

Minireview Article

Visual Impacts of the Digital Environment

Abstract

Digital devices, mainly smartphone with its time-consuming applications and 24/7 internet connection, are accused of being the main cause of severe distraction and excessive disruptions with fragmentation of everyday life that adversely interrupts the adults and youth lifestyles, health, education and social or family relationships. The medical students underwent objective ophthalmic examination to confirm or exclude CVS diagnosis based on Iqbal's four major criteria for accurate CVS diagnosis. Strict reduction or cessation of exposure to blue light for 4 weeks results in spontaneous resolution of cone adaptation/saturation thus eventually the retina regains its normal foveal functions and responses with normal visual acuities and performances.

Keywords:

Light emitting diodes; Blue light; Computer vision syndrome; Digital eye strain; Multifocal electroretinogram; Screen-induced foveal dysfunction; Retinal phototoxicity; Retina; Digital environment, digital screens and electronic devices; Iqbal's criteria and instructions.

Introduction

The innovative digital technology has occupied an enormous time of the individuals' daily activities and dramatically affected the modern lifestyle [1-5]. The digital environment means watching and interacting with several types digital devices for extended periods through the

entire day [1, 2]. The average screen-time in the digital environments reaches up to eight hours daily for the American adult interacting with ≥ 10 different types of digital screens [6, 7]. Within the digital environments, the routine exposure to various digital devices and electronic screens for several daily screen-hours has elicited various visual, ocular surface and extraocular symptoms and complaints known as the computer vision syndrome (CVS) or digital eye strain (DES) [1, 3-5]. The American Optometric Association (AOA) defined the computer vision syndrome (CVS) as follows “Computer vision syndrome, also referred to as digital eye strain, describes a group of eye- and vision-related problems that result from prolonged computer, tablet, e-reader and cell phone use” [8]. Meanwhile, the Tear Film & Ocular Surface Society (TFOS) considered DES as a more appropriate and specific term than CVS and further redefined DES as “the development or exacerbation of recurrent ocular symptoms and/or signs related specifically to digital device screen viewing” [1]. CVS is a multifactorial syndrome that affects more than one human system and its sequelae extend beyond the eye [3-5].

The CVS main visual symptoms are visual blur, eye strain/fatigue, seeing unclear objects post-screen use, glare/seeing halos of light around objects, feeling diminution of vision, double vision/diplopia, difficulty in refocusing the eyes, near vision discomfort/difficulty and increased sensitivity to light [1, 3-5, 8-16]. The CVS main ocular surface symptoms are dry eye, eye redness, itching/eye rubbing, watery eye, eye irritation/discomfort, foreign body sensation, burning sensation, heavy eyelids and frequent blinking [1, 3-5, 8-16]. The CVS main extraocular symptoms are headache, neck/shoulder/back pain, joint pain in fingers and wrists, inability to hold objects well, difficulty to write using a pen, sleep disturbances/insomnia and inattention [3-5, 8-16]. Other serious manifestations; mainly behavioral and mental health issues, such as depression, stress, anxiety, tendency to suicide and midnight hunger with weight gain have also

been linked to CVS sequelae [3-5, 16-20]. However, both CVS visual and ocular surface symptoms could be attributed to accommodation disturbances, dry eye disease (DED), binocular vision dysfunction and contact lens wearing [1, 3, 4].

Digital devices, mainly smartphone with its time-consuming applications and 24/7 internet connection, are accused of being the main cause of severe distraction and excessive disruptions with fragmentation of everyday life that adversely interrupts the adults and youth lifestyles, health, education and social or family relationships [2, 3, 21]. Therefore, such subjects may encounter serious troubles in their lives such as low productivity, poor creativity and weak academic performance [2, 21]. However, it seems that the problem is not in the digital device or the smartphone itself but the way people handle it and misuse it. In other words, digital devices are not responsible for exacerbation of CVS but the way of its usage is the actual problem [2-5]. The main risk factors and incorrect practices of the individuals' screen-styles are improper or too close eye-screen distance, screen edge at/above horizontal eye level, improper gaze angle (e.g. when lying down or in beds), improper or poor lighting conditions, screen-glare, poor screen-resolution or design, uncomfortable seating postures, watching screen in the dark, small screen-size, excessive screen brightness, small-font size, texting with both thumbs, prolonged screen-hours (average daily screen-hours exceeds five hours) and associated uncorrected refractive errors [3-5]. These risk factors constitute the digital screen or smartphone misuse or abuse practices that are responsible for development, exacerbation and aggravation of CVS [3-5].

Methodology:

In our three published studies by Iqbal et al. [3-5], all medical students responded to the subjective valid and reliable computer vision syndrome form-3 (CVS-F3) questionnaire (.742

Cronbach's alpha reliability coefficient, .773 Guttman Split-Half Coefficient and 82% construct validity rate with the Pearson's correlation validity coefficient) that was designed to be ideal for University students [3-5]. The medical students underwent objective ophthalmic examination to confirm or exclude CVS diagnosis based on Iqbal's four major criteria for accurate CVS diagnosis [3-5, 9-13]. The complete ophthalmic examination included both uncorrected and corrected distance visual acuities (UDVA and CDVA; respectively) measurements, testing pupillary reflexes, DED tests, intraocular pressure measurement, subjective and cycloplegic refraction measurements, slit-lamp and dilated fundus examinations [3-5]. The exclusion criteria were amblyopia, strabismus, accommodation-convergence imbalance, near vision abnormalities, anisometropia greater than 2 diopters (D), myopia >6 D, hyperopia >4 D, astigmatism >4 D, eye or retinal pathology, current eye or systemic diseases and previous eye or systemic surgeries [3-5].

