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3 **Intra- and inter-model variability of light detection using a**
4 **commercially available light sensor**

5

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16 4.Myopia

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25 **ABSTRACT**

26 Purpose

27 The veracity of claims made by researchers and clinicians when reporting the impact
28 of lighting on vision and other biological mechanisms is, in part, reliant on accurate
29 and valid measurement devices. We aim to quantify the intra- and inter-watch
30 variability of a commercially available light sensor device which has been widely
31 used in vision and other photobiological research.

32

33 Methods

34 Intra- and inter-watch differences were investigated between four Actiwatch
35 Spectrum Pro devices. The devices were used to obtain measurements on two
36 separate occasions, under three different controlled light conditions; the Gretag
37 Macbeth Judge II lightbox was used to produce Simulated Daylight (D65), Illuminant
38 A (A) and Cool White Fluorescent (CWF) lighting.

39

40 Results

41 Significant inter-watch differences were noted when considering tricolour (red, green,
42 blue) and the white sensor outputs under each of the three illuminants ($p < 0.01$). A
43 significant interaction was also found between tricolour sensor and watch used
44 ($p < 0.01$).

45 Intra-watch differences were noted for the tricolour and for the white sensor outputs
46 under the three illuminants (≤ 0.05), for all but one watch which showed no significant
47 intra-watch difference for the white 'sensor output' under the D65 illuminant.

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49 Conclusion

50 Use of spectral sensitivity devices is an evolving field. Before drawing causal
51 relationships between light and other biological processes, researchers should
52 acknowledge the limitations of the instruments used, their validation, and the
53 resultant data. The outcomes of the study indicate caution must be exercised in
54 longitudinal data collection and the mixing of watches amongst study participants
55 should be avoided.

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59 **INTRODUCTION**

60 Wearable accelerometers and light sensors have been widely adopted for clinical
61 and research purposes. [1,2,3,4,5,6] Studies exploring circadian entrainment; sleep
62 quality; and physical activity have been particularly embrasive of such technologies.
63 [7,8,9,10,11,12] More recently, vision researchers have made greater use of light
64 and activity monitors. [13,14,15,16] The appeal of such devices is unsurprising; they
65 offer a seemingly objective, and largely unobtrusive, method of recording data whilst
66 reducing reliance on more subjective recall methods such as questionnaires or
67 interviews.

68
69 There is a compelling link between onset of myopia, shortsightedness, and time
70 spent outdoors, [17] the study of which relies upon accurate and valid monitoring of
71 lighting exposure. Whilst a range of light sensors have been employed for such
72 photobiological studies, models from the Phillips Respironics' Actiwatch (Philips
73 Healthcare, Best, NL) range have proven to be particularly popular for vision related
74 studies within child and adult cohorts.

75
76 While the research interest in various iterations of the Actiwatch has led to numerous
77 validation and evaluative studies relating to the accelerometry aspect,
78 [18,19,20,21,22,23,24,25] there are comparatively fewer studies validating the light
79 detection features. [26,27,28,29,30]

80 Given the paucity of data and the growing interest in light detection research, this
81 study aims to investigate intra- and inter-model variability in light detection of a
82 commercially available device from Philips Respironics: the Actiwatch Spectrum Pro.

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85 **METHODS**

86 Four static Actiwatch Spectrum Pro devices were used to obtain measurements in a
87 controlled lighting environment. The Phillips Actiware software was used to set
88 epoch length to 15 s for ~5 min exposure periods, from which 13 consecutive
89 sampling points (i.e. ~3 min' worth) were extracted for analysis. In general, the data
90 were only extracted for analysis after at least ~1 min of recording to minimise any
91 erratic measurements due to potential sensor adaptation or otherwise.

92

93 The included data were also checked to ensure the 'Activity' outputs were recorded
94 as zero for the period during which light data were extracted i.e. the watch had not
95 moved or fallen during the process of recording.

96

97 To investigate intra- and inter-watch differences, the four Actiwatch Spectrum Pro
98 devices were used to obtain measurements on two separate occasions, under three
99 different controlled light conditions. A Gretag Macbeth Judge II lightbox (Gretag
100 Macbeth, New Windsor, New York, USA) was used to simulate the lighting
101 conditions: Simulated Daylight (D65), Illuminant A, and Cool White Fluorescent
102 (CWF) lighting, which have published colour temperatures of 6500 K, 2856 K,
103 and 4150 K. All watches were affixed such that the watch was statically face-up and
104 in the horizontal plane within the lightbox.

