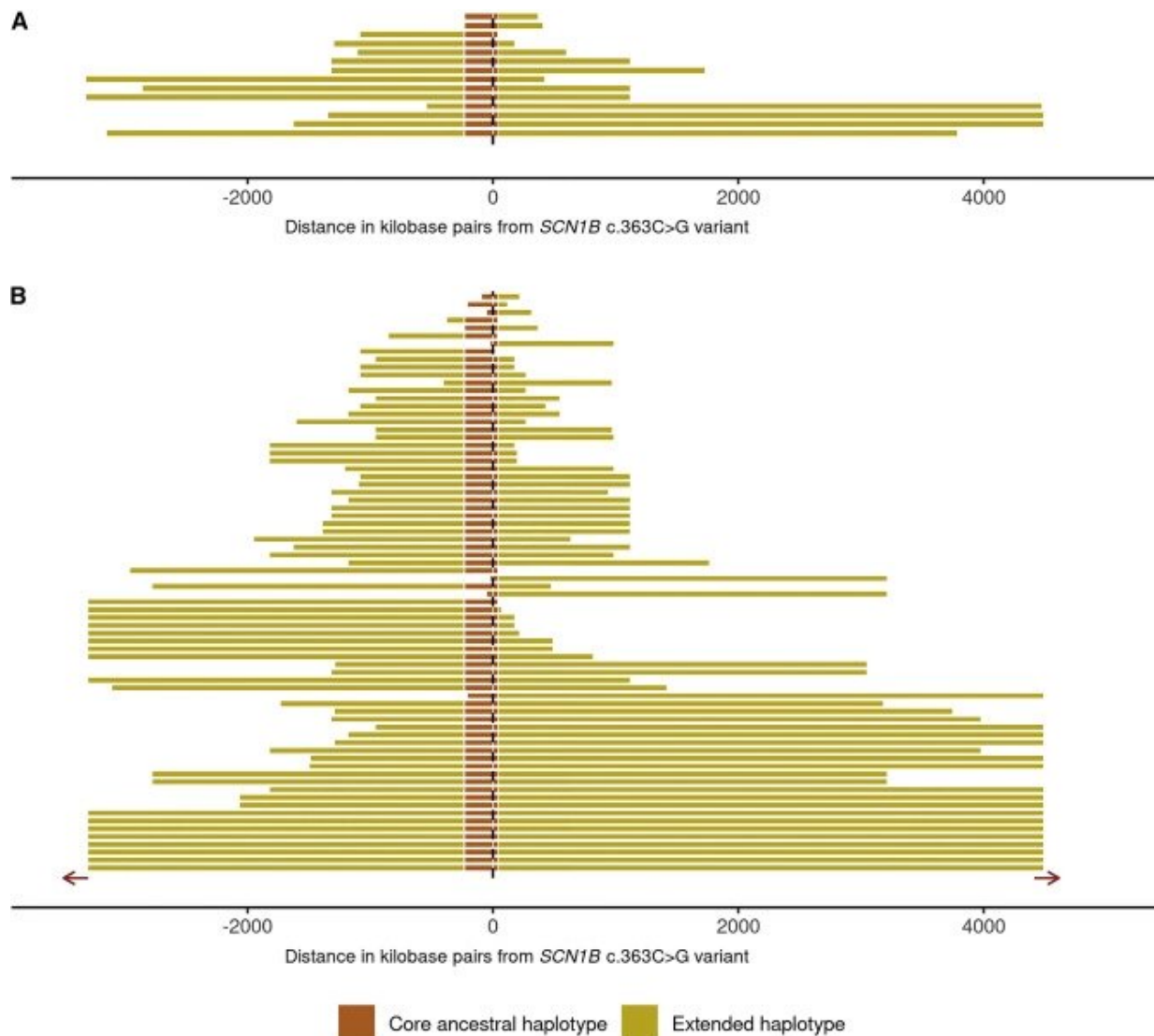


One type of epilepsy traced to a mutation in a single person 800 years ago

October 26 2022, by Bob Yirka



Representation of shared haplotype regions around the *SCN1B* locus. (A) Shared haplotype regions identified in the 14 epilepsy-affected families. Family

identifiers from top to bottom are M, D, C, I, L, E, N, A, K, J, B, H, G, F. Location of the SCN1B c.363C>G variant is shown by the dotted line. Dark orange represents the 0.5 cM core ancestral region shared by all 14 families. Yellow represents the regions shared by at least two epilepsy-affected families. (B) Haplotype regions in 74 UK Biobank samples carrying the SCN1B c.363C>G variant that are shared with the 14 epilepsy-affected families. Red arrows indicate that some individuals may share longer haplotypes than displayed in the plots. Credit: *The American Journal of Human Genetics* (2022). DOI: 10.1016/j.ajhg.2022.10.004

A team of researchers affiliated with several institutions in Australia and the U.K. has found evidence that suggests one type of epilepsy people carry today can be traced back to a mutation that occurred in a single person approximately 800 years ago. In their paper published in *The American Journal of Human Genetics*, the group describes finding the genetic variant responsible for the disease in the U.K. Biobank.

Epilepsy is not just one [disease](#), it is a group of diseases that have shared symptoms. The most common symptom is the periodic onset of a burst of abnormal brain activity, which presents as seizures. One type of the disease is called febrile [epilepsy](#)—it is notable because the seizures it causes are accompanied by a fever. Prior research has shown that it can be traced to the SCN1B c.363C>G [variant](#). It is less severe than other types of epilepsy; some people with the variant do not even know they have it.

In this new effort, the researchers sought to learn if the variant responsible for febrile epilepsy had arisen in multiple people over time or if it could be traced back to just one person. To find out, they traced the lineage of 14 people with the variant. Finding nothing that suggested multiple people had originally had the variant, they expanded their search to include data from the U.K. Biobank and found 74 people with

the variant. A closer look showed that all of them had the same sort of patterns in their variants, which together made them a haplotype.

Because it is unlikely that all of the people they found with the variant had the same haplotype without having a shared ancestry, the variant probably arose in a single person and was passed down over several generations. A closer look at the variant suggests it arose in a person who lived approximately 800 years ago, probably in Britain.

The researchers note that it is quite rare for such a variant to survive for so long in a population. But they also note that because its symptoms are not severe enough to prevent people with the variant from procreating, there is no mechanism for weeding it out.

More information: Bronwyn E. Grinton et al, A founder event causing a dominant childhood epilepsy survives 800 years through weak selective pressure, *The American Journal of Human Genetics* (2022). [DOI: 10.1016/j.ajhg.2022.10.004](https://doi.org/10.1016/j.ajhg.2022.10.004)

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Citation: One type of epilepsy traced to a mutation in a single person 800 years ago (2022, October 26) retrieved 30 December 2022 from <https://medicalxpress.com/news/2022-10-epilepsy-mutation-person-years.html>

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