping genes in GWAS Catalog database. When zooming into the genes underlying these 12 phenotype pairs, a list of 62 different genes was generated (Table S5 in File S1). Among them, 33 genes contribute to two pairs of phenotypes, showing they are involved in three or four unique phenotypes. However, most of those implicated (51 out of 62) have already been reported as pleiotropic genes to the studied 24 phenotypes in the GWAS Catalog. Except two well-known genes from ALZ-HDL (*TOMM40/APOC1* in Table 4), the remaining nine genes were reported as pleiotropic but for different phenotype pairs (Table S5 in File S1). Further larger-scale cross-phenotype GWAS studies are needed to confirm their importance in our reported pairs.

Discussion

Our study follows up the recent report (Bulik-Sullivan *et al.* 2015) on genetic correlations among complex phenotypes by focusing on genic variants of those phenotypes. We demonstrate that gene and gene-set analysis may provide new insights on cross-phenotype connections through integration of three complementary analyses that involved SNPs, genes, and gene-sets on the same dataset. The complete investigation on genetic sharing enables us to form a better picture of how common biological elements contribute to disease comorbidities and trait manifestations.

Genetic correlations that reflect genetic relationship between different phenotypes can be estimated by earlier family-based study design in behavior genetics, whole genome-wide SNP genotypes, and, most recently, GWAS summary statistics (Gratten and Visscher 2016). The potential causes can be ascribed to real biological pleiotropy, mediated phenotype pathways, genetic heterogeneity and spurious bias from study designs (shared controls or ascertainment bias) (Solovieff *et al.* 2013). Our estimation of the genetic overlaps at three different levels, using GWAS summary statistics only, reflects a portion of overall biological pleiotropy among multiple phenotypes, hence provides only partial explanation to the genetic correlation observed in Bulik-Sullivan *et al.* (2015). In contrast to previous cross-phenotype genetic evidence from overall heritability or genetic correlation estimations (International Schizophrenia Consortium 2009; Cross-Disorder Group of the

Psychiatric Genomics Consortium 2013; Parkes *et al.* 2013), our study gives more specific biological clues (in terms of genes and pathways) to shared disease etiologies. Though joint GWAS analyses across multiple phenotypes have enabled finding susceptibility loci, genes, and gene-sets that affect different related phenotypes (Ramanan and Saykin 2013; Network and Pathway Analysis Subgroup of Psychiatric Genomics Consortium 2015), all of these studies employed close-related phenotypes in the same disease category, for instance, psychiatric disorders and autoimmune disorders. Inspired by the findings of Bulik-Sullivan *et al.* (2015), *i.e.*, that widespread and unexpected genetic correlation exists across 24 phenotypes with different pathogenicity mechanism, our multilevel gene-centric pipeline has confirmed significant genetic sharing for those well-connected phenotypes pairs (lipid traits and inflammatory bowel diseases) (Lees *et al.* 2011; Global Lipids Genetics *et al.* 2013); more importantly, our analyses gave new insights on how these phenotypes link to each other through genetics. This new knowledge gained from our systematic adoption of gene/gene-set approaches provides a new perspective to explain complex disease etiologies (Elbers *et al.* 2009; Manolio *et al.* 2009; Solovieff *et al.* 2013). Usually for polygenic traits that involve many genetic hits with small effect size, disease connections could be due mainly to contribution of multiple different SNPs in one gene or multiple genes in a gene-set. Under this scenario, the advantage of our GWAS gene or gene-set analyses, which combine original SNPs with marginal statistical evidence in the same biological unit, are expected to be fully expressed (Wang *et al.* 2010; Ramanan *et al.* 2012). In addition, our new gene-set association approach “LDRT,” adopting a cut-off free competitive test (de Leeuw *et al.* 2016), can analyze all genic SNPs together, and hence provide additional information that may lose in traditional gene-set enrichment analyses as implemented in DEPICT (Pers *et al.* 2015), from which we observed unbalanced numbers (ranging from 0 to 927) of significant gene-sets within each phenotype and only significant sharing between HDL and TG, or CD and RA across those 24 phenotypes (Table S6 in File S1).

On the other hand, our pipeline has generated a few unexpected results. We are missing some well-known genetically correlated phenotypes in psychiatric diseases (such as SCZ-BPD and AN-SCZ) (International Schizophrenia Consortium *et al.* 2009; Bulik-Sullivan *et al.* 2015). This could be due to information loss when retrieving only genic SNPs (see

Figure 2 Cross-phenotype clustering and linking. Phenotypes were clustered and linked according to different levels of sharing: (A) hierarchical clustering of phenotypes with genetic correlation (r) as distance measurement ($1-r^2$). (B) phenotypes were connected to each other within or across different clusters. Edges with different colors and thickness are noted with different numbers: (1) internal linking at gene level in cluster 2; (2) internal linking at two levels (SNP and gene, or gene and gene-set) in cluster 2; (3) internal linking at three levels in cluster 2; (4) internal linking at three levels in cluster 3; (5) cross-cluster linking at gene level; (6) cross-cluster linking at two levels; (7) cross-cluster linking at all three levels; (8) cross-cluster linking at gene-set level only. Cluster 1: ALZ, HH, IHC, BL, and BW; Cluster 2: LDL, ASD, SMO, EDU, AM, BMI, CO, CHD, HDL, TG, FG, and T2D; Cluster 3: AN, RA, CD, UC, MDD, BPD, and SCZ.

