

## Multi-omic analysis along the gut-brain axis points to a functional architecture of autism

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## Acknowledgements

We would like to thank Allan Packer, Paul Wang, Natalia Volfovsky, Kelsey Martin and John Spiro for their critical review of the manuscript. We'd also like to add Kevin Liu, Hannah Sherman and Xue-Jun Kong for insightful discussions. Y.S. and T.D.L. are supported by the Wellcome Trust (WT098051).

## Contributions

J.T.M. and G.T.-O. conceived and designed the study, developed the software, analyzed the data, interpreted the results and wrote the manuscript; R.B. contributed to study design, data analysis, result interpretation and manuscript editing; R.H.M. contributed to study design, data analysis and manuscript editing; R.J.X. and S.K.M. contributed to study design and manuscript editing; G.R. and B.C. contributed to software development and manuscript editing; D.-M.J. and Y.S. contributed to data analysis and manuscript editing; K.B., B.D.N., M.F.Z., M.D., O.V.A., A.S.K., A.N., M.M., M.W., D.N.F., E.L., W.Z., V.F., V.N.D., D.P.W., M.E.B., R.K., J.G., S.M.D. and T.D.L. provided access to data and contributed to manuscript editing.

## Conflict of Interest

- 1 R.H.M. is Scientific Director at Precidiag Inc.; T.D.L. is co-founder and Chief Scientific Officer of Microbiotica; S.K.M. is a co-founder and has equity in Axial Therapeutics; R.B. is currently Executive Director of
- 2 Prescient Design, a Genentech Accelerator; G.T.-O. is a Consultant-in-Residence at the Simons Foundation.
- 3

## 4 Abstract

5 Autism is a highly heritable neurodevelopmental disorder characterized by heterogeneous cognitive, behav-  
6 ioral and communication impairments. Disruption of the gut-brain axis (GBA) has been implicated in  
7 autism, with dozens of cross-sectional microbiome and other omic studies revealing autism-specific profiles  
8 along the GBA albeit with little agreement in composition or magnitude. To explore the functional architec-  
9 ture of autism, we developed an age and sex-matched Bayesian differential ranking algorithm that identified  
10 autism-specific profiles across 10 cross-sectional microbiome datasets and 15 other omic datasets, including  
11 dietary patterns, metabolomics, cytokine profiles, and human brain expression profiles. The analysis uncov-  
12 ered a highly significant, functional architecture along the GBA that encapsulated the overall heterogeneity  
13 of autism phenotypes. This architecture was determined by autism-specific amino acid, carbohydrate and  
14 lipid metabolism profiles predominantly encoded by microbial species in the genera *Prevotella*, *Enterococ-*  
15 *cus*, *Bifidobacterium*, and *Desulfovibrio*, and was mirrored in brain-associated gene expression profiles and  
16 restrictive dietary patterns in individuals with autism. Pro-inflammatory cytokine profiling and virome asso-  
17 ciation analysis further supported the existence of an autism-specific architecture associated with particular  
18 microbial genera. Re-analysis of a longitudinal intervention study in autism recapitulated the cross-sectional  
19 profiles, and showed a strong association between temporal changes in microbiome composition and autism  
20 symptoms. Further elucidation of the functional architecture of autism, including of the role the microbiome  
21 plays in it, will require deep, multi-omic longitudinal intervention studies on well-defined stratified cohorts  
22 to support causal and mechanistic inference.

## 23 Introduction

24 Autism spectrum disorder (ASD) encompasses a broad range of neurodevelopmental conditions defined by  
25 heterogeneous cognitive, behavioral and communication impairments that manifest early in childhood [1]. To  
26 date, over a hundred genes have been identified as putatively associated with ASD, with some genotypes now  
27 having a standardized clinical diagnosis [2]. However, most of the genetic variants are still associated with  
28 heterogeneous phenotypes, making it difficult to identify molecular mechanisms that might be responsible for  
29 particular impairments [3]. Some studies have also looked at the presence of abnormalities in different brain  
30 regions in children with ASD [4, 5]. However, whether such neuroanatomical features could mechanistically  
31 determine autism, and whether environmental factors could induce analogous ASD-like symptoms, remains  
32 unresolved [1].

33 In addition to risk factors, one comorbidity that has been linked to ASD with high confidence is the  
34 occurrence of gastrointestinal (GI) symptoms, such as constipation, diarrhea, or abdominal bloating, but  
35 causal insights remain elusive [6, 7, 8]. Mechanistically, much research has been focused on the interplay  
36 between the GI system and processes controlled by the neuroendocrine, neuroimmune, and autonomous  
37 nervous systems, all of which converge around the GI tract and together modulate the gut-brain axis (GBA)  
38 [9, 10, 11].

39 The GBA facilitates bidirectional communication between the gut and the brain, contributing to brain  
40 homeostasis and helping regulate cognitive and emotional functions [9, 12]. Over the past decade, research  
41 on the factors modulating the GBA has revealed the central role played by the gut microbiome—the trillions  
42 of microbes that colonize the gut—in regulating neuroimmune networks, modifying neural networks, and  
43 directly communicating with the brain [13]. Dysregulation of the gut microbiome and the ensuing disruption  
44 of the GBA are thought to contribute to the pathogenesis of neurodevelopmental disorders including autism,  
45 but the underlying mechanisms and the extent to which the microbiome explains these dynamics is still  
46 unknown [14, 15, 16, 17].

47 Several dozen autism gut metagenomics studies have revealed many, albeit inconsistent, variations in  
48 microbial diversity in individuals with ASD compared with neurotypical individuals [18, 19, 17]. Similarly,  
49 metagenome-based functional reconstructions and metabolic analyses have also shown strong, albeit incon-  
50 clusive differences between ASD and neurotypical individuals [20, 21, 22]. Comparative analyses at other  
51 omic levels have further shown little agreement across studies [23] raising the question of whether the re-  
52 sults obtained so far reflect intrinsic biological differences among cohorts, insufficient statistical power, or  
53 experimental biases that preclude meaningful comparisons [24].

54 A wide range of factors could explain the disagreement across studies, including confounding variation  
55 due to batch effects, the application of inappropriate statistical methodologies, and the vast phenotypic  
56 and genotypic heterogeneity of ASD. Batch effects can be caused by many factors including misspecified  
57 experimental designs, technical variability, geographical location, and demographic composition, and several  
58 algorithms have been proposed to correct for them, but a lack of standardized statistical methods further  
59 complicates interpretation [25, 26, 27, 28]. Microbiome datasets, like other omic datasets, are compositional,  
60 and failure to account for the compositional nature of sequencing counts can lead to high false positive and  
61 false negative rates when identifying differentially abundant microbes [29, 30, 31]. Microbiome analysis in  
62 ASD is further confounded by the phenotypic and genotypic heterogeneity of the disorder, which is known to  
63 be critical for stratifying ASD subtypes and constructing reliable diagnostics but is typically not measured  
64 or controlled for [32, 33, 1].

65 Understanding functional architecture—the network of interactions among different omic levels that  
66 determines individual phenotypes—of complex neurodevelopmental disorders such as autism, requires an  
67 accurate and comprehensive characterization of the different omic levels contributing to it [34]. Traditionally  
68 focused on the human genomic, metabolic, and cellular components of phenotype determination, mounting  
69 evidence of the role the GBA plays in phenotype determination through bidirectional modulatory mechanisms  
70 raises the need for considering the metagenomic and metabolic contributions of the microbiome as potential  
71 key components of the functional architecture of autism [35, 36].

72 To identify autism-specific omic profiles while reducing cohort-specific confounding factors, we have  
73 devised a Bayesian differential ranking algorithm to estimate a distribution of microbial differentials, or  
74 relative log-fold changes, [31] across multiple potential ASD subtypes implicit in 25 omic datasets (Table 1).  
75 A key feature of this approach was to match individual study participants by sex and age within each study  
76 to adjust for confounders in childhood development and cohort-specific batch effects. The preponderance of  
77 autism among males is well documented and several potentially sex-dependent mechanisms to explain this  
78 phenomenon have been proposed [37]. Furthermore, the development of the microbiome during childhood  
79 is a hallmark of microbiome dynamics in the human gut [38, 39, 40, 41, 42]. Our analysis provides insights  
80 into the complexity of the interplay among multiple omic levels in ASD, highlights the inherent limitations  
81 of cross-sectional studies for understanding the functional architecture of autism, and provides a framework  
82 for further studies aimed at better defining the causal relationship between the microbiome and other omic  
83 levels and ASD.

## 84 Results

85 The structure of our analysis consisted of a multi-cohort and multi-omic meta-analysis framework that al-  
86 lowed us to combine independent and dependent omic data sets in one integrated analysis [43, 44]. To  
87 minimize issues of compositionality and sequencing depth [45], we modeled overdispersion using a negative  
88 binomial distribution for modeling sequencing count data [46] (Box 1 “TACKLING METAGENOMIC UN-  
89 KNOWNs”). Our differential ranking approach incorporated a case-control matching component consisting  
90 of individually pairing ASD children with age- and sex-matched neurotypical control children within each  
91 study cohort, allowing us to adjust for confounding variation and batch effects (Supplemental methods). Fi-  
92 nally, we cross-referenced the 16S-based microbial differential ranking analysis from eight age-sex matched  
93 cohorts against 15 other omic datasets to contextualize the potential functional roles these microbes could  
94 play in autism (Figure 1).

### 95 Age- and sex-matching increases informational content of cross-sectional ASD 96 datasets

97  
98 We compared the age- and sex-matched differential ranking analysis to the standard group-averaged dif-  
99 ferential ranking analysis across eight out of the ten 16S studies [47, 48, 49, 21, 50, 51, 52, 53]. Age- and  
100 sex-matched differential analysis outperformed standard group averaging with respect to  $R^2$ , and its overall  
101 performance strictly improved as more studies were added (Figure S1). This performance boost reflected a  
102 reduction in model uncertainty with larger cohorts that was indicative of overlapping differentially abundant  
103 taxa across studies and of reduced confounding variation.

### 104 Global differential ranking analysis reveals a distribution of significant ASD- 105 microbiome associations

106  
107 A global, age- and sex-matched differential ranking analysis of the eight 16S datasets selected for this study  
108 revealed a clear partitioning of microbial differences with respect to ASD and cohort membership (Figure  
109 2a, Figure S2). The distribution of the overall case-control differences showed a strong ASD-specific signal  
110 driven by 142 microbes more commonly found in ASD children and 32 microbes more commonly found in  
111 their control counterparts (Table S1). The variability observed is most likely due to confounding factors such  
112 as cohort demographics and geographic location, with the eight cohorts originating from Asia, Europe, South  
113 America, and North America. Analogous global differential ranking trends could be observed for the virome,  
114 shotgun metagenomics sequencing (SMS), and RNA-seq datasets (Figure S3). To determine whether these  
115 highly significant microbiome signals ( $p$ value<0.0025) could be used to distinguish ASD subjects from their  
116 age- and sex-matched control counterparts, we trained random forest classifiers on train/validation/test splits  
117 on data derived from 16S—targeted sequencing of the microbial 16S ribosomal RNA gene—and SMS—whole  
118 genome sequencing of microbial communities. Despite the strong microbiome effect size, we faced difficulties  
119 fitting generalizable classifiers. Our best classifiers had an average cross-validation accuracy of about 75%  
120 (Figure 2b), falling within the range of 52%–90% classification accuracy observed in previous studies [50,  
121 49, 54]. We suspect that the vast heterogeneity across cohorts hampered classification performance. While  
122 cohort size did not impact predictability (Figure 2c), some cohorts with skewed sex ratios or age ranges did  
123 exhibit lower classification performance (Figure. 2d-e). In the Zurita et al. cohort, sex-specific factors could  
124 confound classification (four girls and 56 boys) [48], and in the Kang et al. cohort, age-associated microbiome  
125 development factors could hamper classification accuracy (all children were 10 years or older). In addition, all  
126 subjects in the Kang et al. cohort had known GI symptoms [55], further compromising classifier performance  
127 because none of the other studies controlled for this variable. As a result, and analogous to the phenotypic  
128 and genotypic heterogeneity observed in ASD, the microbiome composition of ASD children also exhibits  
129 high heterogeneity, precluding the identification of a homogeneous universal ASD microbial profile and the  
130 construction of generalizable classifiers.

