

2 **Initial fixation placement in face images is driven**
3 **by top-down guidance**

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7 **Abstract** The eyes are often inspected first and for longer
8 period during face exploration. To examine whether this
9 saliency of the eye region at the early stage of face inspec-
10 tion is attributed to its local structure properties or to the
11 knowledge of its essence in facial communication, in this
12 study we investigated the pattern of eye movements pro-
13 duced by rhesus monkeys (*Macaca mulatta*) as they free
14 viewed images of monkey faces. Eye positions were
15 recorded accurately using implanted eye coils, while
16 images of original faces, faces with scrambled eyes, and
17 scrambled faces except for the eyes were presented on a
18 computer screen. The eye region in the scrambled faces
19 attracted the same proportion of viewing time and fixations
20 as it did in the original faces, even the scrambled eyes
21 attracted substantial proportion of viewing time and fixa-
22 tions. Furthermore, the monkeys often made the first sac-
23 cade towards to the location of the eyes regardless of image
24 content. Our results suggest that the initial fixation place-
25 ment in faces is driven predominantly by “top-down” or
26 internal factors, such as the prior knowledge of the location
27 of “eyes” within the context of a face.

28 **Keywords** Eye movements · Faces · Eyes · Monkey

29 **Introduction**


30 Visual exploration of the world around us involves a series
31 of saccadic eye movements and fixations, and we tend to

concentrate our fixations on interesting and informative 32
regions in the scene (Yarbus 1967). The choice of the 33
potential fixation targets can be driven by both bottom-up 34
exogenous or external factors and top-down endogenous or 35
internal factors. External factors are image immanent fea- 36
tures, such as local image contrast and local image struc- 37
ture, which transiently attract eye gaze, independent of a 38
particular task. Internal factors, such as an individual’s 39
attentional state, expectation, experience and memory, are 40
top-down and task-dependent (Noton and Stark 1971; 41
Mannan et al. 1997; Henderson 2003). It is argued that in 42
general, the initial saccade to an image is driven predomi- 43
nantly by external factors (Parkhurst et al. 2002; Peters 44
et al. 2005), but can also be biased by internal factors (Hen- 45
derson 2003). 46

As faces can provide visual information about an individ- 47
ual’s gender, age and familiarity, and their expressions pro- 48
vide significant cues to intention and mental state, the 49
ability to recognize these cues and to respond accordingly 50
plays a crucial role in our social communication (Bruce and 51
Young 1998; Emery 2000). Just like humans, rhesus mon- 52
keys are sensitive to faces of conspecifics. They are able to 53
discriminate faces of unfamiliar individuals after only a 54
short exposure to sets of their images (Parr et al. 2000). 55
Viewing of faces is accompanied by longer fixations com- 56
pared with natural scenes (Guo et al. 2006), and is typically 57
associated with a stereotypical eye scanning patterns (Keating 58
and Keating 1982; Nahm et al. 1997; Guo et al. 2003; 59
Gothard et al. 2004; Ghazanfar et al. 2006). Specifically, the 60
eye region in neutral, expressive or vocalizing faces is often 61
the first destination of the saccade and attracts a dispropor- 62
tionate share of fixations compared with other local facial 63
features, suggesting its dominant saliency in the faces. 64

However, it remains unclear whether this interest in 65
eyes, especially at the earliest stage of face exploration, is 66

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67 attributable solely to its local structure properties, or may
 68 derive from the knowledge/memory of its location and
 69 essence in facial communication. To address this question,
 70 in this experiment we systemically manipulated the local
 71 image structures of inner face components (i.e. eye region),
 72 and compared rhesus monkeys' eye scanning patterns when
 73 viewing original monkey face and modified face images.
 74 Our results suggested that the top-down guidance (i.e. prior
 75 knowledge of the location of the eyes in the faces) plays a
 76 crucial role in the saliency of the eye region during early
 77 stage of face exploration.

