

---

---

## Transient Smartphone “Blindness”

**TO THE EDITOR:** Transient monocular vision loss is a common clinical presentation, and the cause is not always thromboembolic.<sup>1</sup> We present two cases in which careful history taking established a benign cause (for the case histories, see the Supplementary Appendix, available with the full text of this letter at NEJM.org).

A 22-year-old woman presented with a several months' history of recurrent impaired vision in the right eye that occurred at night. The results of ophthalmic and cardiovascular examinations were normal. Vitamin A levels and the results of magnetic resonance angiography, echocardiography, and a thrombophilia screening were also normal.

The second case involved a 40-year-old woman

who presented with a 6-month history of recurrent monocular visual impairment on waking, lasting up to 15 minutes. The results of investigations for a vascular cause were again normal. Aspirin therapy had been commenced.

When the patients were seen in our neuro-ophthalmic clinic, detailed history taking revealed that symptoms occurred only after several minutes of viewing a smartphone screen, in the dark, while lying in bed (before going to sleep in the first case and after waking in the second). Both patients were asked to experiment and record their symptoms. They reported that the symptoms were always in the eye contralateral to the side on which the patient was lying.

We hypothesized that the symptoms were due

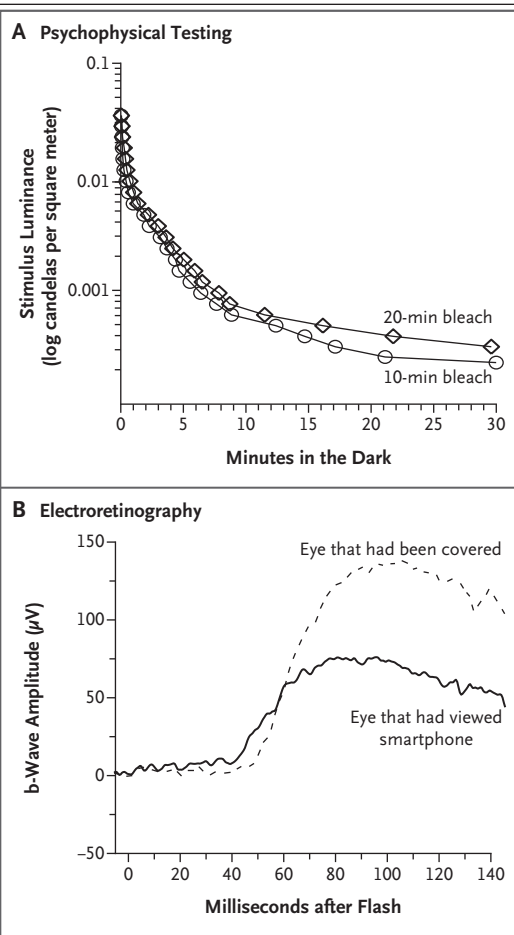
**Figure 1. Diminished Retinal Sensitivity after Smartphone Viewing.**

In Panel A, the points plot visual threshold as a function of time after 10 or 20 minutes of smartphone viewing. The y axis plots the minimum intensity of light that the participant was able to see in the dark. Initially, the participant required a higher-intensity stimulus, indicating low sensitivity; after approximately 20 minutes, the participant was able to see stimuli 100 times dimmer. In Panel B, the two traces show averaged electroretinographic responses to a dim flash of light that was delivered within a few minutes after 20 minutes of monocular smartphone viewing. The response amplitudes are very different, indicating that the eye that had viewed the smartphone had much lower retinal sensitivity than the eye that had been covered (this interocular difference is what the patients perceived as transient monocular blindness). After approximately 20 minutes, responses from both eyes were very similar (see the Supplementary Appendix).

to differential bleaching of photopigment, with the viewing eye becoming light-adapted while the eye blocked by the pillow was becoming dark-adapted. Subsequently, with both eyes uncovered in the dark, the light-adapted eye was perceived to be “blind.” The discrepancy lasted several minutes, reflecting the time course of scotopic recovery after a bleach.<sup>2-4</sup>

In a study approved by a research ethics committee, two of the authors monocularly viewed a smartphone screen at arm’s length and quantified the time course of recovery of sensitivity in the dark both psychophysically and electrophysiologically (Fig. 1). Visual sensitivity was appreciably reduced after smartphone viewing, taking several minutes to recover, and this reduction in sensitivity was measurable at the level of the retina (Fig. 1B).

Although most people view screens binocularly, people frequently use smartphones while lying down, when one eye can be inadvertently covered. Smartphones are now used nearly around the clock, and manufacturers are producing screens with increased brightness to offset background ambient luminance and thereby allow easy reading. Hence, presentations such as we describe are likely to become more frequent. Our cases show that detailed history taking and an understanding of retinal physiology can reassure both patient and doctor and can avoid unnecessary anxiety and costly investigations.



Ali Alim-Marvasti, M.R.C.P.

National Hospital for Neurology and Neurosurgery  
London, United Kingdom

Wei Bi, Ph.D.

City University London  
London, United Kingdom

Omar A. Mahroo, Ph.D.

King's College London  
London, United Kingdom

John L. Barbur, Ph.D.

City University London  
London, United Kingdom

Gordon T. Plant, M.D.

Moorfields Eye Hospital  
London, United Kingdom  
plant@globalnet.co.uk

Supported by a Fight for Sight (United Kingdom) grant (to Dr. Mahroo).

Disclosure forms provided by the authors are available with the full text of this letter at NEJM.org.

1. Petzold A, Islam N, Hu HH, Plant GT. Embolic and non-embolic transient monocular visual field loss: a clinicopathologic review. *Surv Ophthalmol* 2013;58:42-62.
  2. Hecht S, Haig C, Chase AM. The influence of light adaptation on subsequent dark adaptation of the eye. *J Gen Physiol* 1937; 20:831-50.
  3. Lamb TD, Pugh EN Jr. Dark adaptation and the retinoid cycle of vision. *Prog Retin Eye Res* 2004;23:307-80.
  4. Cameron AM, Mahroo OA, Lamb TD. Dark adaptation of human rod bipolar cells measured from the b-wave of the scotopic electroretinogram. *J Physiol* 2006;575:507-26. DOI: 10.1056/NEJMc1514294
- 
-