Furthermore, the medical students underwent multifocal electroretinogram (mfERG) examination. We used the mfERG device (RETIscan; Roland Instruments, Wiesbaden, Germany) in accordance with the standard protocol for mfERG of the International Society for Clinical Electrophysiology of Vision (ISCEV). The mfERG stimulus used in our studies was 61 hexagons in dilated subjects with system age-matched norms. The protocol adhered to ISCEV standards and our cut-off values were the normal ranges provided by the ISCEV standard protocol. Eventually, we documented the first foveal peak and amplitude density (P1 AD) in all the mfERG Rings and Quadrants.

Outcomes: retinal impacts of the digital environment

We have exhibited that the digital environment including the digital devices and electronic screens that contains light emitting diodes (LEDs) that emits blue light, affects the macular integrity as we have already documented the existence of the screen-induced foveal dysfunction (SFD) in our three published studies [3-5]. We are the first ophthalmic team that investigated the mfERG foveal changes elicited by the exposure to blue light emitting-screens. These mfERG changes exhibited the reduction in foveal responses representing the foveal dysfunction that was associated with corresponding reduction in visual performances and acuities. The SFD was recorded in the university students diagnosed as positive CVS-cases who watching digital devices for prolonged screen-hours (>5 average screen-hours) with extensive exposure to various types of blue light emitting-screens such as laptops, smartphone, pads/tabs and/or desktop devices [3-5]. Interestingly, most of these positive CVS-cases were medical students who were involved in the University mandated computer system use program.

Our studies included two groups; the control and the CVS groups. The control groups involved medical students that had no-CVS diagnosis, spending less than three daily screen-hours on average, exhibited normal mfERG findings that revealed within normal preserved foveal peak and mfERG Quadrants and Rings were within ISCEV standard protocol normal ranges. On the other hand, CVS groups included medical students that had positive CVS diagnosis that was based on Iqbal's four major criteria for accurate CVS diagnosis [3-5, 9-12], spending more than five daily screen-hours on average, exhibited abnormal mfERG findings with statistically significant foveal amplitude reduction in P1 AD in most of the mfERG Quadrants and Rings below ISCEV standard protocol normal ranges [3-5]. In comparison with the control groups, the CVS groups exhibited a statistically significant foveal amplitude reduction in the uncorrected and the corrected distance visual acuities (UDVA and CDVA; respectively) [3-5].

Furthermore, we discovered that the SFD is a potential reversible phenomenon [5]. We recorded both the mfERG changes and associated visual acuities before and 4 weeks following strict reduction of the screen-time to ≤ 1 screen-hour daily in both the control and the CVS groups [5]. Thereafter, the medical students in the CVS group exhibited remarkable statistically significant improvements in mfERG foveal responses near to normal ranges with correlated improvements in both UDVA and CDVA [5]. We also documented a positive correlation between the differences of average daily screen-hours reduction and the differences in mfERG Quadrants and Rings P1 AD [5, 10]. Therefore, the lower the daily screen-hours with less exposure to the blue light emitted from digital screens, the more the improvements in the foveal responses [10].

Based on our outcomes, we have defined the term screen-induced foveal dysfunction (SFD) as **“the multifocal electroretinogram reduced foveal responses below standard normal ranges that are mostly associated with reduced visual acuities and performances in computer vision syndrome positive-cases”** [3-5, 9-12]. Therefore, SFD could be discovered in positive CVS-cases and is mostly associated with blurring of vision, feelings of diminution of vision, visualization of unclear objects especially post-screen use, feeling the diminution of vision, complaining of annoying halos of light around objects with subsequent reduction in visual performances. In addition, we think that the SFD is a potential type of retinal phototoxicity that could be attributed to excessive exposure to blue light emitting LEDs, encountered in the manufacture of modern digital screens and electronic devices, with subsequent photochemical injury [3-5, 9-12]. Furthermore, we have discovered that SFD is a temporary retinal phototoxicity phenomenon that has short-term adverse impacts on normal foveal functions and intact macular integrity. Moreover, SFD might be reversed by restrict reduction of the screen-time thus minimizing the retinal exposure to blue light emitting screens [9-12]. Meanwhile, we

unfortunately don't know underlying pathophysiological mechanisms of SFD; however, it might be caused by the macular cone/bipolar cell dysfunction due to the cone adaptation and/or saturation resulting from the excessive levels of blue light with a potential level of retinal phototoxicity resulting in a photochemical injury inducing SFD. Strict reduction or cessation of exposure to blue light for 4 weeks results in spontaneous resolution of cone adaptation/saturation thus eventually the retina regains its normal foveal functions and responses with normal visual acuities and performances.

Similar to our outcomes, Cougnard-Gregoire et al. [14] concluded that the potential toxicity of long-term cumulative exposure to blue light emitting LEDs and the dose-response effect are currently unknown. In agreement with our results, Li et al [22] reported the mfERG outcomes that ≥ 8 daily hours viewing of the screens reduced the retinal photoreceptor cells amplitude in the parafoveal region of the macula with delayed peak time. They also stated that the long-term exposure to blue light is a cause of structural and functional damage of the retinal tissue [22].

Finally, our studies recommended that the higher educational authorities should re-plan the mandated computer system use program and consider other alternatives. We also recommend that further future studies including mfERG investigations regarding this topic.

Ethics Statements

The three studies by Iqbal et al. gained the approval of the Medical Research Ethics Committee (MREC) in Faculty of Medicine, Sohag University, Egypt. All three studies were registered as clinical trials at the ClinicalTrial.gov (ID: NCT04398212 and NCT04405648) and the Pan African Clinical Trial Registry (PACTR201811618954630). All studies were conducted in

accordance with the tenets of the Declaration of Helsinki. All participants signed an informed consent prior to enrolment in the studies.

Supplementary material and/or additional information

None.

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