78 Methods

79 Subjects

80 Two male adult rhesus monkeys (*Macaca mulatta*, 4.5–
 81 6.0 kg) were used in this study. Initially they were trained
 82 to fixate a spot on a computer screen for several seconds in
 83 a dimming fixation detection task (Guo et al. 2003). For the
 84 purpose of recording eye movements, a scleral eye coil and
 85 head restraint were then implanted under aseptic condi-
 86 tions. Throughout the period of the recordings, the animal's
 87 weight and general health were monitored daily. All proce-
 88 dures complied with the "Principles of laboratory animal
 89 care" (NIH publication no. 86-23, revised 1985) and UK
 90 Home Office regulations.

91 Stimuli and apparatus

92 Digitized grey scale images were presented through a VSG
 93 2/3 graphics system (Cambridge Research Systems) and
 94 displayed on a high frequency non-interlaced gamma-cor-
 95 rected color monitor (110 Hz frame rate, Sony GDM-
 96 F500T9) with the resolution of 1,024 × 768 pixels. At a
 97 viewing distance of 57 cm the monitor subtended a visual
 98 angle of 40° × 30°. The mean luminance of uniform grey
 99 background was kept at 6.0 cd/m².

100 Twenty neutral monkey (*Macaca mulatta*) face images
 101 were used as stimuli. All images (512 × 512 pixels, 256
 102 grey-levels) were gamma-corrected. For each original face

103 image, we created two scrambled versions (scrambling eye
 104 region only, scrambling whole face except for eye region)
 105 with the same first- and second-order statistics (image prop-
 106 erties determined by the amplitudes of the Fourier spec-
 107 trum) but different higher-order correlations (image
 108 properties determined by the phases of the Fourier spec-
 109 trum). This was done by computing the Fourier transform
 110 over the scrambled facial features and randomizing the
 111 phase spectrum (0–2π) in the frequency domain. The
 112 Fourier amplitude spectrum of the images was not affected
 113 by this procedure. Without higher-order statistical structures
 114 corresponding to the sparse distributions of local features,
 115 these scrambled image regions lack any visual objects and
 116 have a cloud-like appearance (see Fig. 1 for examples),
 117 although they have the same mean luminance and root-mean-
 118 square contrast as the corresponding facial features (Guo
 119 et al. 2005).

120 In total, three different classes of images were presented
 121 to monkeys: (1) 20 original face images, (2) 20 face images
 122 with scrambled eyes (eyes scrambled); (3) 20 scrambled
 123 faces except for eye regions (eyes only). All images were
 124 displayed once in a random order at the center of the screen
 125 with a resolution of 512 × 512 pixels (20° × 20°).

126 During the experiments the monkey was seated in a pur-
 127 pose-built primate chair with head restrained, and viewed
 128 the display binocularly. To calibrate eye movement signals,
 129 a small red fixation point (FP) (0.2° diameter, 7.8 cd/m²
 130 luminance) was displayed randomly at 1 of 25 positions
 131 (5 × 5 matrix) across the monitor. The distance between
 132 adjacent FP positions was 5°. The monkey was trained to
 133 follow the FP and maintain fixation for 1 s. After the cali-
 134 bration procedure, the trial was started with a FP displayed
 135 on the center of monitor. If the monkey maintained fixation
 136 for 500 ms, the FP disappeared and a face image was pre-
 137 sented for 10 s. During the presentation, the monkeys pas-
 138 sively viewed the images. No reinforcement was given
 139 during this procedure, neither were the animals trained on
 140 any other task with these stimuli, which could have poten-
 141 tially affected the structure of their behaviour. It was con-
 142 sidered that with their lack of training, and in the absence of
 143 instrumental responding, their behavior should be as natural
 144 as possible.

