

# Flashing lights and epileptic spasms: should we be routinely performing intermittent photic stimulation in infants?

Marvin H. Braun<sup>1</sup>, Naureen Jooma<sup>2</sup>, Morris H. Scantlebury<sup>3</sup>

<sup>1</sup> Robert Haslam Fellow in Pediatric Neurology, Department of Pediatric Neurology, Alberta Children's Hospital, Calgary, AB, Canada

<sup>2</sup> Clinical Neurophysiology, Alberta Children's Hospital, Calgary, AB, Canada

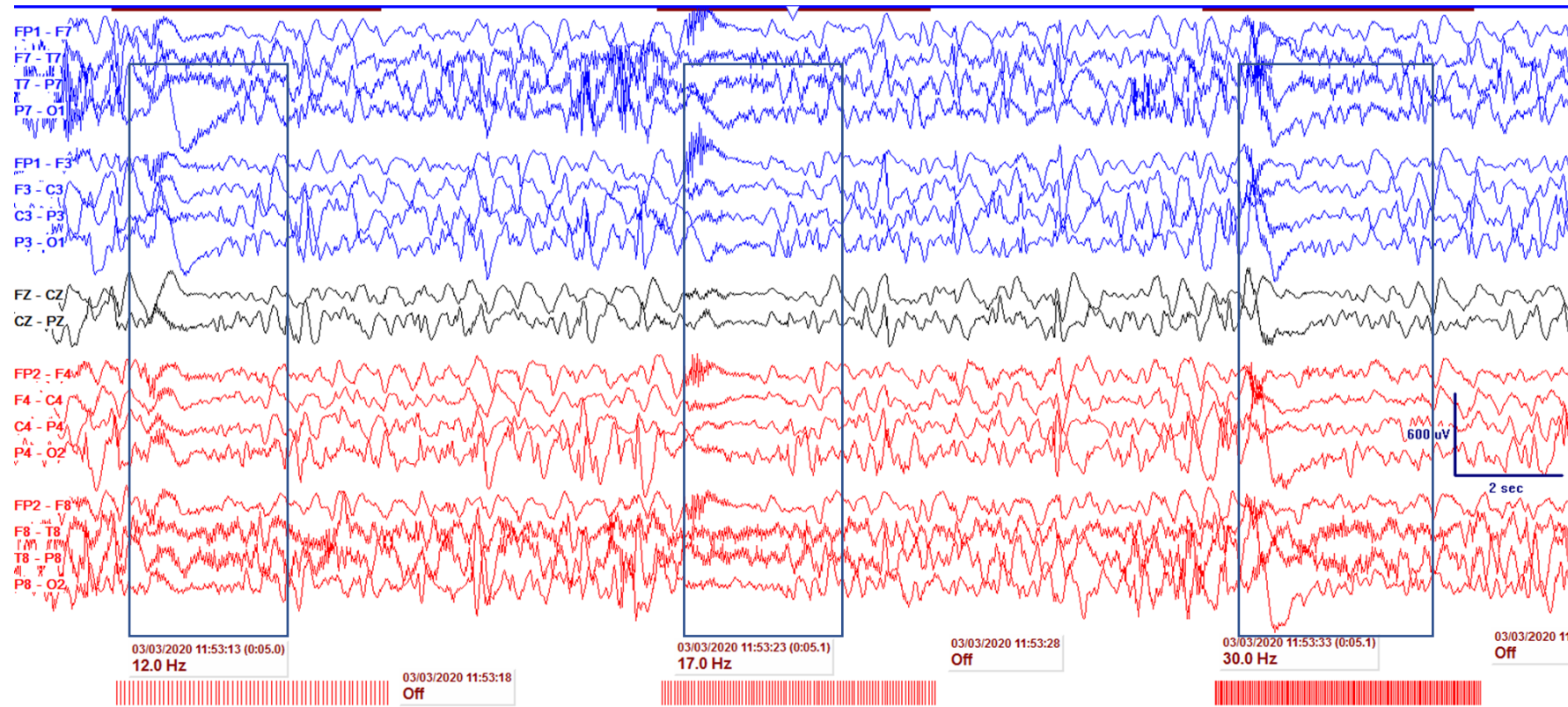
<sup>3</sup> Departments of Pediatrics, Clinical Neuroscience. Member, Alberta Children's Hospital Research Institute (ACHRI) & Hotchkiss Brain Institute (HBI), Alberta Children's Hospital, Calgary, AB, Canada

# Introduction

- Photosensitivity is an abnormal cortical response to light stimulation and is tested for on routine EEGs with intermittent photic stimulation.
- It can result in abnormal cortical responses alone (photoparoxysmal responses) or clinical seizures (photoconvulsive response).
- Most commonly associated with generalized epilepsy.
- Not routinely tested for in young (<1 year old) patients.

# Case Report

- Patient with **ALG13** mutation (c.320A>G [p.Asn107Ser]), an X-linked dominant epileptic encephalopathy gene which results in a severe development and epileptic encephalopathy.
- Developed infantile spasms (IS) at 4 months of age and began treatment at 6 months.
- Had recurrence of IS at 13 months of age and on her subsequent 5 EEGs had reproducible photoconvulsive (with IPS while awake) or photoparoxysmal (with IPS while asleep) responses.
- With escalation of therapy, the strength of the IPS declined and then disappeared.



Representative EEG of intermittent photic stimulation triggering electroclinical epileptic spasms, showing spasm onset with diffuse high-amplitude slow waves followed by diffuse attenuation and low-amplitude fast activity. Boxes demarcate the spasm. Spasms were triggered by 12, 17 and 30-Hz stimulation.

# Discussion

- Photic-induced myoclonic seizures have been reported in trisomy 13 and benign myoclonic seizures of infancy, but spasms induced by IPS have never been reported.
- The findings may be an isolated response of this single patient or a phenotypic feature of this *ALG13* mutation.
- However, since it has never been consistently looked for, it may be a more general response in populations of patients with infantile spasms.
- Therefore, we recommend IPS as part of routine EEG for patients with IS, regardless of age.