

1 Scientific maps should reach everyone: a
2 straightforward approach to let colour blind
3 people visualise spatial patterns

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Abstract

Maps represent powerful tools to show the spatial variation of a variable in a straightforward manner. A crucial aspect in map rendering for its interpretation by users is the gamut of colours used for displaying data. One part of this problem is linked to the proportion of the human population that is colour blind and, therefore, highly sensitive to colour palette selection. The aim of this paper is to present a function in R - `cblind.plot` - which enables colour blind people to just enter an image in a coding workflow, simply set their colour blind deficiency type, and immediately get as output a colour blind friendly plot. We will first describe in detail colour blind problems, and then show a step by step example of the function being proposed. While examples exist to provide colour blind people with proper colour palettes, in such cases (i) the workflow include a separate import of the image and the application of a set of colour ramp palettes and (ii) albeit being well documented, there are many steps to be done before plotting an image with a colour blind friendly ramp palette. The function described in this paper (`cblind.plot`), on the contrary, allows to (i) automatically call the image inside the function without any initial import step and (ii) explicitly refer to the colour blind deficiency type being experienced, to further automatically apply the proper colour ramp palette.

60 1 Introduction

61 Maps are widely used to convey geographical information. Various map types
62 exist and are used in activities from day-to-day tasks such as route planning
63 to furthering scientific studies of such topics as climate change and the spread
64 of invasive species. Despite their widespread use, maps can be problematic
65 information resources. Any map is a generalisation, hence an imperfect repre-
66 sentation of the features shown. A map is also typically interpreted visually,
67 and hence different people may come to dissimilar conclusions about the
68 mapped features. Standard procedures and good practices for map making
69 exist but may not always be followed or may not fully address concerns.
70 Moreover, maps can be produced in a way that may, accidentally or deliber-
71 ately, deceive and lie to the reader (Monmonier, 2018). Indeed, it has been
72 suggested that maps are “the most used and least understood documents of
73 modern civilisation” (Brown, 1953, cited in Maling, 1989, p.144, and, more
74 recently, in Foody, 2021).

75 Any map is a generalisation and hence an imperfect representation of the
76 features shown. A map is also typically interpreted visually and hence differ-
77 ent people may come to dissimilar conclusions about the mapped features.
78 Standard procedures and good-practices for map making exist but may not
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80 be produced in a way that may, accidentally or deliberately, deceive and lie
81 to the reader (Monmonier, 2018).

82 Hence, showing the variation of a variable over space in a map considering
83 the whole gamut of colours is not a simple matter in its very nature (Pointer,
84 1980). In most cases maps full of colours are used, to catch the eye of the
85 reader in a straightforward manner.

86 The misuse of colour could be dangerous, by producing a misleading
87 perception of the achieved results. As an example, yellow is expected to
88 catch the human retina more than other colours. If used in the wrong palette
89 position, it would lead the eye to assign more importance to that particular
90 range. For instance, using it in the middle of the palette colour ramp would
91 lead the reader to see such values as maxima (Cramieri et al., 2020).

92 One common problem with contemporary mapping is that software pack-
93 ages often offer a range of colour palettes for data display, and these may
94 vary in suitability for both a mapping task and the target audience. One part
95 of this problem is linked to the proportion of the human population that
96 is colour blind and, therefore, highly sensitive to colour palette selection.
97 *Colour vision impairments*, also known as *colour vision deficiency* (CVD,
98 Simunovic, 2010) (hereafter even referred to as *colour blindness*), should se-
99 riously be taken into account. In fact, the impossibility to see some of the

100 displayed colours by some people does not allow them to appreciate the dif-
101 ferences between minima and maxima on a map. Furthermore, this might
102 seriously impact scholars' and students' scientific learning affected by such
103 deficiencies having no proper access to graphing parts of articles and/or books
104 (Albany-Ward & Sobande, 2015).

