Supplemental data
at Neurology.org/ng

Ziv Gan-Or, MD, PhD

GBA P.T369M SUBSTITUTION IN PARKINSON DISEASE: POLYMORPHISM OR ASSOCIATION? A META-ANALYSIS

The lysosomal enzyme glucocerebrosidase (GCase), encoded by GBA, has an important role in Parkinson disease (PD). GBA mutation carriers have an increased risk for PD, earlier age at onset, faster progression, and various nonmotor symptoms including cognitive decline, REM sleep behavior disorder, hyposmia, and autonomic dysfunction. Furthermore, GCase enzymatic activity is reduced in the peripheral blood and brain of noncarrier, sporadic PD patients. Biallelic GBA mutations, which have been classified as “severe” or “mild,” may cause Gaucher disease (GD), a lysosomal storage disorder. Mild mutations may lead to GD type 1, and 2 severe mutations result in neuronopathic GD (type 2 and type 3). There are 2 GBA variants, p.E326K and p.T369M, which do not cause GD in homozygous carriers, but may modify GCase activity and GD phenotype. It is now clear that p.E326K is a risk factor for PD, but whether p.T369M is associated with PD is still controversial. In some studies, the p.T369M substitution was associated with PD, while in others it had similar or increased frequency in controls. Of interest, it was recently demonstrated that the GBA p.T369M substitution was associated with reduced enzymatic activity in patients with PD and controls compared with that in noncarriers (7.64 vs 11.93 μmol/L/h, p = 0.001). Of interest, it was even lower than the average enzymatic activity of the p.E326K substitution, which was 9.81 μmol/L/h. Because clinical trials on GBA-associated PD are ongoing, and because treatment specifically targeting GBA is likely to be available in the future, it is important to determine whether the GBA p.T369M substitution is associated with PD.

Herein, we performed a meta-analysis of all studies that reported this variant in case-control cohorts. PubMed was searched in May 2016 for all articles that include the words “Parkinson” or “Parkinson’s” and either “GBA” or “glucocerebrosidase,” and 391 articles were identified. These articles were screened to identify all case-control studies that included data on GBA p.T369M (flowchart in figure e-1 at Neurology.org/ng). Initially, 12 independent studies were identified, and data from 2 additional cohorts that were not independently published were retrieved from a previous meta-analysis of the association of all GBA mutations in PD. A total of 4,920 patients with PD and 3,500 controls were analyzed (table e-1, prior to exclusion of outlier studies). The “Metafor” R package was used, with a fixed-effect model and continuity correction of 0.5 for studies that had zero GBA p.T369M carriers in either the PD patient group or controls. The Cochran-Mantel-Haenszel test was used for pooling the allele frequencies and calculating the odds ratios (ORs), and the Tarone test was used to examine heterogeneity.

In the meta-analysis, the GBA p.T369M allele was associated with an increased risk for PD (OR = 1.78, 95% confidence interval [CI] = 1.25–2.53, p = 0.002, figure 1A); however, the heterogeneity was statistically significant (Tarone p = 0.01). The heterogeneity was mainly introduced by 2 cohorts with the highest and lowest extreme values; one had 10 GBA p.T369M carriers in the controls (allele frequency of 0.021) and none in patients, and the second had 17 carriers among the patients with PD (allele frequency of 0.018) but none in controls (additional details in table e-1). After removing these outliers, a total of 4,320 patients and 2,925 controls remained, and the heterogeneity was eliminated (Tarone p = 0.28). The p.T369M allele frequency was 0.011 in patients and 0.0066 in controls, with an OR of 1.74 (95% CI = 1.19–2.55, p = 0.0059, figure 1B). The allele frequency of p.T369M among controls in the current meta-analysis (0.0066) is similar to the allele frequencies in the 1000 Genomes Project (0.005) and Exome Aggregation Consortium (0.0065). Although fixed-effect model is probably a better estimate of the risk in this case, random-effect model was also performed, demonstrating only a trend (OR = 1.55, 95% CI = 0.96–2.50, p = 0.07 for all 14 studies, and OR = 1.54, 95% CI = 0.99–2.40, p = 0.056 after removing the 2 outliers).

These results suggest that the GBA p.T369M substitution may be a risk factor for PD, with a minor effect, similar to that of common genetic variants identified in genome-wide association studies. Therefore, the risk for PD among GBA p.T369M substitution carriers is only slightly elevated as compared to the general population, and the vast majority of carriers will not develop PD. Furthermore, additional studies that include the GBA p.T369M substitution are necessary to reach a better understanding of the clinical relevance of this substitution.
estimation of its risk. This observation may have some important implications, especially once GBA-specific treatment will become available. Based on the current results and the reported reduced GCase activity in carriers of GBA p.T369M, patients with PD with this substitution may be considered for clinical trials of GBA-associated treatments.

The observation that variants not known to cause GD are associated with PD may imply that the mechanisms underlying the 2 conditions are different. However, it is also possible that the same mechanism is involved, yet the effect of this variant is too mild to cause GD, but together with other genetic and/or environmental factors, it enhances the risk for PD. Further studies aimed at understanding the mechanisms underlying GBA-associated PD may elucidate the role of this alteration in PD pathogenesis.