### 131 Children with ASD exhibit significant individual differences at several omic levels

132  
133 Differential ranking analysis of three core omic levels—microbiome (16S and SMS) and human transcriptome  
134 (RNAseq)—revealed strong and highly significant differences between ASD subjects and their neurotypical

135 counterparts (p-value < 0.0025) (Figure 2f; Table S2; Table S3). Two additional omic levels—the metabolome  
136 and the virome—didn’t show significance signals (Figure S4, Table S4). Amongst the models that yielded a  
137 statistically significant signal, the 16S and SMS datasets had a larger effect size than the RNAseq datasets  
138 (Figure 2f). While each omic dataset by itself showed strong associations with ASD, a side-by-side comparison  
139 of the 16S and SMS datasets—two datasets that should show high equivalence—revealed a significant lack of  
140 overlap between them, highlighting the outstanding challenge of batch effects in microbiome studies (Figure  
141 S5). The reasons for this discrepancy could be many, but most likely center around sample size—our  
142 study looked at eight 16S datasets versus only three SMS datasets containing 754 individuals and 166  
143 individuals, respectively. Another major challenge when estimating species profiles with metagenomics  
144 reference libraries is assigning a species identification to a read—there are many reads that do not uniquely  
145 map to individual species—and as a result, these multi-mapped reads can give rise to numerous false positive  
146 taxa [56, 57, 58, 59]. Based on this, we decided to focus primarily on the 16S datasets to define a global  
147 differential ranking profile.

## 148 Sibling-matching and unrelated sex- and age-matching show significant discrep- 149 ancies

151 To determine whether sex- and age-matched differentials could be universally predictive, we compared the  
152 16S differentials obtained from the age- and sex-matched cohorts with two sibling-matched cohorts [60] [61].  
153 Interestingly, we observed a significant negative correlation between the differentials extracted from the age-  
154 and sex-matched cohort and the two sibling-matched cohorts, suggesting that ASD-specific microbes in the  
155 sibling-matched studies are enriched in the control group in the age- and sex-matched studies and vice versa  
156 (Fig. 2g). Permanova applied to the age-sex matched cohort revealed a strong age-confounder across cohorts  
157 (pvalue < 0.001), with little confounding variation due to sex (Table S5). In contrast, Permanova applied  
158 to these sibling-matched cohorts revealed that household is a major confounder in both cohorts (pvalue <  
159 0.001), but ruled out age as a confounder and indicated that sex was a confounder only in the David et al.  
160 cohort (pvalue < 0.001). The observed discrepancies point to different sets of confounders possibly affecting  
161 the analysis: in the case of the age- and sex-matched studies, family confounders aren’t typically accounted  
162 for, while sibling-matched studies don’t typically adjust for age confounders. In addition, and while cohorts  
163 such as the one studied in Maude et al. specifically control for the possibility, siblings often exhibit a higher  
164 risk of developing ASD compared to the general population [62].

## 165 Host cytokine concentrations are correlated with microbial abundances

166 Immune dysregulation, ranging from circulating ‘anti-brain’ antibodies and perturbed cytokine profiles to  
167 simply having a family history of immune disorders, has been repeatedly associated with ASD [63, 64].  
168 Recently, for example, Zurita et al. showed that concentrations of the inflammatory cytokine transforming  
169 growth factor beta (TGF- $\beta$ ) are significantly elevated in ASD children. We reanalyzed this dataset, after age-  
170 and sex-matching, and observed that microbial differentials associated with TGF- $\beta$  and IL-6 concentrations  
171 were positively correlated with the global microbial log-fold changes between ASD and control pairs (IL-6 :  
172  $r=-0.435$ ,  $p=0$ ; TGF- $\beta$  :  $r=0.291$ ,  $p=0$ ) (Table S6). To validate the integrity of these microbial profiles with  
173 respect to the cytokine changes, we calculated the log-ratios of these microbial abundances and showed them  
174 to, in turn, be highly correlated with TGF- $\beta$  and IL-6 concentrations (IL-6 :  $r=0.50$ ,  $p=0.0007$ ; TGF- $\beta$  :  
175  $r=0.45$ ,  $p=0.002$ ) (Figure 3 a-d).

176 Four clusters of microbial genera—*Prevotella*, *Enterococcus*, *Bifidobacteria*, and *Desulfovibrio*—were pre-  
177 dominantly associated with the cytokine differentials. Partial mechanistic insights on some of these cytokine-  
178 microbe associations have been previously published. Both *B. longum* and *E. faecalis* have shown anti-  
179 inflammatory activities: *B. longum* downregulates IL-6 in fetal human enterocytes in vitro [65] and *E.*  
180 *faecalis* has been observed to upregulate TGF- $\beta$  in human intestinal cells [66]. *P. copri* associations with  
181 different cytokines have also been observed in multiple disease contexts [67]. Similarly, *Bifidobacteria* and  
182 *Prevotella* both co-occurred with phages enriched in ASD or in neurotypical children (Figure S6, Table  
183 S7), but while microbes have previously been reported to mediate viral infections [68, 69], the mechanistic  
184 underpinnings of these interactions with the host’s immunity remain poorly understood [70, 71, 72].

## 185 The microbiome metabolic capacity is reflective of the human brain-associated 186 metabolic capacity in ASD

188 To determine potential crosstalk between the human brain and the microbiome physiology, we compared  
189 the metabolic capacities encoded by the microbial metagenome—combining the individual metabolic capac-  
190 ities of thousands of different microbes—and the differentially expressed human genome in the brain, two  
191 omic levels representing entirely different biological contexts. We observed that over 100 human metabolic  
192 pathways differentially expressed in the brain tissues of ASD individuals had analogous microbial path-  
193 ways differentially abundant in the microbiome of children with ASD, suggesting a potential coordination  
194 of metabolic pathways across omic levels in ASD (Fig. 3e). Pathways related to amino acid metabolism,  
195 carbohydrate metabolism and lipid metabolism were disproportionately represented among the overlapping  
196 genes (Table S8).

## 197 **The microbiome metabolic capacity reflects restrictive diet patterns in children** 198 **with ASD**

199 Autistic traits in early childhood have been shown to correlate with poor diet quality later in life, however,  
200 little is known about how diet quality is directly linked to autistic traits [73]. Here, we re-analyzed the paired  
201 microbiome and dietary survey data from Berding et al.. A microbiome-diet co-occurrence analysis revealed  
202 startlingly similar amino acid, carbohydrate and lipid metabolism association patterns to those observed  
203 in the microbiome-brain metabolic capacity analysis (Table S9). Interestingly, both microbes enriched in  
204 ASD and in control subjects co-occurred primarily with amino acid dietary compounds (Figure 3f, Figure  
205 S7). Autistic children were less likely to consume foods high in glutamic acid, serine, choline, phenylalanine,  
206 leucine, tyrosine, valine and histidine, all compounds involved in neurotransmitter biosynthesis. Even though  
207 the metabolomic analysis did not yield statistically significant signals after FDR correction, the metabolites  
208 that showed the strongest signal included glutamate and phenylalanine, consistent with the microbiome-  
209 diet analysis [74, 75, 76]. Disruptions in the biosynthesis of these neurotransmitter molecules have been  
210 implicated in a wide variety of psychiatric disorders, and a recent blood metabolomics study has shown the  
211 potential of using branched chain amino acids to define autism subtypes [33]. Due to the incompatibility  
212 between the molecular features across datasets, it was not possible to combine any of the metabolomics  
213 datasets to boost the statistical power, which remains a major limitation of metabolomics technologies at  
214 present (see Methods).

## 215 **Differential microbial rankings show disease-specific correlations**

216 One major challenge in determining microbiome-disease associations is identifying correlations specific to a  
217 particular condition and not generally present across diseases [77, 78]. To determine how specific to ASD our  
218 global differential microbiome profile was, we cross-referenced it against differential ranking results obtained  
219 from an Inflammatory Bowel Disorder (IBD) dataset [79] and a Type 1 Diabetes (T1D) dataset [80]. IBD  
220 shares some comorbidities with ASD [81, 82], while no direct correlation between ASD and T1D has been  
221 reported to date. The analysis revealed a notable overlap between microbes enriched in ASD and IBD, and  
222 this overlap was stronger than both the overlap between IBD and T1D and between ASD and T1D (Figure  
223 S8). Whether this ASD-IBD overlap is suggestive of a common microbial profile or is confounded by the  
224 restrictive dietary nature of these two clinical conditions is currently unclear. Higher resolution and properly  
225 designed clinical studies will have to be performed to get to a mechanistic understanding of the potential  
226 microbiome connection between these two conditions.

## 227 **ASD microbiome profiles weaken after fecal matter transplant consistent with** 228 **reported behavioral improvement**

229 While the preceding cross-sectional analyses showed significant associations among several omic levels (vi-  
230rome, microbiome, immunome) or diet and ASD, insights into causality are still limited. By contrast,  
231 longitudinal intervention studies provide an opportunity to obtain stronger insights into causality. To test  
232 this, we re-analyzed data from a two-year, open label fecal matter transplant (FMT) study with 18 children  
233 with ASD [83]. In this study, the children were subjected to a two-week antibiotic treatment and a bowel  
234 cleanse followed by two days of high dose FMT treatment and eight weeks of daily maintenance FMT doses.  
235 Based on one of the most common evaluation scales for ASD, the Childhood Autism Rating Scale (CARS),  
236 significant improvements were achieved after the ten week course of treatment. Two months later the initial  
237 gains were largely maintained, and a two-year follow-up showed signs of further improvement in most of the  
238 patients. The results are consistent with a potential role of the microbiome in improving autism symptoms,  
239 but how the underlying changes in microbiome composition related to those seen in other studies remained  
240 unknown.

241 Here, we re-analyzed the original raw data in the context of the ASD profiles revealed by our cross-  
242 sectional differential ranking analysis (Table S10). All microbes associated with ASD in the 18 children prior  
243 to the FMT treatment had been identified as ASD-associated microbes in our age- and sex-matched cross-  
244 sectional analysis, recapitulating 74% of the cross-sectional profile. Immediately following FMT treatment,  
245 the abundances of the ASD-associated microbes decreased in all 8 children (Figure 4). The two-year follow-  
246 up analysis revealed that all the ASD-associated microbes, mostly *Enterococcaceae*, continued to be depleted.  
247 Consistent with the findings of Kang et al., we also observed *Desulfovibrio sp.*, and *P. copri* increase over  
248 the two year period, while *Bifidobacteria sp.* could be found both among depleted and enriched species and  
249 other *Prevotella sp.* were depleted, pointing to a potentially wide functional diversity within these genera  
250 not noted in the original study.

## 251 **Discussion**

252 The functional architecture of ASD, and in particular the potential role the microbiome plays in modulating  
253 the GBA in the context of autism, remains poorly understood due to disagreements among existing micro-  
254 biome and other omic studies. Our Bayesian model highlighted a distribution of highly significant microbial  
255 differentials obtained from individual age- and sex-pairings between children with ASD and neurotypicals,  
256 and parallel analyses at the immunome, human transcriptome, and dietome levels revealed strong associ-  
257 ations among omic levels. The virome and the direct metabolome signals, while present, were markedly  
258 weaker than the other omic signals. The inferred ASD-specific metabolic profiles from the microbiome and  
259 the human transcriptome, on the other hand, showed a high and significant degree of overlap in microbial  
260 and human pathways expressed in the gut and in the brain, respectively. The metabolic connection implied  
261 by this overlap, which included differentially enriched carbohydrate and amino acid metabolic pathways in

262 ASD, is a remarkable observation given the fundamental difference between the gut and brain physiologies,  
263 which would a priori suggest a reduced overlap in metabolic capacities. The microbiome-diet co-occurrence  
264 analysis also highlighted a reduced intake of amino acids and carbohydrates linked to specific microbiome  
265 profiles in ASD children. These metabolic and dietary imbalances, particularly regarding glutamate levels,  
266 were further apparent, albeit weakly, in the serum, fecal and urine metabolomes we analyzed. This multi-  
267 scale overlap we observed along the GBA points to the existence of a functional architecture of ASD driven  
268 by the metabolic potential at the genomic and metagenomic levels.