105 Colour blind deficiencies are basically represented by three main colour
106 misperceptions: (i) protanopia, i.e., the inability to perceive the red color
107 (560 nm); (ii) deuteranopia, i.e., the inability to recognize the green color
108 (530 nm); (iii) tritanopia, i.e., the inability to distinguish the blue color (420
109 nm, Figure 1, Viénot et al., 1995; Gordon, 1998; Gegenfurtner, 2003). Box
110 1 explains every deficiency type in detail. This leads to the impossibility of
111 recognizing some types of colour ramps in which there is a gradient from blue
112 to green/yellow to red. For instance, this is the case of frequently (mis)used
113 rainbow colour palettes (Golebiowska & Coltekin, 2020; Stoelzle et al., 2021),
114 which are part of a bunch of papers in the scientific literature (e.g., Mesgaran
115 et al., 2014; Gardner et al., 2019; Ellis-Soto et al., 2021; Feilhauer et al., 2021;
116 Rocchini et al., 2021).

117 Inherited colour blindness affects more than 5% of human population
118 (i.e., 8% of males and 0.5% of females, being inherited as X-linked reces-
119 sive disease) mainly because of founder events and genetic drift (Simunovic,
120 2010; Birch, 2012). There is a wide literature in different fields of research
121 presenting problems for colour blind people: from the hampering of medi-
122 cal profession (Spalding, 1999) to students proper learning (Ramachandran,
123 2014), and from road accidents (Cole et al., 2002) until unintentional injuries
124 (Cumberland et al., 2004). On the contrary, this phenomenon is somewhat
125 disregarded in scientific rendering in form of maps. In order to implement
126 routines to help colour blind people, website examples exist to (i) choose
127 colour ramp palettes (seaborn: [https://seaborn.pydata.org/tutorial/
128 color_palettes.html](https://seaborn.pydata.org/tutorial/color_palettes.html), ColorBrewer: <https://colorbrewer2.org/>, Har-
129 rower & Brewer, 2003) or (ii) create them (colorshemesdesigner:
130 <https://paletton.com/#uid=1000u0k1111aFw0g0qFqFg0w0aF>) with an on-
131 line platform dedicated to the creation of colour schemes. However, they seem
132 to be too much complicated, in our opinion, for colour blind people.

133 Furthermore, packages in R are devoted to test for colour blindness
134 (`colorblindcheck` package, Nowosad, 2021) or to make use of colour ramp
135 palettes for colour blind people (`viridis`, Garnier et al., 2021). However,
136 as far as we know, no analytical and straightforward function exists that
137 allows to input an image and plot it for colour blind people simply and
138 straightforwardly. This paper aims to present a package `cblindplot` and its
139 main function - `cblind.plot()` - which enables colour blind people to just
140 enter an image in a coding workflow, specify their colour blind deficiency

141 type, and immediately get a colour blind friendly plot as output.

142 **2 The function: step by step**

143 **2.1 Code**

144 We implemented our idea in the `cblindplot` package (<https://github.com/ducciorocchini/cblindplot>). Its main function `cblind.plot()` updates
145 the color palette on an input image and returns a new visualization along
146 with a new color legend. The function has four basic steps. In the first step,
147 an image is imported into R. Then, a Principal Component Analysis (PCA)
148 is performed on the input image, and only the first principal component is
149 extracted. In the third step, a new color palette is applied to the single
150 dimension derived from PCA. Lastly, the plot with a meaningful legend for
151 color blind people is returned.
152

153 The user is asked to just choose the input image and a type of color vision
154 deficiency: "protanopia", "deuteranopia", or "tritanopia":

```
155 cblind.plot <- function(im,  
156                          cvd = c("protanopia", "deuteranopia", "tritanopia"),  
157                          r = 1, g = 2, b = 3,  
158                          crop_manual = FALSE,  
159                          select_class = FALSE
```