Table S2 in [File S1](#) for details) from overall genome-wide association studies, which contained more noncoding variants important to both phenotypes at a regulation or epigenetic level (Tsankova *et al.* 2007; Maurano *et al.* 2012). The

missing of other correlated phenotypes can be ascribed to false negative factors in our pipeline, including too stringent a cut-off for selecting significant genes or gene-sets, limited numbers of pathways used, and incomplete coverage of

Table 4 Annotation of 12 novel phenotype pairs

Phenotype pair	Total genes (levels) ^a	Epidemiological evidence	LDSC evidence ^b	Catalog evidences ^c
ALZ-HDL	3 (Gene)	Casserly and Topol (2004)	F; across clusters	6*; <i>TOMM40</i> , <i>APOC1</i>
AN-CHD	1 (Gene)	Casiero and Frishman (2006)	F; across clusters	0; NA
AN-LDL	1 (Gene)	Weinbrenner <i>et al.</i> (2004)	F; across clusters	0; NA
AM-CD	18 (Gene)	Ballinger <i>et al.</i> (2003)	F; across clusters	1; 0
AM-UC	24 (Gene)	Ballinger <i>et al.</i> (2003)	F; across clusters	2; 0
BMI-SCZ	6 (Gene)	Zammit <i>et al.</i> (2007)	T; across clusters	2; 0
BMI-TG	11 (Gene)	Locke <i>et al.</i> (2015)	T; within cluster	1; 0
CD-FG	5 (Gene)	Bregenger <i>et al.</i> (2006)	F; across clusters	1; 0
CD-LDL	8 (Gene)	Agouridis <i>et al.</i> (2011)	F; across clusters	1; 0
CD-TG	8 (Gene-set)	Agouridis <i>et al.</i> (2011)	F; across clusters	2; 0
HH-SCZ	8 (Gene)	Zammit <i>et al.</i> (2007)	F; across clusters	9*; 0
T2D-TG	2 (Gene)	Locke <i>et al.</i> (2015)	T; within cluster	1; 0

^a Gene-sets was considered by counting genes in gene-sets with GATES $P < 0.05$; cells with gene-set sharing were underscored.

^b F, for near-zero genetic correlation; T, for nonzero genetic correlation; within clusters or cross clusters was given in Figure 2.

^c Overlapping genes were counted from GWAS Catalog reported genes for each phenotype pair, with "*" showing significance after multiple testing; only hits replicated at both phenotypes in Catalog were given. NA, not applicable.

human genes in the gene-set collection. In addition, we report a few phenotype pairs that have inconsistent evidence from LDSC genetic correlation analyses or the GWAS Catalog record. Though they are subjected to possible false positives arising from our methodology (insufficient correction of gene-level dependence), there are other explanations for this inconsistency. Some phenotype pairs (for instance HDL-LDL) originated from the difference on how to handle effect direction at SNP level—LDSC correlation considered both positive and negative effects (Bulik-Sullivan *et al.* 2015), but we treat them equally with no signs. For the newly found seven pairs of phenotype: the link between AM and inflammatory bowel diseases (CD and UC) have been established previously through epidemiological survey (Ballinger *et al.* 2003), hormone change (Khalili *et al.* 2012), or candidate gene study (Yang *et al.* 2007). CD-LDL and CD-TG have also been reported with significant genetic sharing by another study using a different approach (Andreassen *et al.* 2015), while CD-FG may also reflect the connection between inflammation and metabolic traits; AN-CHD and AN-LDL sharing were driven by the same *ABO* gene, which was recently reported as an important element to multiple phenotypes (Pickrell *et al.* 2015). These connections may have not been captured by genetic correlation analyses, and wait confirmation by future cross-phenotype findings in GWAS Catalog.

The effectiveness and new perspective provided from our GWAS gene or gene-set investigation can serve as proof-of-principle for future cross phenotype studies. Genetic sharing among those 24 phenotypes provided new clues that a combination of gene-wise *P*-values of two related disorders, or even distinct two types of diseases or traits, could greatly increase power for susceptibility gene identification (Barabasi *et al.* 2011; Ramanan and Saykin 2013). Since only summary statistics are needed, our gene or gene-set analyses can easily extend to rare variants from next generation sequencing (Cirulli and Goldstein 2010; Gudbjartsson *et al.* 2015), which would generate another set of gene-level summary statistics that can be combined with those from common variants targeted by GWAS (Curtis 2012; Luo *et al.* 2012). Without doubt, more genetic knowledge on complex disorders will be gained when adopting this strategy on larger-scale datasets in the future. Ultimately, advanced gene/gene-set analyses based on complete variant-level information will play a key role in precision medicine by providing biological guidance on drug development and clinical diagnosis for one disease, or one category of diseases (Barabasi *et al.* 2011; Ramanan and Saykin 2013; Solovieff *et al.* 2013).

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