269 While the differential distributions we determined were highly significant, the global analysis did not  
270 provide reliable ASD classifiers or uncover universal microbial ‘smoking guns’ linked to autism. However,  
271 several microorganisms consistently detected across omic levels pointed to potentially interesting functional  
272 connections. For example, our analysis suggested that *B. longum* exhibited a down-regulation of IL-6, which  
273 has been observed across a number of in vitro and cohort studies [84, 85]. The diet co-occurrence analysis also  
274 showed a strong association between *P. copri* and carbohydrate depletion in ASD. The population dynamics  
275 of *P. copri* have been reported to be driven primarily by carbohydrates in the diet [67]. Multiple other  
276 microbes, including several *Bifidobacteria*, *Enterococcus* and *Desulfovibrio* species, stood out in the immune  
277 and viral analyses. In the FMT study, the relative proportions of several *Prevotella*, *Bifidobacteria*, *Desul-*  
278 *fovibrio* and *Enterococcus* species also showed strong associations with ASD symptoms, further suggesting  
279 a causal role for these microorganisms in shaping autism symptoms.

280 Despite our inability to determine actual metabolomic profiles at this point (see Methods), our metabolite  
281 analysis based on microbiome- and brain-derived metabolite inferences as well as the diet-derived metabolite  
282 data reveals a picture of a unifying and distinct ASD functional architecture. With the brain, the immunome  
283 and diet as major effectors, the multi-factorial complexity of ASD is reduced to a multi-scale set of inter-  
284 actions centered around human and bacterial metabolism that in turn determines phenotypic, genomic and  
285 metagenomic attributes via multiple feedback loops (Figure 5). The association of specific genotypes with  
286 ASD has been clearly established [2]; the pivotal role of the immune system in mediating the communication  
287 between the gut microbiome and the human brain as well as other peripheral systems is also firmly estab-  
288 lished [86]; further, the central role of the microbiome in mediating diet-derived nutrient mobilization has  
289 been extensively documented [87]; and several hard-wired feedback loops among these effectors such as the  
290 hypothalamus-mediated regulation of appetite and diet, have also been described [10].

291 A major limitation of our meta-analysis is the lack of consistent behavioral, genotypic or electronic  
292 healthcare record data that would have allowed subtyping of the ASD subjects in light of environmental  
293 confounders. Furthermore, we cannot definitely recommend age- and sex-matching over sibling matching  
294 based on our analysis. Age remains a major confounding factor in early childhood microbiome development  
295 and controlling for this is key for understanding microbial fluctuations [88]. On the other hand, sibling-  
296 matching may help control environmental factors, but mostly rules out the ability to age-match subjects,  
297 thus potentially introducing the age confounder [89]. And while our approach revealed strong associations  
298 among the microbiome, other omic levels, and ASD, the vast heterogeneity in behavioral patterns and in  
299 genotypes is a major obstacle in constructing diagnostics and treatments for ASD symptoms [90, 32].

300 Our analysis has further exposed the limitations of cross-sectional cohort studies and the need for lon-  
301 gitudinal intervention studies to further our understanding of the functional architecture of ASD. Building  
302 realistic causal models of autism needs to take into account the multi-factorial complexity underlying differ-  
303 ent ASD subtypes, which will require a concerted effort to simultaneously analyze several omic levels and at  
304 clinically relevant time scales. For instance, understanding the engraftment dynamics of FMT and its func-  
305 tional implications on the recipients’ gut microbiomes requires frequent initial sampling of the microbiome,  
306 immunome and metabolome, but tracing any behavioral changes over time requires less frequent sampling  
307 over periods of up to several years in combination with reliable behavioral, medical and dietary surveys  
308 [91, 92]. Collecting and integrating such multi-scale omic datasets presents unique logistical and analytical  
309 challenges.

310 Managing data acquisition and access will require coordinating multiple sites and potentially centralizing  
311 some aspects of sample processing. Recent initiatives such as The Environmental Determinants of Diabetes in  
312 the Young (TEDDY) study, an international long-term, multi-center initiative to link specific environmental  
313 triggers to particular Type 1 diabetes-associated genotypes, provide a blueprint for similar approaches in  
314 ASD [93]. A key component of such an initiative would be the establishment of standardized sampling  
315 and processing protocols that would minimize technical confounders, one of the top confounders at most  
316 omic levels. For instance, our analysis showed major batch effects when comparing 16S and SMS datasets  
317 across cohorts ( $r=-0.023$ ,  $p=0.48$ , Figure S5b) as well as within a cohort ( $r=0.17$ ,  $p=1e-5$ , Figure S5b). And  
318 while there are extensive efforts underway to calibrate microbiome datasets [94], other omic levels such as  
319 the metabolome [26] present even more fundamental technical issues that make it imperative to develop  
320 concerted strategies to be able to include them in an integrated analysis.

321 In addition to the considerable variations in statistical properties across datasets, interactions among  
322 omic levels are mostly underdetermined, making the construction of informative models a major challenge.  
323 Determining the necessary biologically relevant and unbiased assumptions is a non-trivial process and can  
324 inadvertently lead to model mis-specifications resulting in misleading conclusions. As pointed out recently  
325 for genetic, environmental and microbiome models in ASD [95, 96], addressing these issues will be critical  
326 to inferring causal mechanisms from population-scale studies. In addition, and given the vast heterogeneity  
327 of ASD, designing cohort studies that minimize confounding factor effects will be key to furthering our  
328 understanding of autism. For example, while our analysis could not identify ASD subtypes implicated in GI  
329 symptoms, we have determined stronger associations between gut microbes, host immunity, brain expression  
330 and dietary patterns than previously reported, highlighting the potential for boosting the statistical power  
331 and biological insight with comprehensive omic analyses. We conclude that multi-omic longitudinal inter-  
332 vention studies on appropriately stratified cohorts, in combination with comprehensive patient metadata,  
333 provide an optimal approach to advance our understanding of the etiology of autism to the next level.

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## 730 Methods

### 731 Search strategy and inclusion criteria

732

733 We performed a systematic search for published and/or publicly deposited or not yet published and/or  
734 publicly available human microbiome, metabolome, immunome, transcriptome AND autism/ASD datasets  
735 in several NCBI databases (PubMed, SRA, and BioProject), UCSD’s MassIVE resource, the PsychENCODE  
736 consortium, the American Gut Project, and from individual research groups worldwide. About half of 70+  
737 studies we identified were already deposited on public data repositories or were made directly available to  
738 us by the research groups.

739 Most studies consisted of heterogenous—no genotype or phenotype stratification—ASD and neurotypical  
740 age- and sex-matched cohorts and had one or two datasets (microbiome [16S, shotgun metagenomic se-  
741 quencing (SMS)], metabolome [urine/serum/fecal], immunome [cytokines], transcriptome [RNAseq], dietary  
742 survey, behavioral survey) associated with them, with only a few studies having three or more omic datasets  
743 associated with them (Table 1). We adopted a multi-cohort and multi-omics meta-analysis framework that  
744 allowed us to combine independent and dependent omic data sets in one overall analysis[43, 44]. In total,  
745 we analyzed 597 ASD-control pairs. To reduce the batch effects and noise associated with primer choice in  
746 the 16S datasets, a major confounder in microbiome analyses, we restricted the 16S datasets to include only  
747 those targeting the variable region V4 of the bacterial ribosomal RNA, a region exhibiting higher hetero-  
748 geneity and lower evolution rates than other variable regions[97, 98, 99]. Our analysis included 16S datasets  
749 obtained targeting the V4 region exclusively, the V3-V4 region, or the V4-V5 region.

750 The final metabolomic meta-analysis we present here consists of the combined analysis of only four  
751 independently preprocessed, normalized, and analyzed metabolomic datasets. Despite several more ASD-  
752 related datasets being available, the disparity in mass spectrometric technologies used to generate them,  
753 which results in the detection of different subsets of metabolites, precluded their side-by-side comparison  
754 (Table 1). For example, targeted mass spectrometry enables the precise determination of concentrations for

755 a finite number of metabolites, whereas untargeted mass spectrometry detects up to two or three orders of  
756 magnitude more metabolites but is compositional in nature and thus does not yield absolute abundances.  
757 Furthermore, batch effects due to sample-processing such as differences in reagents, sample storage and  
758 mass-spectrometry instruments can introduce unwanted variation in both the abundances and the detected  
759 molecular features [26]. One additional obstacle we encountered was the proprietary nature of many of the  
760 metabolomic datasets that made it impossible to access the raw data and run standardized workflows.

761 Of the 40 transcriptomic datasets that were available in recount3 [100], the vast majority were obtained  
762 from studies with model animals, and only four of them had been obtained from postmortem processing  
763 of brain samples from autistic and neurotypical individuals. These four datasets collected different brain  
764 tissue types, including from the amygdala, the prefrontal cortex, the anterior cingulate and the dorsolateral  
765 prefrontal cortex.

## 766 Data processing

767  
768 16S amplicon and shotgun metagenomics samples were downloaded from the SRA. The 16S amplicon samples  
769 were processed using Deblur and subsequently mapped to bacterial whole genomes captured in the Web of  
770 Life using Woltka[101]. This is done in order to make the amplicon data comparable to shotgun metagenomics  
771 data. Bacterial abundances were extracted from shotgun metagenomics samples using Woltka and Bowtie2.  
772 Viral abundances were extracted from shotgun metagenomics samples using GPD and BWA. RNA expression  
773 data were obtained directly from recount3 [100]; the four metabolomics datasets were provided by the authors.

774 The approach of mapping both 16S amplicon sequences and SMS samples to a common set of microbial  
775 reference genome provided a consistent taxonomic annotation between the different 16S amplicon types and  
776 SMS datasets. However, there are notable limitations in taxonomic resolution in all of these datasets. For  
777 instance, multiple *Bifidobacterium* species that are associated with non-human microbiomes were found in-  
778 cluding *B. asteroides* (honeybee), *B. callitrichos* (marmoset), *B. choerinum* (pig) and *B. sanguine* (tamarin).  
779 We observed that these 16S amplicons were multi-mapped across many different *Bifidobacterium* genomes,  
780 which highlights the lack of species level resolution highlighted in previous studies [102]. Similarly, taxo-  
781 nomic profiles obtained from shotgun sequencing are known to have elevated false positive rates taxonomic  
782 identifications due to high genome similarity between microbial reference genomes [59].

783 To enable age- and sexmatching, a bipartite matching between ASD and neurotypical subjects was  
784 performed using age and sex covariates. Subjects that could not be matched were excluded from the meta-  
785 analysis. Amongst the 16S and SMS datasets, there were multiple longitudinal datasets. To integrate these  
786 datasets into the cross-sectional analysis, we only picked the first time point for each subject.

## 787 Differential ranking analysis

788  
789 One of the most common approaches to evaluating microbiome and other omic studies consists of determining  
790 differences in the abundances of microbial taxa, human metabolites or other omic features between cases and  
791 controls [103]. Such differential abundance analysis is typically performed by computing the log fold changes  
792 between the case and control groups[104, 46, 105]. However, confounders such as sex-, age-, and geography-  
793 related batch effects, compositionality, high-dimensionality, over-dispersion, and sparsity, prevented a reliable  
794 estimation of differential abundances and thus compromised the side-by-side comparison of these differential  
795 abundances across studies in the manner of a traditional meta-analysis [106, 107]. Here, we set out to  
796 overcome these inherent limitations of traditional meta-analyses by developing a generalizable approach  
797 for controlling for select confounders that would help reveal a comprehensive picture of ASD-specific omic  
798 signals.

