

Cranwell MB, Pearce B, Loveridge C, Hurlbert AC. [Performance on the Farnsworth-Munsell 100-Hue Test is significantly related to non-verbal IQ.](#) *Investigative Ophthalmology and Visual Science* 2015, 56(5), 3171-3178.

Copyright:

This is an Accepted Manuscript of an article published by Association for Research in Vision and Ophthalmology in *Investigative Ophthalmology and Visual Science* in May 2015, available online: <http://dx.doi.org/10.1167/iovs.14-16094>

DOI link to article:

<http://dx.doi.org/10.1167/iovs.14-16094>

Date deposited:

21/07/2015

Embargo release date:

01 November 2015

1 Performance on the Farnsworth-Munsell 100-Hue Test is significantly related to non-verbal IQ

2 Matthew B. Cranwell, Bradley Pearce, Camilla Loveridge, & Anya C. Hurlbert

3 Corresponding Author:

4 Matthew B. Cranwell

5 Institute of Neuroscience

6 Henry Wellcome Building

7 Newcastle University

8 Framlington Place

9 Newcastle Upon Tyne

10 United Kingdom

11 NE2 4HH

12 m.b.cranwell@ncl.ac.uk

13

14

15 Please cite as: Cranwell, M.B., Pearce, B., Loveridge, C. & Hurlbert, A.C. (2015). Performance on the

16 Farnsworth-Munsell 100-Hue Test is significantly related to non-verbal IQ. *Investigative*

17 *Ophthalmological & Visual Science*, 56, 3171-3178. doi: 10.1167/iovs.14-16094

18

19

20

21

22

23

24

25 **Abstract**

26 **Purpose.** The Farnsworth-Munsell 100-Hue test (FM100) is a standardised measure of chromatic
27 discrimination, based on coloured cap-sorting, which has been widely used in both adults and
28 children. Its dependence on seriation ability raises questions as to its universal suitability and
29 accuracy in assessing purely sensory discrimination. This study investigates how general intellectual
30 ability relates to performance on both the FM100 and a new computer-based chromatic
31 discrimination threshold test, across different age groups in both typical and atypical development.

32 **Methods.** Participants were divided into two main age groups, child (6-15 years) and young adult
33 (16-25 years), with each group further subdivided into typically developing (TD; 3 groups; TD 6-7y,
34 TD 8-9y, TD Adult) individuals and atypically developing individuals, all but one carrying a diagnosis
35 of Autism Spectrum Disorders (ASD; 2 groups; ATY child 7-15y, ASD Adult). General intelligence was
36 measured using the Wechsler Abbreviated Intelligence Scale and Wechsler Intelligence Scale for
37 Children. All participants completed the FM100. Both child groups also completed a computer-based
38 chromatic discrimination threshold test, which assessed discrimination along cone-opponent (“red-
39 green”, “blue-yellow”) and luminance cardinal axes using a controlled staircase procedure.

40 **Results.** FM100 performance was better in adults than in children. Furthermore, performance
41 significantly positively correlated with non-verbal IQ for all child groups and the young adult ASD
42 group. The slope of this relationship was steeper for the ASD than TD groups. Performance on the
43 chromatic discrimination threshold test was not significantly related to any IQ measure. Regression
44 models reveal that chromatic discrimination threshold, although a significant predictor of FM100
45 performance when used alone, is a weaker predictor than NVIQ used alone or in combination.

46 **Conclusions.** The results indicate that FM100 performance is not purely a measure of colour
47 discrimination but instead also reflects general non-verbal ability. Other measures of chromatic
48 discrimination ability are therefore required for its accurate assessment, particularly in early or
49 atypical development.

50

51

52

53

54

55 Introduction

56 Colour perception is a major contributing factor to a wide range of different behavioural tasks,
57 including, for example: visual search¹, object recognition² and evaluation of material properties (e.g.
58 ripeness of fruit or healthiness of skin³). The low-level ability to discriminate between colours
59 (chromatic discrimination ability) also underlies development of higher-level abilities and behaviours
60 such as colour naming⁴, affective responses to colour⁵ and colour memory⁶. In recent years, colour
61 perception has been increasingly studied in developmental disorders such as autism spectrum
62 disorder (ASD)⁷⁻⁹, Attention Deficit Hyperactivity Disorder^{10,11}, and Williams Syndrome¹², as the
63 extent of atypical sensory processing across all visual domains has become more evident in these
64 disorders. It is therefore increasingly important to ensure that the diagnostic tools used to assess
65 sensory processing, and in particular colour perception, are both sensitive and specific in isolating
66 atypicalities in sensory processing. This requirement is complicated by the hypothesised relationship
67 between sensory processing and intelligence^{13,14}; according to the original hypothesis of Galton and
68 Spearman¹³⁻¹⁵, higher intelligence is associated with better sensory discrimination abilities. The
69 support for this hypothesis has been mixed¹⁵⁻¹⁹, with low correlations found between general
70 intelligence and some measures of sensory discrimination (including colour perception), and other
71 more recent evidence¹⁹ demonstrating a strong link between IQ and performance on a visual
72 motion discrimination task. Nonetheless, the putative relationship makes it vital to ensure that tests
73 of sensory processing are not confounded by direct contributions of general ability to performance.

74 The Farnsworth-Munsell 100-Hue Test²⁰ has been widely used by clinicians and visual scientists
75 as a measure of chromatic discrimination ability²¹ in both typically developing individuals with
76 normal colour vision²² and in individuals with congenital or acquired colour vision deficits²³⁻²⁶ or
77 developmental disorders^{7,8,10,12}. The FM100 test involves arranging a set of individual coloured caps
78 of similar lightness and saturation in order between the hues of two fixed caps (e.g. blue and green),
79 so that a smooth colour gradient is formed, with the hue differences between neighbouring caps as
80 small as possible. The FM100 has been used in a wide range of ages, from early childhood (5 years)
81 to elderly, and has the advantage of age-expected norms^{22,27}. It is also useful in identifying
82 congenital and acquired retinal diseases and as a measure of lens yellowing during normal aging²⁸.

83 Other laboratory measurements of chromatic discrimination ability^{29,30}, for example using
84 detection tasks for forms defined by chromatic contrast embedded in backgrounds with dynamic
85 random luminance variation, conclude that chromatic discrimination ability changes over the life
86 span with sensitivity peaking in late adolescence/the twenties^{31,32} and subsequently decreasing
87 throughout the remaining adulthood^{22,27,33}. The improvement in FM100 performance with age up to

88 early adulthood, evident in studies that establish age-dependent norms^{22, 27}, is consistent with this
89 age-related enhancement of chromatic discrimination ability. Yet other studies suggest that this
90 change in FM100 performance may be due at least partly to improvements in the ability to perform
91 seriation tasks^{34, 35}. A recent study concludes that the general ability to perform a seriation task has
92 sufficiently developed by the age of five that its further development does not explain the
93 improvement in FM100 with age, but also that there are other non-colour-specific factors related to
94 the difficulty of the discrimination which affect FM100 performance^{36, 37}. It has also been observed
95 that the Roth-28 (a shortened version of the FM100 using every third cap) is a time-consuming test
96 that requires attention and a degree of visuo-motor competence, and is subject to learning and
97 practice effects³⁸, and therefore the question has been raised as to whether other tests of colour
98 discrimination might be more useful or appropriate³⁹. Although a previous study concludes that the
99 specific correlation between the FM100 and general intelligence is low¹⁶, we have found (in a
100 preliminary study to that reported here) that performance on the FM100 is related to non-verbal
101 ability (NVIQ) in adults⁹, and more strongly so in individuals diagnosed with ASD. Taken together,
102 these results argue strongly for dissociating the various factors contributing to performance on the
103 FM100 – including age, attentional ability, general intelligence, and chromatic discrimination ability -
104 before accepting it as an unequivocal measure of chromatic discrimination ability suitable for
105 widespread diagnostic use.

106 Here, the contribution of general cognitive ability to performance on two chromatic
107 discrimination tests (the FM100 and a computer-based form-discrimination threshold task) is
108 assessed. Specifically, we aim to determine the extent to which the relationship between non-verbal
109 general ability and performance on the FM100 varies with age (e.g. between adults and children)
110 and also with development, between typical and atypical development.

