Dear Editor

An interesting study recently reported by Johnston et al1 investigated the interblink interval between dry and normal eye subjects. The authors concluded that this interval was reduced in dry eyes in comparison with normal eyes, that there was a greater variability in interblink intervals in normal eyes, and that the parameters were useful for diagnosis of dry eye disease.1

Using a method that measured the interblink interval achieved exactly what the authors expected, and enabled better discrimination between normal and dry eye subjects. As the authors state, monitor usage may have impaired their data slightly, and hence a comparative study of the same cohort but using a fixed visual stare would have been both interesting and relevant. The variability seen, as the authors discuss, could be related to an integral homeostatic mechanism. Taylor et al, who investigated and published an interesting paper relating to dopamine’s role in blink rate, suggested the ventromedial part of the caudate nucleus in particular is critically involved.2

We thank Johnston et al for reporting this enlightening study. Should it be repeated or augmented, a comparative study of fixed visual stare to monitor usage in the same participants and also comments on the relationship with dopamine levels would be interesting.

Disclosure

The authors report no conflicts of interest in this correspondence.

References

Author’s response

Patrick R Johnston¹
John Rodriguez¹
Keith J Lane¹
George Ousler¹
Mark B Abelson¹,²
¹Ora Inc, Andover, MA, USA; ²Schepens Eye Research Institute and Harvard Medical School, Boston, MA, USA

Correspondence: Patrick R Johnston
Ora Inc, 300 Brickstone Sq, 3rd Floor, Andover, MA 01810, USA
Tel +1 978 685 8900
Fax +1 978 689 0020
Email pjohnston@oraclinical.com

We thank Lemon and Shah for their insightful comments. The finding that dopamine mediates spontaneous blink through the caudal nucleus provides the mechanism behind altered blink states in diseases such as Parkinson’s disease and schizophrenia. Fatigue has also been linked to impaired dopaminergic transmission,¹,² and it is fascinating to speculate that the changes observed in blinking could be in part centrally mediated. A local role of dopamine might also be proposed. The finding of dopaminergic nerves adjacent to goblet cells suggests a role in the regulation of these mucin-secreting cells which are critical to a stable tear film.³ Dopamine receptors have also been isolated from bovine cornea.⁴

Our field of study is at a crossroads between local, ocular, and central stimuli, as we challenge subjects both by desiccation of the ocular surface and by visual task to assess effects on blinking. The suggestion by Lemon and Shah to assess subjects under fixed-stare conditions is well taken. In fact, we intend to study alterations in blink patterns under various conditions of task and nontask, trying to tease out changes due to ocular dryness and those due to discomfort. It is quite possible that sympathetic dopaminergic stimulation from the stressed cornea and conjunctiva results in blinking, demonstrating its fundamental role in both central and local control of this critical function.

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References