799 To minimize confounder effects, we developed a Bayesian differential ranking algorithm that used bipar-  
800 tite matching to optimize the age- and sex-based pairing of ASD and control subjects within each dataset.  
801 This approach helped both control for potential age and sex confounders and minimize batch effects such as  
802 sample collection method, sample processing protocol, and geographical provenance [108]. These Bayesian  
803 models were fitted via MCMC using Stan [109]. Conceptually, this allowed us to compute log-fold change  
804 differences of microbes between age- and sex-matched subjects, but because we did not have absolute abun-  
805 dance information we could only estimate this log-fold change up to a constant [31] (Supplemental methods).  
806 To determine if there was a significant difference between the age- and sex-matched pairs, we constructed an  
807 effect size metric utilizing our model’s uncertainty estimation (see Supplemental methods for more details).  
808 To show that this model is relevant for biological data, we built a simulation benchmark using the 16S count  
809 data. Specifically, we fitted the Bayesian model on the 16S cross-sectional cohort, and simulated microbial  
810 counts based on those estimated parameters. We showed that the ground truth log-fold changes across all  
811 of the microbes are within the 95% credible intervals estimated by our algorithm. When we evaluated our  
812 Bayesian model fit on the 16S, SMS and RNAseq datasets, our models fits achieved  $R_{hat}$  values below 1.1  
813 and ESS values above 300, indicating that the draws from the posterior distribution are reliable [109].

814 To identify microbes that were ASD-specific or neurotypical specific, we fitted a Gaussian mixture model  
815 on top of the estimated log-fold changes, binning the taxa into three different groups, those taxa hypothesized  
816 to be more abundant in neurotypical controls, those more abundant in ASD children and those that are  
817 equally prevalent in both groups, or are “neutral”.

818 This strategy was inspired by the work done with ANCOM-BC [110]. The major difference in our ap-  
819 proach compared to ANCOM-BC is that our approach assumes a negative binomial distribution for modeling  
820 counts and allows for Bayesian model uncertainty quantification. The reference frame in the cross-sectional  
821 analysis refers to the average abundance of the microbes that are categorized as neutral in Figure 1a. These  
822 same microbes were used to construct a reference frame in the FMT analysis to standardize all of the time

823 points. The FMT analysis used the same matching strategy, but instead of matching on age and sex, the  
824 matchings were performed on the subjects to compare different time points.

825 The heatmap shown in Figure 1 displayed the log-fold changes for each case-control pair. To do this, a  
826 robust CLR transform was performed and all zeros were imputed to the mean abundance for visualization  
827 purposes. The case-control log-fold changes were computed for each pair as highlighted in Figure S7.

## 828 Other methods

829

830 We fitted Random Forests models on nine 16S datasets and on three SMS datasets. We randomly split the  
831 samples into 90/10 training and test splits, performed a 10-fold cross-validation on the training datasets to  
832 obtain optimal model parameters, and computed predictions on the held-out test dataset. PERMANOVA  
833 with Bray-Curtis distances was used to determine if confounding variation due to household, age and sex  
834 were statistically significant in the sibling cohorts.

835 We used MMvec [111] to perform the diet-microbe co-occurrence analysis. Here, microbes were used to  
836 predict dietary intake. This analysis enabled the estimation of conditional probabilities, namely the probabili-  
837 ty of observing a dietary compound given the microbe was already observed. To estimate these conditional  
838 probabilities, MMvec performs a matrix factorization, identifying the factors that explain the most infor-  
839 mation in these interactions. We compared the MMvec microbial factors against the cross-sectional log-fold  
840 changes. We then compared the MMvec dietary factors against t-statistics that measure the differences in  
841 dietary compounds between ASD and neurotypical children.

842 To identify candidate viral-microbe interactions, we ran MMvec on each of the SMS datasets. We then  
843 pulled out the top co-occurring viral taxa for each microbe that had a conditional log-probability greater  
844 than 1, amounting to 78580 microbe-viral interactions. Then we filtered out the microbe-viral interactions  
845 that were not present in the GPD [70], leaving 31276 microbial-viral interactions.

846 We used Songbird [31] to perform the cytokine-microbe analysis via a multinomial regression that used  
847 the cytokines to predict microbial abundances. We reported biased microbial log-fold changes with respect to  
848 cytokine concentration differences. Pearson correlation was used to determine the agreement between the 16S  
849 cross-sectional microbial differentials and the microbe-cytokine differentials. To directly link these microbial  
850 abundances to the cytokine concentrations, we computed log-ratios, or balances, of microbes for each sample.  
851 For example, for IL-6 the numerator consisted of the top 30 microbes that are estimated to increase the most  
852 in abundance when IL-6 concentration increased, and the denominator consisted of the bottom 30 microbes  
853 which are estimated to be the most decreased when IL-6 concentration increases. Once these partitions are  
854 defined, the balances for each sample are computed by taking the log-ratio of the average abundance of the  
855 numerator group and the denominator group. See Morton et al 2017 for more details behind balances [112].  
856 Pearson correlation between these balances and the cytokine concentrations are then computed to measure  
857 the agreement between the microbial abundances and the cytokine concentrations.

858 To identify key microbial genes, we performed a comparative genomic analysis in which we binned  
859 the microbial genomes into those associated with ASD and those associated with control subjects. Using a  
860 binomial test, we were able to determine if a particular gene was more commonly observed in ASD-associated  
861 microbes than by random chance. Significant microbial genes and RNA transcripts were subsequently  
862 mapped to KEGG pathways. To directly compare the two contrasting omics levels and gauge metabolic  
863 similarity, we retrieved all the molecules involved in both the microbial and human pathways and calculated  
864 their intersection. Since the metabolomics datasets were not directly comparable, we performed Wilcoxon  
865 tests on age- and sex-matched metabolomics samples within each cohort separately. While our analysis  
866 revealed multiple metabolites that were below the 0.05 p-value threshold, none of these metabolites passed  
867 the FDR corrected threshold.

## 868 Software Availability

869

870 Software implementation of our Bayesian age-sex matched differential ranking algorithm can be found at  
871 <https://github.com/flatironinstitute/q2-matchmaker>

872 We want to acknowledge Matplotlib [113], Seaborn [114], Scipy [115], Numpy [116], Xarray [117], Arviz [118],  
873 Scikit-learn [119], biom-format [120] and Scikit-bio [121] for providing the software foundation that this work  
874 was built upon.

875 **Tables and Figures**

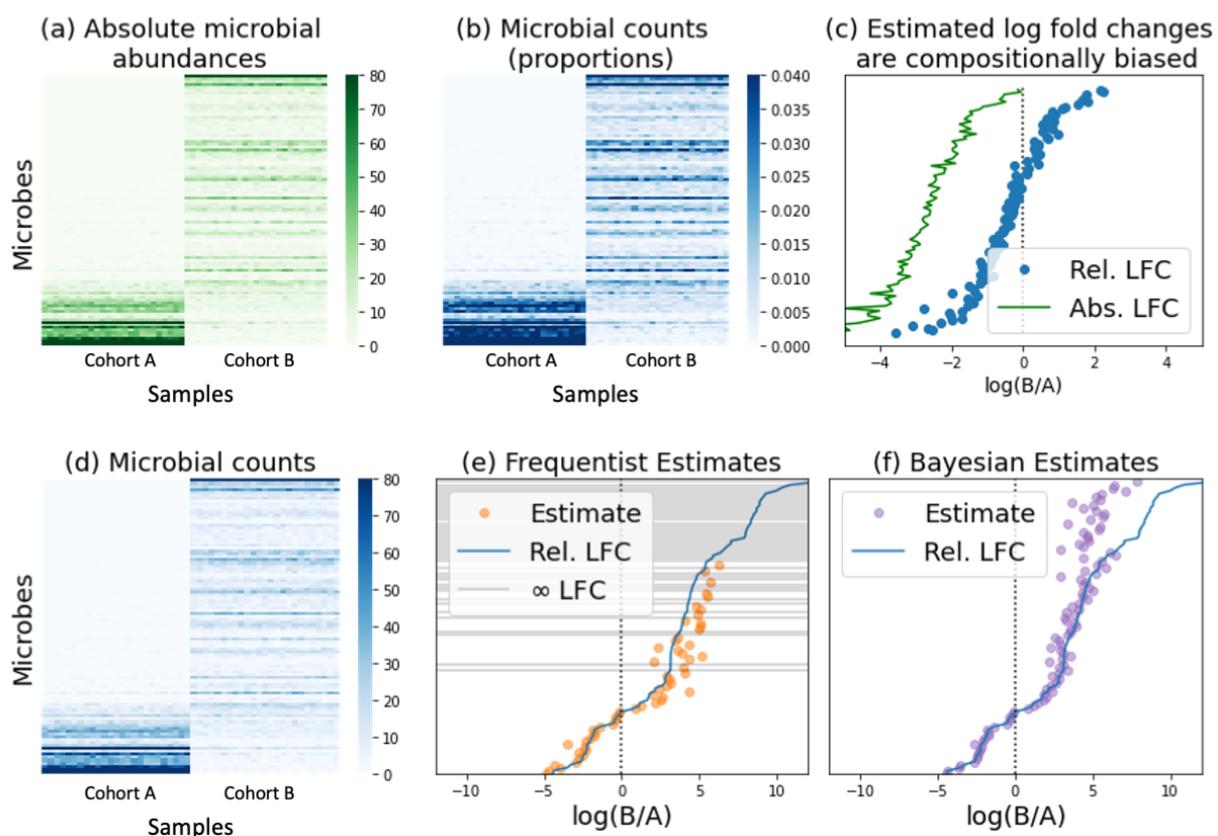


Figure BOX 1: **TACKLING METAGENOMIC UNKNOWNNS**

Metagenomic sequence data present unique quantification challenges due to a lack of total microbial load measurements, which precludes the determination of absolute microbe abundances, and to limitations brought about by sampling and sequencing depth limitations, which result in an incomplete representation of the metagenome. We devised a Bayesian differential ranking algorithm to address both these challenges, the compositional challenge and the zero-inflation challenge.

**The compositional challenge:** Most sequencing count datasets lack absolute abundance information in the form of cells, colony forming units, or transcripts per volume. This limitation preempts the reliable estimation of log fold changes and is a defining characteristic of compositional data that can lead to excessive false positives or false negatives depending on the magnitude of the change in absolute abundances [31, 45]. As illustrated in panels a) through c), microbial counts (a) are typically converted into proportional abundances (b) that are then used to compute log-fold ratios. Fold change calculations adopt the general formula  $\frac{B}{A} = \frac{N_B p_B}{N_A p_A} = \frac{p_B}{p_A} \times \frac{N_B}{N_A}$ , where  $A$  and  $B$  represent the two samples being compared,  $p_A$  and  $p_B$  represent the microbial proportions in  $A$  and  $B$ , and  $N_A$  and  $N_B$  represent the total number of microbes in  $A$  and  $B$ , also known as the ground truth. A key limitation of sequencing count data is their lack of proportionality to the corresponding absolute abundances in the original samples due to sequencing depth constraints [122]. Our inability to observe  $N_A$  and  $N_B$  introduces a bias that ultimately prevents us from performing FDR correction to identify differentially abundant microbes [123]. This bias depends on the change in microbial population size, with large population shifts leading to increased false positive and false negative rates, and an overall skewed representation of the ground truth (c).

**The zero-inflation challenge:** Sampling errors and shallow sequencing lead to disproportionately high numbers of zero counts, especially for microbes present in low abundances (d). Multinomial, Poisson and Negative Binomial distributions have been used to explicitly handle zero counts [46]. However, estimating log-fold differentials remains problematic when microbes are not observed in any of the samples in one group since  $\log 0$  is  $-\infty$  and thus the true log-fold change of a zero-count microbe can not be determined (e). Bayesian inference avoids this problem by introducing a prior that prevents nonsensical log-fold change estimates (f). Specifically, this introduces a rounded-zero assumption whereby all microbes have a non-zero chance of being observed. Panel h highlights what these log-fold changes would look like using a Dirichlet prior, where every microbe has the same probability of being observed before collecting data.