160 The `cvd` is a meaningful argument directly related to the type of colour
161 blindness (Box 1 and Figure 1): `cvd=c("protanopia", "deuteranopia",`
162 `"tritanopia")`. They will directly link to the `cividis`, `viridis` and `magma`,
163 respectively, palettes that are going to enhance the main colours seen by peo-
164 ple affected by such deficiency types. We chose such colour schemes follow-
165 ing Viénot et al. (1995), who developed a simulation of the reduced colour
166 gamut of colour defective people, explicitly considering the three aforemen-
167 tioned colour blind deficiency types (see also Box 1 and Figure 1). The
168 `cblind.plot()` function also allows to change the order of the RGB bands
169 (by default, the first image band relates to the red colour, the second to the
170 green colour, and the third to the blue colour), manually crop the input image
171 (the `crop_manual()` argument), and the `select_class()` makes it possible
172 to select only certain colours in the image for the further processing.

173 The complete code of the `cblindplot` package is available at [https://](https://github.com/ducciorocchini/cblindplot)
174 github.com/ducciorocchini/cblindplot, while we provide an empirical
175 example of its application in the next section.

176 2.2 Empirical example

177 In order to test the function we decided to directly get in the game by making
178 use of a (wrong) plot that we published in Rocchini et al. (2021), where
179 a rainbow colour palette was used to show the variability in space of the
180 Similaun Glacier (Italy, Figure 2). In that case, a rainbow colour ramp
181 palette was used passing from: (i) low spatial variability, represented by blue
182 and green, related to a small lake at north-east and snow at north-west,
183 respectively; (ii) medium variability (yellow) related to woodlands and high
184 elevation grasslands; and (iii) high variability (red), related to crevasses and
185 cracks of the calcareous rock composing the glacier. All of this description
186 makes no sense for colour blind people.

187 Hence, the issue can be solved in a straightforward manner by storing a
188 screenshot of the image and import it in R by the `cblind.plot` function.
189 No previous import is required but a direct call is done into the function
190 under the argument `im`. The three bands composing the RGB of the image
191 are generally mounted as: red in the first band, green in the second, and
192 blue in the third one; this is the order used by the `cblind.plot` function to
193 import them by default, which can be changed at any time. As previously
194 stated, the output is straightforward once the user declares her/his deficiency
195 type. For protanopia (Box 1 and Figure 1) a `viridis` colour ramp palette
196 from the `viridis` package is applied to enhance the contrast between blue
197 and yellow which can easily be seen by people affected by such a deficiency,
198 for deuteranopia a `cividis` colour ramp palette is used to smoothly pass
199 from blue to green to yellow, all colours that can easily be seen, and for
200 tritanopia a `magma` colour ramp palette is used to avoid the use of pure
201 blue. As previously stated, this is in line with the simulation screen figure of
202 the 'Jardin des Plantes' (photo: Jean Le Rohellec; Grande Galerie, FNAC)
203 seen by colour-blind people provided by Viénot et al. (1995), associating a
204 smooth blue-to-yellow colour ramp to protanopia-affected people, sharp blue-
205 to-yellow colour ramp to deuteranopia-affected people, to deep blue-to-yellow
206 colour ramp to tritanopia-affected people. We suggest the reader to compare
207 colours used in Figure 2 for the different deficiency types with that published
208 by Viénot et al. (1995)¹. Following Figure 1 for protanopia and deuteranopia
209 the colour ramp palettes can be used interchangeably (Rigden, 1999).