111

112

113

114

115

116

117

118 **Methods**

119 a) Participants

120 Ninety-two participants took part in the study, split across five different participant groups
121 on the basis of chronological age (adult/child) and the typicality of their development (typically
122 developing/atypically developing). All adult participants gave informed consent. For the child groups,
123 parental consent was obtained and assent was given by the participant prior to the start of the
124 experiment in line with the Declaration of Helsinki. Ethical approval was given from the Faculty of
125 Medical Sciences Ethics Committee, Newcastle University (Approval Code:00618/2012; for child
126 participants) and from the Newcastle University Psychology Ethics Committee (Approval Code
127 060004; for adult participants) .

128 There were two adult groups: a typically developing group (TD adult; n=28, mean age= 20.07
129 years, male=15, female=13) and an atypically developing group diagnosed with ASD (ASD Adult;
130 n=18, mean age=18.11, male=15, female=3). There were no psychiatric or developmental disorder
131 diagnoses in the TD Adult group. Intelligence was assessed using the Wechsler Abbreviated Scale of
132 Intelligence (WASI)⁴⁰. Verbal IQ (VIQ) was determined by Vocabulary and Similarities subtests, and
133 Performance or Non-Verbal IQ (NVIQ) was measured using Block Design and Matrix Reasoning
134 subtests.

135 Three groups of children were recruited from local mainstream and special needs schools:
136 two typically developing (TD) groups, 12 children aged 6-7 years old (TD 6-7y; male=5, female=7) and
137 19 children aged 8-9 years old (TD 8-9y; male=5, female=14); and one group of 17 children
138 diagnosed with ASD plus one child with Williams Syndrome (ATY child; male=12, female=6). There
139 were no developmental disorders reported for any children in the two TD groups. Four sub-tests of
140 the Wechsler Intelligence Scale for Children (WISC⁴¹) Fourth Edition were administered as measures
141 of IQ: Block Design, Picture Concepts, Vocabulary, and Similarities. NVIQ was determined by the
142 combined scores on Block Design and Picture Concepts subtests. VIQ was calculated from the scores
143 on Vocabulary and Similarities subtests. (See Table 1 for included participant demographics). All
144 participants were screened for colour deficiencies using the Neitz Test of Color Vision⁴² ; all included
145 participants were classified as normal trichromats.

146 All participants were tested on the IQ tests described above and the Farnsworth-Munsell
147 100-Hue test (FM100) described below. Additionally, the three child groups (TD 6-7y, TD 8-9y, and
148 ATY child) were tested on the chromatic contrast discrimination threshold test, described below.
149 This test was developed for a children's colour perception test battery partly in response to the

150 observed relationship between performance on the FM100 and non-verbal ability (NVIQ) in adults⁹
151 and was therefore not available at the time of the adult testing.

152 Results from six participants were removed from the FM100 analysis, for the following
153 reasons: one ASD adult, one TD 6-7y child and two ATY child participants due to Total Error Scores of
154 over 500, implying poor task comprehension²²; one TD 8-9y child and one ATY child due to task non-
155 completion (time constraints); one TD 8-9y child for deuteranomaly, as evidenced by Neitz Test
156 screening and the FM100 error pattern. Table 1 reports indices for the remaining participants,
157 whose FM100 results were included in analysis. On the chromatic contrast discrimination threshold
158 test (see table S1), results were excluded from two ATY child participants due to task non-
159 completion (time constraints), and from the one TD 8-9y child due to deuteranomaly.

160 **Table 1.** Participant group demographics, showing only participants whose results were included in
161 the FM100 analysis (see text for explanation). Chronological age is shown in years. Verbal and Non-
162 Verbal IQ are shown as standardised scores. In both adult groups, IQ was assessed using the WASI. In
163 child groups (TD 6-7y, TD 8-9y and ATY child) IQ was assessed using the WISC Fourth Edition.
164 Standard deviations are shown in brackets.

165

Group	Chronological Age	Verbal IQ	Non-Verbal IQ
TD Adult (n=28)	20.07 (2.04)	96.36 (10.47)	108.89 (9.23)
ASD Adult (n=17)	18.12 (1.22)	86.35 (19.05)	93.59 (18.94)
TD 6-7y (n=11)	6.37 (0.17)	111.45 (10.78)	109.27 (10.11)
TD 8-9y (n=17)	8.95 (0.31)	121.24 (11.84)	112.24 (13.79)
ATY Child (n=15)	12.06 (2.48)	71.4 (22.82)	85.53 (20.21)

166

167 b) Farnsworth-Munsell 100-Hue Test (FM100)

168 The Farnsworth-Munsell 100-Hue test²⁰ is a measure of chromatic discrimination. It consists
169 of 85 coloured caps split across four trays. The caps vary only in hue, with lightness and saturation
170 kept constant. Each tray has 21 removable intermediate caps (with the exception of the first tray
171 where there are 22 caps) whose hues vary smoothly between those of the two fixed caps at either
172 end. Standard administration procedures were followed: For each tray, the intermediate caps are
173 removed from the tray and placed in a random arrangement while the participant looks away. The

174 participant is then asked to view and place the intermediate caps in the correct order in the tray
175 between the two fixed caps, with as little difference in hue between neighbouring caps as possible.
176 Standard prompts of, “Which colour is most like the one at the end?”, were used to ensure that the
177 task was understood correctly. The trays were completed in different orders between participants.
178 The order in which the participant placed the caps was recorded by the experimenter. The task was
179 completed under simulated daylight illumination of colour temperature 6500K (D65) produced by a
180 VeriVide D65 “Artificial Daylight” lamp.

181 Standard scoring procedures were followed. Error scores for each tray position are
182 calculated on the basis of the differences between its chosen cap and the two neighbouring caps,
183 generating a baseline score of 2 for each cap when in perfect order. Error scores for caps at the end
184 of each tray were calculated using the neighbouring cap in the same tray and the first cap of the next
185 tray, so that all caps are considered on a continuum around the colour circle. The Total Error Score
186 (TES) is computed by first subtracting the baseline score from each tray position error score and then
187 summing all 85 individual error scores. Specific anomalies of colour vision are revealed by specific
188 error patterns (clustering of cap transposition errors along the protan, deutan or tritan axes).

189

190 c) Chromatic Contrast Discrimination Threshold Test (CCDT)

191 i) Overview

192 The Chromatic Contrast Discrimination Threshold (CCDT) was designed to isolate and assess
193 discrimination within each of the cardinal chromatic mechanisms independently, similarly to the
194 class of contemporary chromatic discrimination tests which include the Colour Assessment and
195 Diagnosis (CAD) test²⁹ and the Cambridge Colour Test (CCT)³⁰. It was developed for use in this and
196 related studies of colour perception in children, with the requirements that it be engaging to
197 children, portable, relatively quick to run, and easily reproduced without specialist equipment. The
198 thresholds measured by the CCDT are comparable to those from the CAD and CCT, although exact
199 comparisons cannot be made because of differences in the specific shape discrimination task used
200 and the background chromaticity and luminance. The CCDT differs from the Farnsworth-Munsell
201 100-Hue test in that it directly measures thresholds for chromatic discrimination around a point of
202 neutral chromaticity, whereas the FM100 does not measure thresholds but instead requires the
203 observers to detect (and then seriate) chromatic differences between colours at a fixed distance
204 from neutral chromaticity. These chromatic differences have been selected to be near threshold for
205 normal observers (note that, in general, discrimination thresholds for hues of roughly equal

206 luminance and saturation will differ from thresholds relative to a neutral chromaticity). Despite the
207 differences between the two types of test, other studies suggest that the age dependence for both is
208 similar³³, supporting the assumption that both rely on the same basic chromatic processing
209 mechanisms.