Fig. 1 Examples of static grey scale monkey face images used in the recording. From left to right: original face image, face image with scrambled eyes, scrambled face except for eye region



145 Eye movement recordings and analysis

146 Horizontal and vertical eye positions were measured using
 147 an 18-in. cubic scleral search coil assembly with 6 min arc
 148 sensitivity (CNC Engineering). Eye movement signals were
 149 amplified and sampled at 500 Hz through CED1401 plus
 150 digital interface (Cambridge Electronic Design). The soft-
 151 ware developed in Matlab computed horizontal and vertical
 152 eye displacement signals as a function of time to determine
 153 eye velocity and position. Fixation locations and durations
 154 were then extracted from the raw eye tracking data using
 155 velocity (less than 0.2° eye displacement at a velocity of
 156 less than $20^\circ/\text{s}$) and duration (greater than 50 ms) criteria
 157 (Guo et al. 2006).

158 **Results**

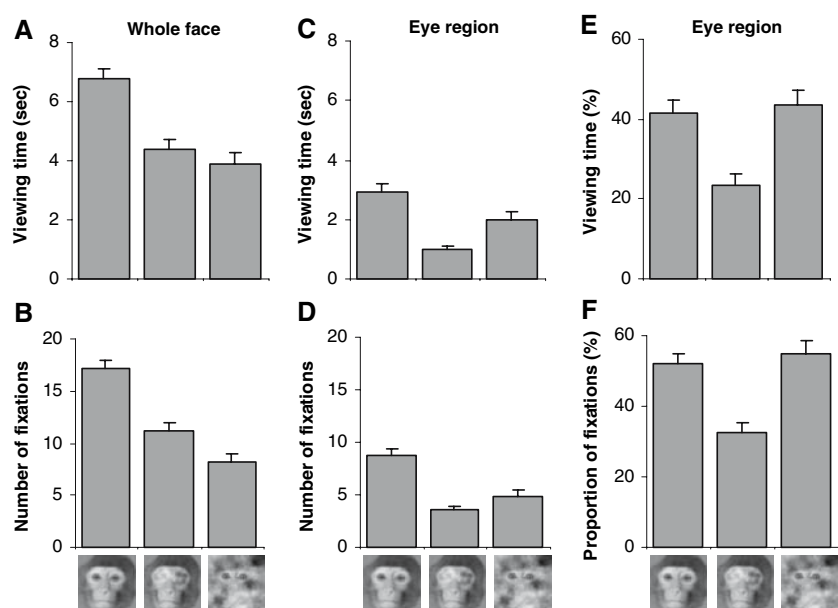
159 Not surprisingly, the original face images were the most
 160 salient to the monkeys. They attracted longer viewing time
 161 (1-way ANOVA, $F_{(2,129)} = 22.36$, $P = 4.6\text{E} - 9$) and more
 162 fixations (1-way ANOVA, $F_{(2,129)} = 30.01$, $P = 1.98\text{E} - 11$)
 163 than the modified face images (Fig. 2a, b). The two mon-
 164 keys spent $68 \pm 3\%$ (mean \pm SEM) of the 10-s image pre-
 165 sentation time viewing the original faces, making
 166 17.11 ± 0.9 fixations across the images. The proportion of
 167 time spent viewing the image decreased to 44 ± 3 and
 168 $39 \pm 4\%$, and the number of fixations declined to
 169 11.25 ± 0.78 and 8.2 ± 0.8 for the eyes scrambled and eyes
 170 only images.

171 Among local facial features, the eye region in original
 172 faces often receives the highest proportion of fixations dur-
 173 ing face exploration (Guo et al. 2003). In this experiment,