105

106 The Actiwatch Spectrum Pro refers to 'white', 'red', 'green', 'blue' outputs. As white
107 light watch outputs are understood to be generated from integration of all three
108 tricolour sensors, white light data were analysed separately. The manufacturer's

109 literature indicates that the Actiwatch Spectrum Pro measures wavelengths between
110 400 and 700 nm, [31] including capturing wavelength sensitivity with respect to the
111 tricolour sensors blue (400-500 nm), green (500-600 nm), and red (600-700 nm) with
112 band widths of ~100 nm. [32]

113

114 **Statistical analysis**

115 A series of paired t-tests were used to investigate intra-watch differences, i.e.
116 differences between runs 1 and 2 for white, red, green, and blue sensor outputs,
117 following exposure to the three illumination conditions. Bias and limits of agreement
118 were generated for each intra-watch combination.

119

120 A mixed design repeated-measures analysis of variance (ANOVA) was used to
121 determine inter-watch differences. The watch used was considered the between-
122 subject factor, with lighting and the tricolour sensors as the within-subject factors; the
123 analysis was repeated for the white sensor with only lighting as the within subject
124 factor.

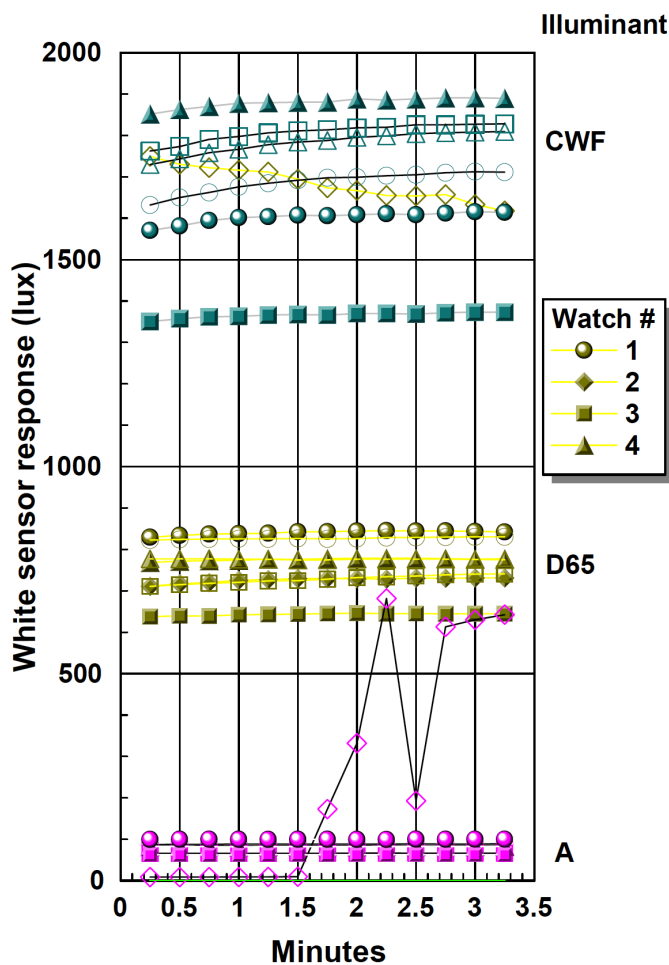
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126 **RESULTS**

127 For watch 2, anomalous measurements e.g. the recording of a zero response, were
128 found under multiple conditions, and anomalous results were recorded for illuminant
129 A (see Fig. 1), thus watch 2 was not included in all analyses.