210 i) Apparatus and Setup

211 Colour stimuli were displayed on a computer screen placed at the back of a black viewing
212 box (36cm x 44.8cm x 62.3cm). Participants rested their heads on a chin rest placed centrally at the
213 front of the box and viewed the screen through an aperture (13.5cm x 9cm) placed 21cm along the
214 box length, from a distance of 62cm. One of two different computer setups was used to control the
215 experiment, depending on the group: for the TD 6-7y and ATY child groups, the experiment ran on
216 a Dell Inspiron Laptop with stimuli displayed on its 14-inch screen; while for the TD 8-9y group the
217 experiment ran on a custom built portable desktop tower, with standard components, running
218 Windows 7 64-Bit edition with a PNY 600 10-bit graphics board with the stimuli displayed on a 10bit
219 23-inch Proart LCD monitor PA 238Q using a display port adapter. The same experimental
220 programme (the Chromatic Contrast Discrimination Threshold test, or CCDT) was used for both
221 setups, written in Matlab (v7.6.0, 2012b, The MathWorks, 2008, 2012), with graphics display
222 functions from Psychtoolbox⁴³ and colorimetric conversion functions from *kccv* (a set of Matlab
223 routines based on standard formulae⁴⁴ tailored for 8- or 10-bit displays appropriately); the 10-bit
224 display used the NVIDIA QUADRO performance drivers. Spectral emission properties of both
225 screens were characterised using a PR-650 spectroradiometer and colorimetric calibration tables
226 were checked regularly using a Minolta CS-100 chromameter and updated when necessary to ensure
227 colorimetric accuracy of the displayed stimuli.

228 ii) Stimuli

229 On each trial, a single coloured arrow (visual angle = 1.83°), pointing either leftwards or
230 rightwards, was presented. The vertical position of the arrow was randomly jittered from trial to trial
231 5.51° above and below the central fixation point (visual angle = 0.92°), on an achromatic grey
232 background (CIE 1931 coordinates: x=0.36, y=0.37; Y=20.46 cd/m² for the 8-bit display; x=0.314,
233 y=0.339; Y=64.8 cd/m² for the 10-bit display). The arrow colour was systematically varied in
234 increments along only one of the three cone-opponent-contrast axes⁴⁵ (L-M or 'Red-Green'; S-
235 (L+M) or 'Blue-Yellow'; L+M or 'luminance'; see supplementary material for further detail). The just-
236 noticeable difference in arrow colour with respect to the background was calculated in ΔE units in a
237 perceptually uniform colour space (CIE L*u*v* space).

238 iii) Design

239 A staircase protocol was used to vary the colour difference between the arrow and
240 background on each trial, stepping through differences on each half of the red-green, blue-yellow
241 and luminance cone-opponent-contrast axes separately, beginning with supra-threshold difference
242 values and moving in progressively smaller increments to difference values that are just reliably
243 detected by the observer. A one-up/two-down procedure was used, in which the participant must
244 be correct twice consecutively to go down the staircase (i.e. testing smaller colour differences)
245 whereas an incorrect answer will take the participant up the staircase (i.e. testing larger colour
246 differences). (Details of step sizes are provided in the supplementary material). Each colour axis was
247 tested in a separate block of trials, with a maximum of 100 trials per colour axis (50 for each colour
248 direction of the axis; e.g. 50 each for “bluer” and “yellower”) and a maximum number of 30 reversals
249 per half-axis, with the two independent staircases for each half-axis randomly interleaved in the one
250 block. Each trial began with a 500ms centrally positioned white fixation dot, followed by the target
251 which appeared for 150ms. The participant had to respond whether the arrow pointed to either the
252 left of right, by pressing the corresponding mouse button. There was no time limit on the response,
253 and the next trial began immediately after the response had been given.

254 iv) Procedure

255 Participants sat in front of the viewing box. Prior to the start of the experiment, the
256 researcher checked that the participant understood the difference between left and right. A short
257 practice set of highly visible arrows (with supra-threshold luminance contrast) was administered to
258 check that the participant understood the experiment. After the practice set the actual experiment
259 began. In a game-like format, participants were presented with a choice for each condition (each
260 depicted by a different storybook image) and performed each condition only once. The order of
261 condition was not counterbalanced but there was no significant difference of order for any of the
262 groups (lowest $p=0.11$). Each colour-axis condition ran until either the maximum number of trials or
263 maximum number of staircase reversals was reached. Once each condition had finished the
264 participant was given a short break before continuing to the next condition. All conditions were
265 either completed in one session or over two sessions.

266

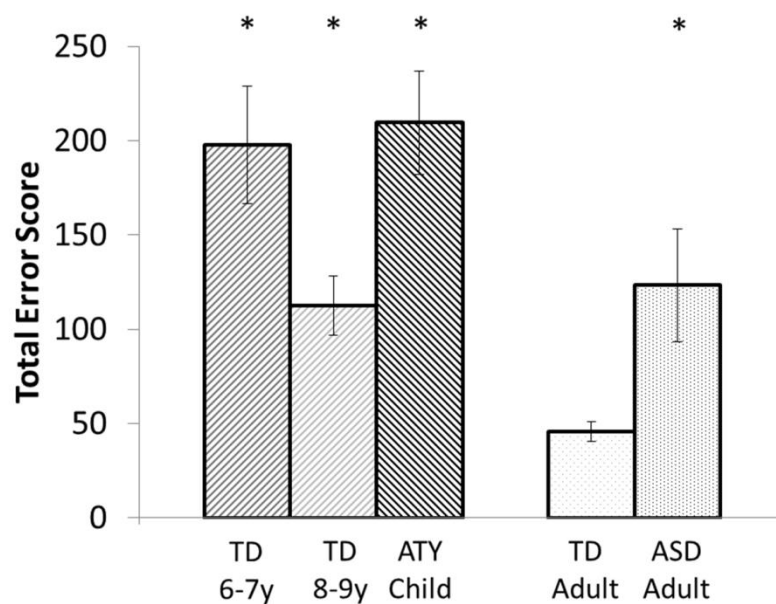
267

268

269 **Results**

270 **Experiment 1: Farnsworth-Munsell 100-Hue Test Results**

271 The error patterns of all included participants revealed no specific anomalies of colour vision. A
272 one-way ANOVA conducted with Total Error Score (TES) as the dependent variable and a between-
273 subjects factor of participant group found a significant difference in TES between groups, $F(4, 83) =$
274 $11.83, p < 0.001$ (see figure 1). Subsequent post hoc t-tests using a Bonferroni-corrected p-value
275 revealed specific group differences: the TD adult group had significantly lower TES (mean TES =
276 45.79) than all other groups (highest $p < 0.001$) and the TD 8-9y group had significantly lower TES
277 (mean TES = 112.47) than the ATY child group (mean TES = 209.47), $t(30) = -3.16, p < 0.05$ (TD 6-7 group
278 mean TES = 197.64 ; ASD adult group mean TES = 123.29). No other specific group TES comparisons
279 revealed significant differences (lowest $p = 0.115$).