174 the eye region in eyes scrambled and eyes only images still
 175 attracted a substantial amount of attention, although the
 176 cumulative viewing time (Fig. 2c) and the number of fixa-
 177 tions (Fig. 2d) were decreased in these two conditions
 178 (viewing time: original face 2.91 ± 0.3 s, eyes scrambled
 179 1 ± 0.13 s, eyes only 1.96 ± 0.32 s, 1-way ANOVA,
 180 $F_{(2,129)} = 12.81$, $P = 8.44\text{E} - 6$; number of fixations: origi-
 181 nal face 8.73 ± 0.65 , eyes scrambled 3.59 ± 0.38 , eyes
 182 only 4.89 ± 0.57 , 1-way ANOVA, $F_{(2,129)} = 24.11$,
 183 $P = 1.27\text{E} - 9$). When the same data in Fig. 2c, d was
 184 expressed as the percentage of face viewing time (Fig. 2E)
 185 and as the proportion of the number of fixations within the
 186 images (Fig. 2f), the eye region in original face and eyes
 187 only images received the same proportion of face viewing
 188 time and fixations (viewing time: original face $41 \pm 3\%$,
 189 eyes only $43 \pm 4\%$, Tukey's least significant procedure,
 190 $P = 0.6$; fixations: original face $52 \pm 3\%$, eyes only $55 \pm 4\%$,
 191 Tukey's least significant procedure, $P = 0.47$). The unrec-
 192 ognisable eyes in eyes scrambled face images attracted less
 193 proportion of face viewing time ($23 \pm 3\%$, 1-way ANOVA,
 194 $F_{(2,129)} = 12.15$, $P = 1.46\text{E} - 5$) and fixations ($32 \pm 3\%$,
 195 1-way ANOVA, $F_{(2,129)} = 14.66$, $P = 1.83\text{E} - 6$).

196 To examine whether there were any differences in the
 197 spatial distribution of sequential fixation placement during
 198 image exploration, we compared the first five fixation
 199 placements in each image (this number was chosen as it
 200 represented the maximum number of saccades for some
 201 images). The probability of fixation placement in the eye
 202 region as a function of fixation sequence is plotted in Fig. 3.
 203 The eyes had a very higher probability as the first saccade
 204 destination ($>90\%$) once the image was presented, even
 205 when they were unrecognisable in the eyes scrambled
 206 images. For the next four saccades, they had the same

Fig. 2 a, b Cumulative viewing time and number of fixations within original face, eyes scrambled and eyes only images. c, d Cumulative viewing time and number of fixations for the eye region within original face, eyes scrambled and eyes only images. e, f Proportion of cumulative face viewing time and number of fixations for the eye region within original face, eyes scrambled and eyes only images. Error bars indicate standard error of mean



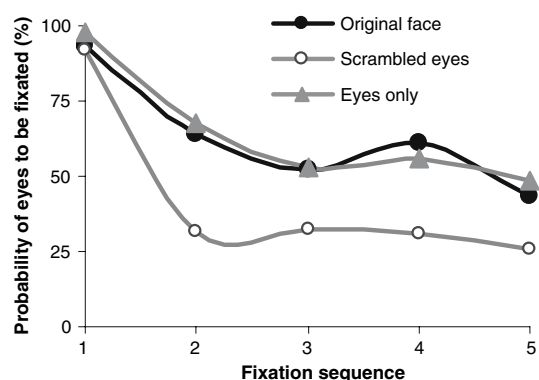


Fig. 3 The probability of the eye region as the destination of first five saccades measured while viewing original face, eyes scrambled and eyes only images

207 probability to be fixated in the original face and eyes only
 208 images (Kolmogorov–Smirnov test, $P > 0.05$), but much
 209 less chance to be inspected in the eyes scrambled images
 210 (Kolmogorov–Smirnov test, $P < 0.05$).

211 Discussion

212 Faces are probably the most important visual stimuli in pri-
 213 mate social communications (Bruce and Young 1998). The
 214 saliency of the face, however, is dependent on appropriate
 215 facial configurations. Disruptions such as inverting faces or
 216 randomly rearranging local facial components would
 217 reduce the amount of viewing time and fixations directed to
 218 the faces (Guo et al. 2003). Here we further observed that
 219 the saliency of the face was decreased with the manipula-
 220 tion of local facial structures by scrambling eyes or non-eye
 221 regions (Fig. 2a, b), suggesting that the selection of the face
 222 as a target for fixation would depend on the prior knowl-
 223 edge concerning the likelihood of the occurrence of the
 224 faces (Carpenter and Williams 1995), such as faces pre-
 225 sented in a given orientation and within a given context.