Organism	Body type	Data type	Num Studies	Num Subject pairs	References
Human	Postmortem brain tissue	RNAseq	4	49	(Velmeshev et al. 2019; Wright et al. 2017; Herrero et al. 2020) SRP072713
Human	Serum	Immune markers	1	22	(Zurita et al. 2020)
Human	Serum	Metabolome	2	50	(Needham et al. 2021; Kuwabara et al. 2013)
Human	Urine	Metabolome	1	26	(Noto et al. 2014)
Human	NA	Dietary survey	1	26	(Berding and Donovan 2019)
Human	NA	Behavioral survey	1	28	(Kang et al. 2019)
Microbial	Fecal	Metabolome	2	43	(Needham et al. 2021; Kang et al. 2018)
Microbial	Fecal	16S amplicon	10	346	(Berding and Donovan 2019; Zurita et al. 2020; Dan et al. 2020; J. Zhu et al. 2021; Fouquier et al. 2021; Zou et al. 2020; Kang et al. 2019, 2017; Son et al. 2015; David et al. 2021) SRP299486
Microbial	Fecal	Shotgun metagenomics	3	83	(Averina et al. 2020; M. Wang et al. 2019; Dan et al. 2020)

Table 1: ASD omic datasets included in this study All sequencing datasets were retrieved from the SRA. [124].

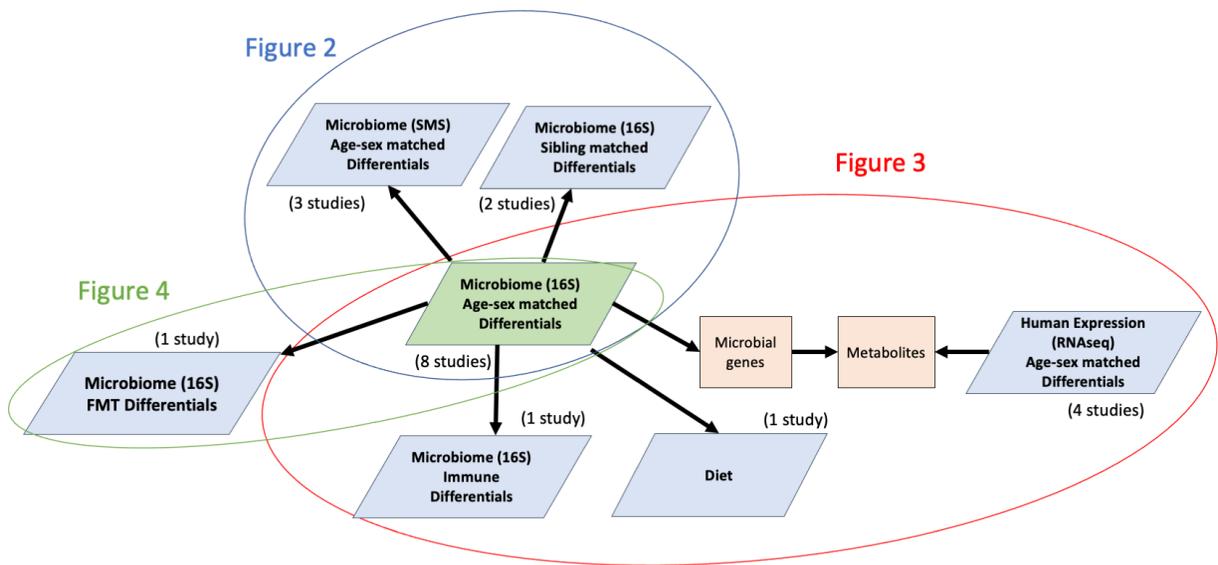


Figure 1: The structure of our meta-analysis across multiple omics levels. For Figure 2, microbial differential abundance on 16S data from age-sex matched cohorts was cross-referenced against sibling matched cohorts and SMS data from other age-sex matched cohorts. For Figure 3, these same 16S microbial differential abundances were cross-referenced against cytokine profiles, dietary surveys and pathways from RNAseq. For Figure 4, the differentially abundant microbes from the age-sex matched analysis was cross-referenced against the Kang et al FMT trial.

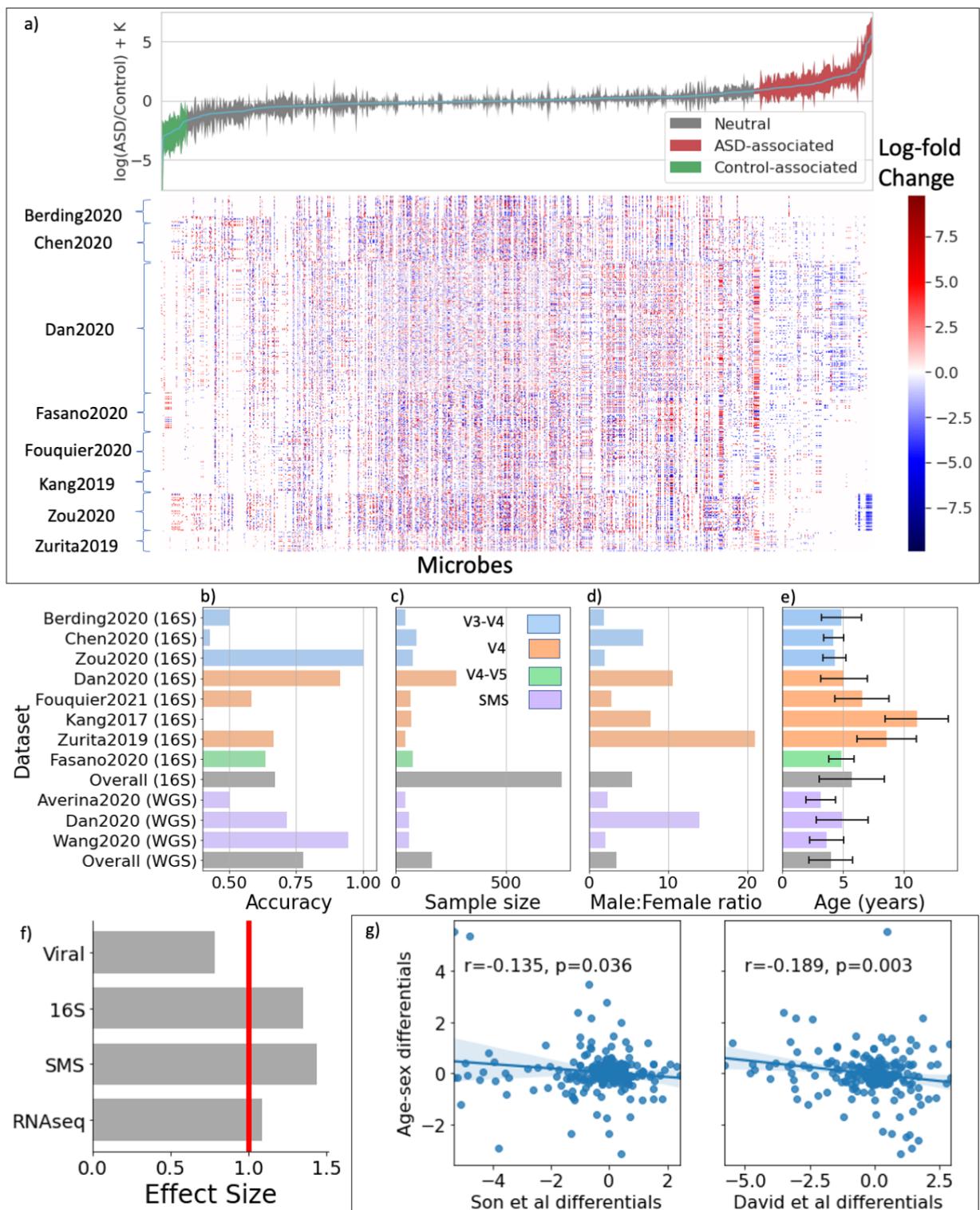


Figure 2: Differential ranking analysis across omics levels. (a) Global microbial 16S log-fold changes between age- and sex-matched ASD and control individuals. Error bars represent the 95% credible intervals. Heatmap showing all CLR transformed microbial differentials for each age- and sex- matched ASD-control pair across all cohorts. (b-e) Held-out random forests ASD classification accuracy, sample size, male:female ratio and age distributions across all 16S and shotgun metagenomics datasets analyzed in this study. V3-V4, V4, and V4-V5 refer to the variable region of the bacterial ribosomal RNA analyzed; SMS refers to shotgun metagenomic sequencing. (f) Effect sizes of different omics levels: viral, 16S, SMS, and RNAseq. (g) Comparisons of sibling-matched differentials from two different studies to the global age- and sex-matched differentials.

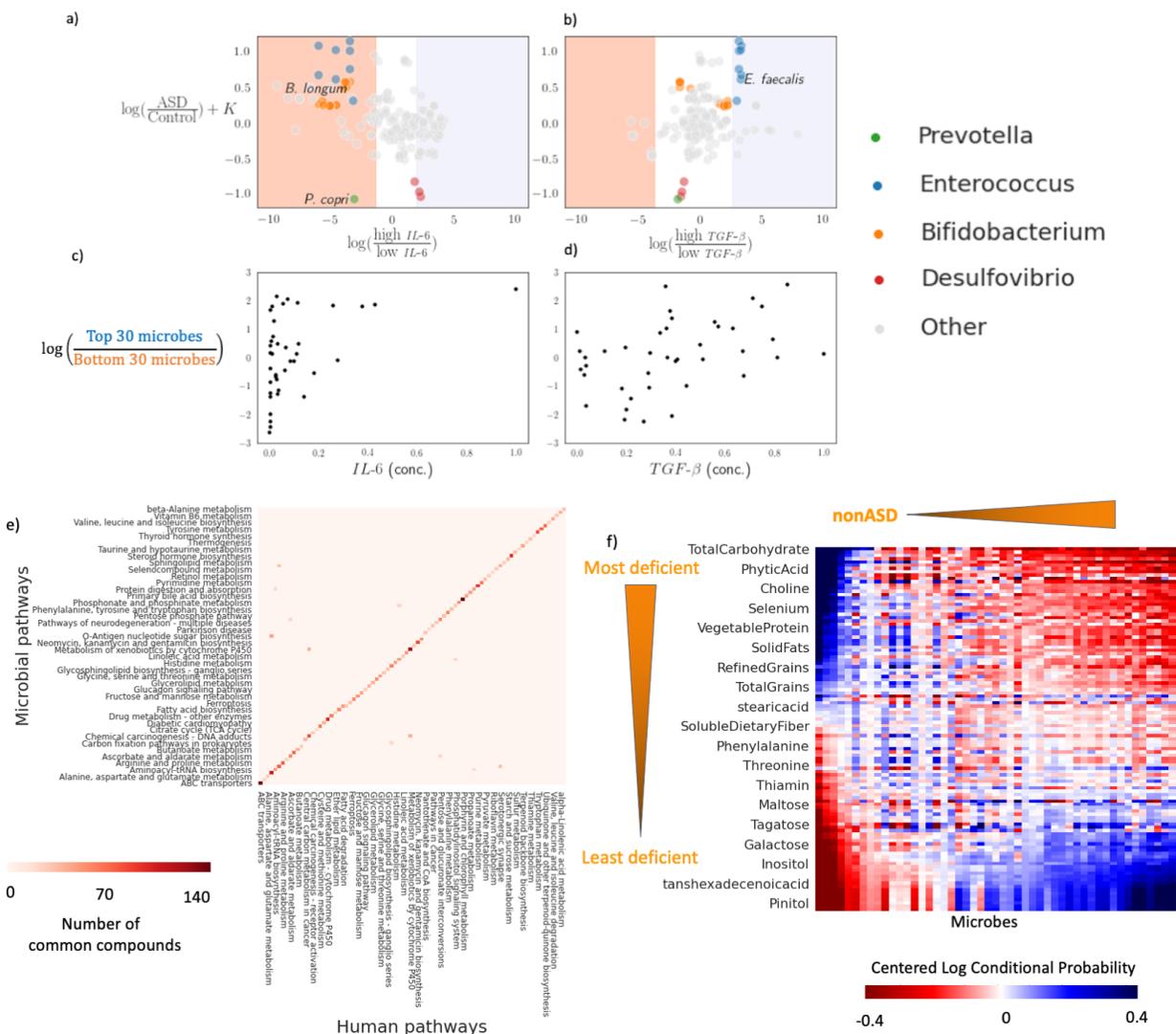


Figure 3: Characterizing the associations between differentially abundant microbes in ASD and cytokines, gene expression in the brain, and dietary patterns. (a-b) Comparison of microbial differentials obtained from age- and sex-matching and cytokine analysis. LFC denotes log-fold change of microbial abundances with respect to a specific cytokine. (c-d) Microbial log-ratios constructed from 30 top and bottom most differentially abundant microbes corresponding to each cytokine. (e) Heatmap showing the overlap of molecules between ASD-enriched pathways in the microbiome and in the brain. (f) Co-occurrence analysis between diet and microbes in ASD.