280

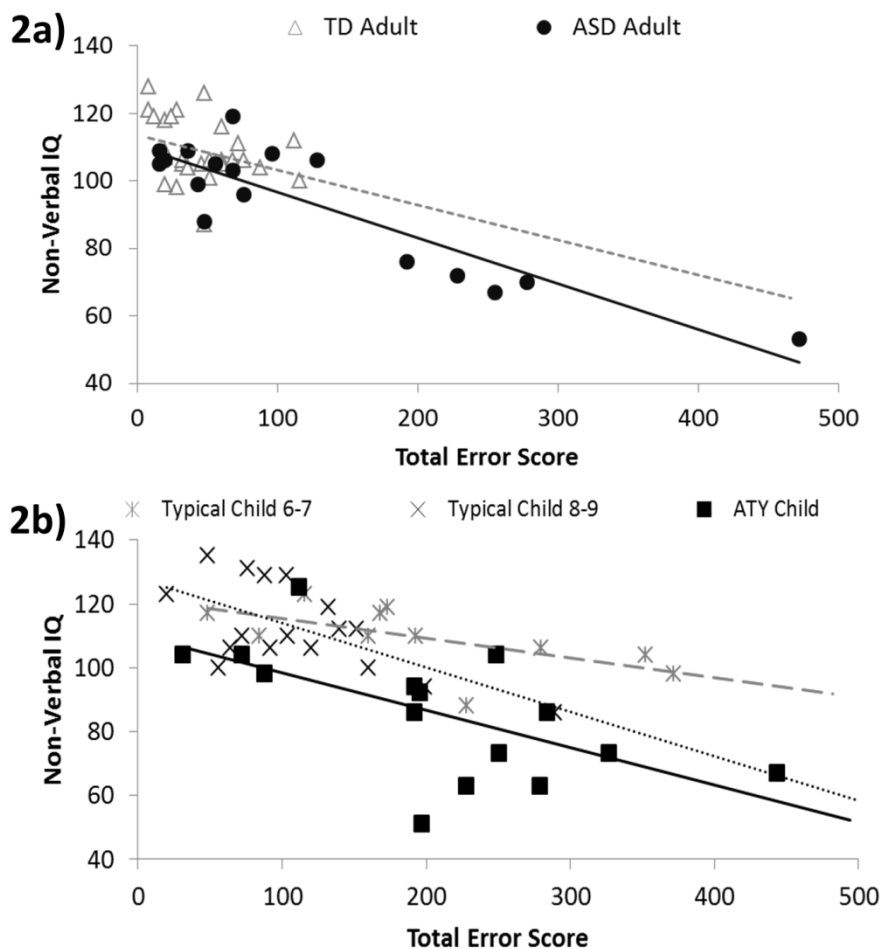
281

282 Figure 1 – Total Error Scores (TES) for the FM100 for each participant group. Higher TES reflects
283 poorer performance on the test. Errors bars are +/- one standard error. The * denotes a significant
284 difference from the expected age-appropriate norms reported by Kinnear & Sahraie (2002).

285

286 a) Correlation of IQ with TES

287 Correlations were calculated for each group between TES and standardised scores on VIQ and
 288 NVIQ subscales of respective IQ tests. For VIQ and TES correlations, significant correlations were
 289 found for ASD adult ($r=-0.68, p<0.005$), and ATY child groups ($r=-0.58, p<0.05$). No other VIQ/TES
 290 correlations were significant (lowest $p=0.33$). For NVIQ correlations with TES there were significant
 291 correlations for ASD adults ($r=-0.89, p<0.001$), TD6-7y ($r=-0.63, p<0.05$), TD 8-9y ($r=-0.65, p<0.005$),
 292 and ATY child groups ($r=-0.62, p<0.05$) (see Figure 2). A trend towards significance was observed for
 293 the TD adult group ($r=-0.32, p=0.095$).



294

295

296 Figure 2 – Individual standardised NVIQ scores plotted against FM100 Total Error Scores. Lines
 297 are least-squares best fits for each group. In both graphs, ASD groups are represented by filled
 298 shapes and solid lines. (a) Adult groups: The TD adult group are denoted by unfilled triangles and
 299 the ASD adult group by filled black circles. (b) Child groups: The TD 6-7y group are denoted by
 300 grey asterisks*, the TD 8-9y group by grey crosses, and the ATY child group by filled black
 301 squares.

302

303 b) Relation to previously reported age-norms

304 Comparisons between the TES found in this sample and expected age-norms²² were made.
305 Difference scores between observed and expected scores for every participant were calculated, with
306 a negative score indicating worse performance relative to the expected age score and a positive
307 score better performance (see figure 1). One-sample t-tests were conducted against a value of 0 to
308 reveal any significant deviations in each participant sample relative to expected age norms.
309 Significantly better than expected performance was found in the TD 6-7y, $t(10)=2.81$, $p<0.05$ and TD
310 8-9y, $t(16)=2.4$, $p<0.05$, groups and significantly worse than expected performance for the ASD adult,
311 $t(16)=2.65$, $p<0.05$, and ATY child, $t(14)=-3.65$, $p<0.005$, groups. No significant differences with age-
312 expected norms were found for the TD adult group, $t(27)=1.02$, $p=0.32$.

313 a) Discrimination thresholds on chromatic and luminance axes

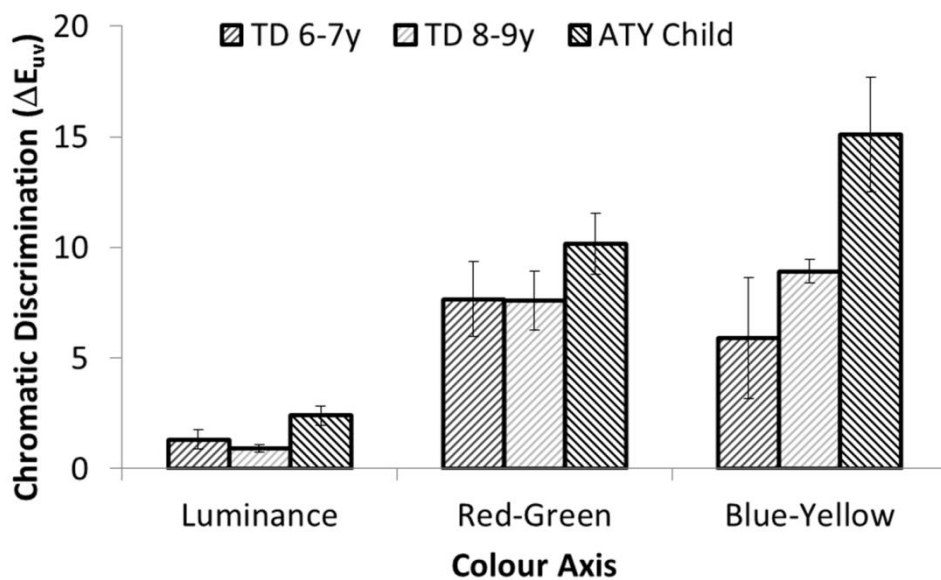
314 Individual staircases for each colour half-axis were analysed for convergence and excluded
315 from further analysis if either of two conditions were met: (a) no reversals in the final 20% of trials or
316 (b) final stimulus contrast less than or equal to zero, relative to background chromaticity. If the
317 staircase for one half-axis did not meet these conditions, the staircase in the other half-axis was also
318 discarded from analysis. For any one axis (Red-Green (R-G), Blue-Yellow (B-Y), or luminance (LUM)),
319 a maximum of 6 participants' staircases were excluded on these criteria for the ATY child group, and
320 a maximum of 3 were excluded for the TD 6-7 child group. All staircases in the TD 8-9y group were
321 accepted. All participants described in Table S1 in the supplementary material contributed
322 acceptable data for at least one axis.

323 Threshold contrasts for each of the six half-axes were calculated in ΔE_{uv} units, as described
324 above. The two thresholds for the two directions for each condition were then averaged together to
325 give an overall threshold for each cardinal axis (e.g. blue and yellow individual thresholds were
326 averaged together for a B-Y axis threshold). To normalise the threshold distributions for each colour
327 axis, ΔE_{uv} thresholds were converted to a logarithmic scale.

328 To compare thresholds between colour axes, data collected from the 8-bit display (TD 6-7y
329 and ATY child groups) and 10-bit display (TD 8-9y child group) systems were analysed separately,
330 given the difference in background luminance between the systems, which affects the absolute
331 magnitude of the thresholds. For the former, a two-way ANOVA with participant group (TD 6-7y/
332 ATY child) and colour axis as fixed factors and threshold (ΔE_{uv}) as dependent factor revealed a
333 significant main effect for colour axis, $F(1,56)=15.78$, $p<0.001$. Post-hoc tests revealed that

334 thresholds were significantly lower for the luminance than both R-G and B-Y colour axes for both TD
 335 6-7y (highest $p < 0.001$) and ATY child groups (highest $p < 0.001$). There was no significant difference
 336 between R-G and B-Y thresholds for either group (lowest $p = 0.329$). A significant main effect was also
 337 observed for group, $F(1,56) = 15.781$, $p < 0.001$. Post-hoc tests conducted for each colour axis further
 338 revealed significant differences between groups on the B-Y axis, $t(19) = 2.93$, $p < 0.05$, with thresholds
 339 significantly higher for the ATY vs TD 6-7y child groups, but not for luminance ($p = 0.22$) or R-G axes
 340 ($p = 0.19$).