226 Among local facial components, eye region is the most
 227 attended feature. During face exploration, both human and
 228 non-human primates demonstrated an exaggerated interest
 229 in the eye region of the faces of conspecifics (Yarbus 1967;
 230 Keating and Keating 1982; Nahm et al. 1997; Guo et al.
 231 2003; Gothard et al. 2004; Ghazanfar et al. 2006). This
 232 preferential interest in the eyes remained when the eyes or
 233 the rest of the facial structures were scrambled. The eye
 234 region in the scrambled faces attracted the same proportion
 235 of viewing time and fixations as it did in the original faces,
 236 even the unrecognisable eyes in the eyes scrambled images
 237 attracted substantial proportion of viewing time (~23%)
 238 and fixations (~32%, Fig. 2e, f). Taken together, it seems
 239 that both the intrinsic structure (e.g. local contrast or local
 240 edges) of the eye region and the knowledge of its location

241 and essence in facial communication contribute to its
 242 saliency during face exploration. However, the declined
 243 cumulative viewing time and fixation numbers towards the
 244 eye region in the eyes scrambled and eyes only images
 245 (Fig. 2c, d) suggest that the eyes are better processed in
 246 concert with other facial features.

247 The eye region is often the first fixation target following
 248 the appearance of the faces (Guo et al. 2003). It is argued
 249 that in general, the initial saccade to an image is driven pre-
 250 dominantly by “bottom-up” process or external factors
 251 such as local image contrast (Parkhurst et al. 2002; Peters
 252 et al. 2005). From this perspective, the saliency of the eye
 253 region at the earliest stage of face inspection could be
 254 attributed to its local structure properties (i.e. the eye region
 255 has relatively higher local contrast in grey scale images).
 256 However, in our test condition of face images with scram-
 257 bled eyes, the eye region was also inspected first even when
 258 it was unrecognisable (Fig. 3), suggesting that the visual
 259 system may retain prior knowledge of the location of
 260 “eyes” within the context of a face from past experience,
 261 and this knowledge could bias the destination of the initial
 262 saccade. In addition, when the face images were inverted or
 263 the position of the eyes were rearranged within the faces,
 264 the time into the trial for the first saccade directed at the
 265 eyes was significantly delayed, indicating the first saccade
 266 within the image was not directed at the eyes although their
 267 local image properties (contrast and structure) were unal-
 268 tered (Guo et al. 2003). Taken together, it seems the initial
 269 fixation placement in a face image is driven predominantly
 270 by “top-down” guidance or internal factors, in particularly
 271 the prior knowledge of the location of “eyes” within the
 272 context of a normal face. Furthermore, it could be the glo-
 273 bal semantic characteristics of the faces that determine the
 274 initial fixation placement rather than the local semantic
 275 characteristics of the eyes.


276 Our results are consistent with previous behavioural,
 277 psychophysical, neurophysiological and neuroimaging
 278 studies on the role of the eyes in social interaction in
 279 humans and non-human primates. The eyes are one of the
 280 first points of contact between infants and mothers, and
 281 play a pivotal role in identity recognition and emotional
 282 communication (Bruce and Young 1998). They often
 283 provide “early warning signals” for rapid assessment and
 284 response to salient and potential harmful events (i.e.
 285 through the process of joint attention), hence may capture
 286 attention involuntarily (Langton et al. 2000; Rauschenberger
 287 2003). While presented alone, the eyes can selectively
 288 activate neurons in superior temporal sulcus and amygdala,
 289 sometimes with the same response amplitude as the presen-
 290 tation of whole face (Emery 2000; Ghazanfar and Santos
 291 2004). Furthermore, the observation that the eyes do not
 292 carry the same relevance for human and monkey infants as
 293 human and monkey adults (Thomsen 1974; Farroni et al.

294 2002) suggests that the sensitivity to the eyes is a learnt
 295 mechanism. Given these considerations, it is reasonable to
 296 assume that the knowledge of the location of eye region
 297 within a face and its social relevance contribute signifi-
 298 cantly to its saliency at the earliest stage of face exploration
 299 even without specific task demands.