¹Notice that in Viénot et al. (1995) the images for the different deficiency types are arranged in a clockwise manner, hence they relate to panels a), b), d) and c) in Figure 2

210 3 Outlook and take home message

211 Providing for the possibility to appreciate coloured maps to everybody by
212 a single function is priceless in our opinion. Doing that in a free and open
213 source environment is mandatory. In fact, open source software would allow
214 the possibility to exactly reproduce analysis by guaranteeing high robustness
215 (Rocchini & Neteler, 2012). Apart for the previously cited `viridis` package,
216 examples exist for oceanography, with the `cmocean` package in R (Thyng et
217 al., 2016, 2020). Additional examples exist making use of Python () lan-
218 guage (Nuñez et al., 2018). However, in such cases, the workflow include
219 a separate import of the image and the application of a set of colour ramp
220 palettes (Garnier et al., 2021). Furthermore, albeit being well documented,
221 there are many steps to be done before plotting an image with a colour blind
222 friendly ramp palette. The function described in this paper (`cblind.plot`),
223 on the contrary, allows to (i) automatically call the image inside the function
224 without any initial import step and (ii) explicitly refer to the colour blind
225 deficiency type being experienced, to further automatically apply the proper
226 colour ramp palette.

227 Additional R packages have been devoted to improve mapping (and colours)
228 of various scientifically sound response variables like biodiversity (Féret & de
229 Boissieu, 2020) or species distributions (Schuetz et al., 2020a,b). Connecting
230 the `cblind.plot` with such packages would be an enormous advantage for
231 color blind people. Hence, the authors of the present paper have contacted
232 the developers of such packages to implement such function in their packages.

233 How the magnitude of spatial variation of data is represented in a map
234 has a high impact on the perception of the main processes shaping it, since
235 different colour signals are processed differently by the human visual system
236 (Rogowitz et al., 1996). From this point of view monotonic palettes based on
237 a monotonic continuous gradient (sensu Stevens, 1966) - like those used in the
238 presented function `cblind.plot` from the `viridis` R package - rather than
239 on abrupt thresholds - like the rainbow colour ramp palette (Borland & Ii,
240 2007) - must be preferred not only referred to colour blindness problems, but
241 also for the common perception of spatial variability. In fact, this is generally
242 based on a monotonic increase in eye stimulus intensity, as well established
243 by pioneering papers on psychological perception of colour variation (Ekman
244 & Sjöberg , 1965; Panek & Stevens, 1966).

245 Based on previous observation, the `cblind.plot` function guarantees to
246 avoid perceptual discontinuities and thus the appearance of false spatial fea-
247 tures (Kovesi, 2015) - in most cases related to noise in the original im-
248 ages/maps - being based on a continuous monotonic colour ramp, i.e. on
249 an incremental and uniform change of colours over the whole output im-

250 age/map. Although one cannot be certain about the real vision of someone
251 else, the provided function is providing a straightforward tool based on well
252 established previous simulations on colour vision about the residual colour
253 information coming out from the above described deficiency types (Brettel,
254 1997). This is just based on the fact that a certain stimulus is perceived dif-
255 ferently in colour blind deficiencies versus normal vision, with a well known
256 reaction from every single deficiency type considered in this paper. That is
257 why we avoided, at the time being, a test with colour blind people, relying
258 rather on medical papers on the matter (Capilla, 2004).

259 Beside problems related to colour blind deficiencies, the rainbow colour
260 map has widely been acknowledged as a fundamental problem-full palette
261 for plotting scalar values to colours. This is mainly because it introduces
262 artifacts and obscure some data by e.g. putting yellow colours as mid values,
263 by finally confusing any user, despite her/his ability to properly see colours,
264 if there is any proper manner to see colours (Moreland, 2009).

265 Colour gamut is of primary importance to synthesise the information
266 contained in a dataset/map. Summarising, there are three main important
267 elements/steps when applying a colour gamut to data rendering: (i) the pos-
268 sibility of using different colour palettes in a software, (ii) the choice made
269 by the map producer about one of the possible palettes, and (iii) how such a
270 choice is perceived by users. From this point of view, awareness of the uncer-
271 tainty and potential limitations of the use of colours in maps may enhance
272 map interpretation and use. While colour-coding guidelines are needed, we
273 are far from having an etiquette on this theme, especially when considering
274 continuous pseudocolour maps, i.e. maps coming out from analytical pro-
275 cesses on geographical data (Reda et al., 2018). The `cblind.plot` function
276 partially solves this issue by providing a direct manner to rescale pseudo-
277 colour values and plot them in a colour-blind free scheme, which is robustly
278 grounded on optical theory. One of the main strengths of the function is that
279 users are not required to exit their analysis process and enter e.g. internet
280 sites but they can integrate the function in their throughput code to properly
281 see output results.