341 Similarly, for data from the TD 8-9y child group, a two-way repeated-measures ANOVA with
 342 a within-subjects factor of colour axis (3 levels) revealed a significant main effect of axis ,
 343 $F(2,34) = 39.46$, $p < 0.001$, with luminance thresholds significantly lower than both R-G, $t(17) = 12.9$,
 344 $p < 0.001$, and B-Y, $t(17) = 6.18$, $p < 0.001$, thresholds. R-G and B-Y thresholds did not significantly differ
 345 from each other, $t(17) = 1.6$, $p = 0.13$.



346

347

348 Figure 3 – Average discrimination thresholds for the luminance, red-green and blue-yellow colour
 349 axes, as determined by the Chromatic Contrast Discrimination Threshold test. The three groups (TD
 350 6-7y, TD 8-9y, and ATY child) are indicated by different diagonal line shading, as shown in the legend.
 351 Error bars indicate standard errors.

352

353 c) Correlations of Thresholds with IQ

354 Correlation analyses were conducted to reveal relationships between VIQ and NVIQ and
355 discrimination thresholds on the three colour axes, using adjusted p-values to control for multiple
356 comparisons. No significant correlations were found in any participant group for either VIQ (lowest
357 $p=0.18$) or NVIQ (lowest $p=0.22$).

358 d) Correlations between FM100 and Chromatic threshold

359 Correlation analyses were conducted to examine the relationship between performance on the
360 FM100 and CCDT tests. Performance was again split via participant group. For the CCDT test, the
361 mean chromatic discrimination threshold was calculated by averaging the B-Y and R-G thresholds.
362 No significant correlation was found for the TD 6-7y child group ($p=0.73$). A significant correlation for
363 the ATY child group was found ($r=0.83$, $p<0.001$), while a trend towards significance was revealed for
364 the TD 8-9y child group, $r=0.44$, $p=0.08$.

365 **Regression Analysis**

366 To further assess the role of possible factors on FM100 performance, a multiple regression
367 was carried out to estimate the extent to which performance on the FM100 might be predicted by
368 NVIQ, chronological age and chromatic discrimination (as measured by the average B-Y and R-G
369 thresholds from the CCDT test. This was done to gain an overall measure of chromatic discrimination
370 to make this analogous to the TES on the FM100). Four predictors were included in the model: NVIQ,
371 chronological age, development typicality (Typical/Atypical) and chromatic discrimination threshold.
372 The analysis included only participants who completed both the FM100 and the CCDT test, from all
373 child groups. Predictors were entered into the regression model using the backward entry method,
374 appropriate in the absence of an *a priori* theory for which predictors would explain the most
375 variance in FM100 performance. From this procedure, two models were generated. Model 1
376 included all of the variables, while Model 2 included all variables except for participant group
377 (development typicality) since this factor was found not to be a significant predictor (see Table 2).
378 Overall, both models significantly explained substantial variance in FM100 performance: Model 1
379 explained 53.2% of the variance, $F(4,30)=8.54$, $p<0.001$ while Model 2 explained 53.1% of the
380 variance, $F(3,31)=11.70$, $p<0.001$. Additional regression models run on data from all participant
381 groups alone and in combination are described in the supplementary materials; in all models, NVIQ
382 is always the highest predictor, and although CCDT thresholds are significant predictors when used
383 solely, they explain much less variance than NVIQ-only models.

384

385 **Table 2.** Backward stepwise regression model for the contributions to FM100 performance of
 386 distinct factors: NVIQ, chromatic discrimination, chronological age and development typicality.
 387 Model 1 includes all of the predictor variables; Model 2 omits only development typicality. The
 388 symbols *, **, *** denote significance at $p < 0.05$, $p < 0.01$ and $p < 0.001$ respectively.

389

Variable	Model 1			Model 2		
	<i>B</i>	<i>SE B</i>	β	<i>B</i>	<i>SE B</i>	<i>B</i>
NVIQ	-3.629	0.953	-0.737***	-3.743	0.859	-0.76**
Chromatic Discrimination	236.143	106.66	0.369*	231.364	103.896	0.361*
Chronological Age	-1.9	0.769	-0.557*	-1.767	0.614	-0.518**
Development	13.69	46.23	0.064			
R ²		0.532			0.531	
F-value		8.539			11.701	

390

391

392 Discussion

393 The major finding in this study is that non-verbal general ability differentially affects
 394 participants' performance on two different chromatic discrimination tasks. The results from
 395 experiment 1 show that performance on the FM100 is significantly associated with non-verbal ability
 396 in all groups except typically developing adults. Furthermore, this association is stronger in children,
 397 and even stronger in atypically developing individuals (the majority with ASD). The results from
 398 experiment 2 demonstrate, conversely, that this association between general ability and colour
 399 perception in the younger age groups does not hold for the computer-based chromatic
 400 discrimination threshold test (CCDT).

401 The results therefore strongly suggest that the FM100 is not an unequivocal measure of
 402 colour discrimination but conflates this with a measure of "non-verbal ability", providing evidence
 403 for previous suggestions that cognitive factors unrelated to chromatic discrimination ability
 404 influence performance on the task^{9,37}. The FM100 performance-NVIQ correlation slopes are steeper
 405 and more significant in the child groups than adult. Furthermore, the results of the regression model
 406 suggest that non-verbal ability is a significant predictor of performance in addition to age and the
 407 ability to discriminate between colours.

408 When considering why the FM100 is associated with general cognitive ability but the
409 chromatic discrimination threshold test is not, the relative task demands are important. Successful
410 performance on the FM100 requires attentional and visuospatial abilities in addition to chromatic
411 discrimination ability. Spatial comparisons are required between the selected and non-selected caps.
412 Attention switching between the local field – in making a comparison between two adjacent caps,
413 and the global field - in overseeing the entire colour gradient - is also essential for good
414 performance. These task demands may be influenced by different factors in each group. In typically
415 developing children competency of global processing does not develop until late into childhood⁴⁶.
416 Individuals with ASD are more likely to process visual information locally rather than globally⁴⁷ and
417 to have difficulty switching between local and global processing⁴⁸. Non-verbal ability may also
418 differentially affect performance between groups. Better performance in the older TD groups may
419 reflect more mature global processing competency than in the younger TD groups. Poorer
420 performance in the ASD groups may be the result of difficulty both in sustaining and switching of
421 attention between local and global fields of the FM100.

422 These results have implications for the FM100 norms that have previously been reported²².
423 These norms implicitly assume that performance is unrelated to IQ and that task demands are
424 consistent between different ages. In the current study, both TD child groups are above average in
425 non-verbal ability and perform significantly better than expected from the Kinnear and Sahraie
426 (2002) norms²². Given that the number of participants in both studies is similar for each respective
427 age group, this comparison calls into question the reliability of the norms for these age groups and
428 more so for the younger age groups for which the number of observers is even smaller (e.g. 9
429 observers for age 5 years)²². These results, combined with those from a preliminary study by
430 Hurlbert and colleagues⁹ instead suggest that caution must be exercised when using FM100 norms,
431 for all ages, but especially for younger children or clinical populations where the NVIQ is lower than
432 average with respect to chronological age, given that the relationship between FM100 performance
433 and NVIQ is stronger for groups of the latter type.

434 In comparison to the FM100, the CCDT has fewer task demands. This test of chromatic
435 discrimination requires attention on a trial-by-trial basis only to identify the direction of an arrow.
436 Like other standardised tests in whose class the CCDT falls (e.g. the CCT³⁰ and the CAD²⁹), the CCDT
437 measures discrimination thresholds along isolated chromatic directions away from a fixed
438 adaptation point. Although other colour discrimination tasks have been adapted for use with
439 children (e.g. the CCT⁴⁹), to our knowledge these have yet to be demonstrated as independent of
440 general ability. Other standardised chromatic discrimination threshold tests call on more complex

441 aspects of visual processing which may introduce additional confounds when used in children: the
442 CCT³⁰, for example, requires participants to identify a global shape composed from local elements
443 while the CAD (or City Colour Vision Test⁵⁰) requires participants to discriminate the direction of a
444 moving stimulus^{29, 50}. The former thus presents a potential confound in distinguishing between
445 deficits in local chromatic discrimination vs global shape processing, while the latter may be unable
446 to dissociate between deficits in motion direction discrimination vs chromatic discrimination, which
447 are known to develop at different rates in children^{33, 51}. The CCDT task used in this study may provide
448 a more direct measurement of chromatic discrimination by being a simple shape identification task,
449 which requires only a coarse binary judgement of left versus right, does not depend on numeracy or
450 literacy skills, and does not involve a trade-off between local and global processing, involving the
451 discrimination of only a single large shape against a uniform background. Performance is more likely
452 to be independent of developmental stage or cognitive ability, allowing for age variations in
453 chromatic discrimination to be more accurately captured. Because participants continue the task
454 only until they reach their own individual threshold, task difficulty also remains constant between
455 individuals even though other factors such as chronological age or chromatic discrimination ability
456 may differ between participants. Because of their shared properties, we would expect CCDT
457 performance to share the same pattern of age dependence as that demonstrated for the CCT and
458 CAD³², and our ongoing studies support this expectation.