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 133 **Fig. 1.** White sensor response of the four watches for illuminants CWF (teal), D65 (yellow),
 134 and illuminant A (purple). Watch 1 denoted by circles, watch 2 by diamonds, watch 3 by
 135 squares, and watch 4 by triangles. Filled symbols represent run 1 and empty symbols run 2.
 136

137 ***Intra-watch differences***

138 A series of paired t-tests showed a significant intra-watch differences for the 'white'
 139 light output under each lighting conditions (D65, A, CWF) for all watches ($p < 0.01$)
 140 except watch 4 which only showed a significant intra-watch difference under the
 141 CWF and the illuminant A lighting conditions, but not D65 ($p > 0.05$). Separately,
 142 paired t-tests showed significant intra-watch differences ($p \leq 0.05$) under each of the
 143 illuminant conditions for the tricolour outputs (R,G,B); all differences remained
 144 significant following application of a Bonferroni correction ($0.05/3 = 0.0167$) except
 145 the red sensor of watch 4 under illuminant A ($p = 0.05$).
 146

147 The bias (i.e. the mean difference between run 1 and run 2) and the limits of
 148 agreement were calculated for outputs under each of the lighting conditions; these
 149 are shown in Tables 1 and 2. Large differences in bias were noted, ranging from
 150 -956.89 to 13.61 for the D65 condition; -255.12 to 11.65 for illuminant A; and
 151 -441.60 to 97.37 for illuminant CWF for the white data outputs. Similarly, the range of
 152 bias was also observed between watches for the tricolour outputs.
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	<i>Watch 1</i>	<i>Watch 2</i>	<i>Watch 3</i>	<i>Watch 4</i>
DAYLIGHT				
<i>Bias</i>	13.61	-956.89	-84.86	-0.39
<i>Standard Dev</i>	3.25	45.89	7.75	3.67
<i>Lower LOA</i>	7.25	-1046.82	-100.04	-7.58
<i>Upper LOA</i>	19.97	-866.95	-69.68	6.80
<i>t-test p value</i>	<0.01	<0.01	<0.01	0.71
% difference in means				
	-1.6	56.9	11.7	0.1
ILLUMINANT A				
<i>Bias</i>	11.65	-255.12	-3.83	-1.50
<i>Standard Dev</i>	0.44	285.88	0.346952	0.42
<i>Lower LOA</i>	10.78	-815.45	-4.50618	-2.32
<i>Upper LOA</i>	12.52	305.20	-3.14613	-0.69
<i>t-test p value</i>	<0.01	0.01	<0.01	<0.01
% difference in means				
	13.2	100	5.8	1.8
CWF				
<i>Bias</i>	-84.69	-8.30	-441.60	97.37
<i>Standard Dev</i>	12.72	0.33	14.85	14.51
<i>Lower LOA</i>	-109.63	-8.95	-470.70	68.93
<i>Upper LOA</i>	-59.75	-7.65	-412.50	125.80
<i>t-test p value</i>	<0.01	<0.01	<0.01	<0.01
% difference in means				
	5	100	24.4	-5.5

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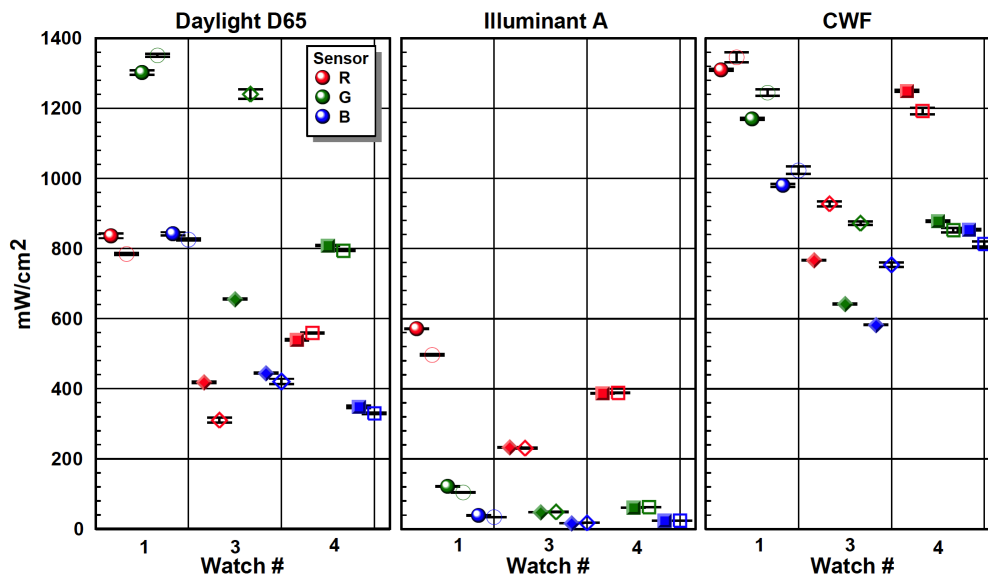
Table 1. Bias (mean difference between runs 1 and 2), standard deviation (SD) of the bias, upper and lower limit of agreement *LOA*, and percentage difference (change from run 1 to 2) is shown for all four watches under each of the lighting conditions for white light outputs only.