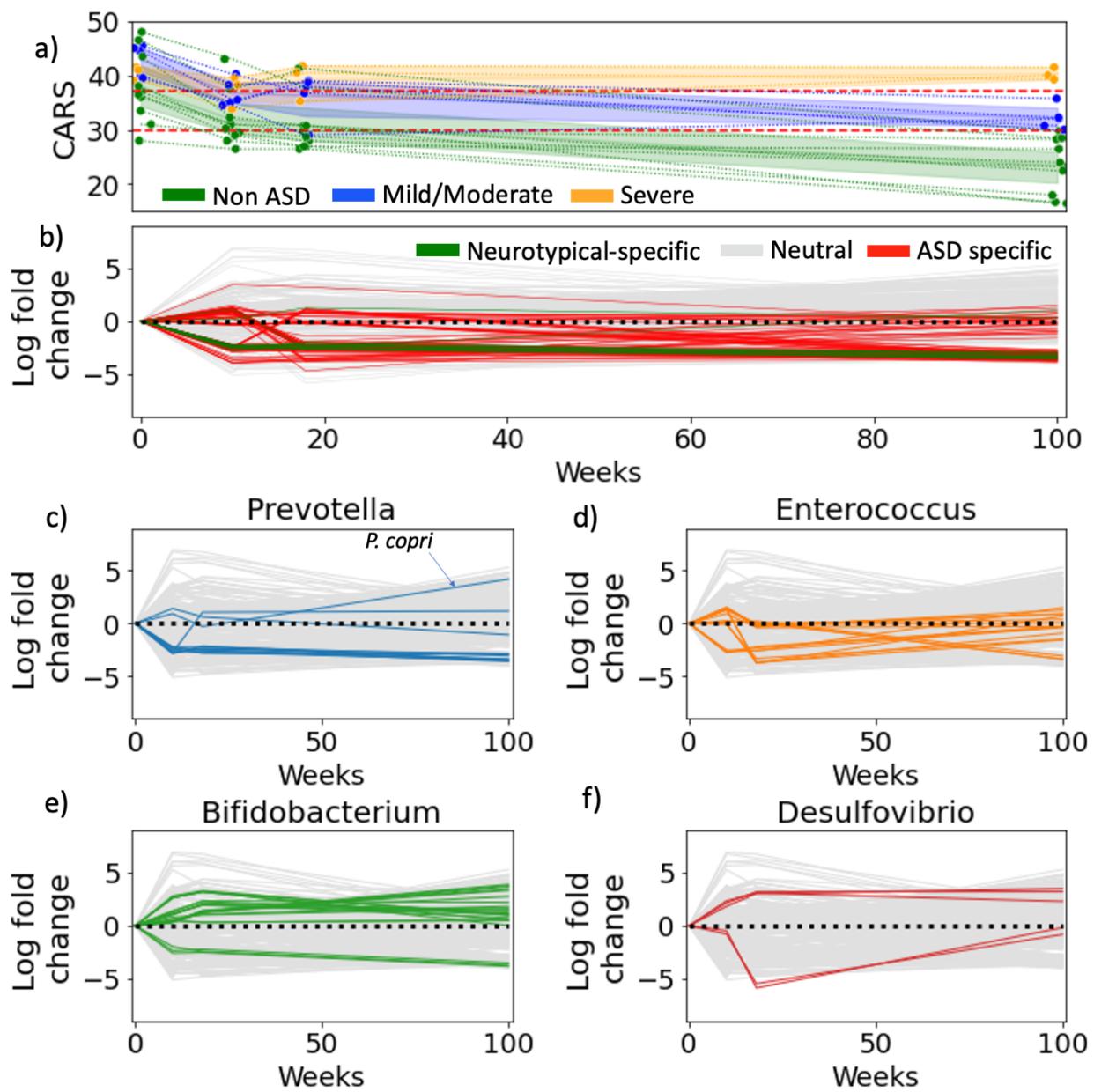


Figure 4: Fecal matter transplants have long-lasting effects on autism gut microbiomes. (a) The improvement of CARS for each ASD child over time. The children are split into 3 groups, non-ASD, mild/moderate and severe based on whether their CARS score fell below 30, between 30-37 or greater than 37 (b) Microbial log-fold changes over time: the time series was generated by calculating log-fold changes between time points for each microbe. ASD-specific microbes highlighted in red were determined in the cross-sectional study. (c-f) Microbial log-fold changes are re-colored with genera highlighted in cytokine comparisons.

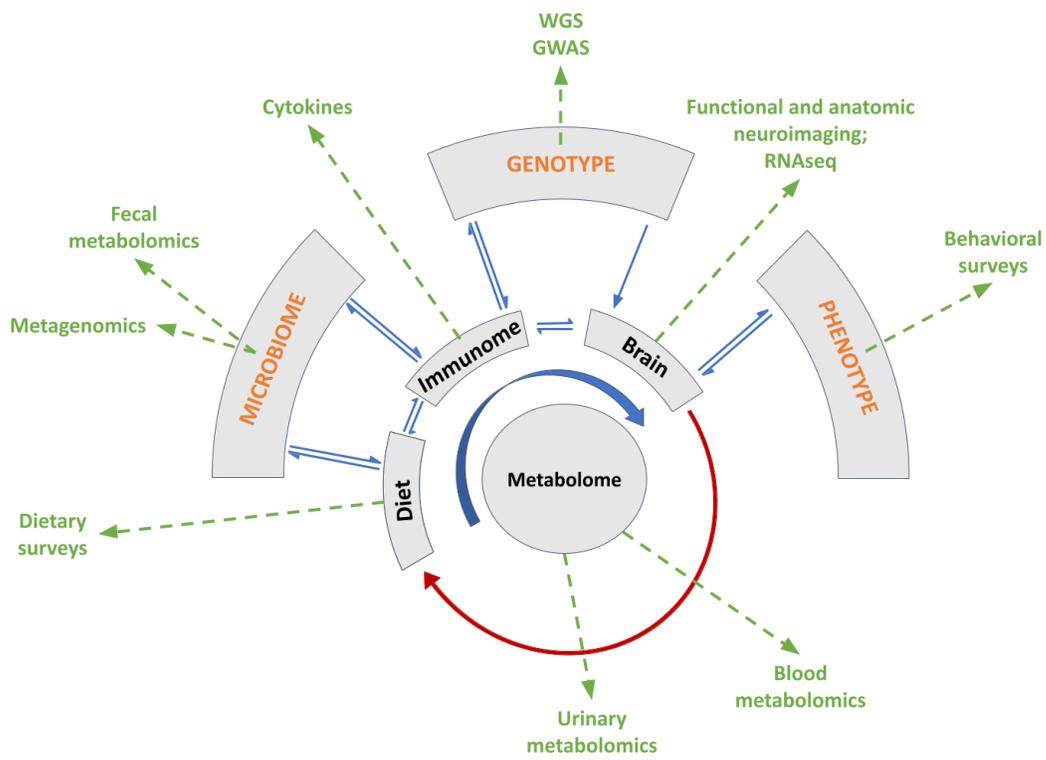


Figure 5: Proposed functional architecture of ASD Hypothesized causal graphs underlying relevant omic levels along the GBA in ASD, and experimental considerations for future studies. Blue arrows denote causal direction, red arrows indicate feedback loops and green arrows indicate measurable data types.

## 876 Supplemental Materials

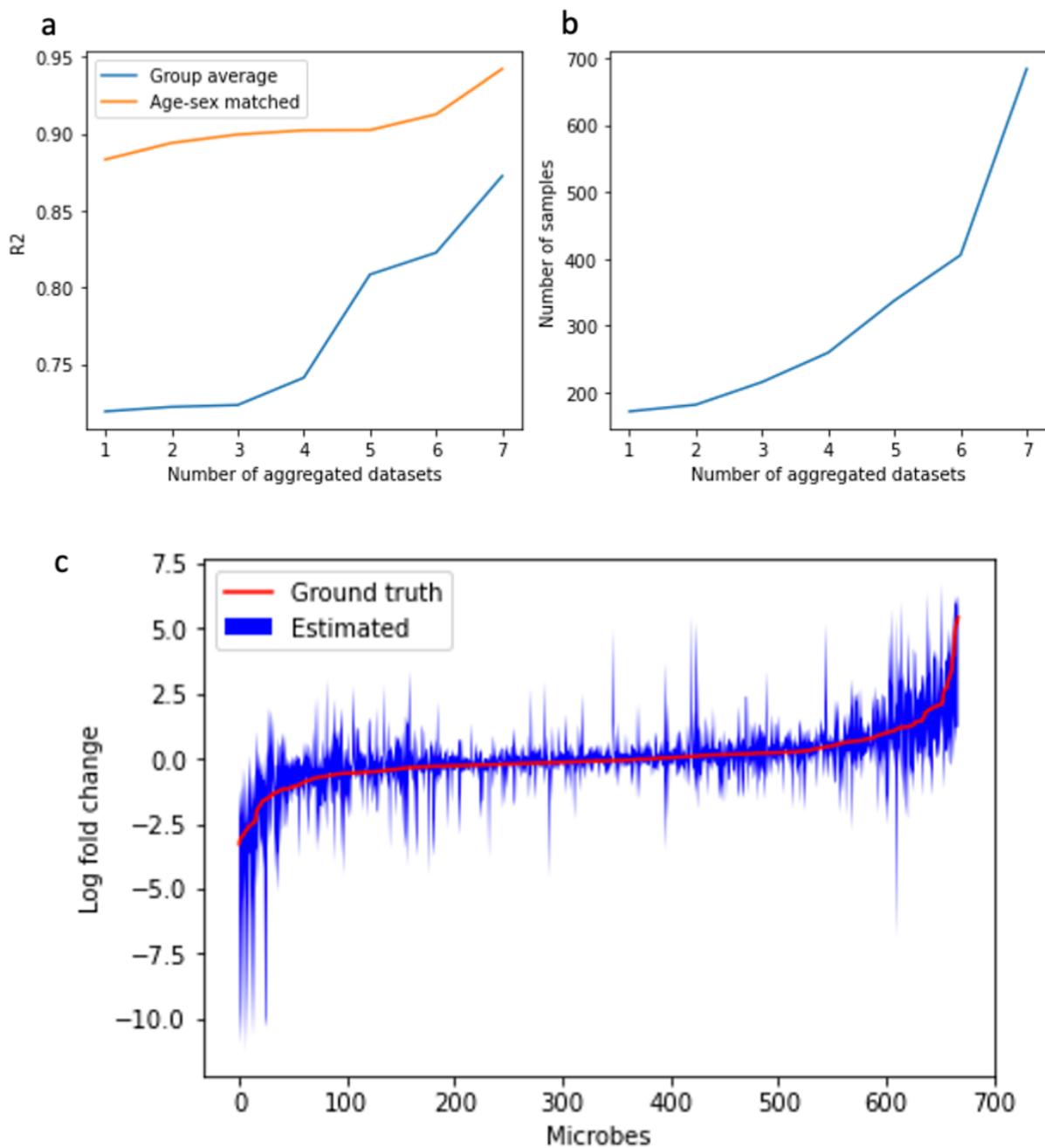


Figure S1: Benchmarks. (a) Comparison of age- and sex-matching approach compared to standard group averaging with respect to dataset size across 7 of the 10 16S studies (excluding Kang et al, David et al and Son et al). (b) Number of samples analyzed. The x-axis represents the number of aggregated datasets, the y-axis on the left panel is the average  $R^2$  metric to measure the model error, and the y-axis on the right panel is the number of samples in the aggregated dataset. (c) Differential abundance estimation derived from a simulated datasets modeled from the cross-sectional cohort from the 8 16S datasets.

