



Erratum: The Genetic Intersection of Neurodevelopmental Disorders and Shared Medical Comorbidities – Relations that Translate from Bench to Bedside

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An erratum on

The Genetic Intersection of Neurodevelopmental Disorders and Shared Medical Comorbidities – Relations that Translate from Bench to Bedside

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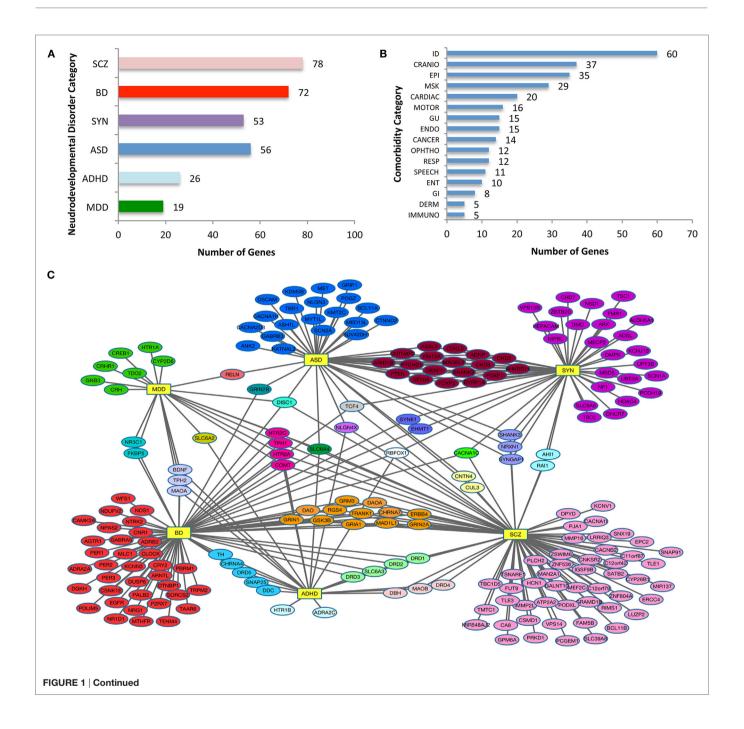
Reason for erratum:

Due to a typesetting error, some information went missing on the y-axis of **Figure 1B** in the original publication.

The publisher apologizes for this error, and the correct version of **Figure 1** appears below. This error does not change the scientific conclusions of the article in any way.

The original article has been updated.

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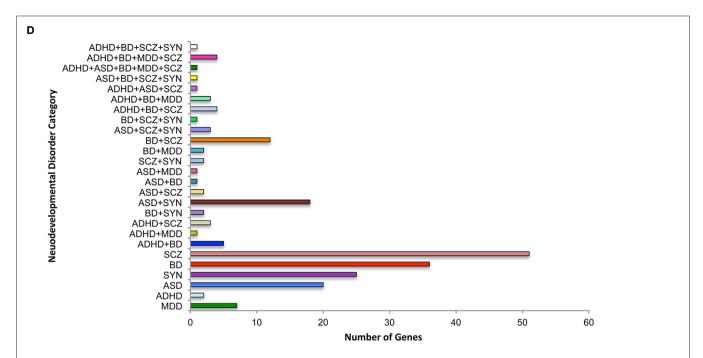


FIGURE 1 | Categorization of neurodevelopmental disorder (NDD) risk genes and their associated comorbidities. (A) NDD categories and the number risk genes for attention-deficit hyperactive disorder (ADHD), autism spectrum disorder (ASD), bipolar disorder (BD), major depressive disorder (MDD), schizophrenia (SCZ), and syndromic neurodevelopmental disorders (SYN). Note that because a single gene may be assigned to more than one category, the total is greater than the 208 genes analyzed. (B) The distribution of NDD risk genes with OMIM-generated, related non-psychiatric conditions. The phenotypic categories listed in (A,B) are not mutually exclusive. (C) Diagram of relationship between NDD risk genes. NDD risk genes depicted with their primary psychiatric associations. Abbreviations for each NDD are as noted for (A), and are shown as hubs (yellow rectangles). Note the connection of NDD genes associated with one or more psychiatric conditions. (D) Histogram with color code that depicts NDD risk genes associated with a single or two or more psychiatric categories, the latter comprising nearly half of the 208 psychiatric NDD genes. NDD abbreviations are noted in Table 1. X-axis indicates number of genes in each combination or single category.

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