	RED				GREEN				BLUE			
DAYLIGHT	Watch 1	Watch 2	Watch 3	Watch 4	Watch 1	Watch 2	Watch 3	Watch 4	Watch 1	Watch 2	Watch 3	Watch 4
Bias	51.54	-777.23	108.38	-19.77	-49.23	-4756.23	-584.77	14.69	16.54	-320.00	23.77	18.62
Standard Dev	4.39	14.78	5.59	3.11	7.60	348.25	14.70	2.10	2.96	11.13	6.73	2.14
Lower LOA	42.93	-806.19	97.43	-25.87	-64.12	-5438.81	-613.58	10.58	10.73	-341.81	10.57	14.42
Upper LOA	60.14	-748.27	119.34	-13.67	-34.34	-4073.65	-555.96	18.80	22.34	-298.19	36.97	22.81
t-test p value	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01
% difference in means	-6.6	46.7	-35.0	3.5	3.6	92.1	47.2	-1.9	-2.0	40.9	-5.7	-5.7
ILLUMINANT A												
Bias	74.69	-246.52	2.46	-0.92	17.23	-253.23	-0.91	-1.40	4.49	-140.52	-1.13	-0.32
Standard Dev	2.39	276.87	1.33	1.50	0.44	292.56	0.23	0.36	0.18	156.01	0.10	0.12
Lower LOA	70.00	-789.18	-0.15	-3.86	16.37	-826.64	-1.36	-2.10	4.13	-446.29	-1.33	-0.55
Upper LOA	79.38	296.13	5.07	2.01	18.09	320.18	-0.46	-0.70	4.85	165.26	-0.93	-0.09
t-test p value	<0.01	0.01	<0.01	0.05	<0.01	0.01	<0.01	<0.01	<0.01	0.01	<0.01	<0.01
% difference in means	-15.1	100	-1.1	0.2	-16.6	100	1.9	2.3	-13.3	100	6.6	1.4
CWF												
Bias	-36.15	-5.94	-160.85	56.92	-75.38	0.00	-230.23	26.15	-43.00	-2.50	-171.62	40.62
Standard Dev	12.61	0.14	6.35	7.51	7.76	0.00	4.97	7.31	7.46	0.10	5.38	6.28
Lower LOA	-60.87	-6.22	-173.29	42.20	-90.60	0.00	-239.97	11.82	-57.62	-2.69	-182.16	28.31
Upper LOA	-11.44	-5.66	-148.40	71.64	-60.17	0.00	-220.49	40.49	-28.38	-2.31	-161.07	52.92
t-test p value	<0.01	<0.01	<0.01	<0.01	<0.01	n/a	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01
% difference in means	2.7	100	17.3	-4.8	6.1	n/a	26.4	-3.1	4.2	100	22.8	-5

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Table 2. Bias (mean difference between runs 1 and 2), standard deviation of the bias, upper and lower LoAs are shown for all four watches under each of the lighting condition Bonferroni correction).

Figure 1 shows the white light sensor outputs for the watches when exposed to the three illuminants, including runs 1 and 2. The watches, except watch 2, show relatively constant recordings during the 3 min. Figure 2 shows the same for the tricolour (R, G, B) outputs with each measurement being the mean recording \pm standard deviation of the time exposure. Visual inspection of the difference vs. mean plots showed that while there was generally no obvious relationship between bias and means for the incandescent lighting conditions (for white or RGB), for the other two lighting conditions there may have been an increase in differences between readings with increasing mean, although this was not always apparent in every case.