459 Although we have here carried out the CCDT only in the younger age groups and therefore
460 demonstrated this independence only for those groups, we would expect the independence to hold
461 for the adult groups also, particularly given their better chromatic discrimination ability, higher
462 absolute IQ, and increased attentional capacity relative to children⁵².

463 It is not possible from this study to address specifically whether sensory processing is related
464 to general ability, as suggested by other studies^{13, 19}. Nonetheless, the results here do highlight the
465 importance of using multiple tests within the same sensory processing domain when assessing this
466 link. For example, the results of the FM100 alone would suggest that there is a link between sensory
467 (chromatic) discrimination and intelligence, while the results of the CCDT test alone would suggest
468 that there is no such relationship. The differential correlation between non-verbal ability and the
469 FM100/CCDT test demonstrate that different tests measuring the same sensory (chromatic)
470 discrimination may give different results depending on the different task demands. In order to more
471 accurately tease apart the possible relationship between intelligence and sensory discrimination,
472 comparisons need to be made on performance from multiple tests measuring the same sensory
473 domain.

474 In summary, we show that general ability affects performance on two different colour
475 discrimination tasks differently: performance on the FM100 is associated with non-verbal ability
476 while the chromatic contrast discrimination task is not. This result impacts upon the appropriateness
477 of the clinical use of the FM100. The results from this study suggest that the use of an appropriate
478 psychophysical task which has fewer task demands and is of equal difficulty across all ages will give a
479 more accurate measure of colour discrimination and ultimately visual function in participants.

480

481 **Acknowledgements**

482 The authors thank the teachers and students at Tasker House, Newcastle College, Barndale House
483 School, South Gosforth First School, Bamburgh School, and Collingwood School, for their help and
484 participation. The authors also acknowledge the research assistance of undergraduate students
485 Charlotte Reay, Max O'Collins and Vikki Ingram. The authors also thank two anonymous reviewers
486 for their very helpful comments on an earlier draft of this manuscript. MBC thanks Dr. Deborah Riby
487 and Professor Ann Le Couteur for their general supervisory support. For research studentship
488 support, we gratefully acknowledge the Estate of David Murray Garside (MBC), the Newcastle Vision
489 Fund (MBC), and the EPSRC (EP/H022325/1).

490

- 491 1. Treisman AM, Gelade G. A feature-integration theory of attention. *Cognit Psychol*
492 1980;12:97-136.
- 493 2. Tanaka J, Weiskopf D, Williams P. The role of color in high-level vision. *Trends in Cognitive*
494 *Sciences* 2001;5:211-215.
- 495 3. Whitehead RD, Re D, Xiao D, Ozakinci G, Perrett DI. You Are What You Eat: Within-Subject
496 Increases in Fruit and Vegetable Consumption Confer Beneficial Skin-Color Changes. *PLoS ONE*
497 2012;7:e32988. doi: 32910.31371/journal.pone.0032988.
- 498 4. Roberson D, Davies I, Davidoff J. Color categories are not universal: Replications and new
499 evidence from a stone-age culture. *J Exp Psychol Gen* 2000;129:369-398.
- 500 5. Palmer SE, Schloss KB. An ecological valence theory of human color preference. *Proc Natl*
501 *Acad Sci U S A* 2010;107:8877-8882.
- 502 6. Ling Y, Hurlbert A. Role of color memory in successive color constancy. *J Optical Soc Am A*
503 2008;25:1215-1226.
- 504 7. Franklin A, Sowden P, Burley R, Notman L, Alder E. Color Perception in Children with Autism.
505 *Journal of Autism and Developmental Disorders* 2008;38:1837-1847.

- 506 8. Franklin A, Sowden P, Notman L, et al. Reduced chromatic discrimination in children with
507 autism spectrum disorders. *Developmental Science* 2010;13:188-200.
- 508 9. Hurlbert A, Loveridge C, Ling Y, Kourkoulou A, Leekam S. Color Discrimination and
509 Preference in Autism Spectrum Disorder. *Journal of Vision* 2011;11:429.
- 510 10. Banaschewski T, Ruppert S, Tannock R, et al. Colour perception in ADHD. *Journal of Child*
511 *Psychology and Psychiatry* 2006;47:568-572.
- 512 11. Kim S, Chen S, Tannock R. Visual function and color vision in adults with Attention-
513 Deficit/Hyperactivity Disorder. *Journal of Optometry* 2014;7:22-36.
- 514 12. Farran EK, Cranwell MB, Alvarez J, Franklin A. Colour discrimination and categorisation in
515 Williams syndrome. *Res Dev Disabil* 2013;34:3352-3360.
- 516 13. Galton F. Inquiries into human faculty and its development. London: Dent; 1883.
- 517 14. Spearman C. "General Intelligence," Objectively Determined and Measured. *Am J Psychol*
518 1904;15:201-292.
- 519 15. Deary IJ. Sensory discrimination and intelligence: Postmortem or resurrection? *Am J Psychol*
520 1994;95-115.
- 521 16. Acton GS, Schroeder DH. Sensory discrimination as related to general intelligence.
522 *Intelligence* 2001;29:263-271.
- 523 17. Deary IJ. Intelligence and auditory discrimination: Separating processing speed and fidelity of
524 stimulus representation. *Intelligence* 1994;18:189-213.
- 525 18. Li S-C, Jordanova M, Lindenberger U. From good senses to good sense: A link between tactile
526 information processing and intelligence. *Intelligence* 1998;26:99-122.
- 527 19. Melnick MD, Harrison BR, Park S, Bennetto L, Tadin D. A strong interactive link between
528 sensory discriminations and intelligence. *Curr Biol* 2013;23:1013-1017.
- 529 20. Farnsworth D. The Farnsworth-Munsell 100-Hue Test for the examination of color
530 discrimination: Manual. Baltimore: Macbeth; 1957.
- 531 21. Dain SJ. Clinical colour vision tests. *Clinical and Experimental Optometry* 2004;87:276-293.
- 532 22. Kinnear PR, Sahraie A. New Farnsworth-Munsell 100 hue test norms of normal observers for
533 each year of age 5–22 and for age decades 30–70. *British Journal of Ophthalmology* 2002;86:1408-
534 1411.
- 535 23. Gunther KL, Neitz JAY, Neitz M. A novel mutation in the short-wavelength-sensitive cone
536 pigment gene associated with a tritan color vision defect. *Vis Neurosci* 2006;23:403-409.
- 537 24. Heywood CA, Gadotti A, Cowey A. Cortical area V4 and its role in the perception of color. *J*
538 *Neurosci* 1992;12:4056-4065.

