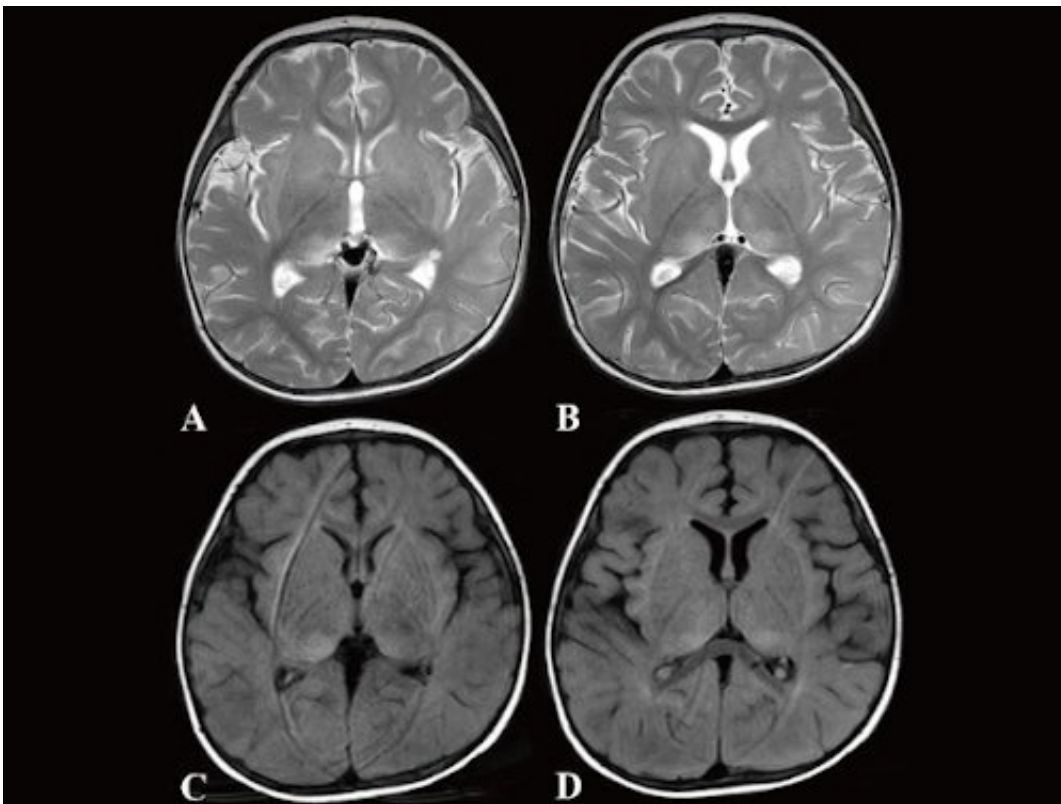


Medical oddity ushers in discovery of unheard-of 'immunity gene' mutations—and a better way to screen them

December 21 2020



Brain MRI of the 10-month old boy with anti-NMDAR encephalitis obtained on day 4. The axial T2 weighted image (T2WI) in A and B and the axial fluid attenuation inversion recovery (FLAIR) image in C and D showed high intensity regions in the bilateral thalamus. New “immunity gene” mutations that researchers discovered to be deleterious were found in the infant. Credit: Satoshi Okada

An infant's odd case of rare encephalitis not typically found in children led researchers to discover never before seen genetic mutations and a more accurate technique to examine the IRAK4 gene responsible for innate immunity.

The IRAK4 gene instructs the production of a protein that plays a key role in the early recognition and response to invading pathogens. Inherited [mutations](#) in this gene can cause an immune system disorder that leaves the body susceptible to recurrent infections of pus-forming pyogenic bacteria. Cases of IRAK4 deficiency are quite rare and only about 10 familial instances were identified so far in Japan.

A Hiroshima University-led research team detected two novel IRAK4 mutations, c.29_30delAT (p.Y10Cfs*9) and c.35G>C (p.R12P), in a 10-month-old boy with anti-N-methyl-d-aspartate receptor (anti-NMDAR) encephalitis and human herpesvirus 6 (HHV6) reactivation. The p.Y10Cfs*9 and p.R12P variations were inherited from his father and mother, respectively. The research team found the mutations by using whole exome sequencing (WES), a method used to exhaustively examine the DNA for genetic disorders.

Anti-NMDAR encephalitis, an [inflammation of the brain](#), happens when the immune system attacks a vital brain receptor, causing psychiatric symptoms, involuntary movement, seizures, autonomic dysfunction, and central hypoventilation. It typically occurs in adult females suffering from ovarian tumor. Only a few cases of anti-NMDAR encephalitis have been so far reported in infants.

The researchers suspect that reactivation of HHV6 may have induced [brain damage](#) that triggered the production of anti-NMDAR antibodies.

"It is well known that anti-NMDAR encephalitis is triggered by HSV1 (herpes simplex virus) infection. The post-infectious autoimmune

process that follows the HSV-induced brain damage is thought to be the cause of anti-NMDAR encephalitis," the researchers said in their study published in the *Journal of Clinical Immunology*.

"The coexistence of anti-NMDAR encephalitis and HHV6 reactivation in this patient may reveal an unknown manifestation associated with IRAK4 deficiency," they said.

More precise method to analyze IRAK4 mutations

Although the researchers have yet to find direct evidence demonstrating a relationship between IRAK4 deficiency and the development of anti-NMDAR encephalitis or HHV6 reactivation, they were able to establish that mutations p.Y10Cfs*9 and p.R12P are damaging.

The researchers developed a novel NF- κ B reporter assay that can precisely evaluate whether an IRAK4 mutation is harmful. The reporter assay allows them to investigate the activation of transcription factor NF- κ B which regulates the body's inflammatory responses.

"Using this system, we confirmed that both novel mutations are deleterious. The current case revealed the possibility that genetics can contribute to characterizing infantile cases with anti-NMDAR [encephalitis](#)," study author Satoshi Okada, a professor at HU's Graduate School of Biomedical and Health Sciences, said.

They resolved potential issues of misevaluation in a previously developed NF- κ B reporter assay by using CRISPR gene-editing technology to remove the IRAK4 naturally expressed in the HEK293 cell—commonly used in studying gene function as it is much easier to transfect than other cell lines.

"HEK293 cells have endogenous IRAK4 expression. Their endogenous

IRAK4 prevents the precise evaluation of the activity of introduced WT or mutant IRAK4," Okada explained.

To date, there are so far 24 known mutations identified in patients with IRAK4 deficiency.

The research team hopes to soon introduce their novel NF-κB reporter assay system to help evaluate IRAK4 gene mutations and achieve early diagnosis.

More information: Shiho Nishimura et al. IRAK4 Deficiency Presenting with Anti-NMDAR Encephalitis and HHV6 Reactivation, *Journal of Clinical Immunology* (2020). [DOI: 10.1007/s10875-020-00885-5](https://doi.org/10.1007/s10875-020-00885-5)

Provided by Hiroshima University

Citation: Medical oddity ushers in discovery of unheard-of 'immunity gene' mutations—and a better way to screen them (2020, December 21) retrieved 19 April 2024 from <https://medicalxpress.com/news/2020-12-medical-oddy-ushers-discovery-unheard-of.html>

This document is subject to copyright. Apart from any fair dealing for the purpose of private study or research, no part may be reproduced without the written permission. The content is provided for information purposes only.