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Fig. 2. Response of red (R), green (G) and blue (B) sensors in watches 1, 3, and 4 under daylight D65, illuminant A and CWF illumination. Two runs are shown; run 1 – filled symbols and run 2 – empty symbols. Error bars represent the standard deviation of measurements taken over the course of ~3mins.

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Inter-watch differences

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A mixed design repeated measures ANOVA, using the first run of measurements, where watch (watch 1, 3, 4) was the between-subject factor, and both lighting and the tricolour sensors served as the within-subject factors, showed a significant overall difference between watches ($p < 0.001$) for the R, G, B outputs. Similarly, a significant inter-watch difference was noted for white outputs too ($p \leq 0.001$). As may be expected there was a significant main effect of lighting on outputs ($p < 0.001$) for both the tricolour sensor and, separately, for white sensor responses; and an interaction between lighting and sensor for the tricolour responses ($p < 0.001$) i.e. some sensors reacted more, and in other cases less, to each illuminant.

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Inter-watch differences were highlighted through significant interactions between illuminant and watch used; this was the case for both tricolour and, separately, white responses ($p < 0.001$). Differences were also found between watches for the tricolour sensors ($p < 0.001$). Outcomes for run 2 (with watches 1, 3, 4) showed the same results.

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DISCUSSION

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Our data show multiple intra-watch differences for measurements obtained using the same watches and lighting conditions on different dates; the data also show a significant inter-watch difference i.e. differences between the outputs of individual watches. The interaction between watch used and sensor (R, G, B) further reinforces the need for calibration of watches prior to their use. [27] Despite the differences, other than the watch which appeared to produce largely anomalous and erratic results (watch 2), our data showed the percentage change between runs 1 and 2 was generally small for the white light response (see Tables 1 and 2).

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As part of their studies, Figueiro et al. (2015) [28] tested six Actiwatch Spectrum devices under various light sources. A measurement range of 20% (from lowest to highest) under high pressure sodium lighting to 9% under 3500 K fluorescent lighting was reported, the range for daylight was approximately 12%. For comparison, excluding watch 2, our range of maximum and minimum measurements across all watches (i.e. the maximum and minimum of all readings across watches 1,3, 4) fluctuated 25% from highest to lowest readings under daylight; 38% under illuminant A; and 29% under the CWF illuminant for the white output. Similarly, others have also reported a high degree of variation between different Actiwatch devices. [27]

While our testing protocol did not evaluate the impact of oblique and direct lighting separately, the use of a lightbox ensured uniform light distribution. Previous work has shown the Actiwatch Spectrum sensors to be sensitive to orientation. [28,29] Price et al. (2012) reported mean percentage of cosine response errors (f) as approximately 25.3%; 33.2%; 32.6%; and 48.6% when the plane of incidence was horizontal and approximately 61.1%; 60.9%; 61.0%; and 64.7% when vertical for the white, red, green, and blue outputs respectively. Figueiro et al. (2015) [28] also reported on spatial sensitivity, for the Actiwatch Spectrum; f errors were 30.7%;39.4%; and 57.2% for the red, green, and blue sensors, respectively. The Actiwatch Spectrum sensors are set back from the watch surface by approximately 2 mm (Price et al. 2012); and encased in an external cover; the positioning of the sensors is understood to limit incident light, [28] particularly for the blue sensor [29].

With respect to future work in ophthalmology

The findings suggest that the Actiwatch Spectrum Pro is a useful tool for characterising light, however, caution must be exercised.

Our work provides some indication of the magnitude of error one might expect when collecting data under different lighting conditions.

Repeatability may be lower for some watches, which will affect the validity of any comparisons between data captured at different time points i.e. longitudinal studies.

The presence of inter-watch differences demonstrates a need to use the same watch for the same individual throughout a study i.e. the watches are not interchangeable. A lack of interchangeability also limits the ability to draw comparisons between datasets from different individuals who have used different watches.

CONCLUSION

In summary, our findings are in general agreement with previous work evaluating the Actiwatch and support the need for calibration. Spectral sensitivity devices appear to be part of an evolving field. Before drawing causal relationships between light and other biological processes, researchers should be clear to acknowledge the limitations of their instruments and understand the potential margins of error which may affect their dataset; it is only then that meaningful differences can be distinguished from noisy data.

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