- 539 25. Ménéage MJ, Papakostopoulos D, Dean Hart JC, Papakostopoulos S, Gogolitsyn Y. The
540 Farnsworth-Munsell 100 hue test in the first episode of demyelinating optic neuritis. *Br J Ophthalmol*
541 1993;77:68-74.
- 542 26. Victor JD, Maiese K, Shapley R, Sidtis J, Gazzaniga MS. Acquired central dyschromatopsia:
543 analysis of a case with preservation of color discrimination. *Clin Vision Sci* 4:183-196.
- 544 27. Knoblauch K, Saunders F, Kusuda M, et al. Age and illuminance effects in the Farnsworth-
545 Munsell 100-hue test. *Appl Opt* 1987;26:1441-1448.
- 546 28. Beirne RO, McIlreavy L, Zlatkova MB. The effect of age-related lens yellowing on
547 Farnsworth–Munsell 100 hue error score. *Ophthalmic and Physiological Optics* 2008;28:448-456.
- 548 29. Birch J, Barbur JL, Harlow AJ. New method based on random luminance masking for
549 measuring isochromatic zones using high resolution colour displays. *Ophthalmic and Physiological*
550 *Optics* 1992;12:133-136.
- 551 30. Regan BC, Reffin JP, Mollon JD. Luminance noise and the rapid determination of
552 discrimination ellipses in colour deficiency. *Vision Res* 1994;34:1279-1299.
- 553 31. Paramei GV, Oakley B. Variation of color discrimination across the life span. *J Optical Soc Am*
554 *A* 2014;31:A375-A384.
- 555 32. Barbur JL, Rodriguez-Carmona M. Colour vision changes in normal aging. In: Elliot AJ,
556 Fairchild MD (eds), *Handbook of Color Psychology*: Cambridge University Press; In Press.
- 557 33. Knoblauch K, Vital-Durand F, Barbur JL. Variation of chromatic sensitivity across the life span.
558 *Vision Res* 2001;41:23-36.
- 559 34. Birch J. *Diagnosis of Defective Colour Vision*. 2nd ed. Oxford: England: Butterworth
560 Heinemann; 2001.
- 561 35. Karpf RJ, Goss AE, Small MY. Naming, Selection, and Ordering of Color (“Hue”) by Young
562 Children. *J Gen Psychol* 1974;90:297-314.
- 563 36. Dain SJ, Ling BY. Cognitive abilities of children on a gray seriation test. *Optometry and Vision*
564 *Science* 2009;86:E701 - E707.
- 565 37. Dain SJ, Ling BY. Cognitive Abilities of Children on a Gray Seriation Test. *Optom Vis Sci*
566 2009;86:E701-E707 710.1097/OPX.1090b1013e3181a1059d1046.
- 567 38. Schroeder A, Kreutz M, Meyer M, Erb C. Influence of learning effects on the results of the
568 cap-sorting test Roth 28-hue (E) desaturated: Influence of learning effects on colour-arrangement
569 tests. *Color Research and Application* 2007;32:16-21.
- 570 39. Foote KG, Neitz M, Neitz J. Comparison of the Richmond HRR 4th edition and Farnsworth-
571 Munsell 100 Hue Test for quantitative assessment of tritan color deficiencies. *J Optical Soc Am A*
572 2014;31:A186-A188.

- 573 40. Wechsler D. *Wechsler abbreviated scale of intelligence*. San Antonio, TX: Psychological
574 Corporation; 1999.
- 575 41. Wechsler D. *Wechsler Intelligence Scale for Children - Fourth Edition*. San Antonio, TX:
576 Harcourt Assessment, Inc; 2003.
- 577 42. Neitz J, Summerfelt P, Neitz M. *The Neitz Test of Color Vision*. Los Angeles: Western
578 Psychological Service; 2001.
- 579 43. Brainard DH. The Psychophysics Toolbox. *Spat Vis* 1997;10:433-436.
- 580 44. Wolf K. The Modulation of Simultaneous Chromatic Contrast. *Institute of Neuroscience*.
581 Newcastle Upon Tyne: University of Newcastle Upon Tyne; 2011:402.
- 582 45. Eskew RT, McLellan JS, Giulianini F. Chromatic detection and discrimination. In:
583 Gegenfurtner KR, Sharpe LT (eds), *Color Vision: From Genes to Perception*. Cambridge, UK: University
584 Press; 1999:345-368.
- 585 46. Kramer JH, Ellenberg L, Leonard J, Share LJ. Developmental sex differences in global-local
586 perceptual bias. *Neuropsychology* 1996;10:402-407.
- 587 47. Plaisted K, Swettenham J, Rees L. Children with Autism Show Local Precedence in a Divided
588 Attention Task and Global Precedence in a Selective Attention Task. *The Journal of Child Psychology
589 and Psychiatry and Allied Disciplines* 1999;40:733-742.
- 590 48. Rinehart NJ, Bradshaw JL, Moss SA, Brereton AV, Tonge BJ. A deficit in shifting attention
591 present in high-functioning autism but not asperger's disorder. *Autism* 2001;5:67-80.
- 592 49. Goulart PRK, Bandeira ML, Tsubota D, Oiwa NN, Costa MF, Ventura DF. A computer-
593 controlled color vision test for children based on the Cambridge Colour Test. *Vis Neurosci*
594 2008;25:445-450.
- 595 50. Barbur JL, Harlow AJ, Plant GT. Insights into the different exploits of colour in the visual
596 cortex. *Proceedings of the Royal Society of London B: Biological Sciences* 1994;258:327-334.
- 597 51. Parrish EE, Giaschi DE, Boden C, Dougherty R. The maturation of form and motion
598 perception in school age children. *Vision Res* 2005;45:827-837.
- 599 52. McAvinue LP, Habekost T, Johnson KA, et al. Sustained attention, attentional selectivity, and
600 attentional capacity across the lifespan. *Attention, Perception, & Psychophysics* 2012;74:1570-1582.

601

Supplementary Information

602 **Farnsworth-Munsell 100-Hue Test Results**

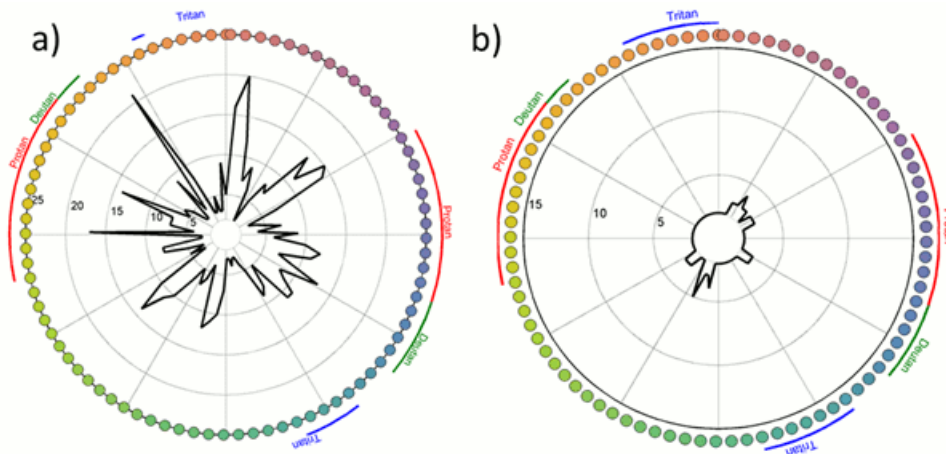


Figure S1. Sample Farnsworth-Munsell 100-Hue error plots for two participants. Error scores at individual cap positions around the hue circle are represented by radial distance of the black contour from the centre. The error scale, shown by labels on concentric grey contours, differs in the two plots. a) Results for a participant from the ASD adult group, who performed below average on the FM100 (FM100 Total Error Score = 472; NVIQ standard score = 53, below average). b) Results for a participant from the TD adult group who performed above average on the FM100 (FM100 Total Error Score = 28; NVIQ standard score = 121, above average).

603

604 **Chromatic Contrast Discrimination Threshold Test (CCDT)**

605 Table S1. Participant group demographics, showing only participants whose results were included in
 606 the CCDT analysis. Chronological age is shown in years. VIQ and NVIQ are shown as standardised
 607 scores. In all groups IQ was assessed using the WISC Fourth Edition. Standard deviations are shown
 608 in brackets.