300 References

301 Bruce V, Young A (1998) In the eye of the beholder. Oxford Univer-
 302 sity Press, New York
 303 Carpenter RH, Williams ML (1995) Neural computation of log likeli-
 304 hood in control of saccadic eye movements. *Nature* 377:59–62
 305 Emery NJ (2000) The eyes have it: the neuroethology, function and
 306 evolution of social gaze. *Neurosci Biobehav Rev* 24:581–604
 307 Farroni T, Csibra G, Simion F, Johnson MH (2002) Eye contact detec-
 308 tion in human from birth. *Proc Natl Acad Sci USA* 99:9602–9605
 309 Ghazanfar AA, Santos LR (2004) Primate brains in the wild: the sen-
 310 sory bases for social interactions. *Nat Rev Neurosci* 5:603–616
 311 Ghazanfar AA, Nielsen K, Logothetis NK (2006) Eye movements of
 312 monkey observers viewing vocalizing conspecifics. *Cognition*
 313 101:515–529
 314 Gothard KM, Erickson CA, Amaral DG (2004) How do rhesus mon-
 315 keys (*Macaca mulatta*) scan faces in a visual paired comparison
 316 task? *Anim Cogn* 7:25–36
 317 Guo K, Robertson RG, Mahmoodi S, Tadmor Y, Young MP (2003)
 318 How do monkeys view faces?—a study of eye movements. *Exp*
 319 *Brain Res* 150:363–374

Guo K, Robertson RG, Mahmoodi S, Young MP (2005) Centre-sur-
 320 round interactions in response to natural scene stimulation in the
 321 primary visual cortex. *Eur J Neurosci* 21:536–548 322
 Guo K, Mahmoodi S, Robertson RG, Young MP (2006) Longer fixation
 323 duration while viewing face images. *Exp Brain Res* 171:91–98 324
 Henderson JM (2003) Human gaze control during real-world scene
 325 perception. *Trends Cog Sci* 7:498–504 326
 Keating CF, Keating EG (1982) Visual scan patterns of rhesus mon-
 327 keys viewing faces. *Perception* 11:211–219 328
 Langton SR, Watt RJ, Bruce II (2000) Do the eyes have it? Cues to the
 329 direction of social attention. *Trends Cogn Sci* 4:50–59 330
 Mannan SK, Ruddock KH, Woodings DS (1997) The relationship be-
 331 tween the locations of spatial features and those of fixations made
 332 during visual examination of briefly presented images. *Spatial Vis*
 333 10:165–188 334
 Nahm FKD, Perret A, Amaral DG, Albright TD (1997) How do mon-
 335 keys look at faces? *J Cogn Neurosci* 9:611–623 336
 Noton D, Stark L (1971) Scanpaths in saccadic eye movements while
 337 viewing and recognizing patterns. *Vis Res* 11:929–942 338
 Parkhurst D, Law K, Niebur E (2002) Modelling the role of salience in
 339 the allocation of overt visual attention. *Vis Res* 42:107–123 340
 Parr LA, Winslow JT, Hopkins WD (2000) Recognizing facial cues:
 341 individual discrimination by chimpanzees (*Pan troglodytes*) and
 342 rhesus monkeys (*Macaca mulatta*). *J Comp Psychol* 114:1–14 343
 Peters RJ, Iver A, Itti L, Koch C (2005) Components of bottom-up
 344 gaze allocation in natural images. *Vision Res* 45:2397–2416 345
 Rauschenberger R (2003) Attentional capture by auto- and allo-cues.
 346 *Psychon Bull Rev* 10:814–842 347
 Thomsen CE (1974) Eye contact by non-human primates toward a hu-
 348 man observer. *Anim Behav* 22:144–149 349
 Yarbus A (1967) Eye movements and vision. Plenum, New York 350

	221	1038	xxxx	Dispatch: 26.6.07		No. of Pages: 5	
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