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398 **Box 1 - Colour blind deficiency types**

399 ▶ **Protanopia:** Inability to perceive the red color. Due to genetic
400 mutations that cause the "L" type retinal cones to fail, or those pho-
401 toreceptors sensitive to large wavelengths (560 nm), which allow the
402 vision of the red color. It has a genetic transmission of the x-linked
403 recessive type, for which it largely affects the male sex.

404 ▶ **Deutanopia:** inability to recognize the green color (the most
405 common). Caused by genetic mutations that result in the failure of the
406 "M" cones, those photoreceptors sensitive to medium wavelengths (530
407 nm), that is, which allow the vision of the green color. It has a genetic,
408 x-linked recessive inheritance, for which it largely affects the male sex.

409 ▶ **Tritanopia:** inability to distinguish the blue color (the rarest).
410 Given by genetic mutations that cause the failure of the "S" cones,
411 photoreceptors sensitive to radiation of short wavelengths (420 nm), or
412 that allow the vision of the blue color. It has an autosomal dominant
413 genetic inheritance, therefore independent of sex.

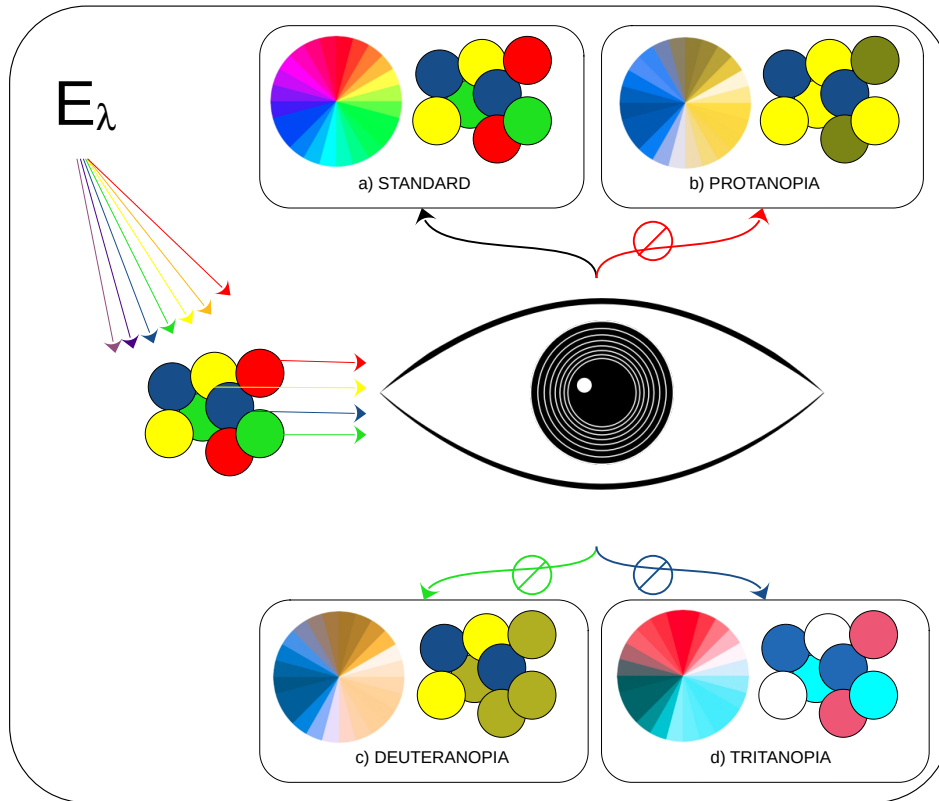


Figure 1: Different manners to perceive colours by the human eye: (a) standard vision; (b) protanopia, i.e., the inability to perceive the red color (560 nm); (c) deuteranopia, i.e., the inability to recognize the green color (530 nm); (d) tritanopia, i.e., the inability to distinguish the blue color (420 nm). Refer to Box 1 for specific information.