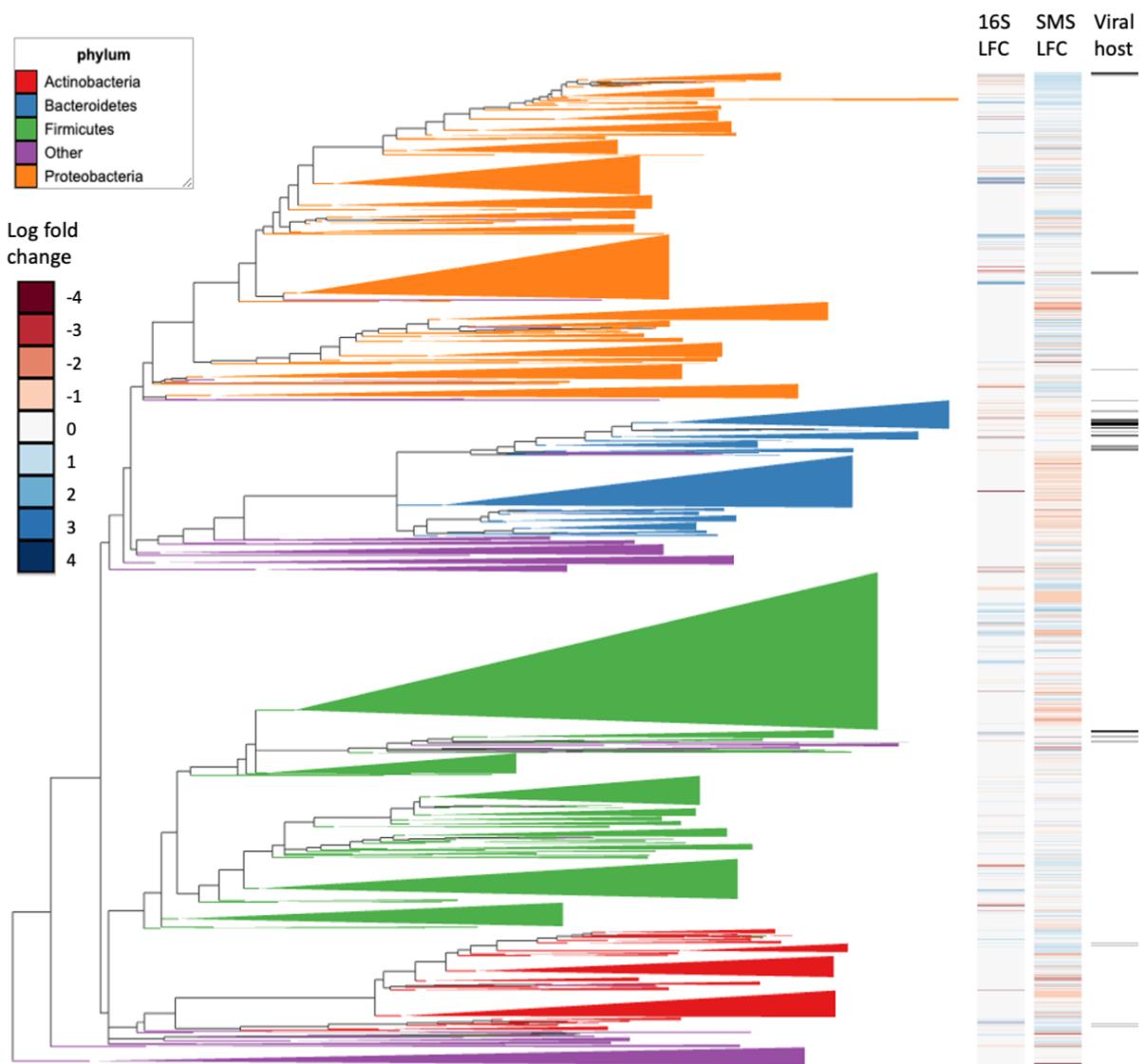


Figure S2: Phylogenetic visualization of microbes with respect to the differentials computed from 16S and SMS differentials. Microbes that were annotated as viral hosts were also highlighted. The phylogenetic visualization was generated using Empress [125].

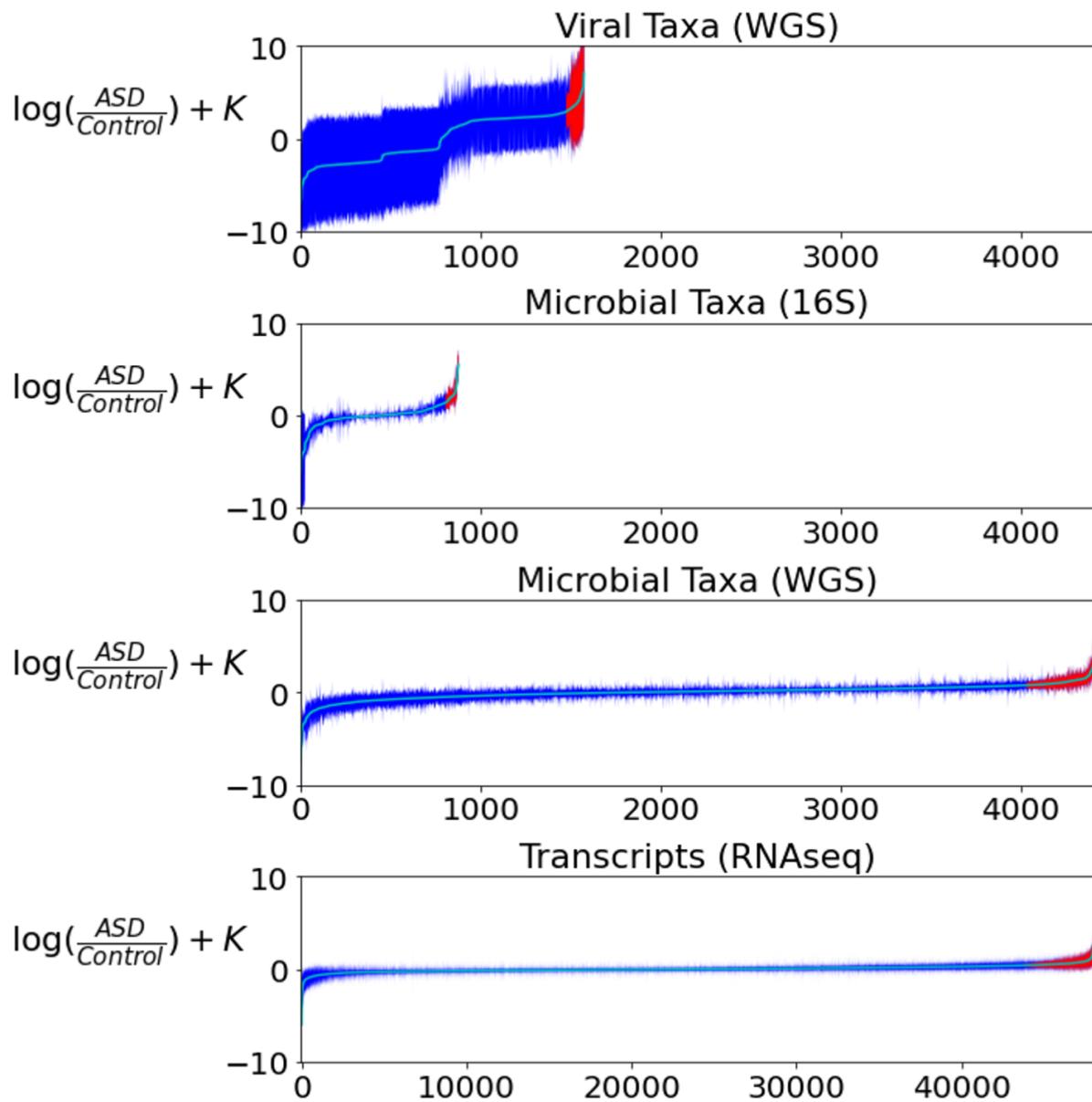


Figure S3: Global differential ranking trends observed for the virome, 16S, SMS, and RNAseq datasets analyzed in this study. The x axis for the virome, 16S and SMS datasets is equivalent to showcase the differences in feature counts; the x axes for the RNAseq dataset is larger by a factor of 10, illustrating the stark difference in number of features of this dataset compared to the other three.

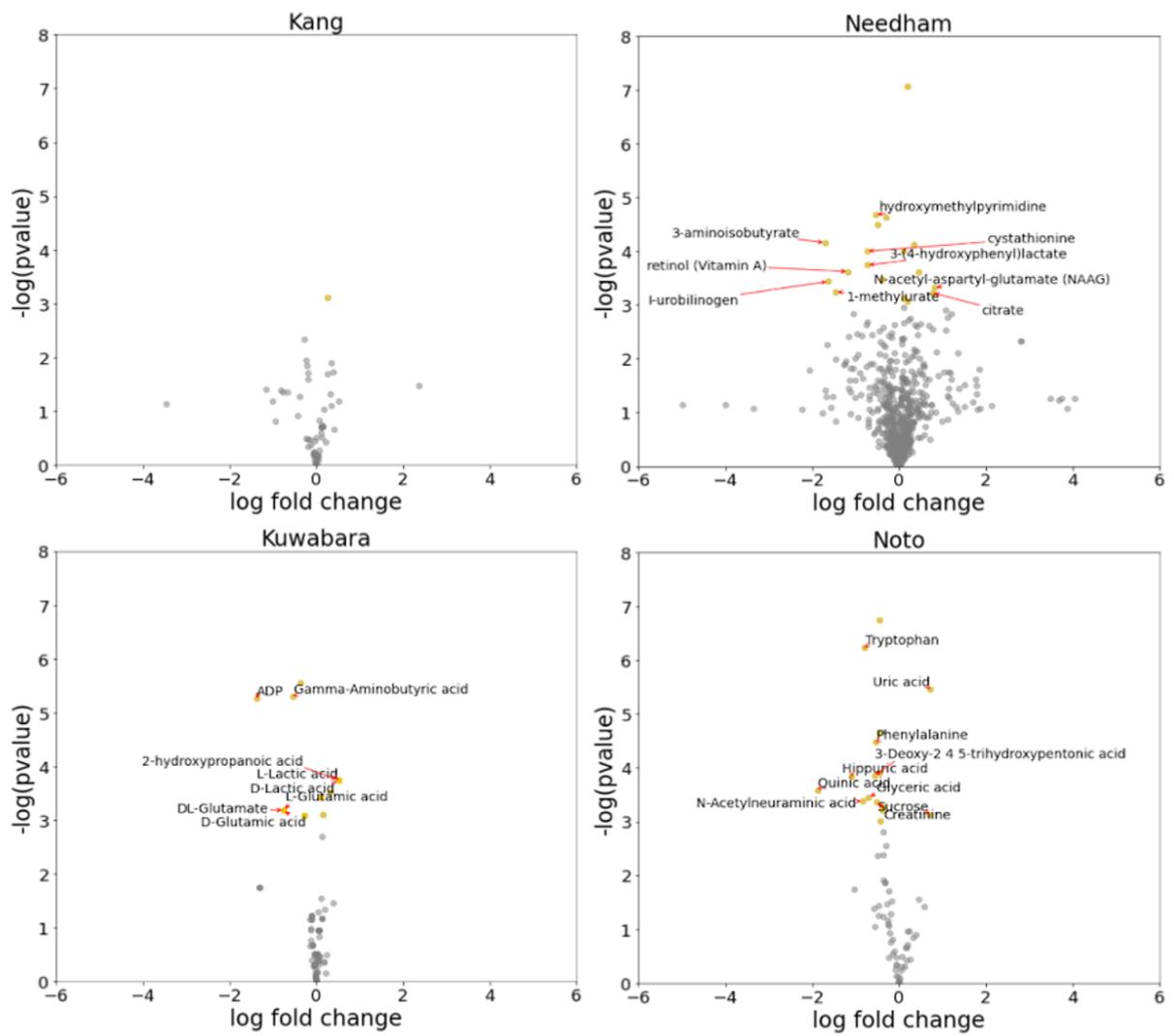


Figure S4: Metabolomics differential ranking analysis across four studies. Paired t-tests were performed to identify differentially abundant metabolites. The metabolites shown in Needham et al consist of both fecal and serum metabolites. None of the metabolites had significant log-fold changes after applying FDR correction.