Group	Chronological Age	VIQ	NVIQ
TD6-7y (n=12)	6.38 (0.17)	113.33 (12.17)	107 (12.45)
TD8-9y (n=18)	8.96 (0.31)	121.94 (11.87)	112.83 (13.61)
ATY Child (n=16)	12.15 (2.42)	71.25 (22.05)	85.13 (19.59)

609

610 Stimuli: The contrast of the arrow stimulus with respect to the background was calculated in cone-
 611 opponent-contrast coordinates following the formulae given in Eskew et al. (1999). The origin of this
 612 coordinate space is given by the cone excitations to a reference white surface (L_0 , M_0 and S_0), in this
 613 case, the uniform grey background, using the Smith-Pokorny (1975) cone fundamentals.¹ Cone

614 excitation values for the arrow are defined with respect to this origin by the ratios: $\Delta L = (L - L_0)/L_0$,
615 $\Delta M = (M - M_0)/M_0$ and $\Delta S = (S - S_0)/S_0$, where L, M, and S are the cone excitations to the arrow
616 stimulus. The cone-opponent contrast coordinates of the arrow stimulus are then calculated for
617 each of three axes:

$$618 \quad LUM = [0.78*\Delta L + 0.37*\Delta M];$$

$$619 \quad RG = [0.7*\Delta L - 0.72*\Delta M + 0.02*\Delta S];$$

$$620 \quad BY = - [0.55*\Delta L + 0.25*\Delta M - 0.8*\Delta S];$$

621 where *LUM* is the luminance axis, *RG* is the “red-green” axis and *BY* is the “blue-yellow” axis.

622 For each half of each axis, a set of displayable colours (numbering on average 75) were calculated
623 whose colour differed from the uniform background only along that axis, in the smallest increments
624 achievable for the given display device. For example, for the “yellower” direction of the *BY* axis and
625 the 10bit display, all ~65 stimuli had effectively the same *LUM* and *RG* coordinates as the grey
626 background while their *BY* coordinates varied systematically from the background *BY* coordinate in
627 steps equal to (on average) $0.3142 \Delta E_{u^*v^*}$, up to a maximum difference of approximately $20 \Delta E_{u^*v^*}$.
628 (The unit $\Delta E_{u^*v^*}$ is defined as a just-noticeable-difference in the perceptually uniform CIE Lu^*v^*
629 space.)

630

631 Staircase Procedure: Six individual staircases, one for each direction of each colour axis (i.e. each
632 half-axis), were completed by each observer. Each staircase drew its stimuli from one of the colour
633 sets calculated as described above. Each colour axis was tested with a separate block of trials, in
634 which the two half-axis staircases were interleaved; e.g. for the *BY* colour axis, the “bluer” and
635 “yellower” staircases were interleaved. Within each half-axis staircase, the arrow colour was varied
636 in 6 decreasing step sizes ranging from approximately $10 \Delta E_{u^*v^*}$ to $0.3 \Delta E_{u^*v^*}$ (for the 10-bit

637 experiment; step-sizes ranged from approximately $10 \Delta E_{u^*v^*}$ to $0.8 \Delta E_{u^*v^*}$ for the 8-bit experiment)
638 in a one-up/two-down procedure. The steps were selected in terms of the indices of the colour
639 stimuli in the colour sets described above, with the smallest step in each case corresponding to an
640 index step-size of 1.

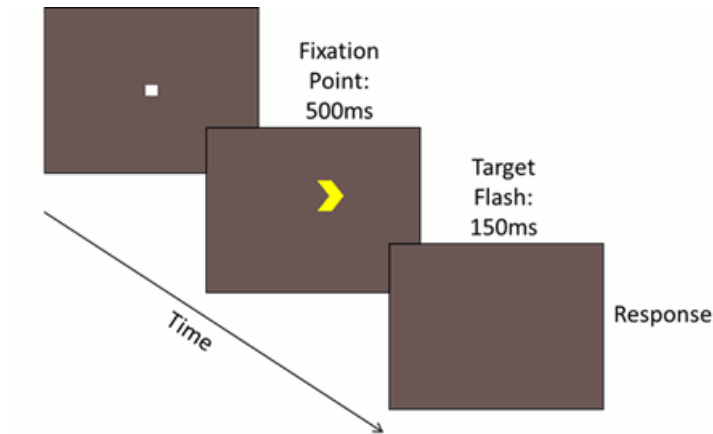


Figure S2. Schematic illustration of stimuli in the CCDT, and the temporal sequence of an individual trial. Sizes of stimuli are not to scale.

641

642

643 **Comparison of 8-bit and 10-bit display systems for CCDT**

644 In this paper, the CCDT was used with two different graphics boards/display systems (8-bit and 10-
645 bit), due to changes in equipment availability over the course of the study. Although the 10-bit
646 system is preferable for widespread use because of its higher chromatic resolution, the two systems
647 are equally effective for the populations studied here. A control experiment in 5 adult individuals,
648 who performed the test on both 8-bit and 10-bit systems in the same day, revealed a significant
649 positive correlation between thresholds on the two systems ($r = 0.88$; $p < 0.001$). All thresholds
650 obtained on the two systems are also comparable to thresholds reported for other standardised
651 chromatic discrimination tests, e.g. the Colour Assessment and Diagnosis (CAD) test¹ and the
652 Cambridge Colour Test (CCT)², in both absolute magnitude and variation across chromatic direction.
653 Note that because thresholds vary across colour space, with both the background chromaticity and
654 luminance, it is not appropriate to make exact comparisons.

655 **Regression Model**

656 Further regression models were run on data from all participants (child and adult groups), using TES
657 as a dependent variable and NVIQ, VIQ, Chronological Age (months), and Development as predictors,
658 excluding CCDT (Backwards entry method). Three iterations of model yielded the following
659 predictions:

660 1) $R^2=64.9\%$, all predictors included

661 2) $R^2=64.9\%$, Development only removed

662 3) $R^2=64.3\%$, Development and VIQ removed (only NVIQ and CA remaining)

663 NVIQ is always the highest predictor, but with slightly lower weighting than in results from Model 2
664 presented in Table 2 of the main paper. A similar final regression model run for the child groups only
665 (with NVIQ and Chronological Age only as predictors after Development and VIQ have been
666 removed) yields R^2 of 47.1%. For the adult groups only, the final regression model explains 66.9% of
667 the variance (with NVIQ only as predictor, after Development, VIQ and Chronological Age have been
668 removed).

669 All models explain more variance when not including CCDT measures. For data from the child groups
670 only, CCDT thresholds (converted to logarithms) are a significant predictor when they are the only
671 predictor but explain much less of the variance (23.4%) than the NVIQ-only model (39.5%). For data
672 from both the child and adult groups, the NVIQ only model predicts slightly more variance (41.4%).
673 The adult only NVIQ only model predicts much of the variance (66.9%).

674 Regression models were also run using the CCDT threshold from each colour axis separately with
675 its corresponding FM100 partial error score (PES), for example, using only the Red-Green PES and
676 Red-Green CCDT thresholds within one regression analysis and the Blue-Yellow PES and Blue-Yellow
677 CCDT thresholds in a separate analysis. The final Red-Green model explained 44% of the variance
678 (final model predictors: Development, Red-Green CCDT threshold and NVIQ). The Blue-Yellow model
679 explained more of the variance (61.3%, final model predictors: Chronological Age, Blue-Yellow CCDT
680 threshold and NVIQ) than the Red-Green model. Despite the difference in overall variance explained

681 between the Red-Green and the Blue-Yellow model, NVIQ was still the highest loading predictor in
682 both models.

683 A final regression analysis was done using Development, Chronological age, NVIQ, Red-Green CCDT
684 threshold and Blue-Yellow CCDT threshold as predictors and FM100 TES as a dependent variable.

685 Five iterations of the model yielded the following predictions:

- 686 1) $R^2 = 54.8\%$, All variables included
- 687 2) $R^2 = 54.8\%$, Development removed
- 688 3) $R^2 = 52\%$, Blue-Yellow CCDT threshold removed
- 689 4) $R^2 = 47.8\%$ Red-Green CCDT threshold removed
- 690 5) $R^2 = 43.3\%$ Chronological Age removed (Only NVIQ remaining).

691

692

693 1 Smith VC, Pokorny J. Spectral sensitivity of the foveal cone photopigments between 400 and 500
694 nm. *Vision Research* 1975; 15:161-171.

695 2 Birch J, Barbur JL, Harlow AJ. New method based on random luminance masking for measuring
696 isochromatic zones using high resolution colour displays. *Ophthalmic and Physiological Optics*
697 1992;12:133-136.

698 3 Regan BC, Reffin JP, Mollon JD. Luminance noise and the rapid determination of discrimination
699 ellipses in colour deficiency. *Vision Res* 1994;34:1279-1299.

700