Author contributions: Ms. Mallett and Mr. Ross: study concept and design; acquisition of data; analysis and interpretation; and critical revision of the manuscript for important intellectual content. Dr. Alcalay: analysis and interpretation and critical revision of the manuscript for important intellectual content. Ms. Ambalavanan: acquisition of data; analysis and interpretation; and critical revision of the manuscript for important intellectual content. Dr. Saidnasky: acquisition of data and critical revision of the manuscript for important intellectual content. Dr. Dion: critical revision of the manuscript for important intellectual content. Dr. Rouleau: critical revision of the manuscript for important intellectual content and study supervision. Dr. Gan-Or: study concept and design; acquisition of data; analysis and interpretation; critical revision of the manuscript for important intellectual content; and study supervision.

Study funding: No targeted funding reported. G.A.R. holds a Canada Research Chair in Genetics of the Nervous System and the Wilder Penfield Chair in Neuroscience. Z.G.-O. is supported by a postdoctoral fellowship from the Canadian Institutes for Health Research (CIHR). Disclosure: Ms. Mallett reports no disclosures. Mr. Ross has received research support from the Canadian Institutes for Health Research (CIHR). Dr. Alcalay has received consultancy fees, speaker honoraria, and travel fees from Sanofi-Genzyme; and has received research support from NIH, the Broad Foundation, the Michael J. Fox Foundation, the Parkinson’s Disease Foundation, and the Smart Foundation. Ms. Ambalavanan reports no disclosures. Dr. Saidnasky has served on the medical advisory board of the National Gaucher Foundation; has served on the editorial board of Molecular Genetics and Metabolism; and has received research support from NIH, NHGRI, and Merck under a Collaborative Research Agreement with NHGRI. Dr. Dion reports no disclosures. Dr. Rouleau has received research support from the CHIR Foundation Scheme, the ALS Society of Canada, and the ALS Association. Dr. Gan-Or has received consultancy fees, speaker honoraria, and travel fees from Sanofi-Genzyme; and has received research support from the Canadian Institutes for Health Research (CIHR), McGill University, and the Michael J. Fox Foundation. Go to Neurology.org/ng for full disclosure forms. The Article Processing Charge was paid by the authors.

This is an open access article distributed under the terms of the Creative Commons Attribution-NonCommercial-NoDerivatives License 4.0 (CC BY-NC-ND), which permits downloading and
GBA p.T369M substitution in Parkinson disease: Polymorphism or association? A meta-analysis
Victoria Mallett, Jay P. Ross, Roy N. Alcalay, et al.
Neurol Genet 2016;2:
DOI 10.1212/NXG.0000000000000104

This information is current as of September 8, 2016
<table>
<thead>
<tr>
<th>Updated Information &amp; Services</th>
<th>including high resolution figures, can be found at: <a href="http://ng.neurology.org/content/2/5/e104.full.html">http://ng.neurology.org/content/2/5/e104.full.html</a></th>
</tr>
</thead>
<tbody>
<tr>
<td>Supplementary Material</td>
<td>Supplementary material can be found at: <a href="http://ng.neurology.org/content/suppl/2016/09/08/2.5.e104.DC1">http://ng.neurology.org/content/suppl/2016/09/08/2.5.e104.DC1</a></td>
</tr>
<tr>
<td>References</td>
<td>This article cites 7 articles, 0 of which you can access for free at: <a href="http://ng.neurology.org/content/2/5/e104.full.html##ref-list-1">http://ng.neurology.org/content/2/5/e104.full.html##ref-list-1</a></td>
</tr>
<tr>
<td>Citations</td>
<td>This article has been cited by 11 HighWire-hosted articles: <a href="http://ng.neurology.org/content/2/5/e104.full.html##otherarticles">http://ng.neurology.org/content/2/5/e104.full.html##otherarticles</a></td>
</tr>
<tr>
<td>Subspecialty Collections</td>
<td>This article, along with others on similar topics, appears in the following collection(s):</td>
</tr>
<tr>
<td></td>
<td>All Genetics <a href="http://ng.neurology.org/cgi/collection/all_genetics">http://ng.neurology.org/cgi/collection/all_genetics</a></td>
</tr>
<tr>
<td></td>
<td>Association studies in genetics <a href="http://ng.neurology.org/cgi/collection/association_studies_in_genetics">http://ng.neurology.org/cgi/collection/association_studies_in_genetics</a></td>
</tr>
<tr>
<td></td>
<td>Parkinson's disease/Parkinsonism <a href="http://ng.neurology.org/cgi/collection/parkinsons_disease_parkinsonism">http://ng.neurology.org/cgi/collection/parkinsons_disease_parkinsonism</a></td>
</tr>
<tr>
<td>Permissions &amp; Licensing</td>
<td>Information about reproducing this article in parts (figures, tables) or in its entirety can be found online at: <a href="http://ng.neurology.org/misc/about.xhtml#permissions">http://ng.neurology.org/misc/about.xhtml#permissions</a></td>
</tr>
<tr>
<td>Reprints</td>
<td>Information about ordering reprints can be found online: <a href="http://ng.neurology.org/misc/addir.xhtml#reprintsus">http://ng.neurology.org/misc/addir.xhtml#reprintsus</a></td>
</tr>
</tbody>
</table>

*Neurol Genet* is an official journal of the American Academy of Neurology. Published since April 2015, it is an open-access, online-only, continuous publication journal. Copyright © 2016 American Academy of Neurology. All rights reserved. Online ISSN: 2376-7839.