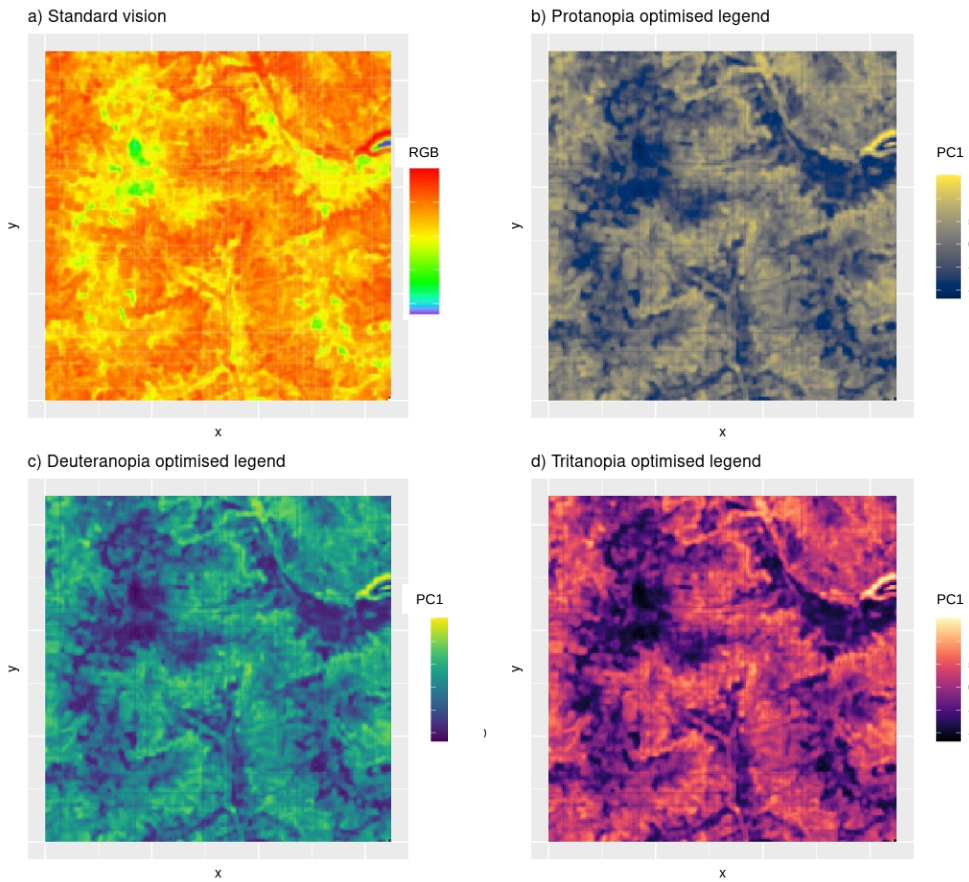


Figure 2: The variability in space of the Similaun Glacier shown by a rainbow colour ramp palette (a) as in Rocchini et al. (2021). Low spatial variability is represented in blue and green (lake), while medium variability is in yellow (woodlands), high variability is in red (crevasses and cracks). Colour blind people cannot ascertain differences between minima (blue) and maxima (red). Hence, different colour ramps are applied considering the different diseases, according to previous tests by Viénot et al. (1995): a smooth blue-to-yellow colour ramp to protanopia-affected people (`cividis` colour ramp in the `viridis` package, sharp blue-to-yellow colour ramp to deuteranopia-affected people (`viridis` colour ramp), to deep blue-to-yellow colour ramp (`magma` colour ramp) to tritanopia-affected people.