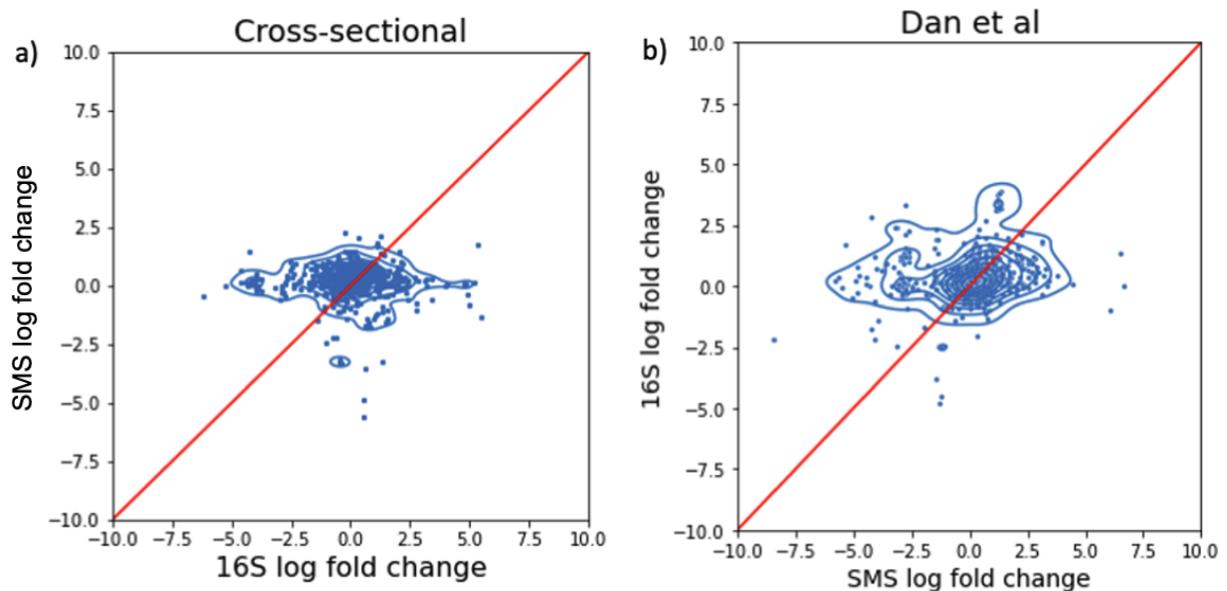


Figure S5: Comparison of log-fold changes computed from 16S and SMS. (a) Comparison of differentials obtained from 16S and SMS on the cross-sectional datasets. (b) Comparison of differentials obtained from 16S and SMS on the same samples from Dan et al.

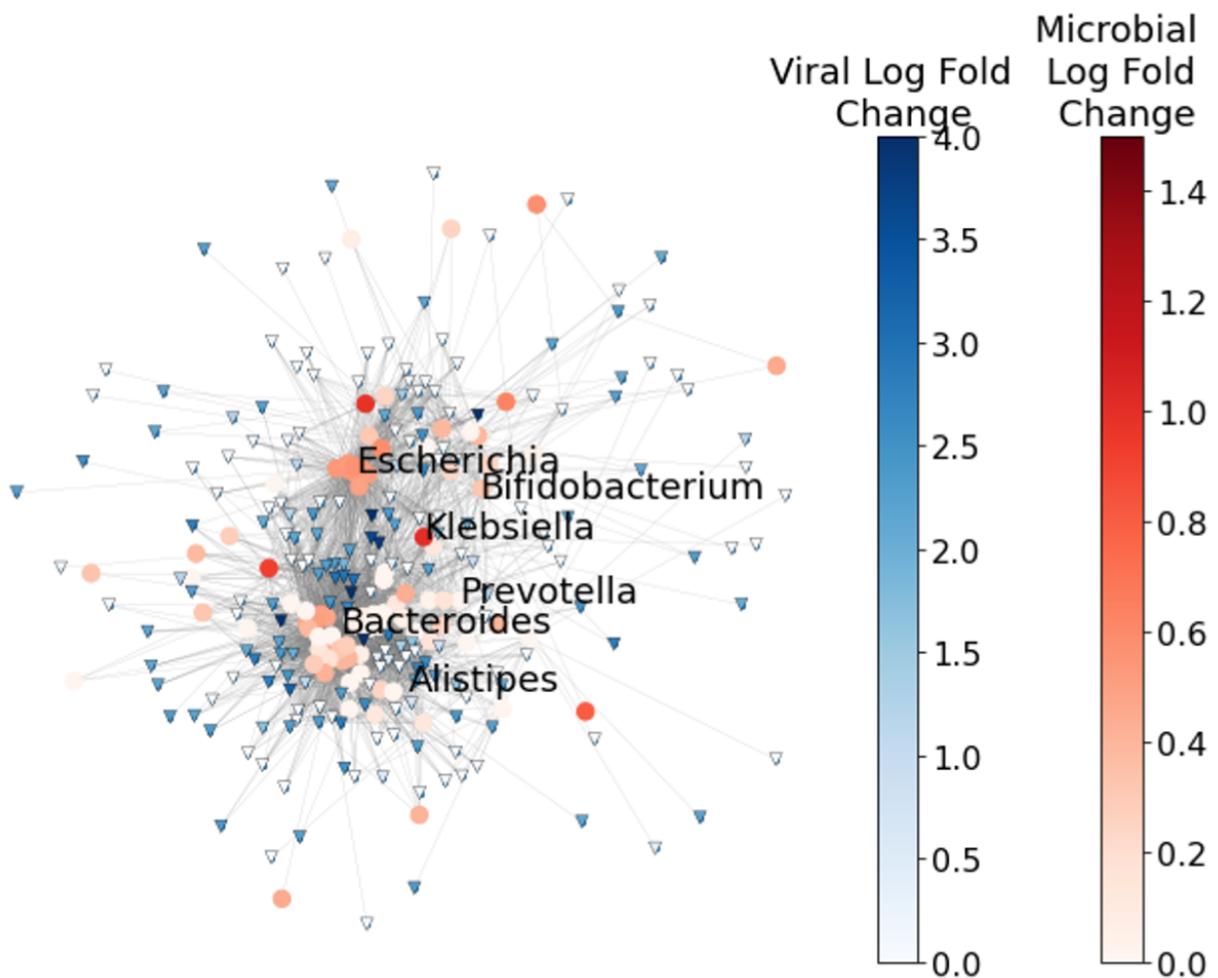


Figure S6: Microbe-viral co-occurrence network estimated using MMvec. Microbes are colored red and viruses are colored blue. Edges are drawn between microbes and viruses if they are highly co-occurring and the interaction was annotated in GPD.

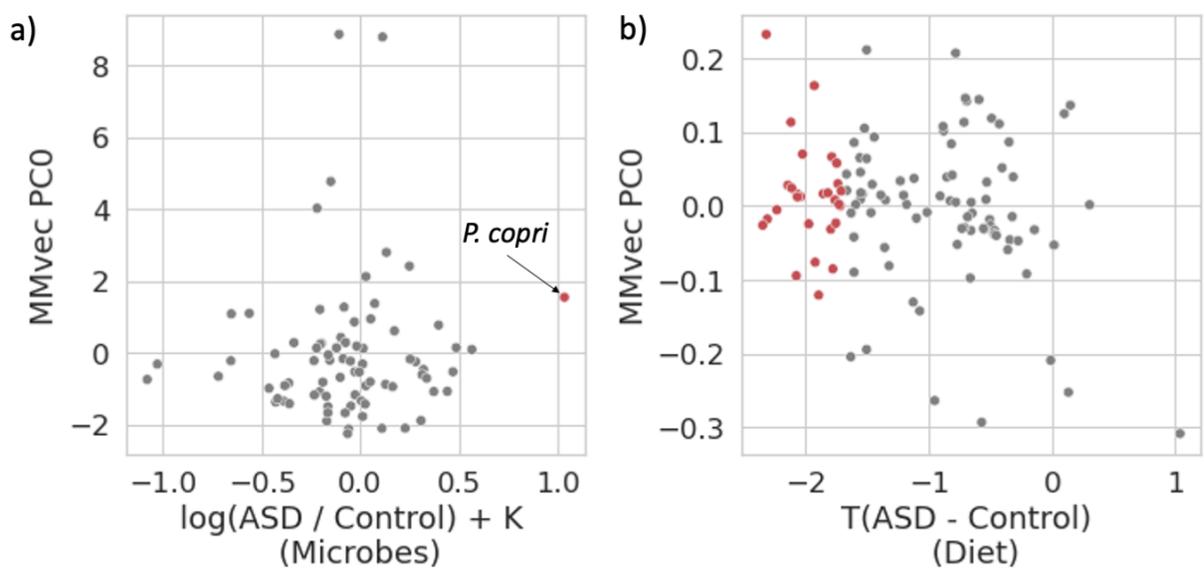


Figure S7: Principal components analysis of microbe-diet interactions. The top principal component explaining the variation in the microbe-diet co-occurrences is compared against the (a) microbial log fold change and (b) dietary differences computed from a t-statistic. Dietary compounds that are significantly significant before FDR correction are highlighted in red.

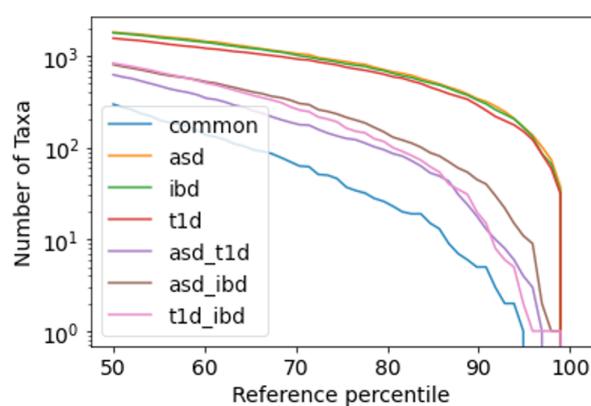


Figure S8: Comorbidity analysis. The number of taxa in common between diseases from differential ranking analysis are shown, with a focus on the intersections between ASD, IBD and T1D.

Table S1: Table of statistics for 16S differentials, including mean log-fold change, standard deviation log fold change, 95% credible intervals and taxonomy for each microbe.

Table S2: Table of statistics for SMS differentials, including mean log-fold change, standard deviation log fold change, 95% credible intervals and taxonomy for each microbe.

Table S3: Table of statistics for RNAseq differentials, including mean log-fold change, standard deviation log fold change, 95% credible intervals and taxonomy for each transcript.

Table S4: Table of statistics for viral differentials, including mean log-fold change, standard deviation log fold change, 95% credible intervals for each virus.

Table S5: Permanova breakdown of sibling matched cohorts looking at the confounding variation due to age, sex and household.

Table S6: Microbial log fold changes due to cytokine differences, including mean log-fold change for each cytokine.

Table S7: Microbe virus co-occurrences, where entries represent the centered log-probability of a microbe and a virus both present for a given sample

Table S8: A list of paired microbe and human pathways in addition to the number of overlapping metabolites

Table S9: Microbe diet co-occurrences, where entries represent the centered log-probability of a microbe and a dietary compound both present for a given subject.

Table S10: Microbial log fold changes between paired time points across all of the subjects in the FMT study.