Dramatic colour changes in a bird of paradise caused by uniquely structured breast feather barbules

Doekele G. Stavenga1, Hein L. Leertouwer1, N. Justin Marshall2
and Daniel Osorio3,*

1Department of Neurobiophysics, University of Groningen, 9747 AG, Groningen, The Netherlands
2Queensland Brain Institute, The University of Queensland, Saint Lucia, Queensland 4072, Australia
3School of Life Sciences, University of Sussex, Brighton BN1 9QG, UK

The breast-plate plumage of male Lawes’ parotia (Parotia lawesii) produces dramatic colour changes when this bird of paradise displays on its forest-floor lek. We show that this effect is achieved not solely by the iridescence—that is an angular-dependent spectral shift of the reflected light—which is inherent in structural coloration, but is based on a unique anatomical modification of the breast-feather barbule. The barbules have a segmental structure, and in common with many other iridescent feathers, they contain stacked melanin rodlets surrounded by a keratin film. The unique property of the parotia barbules is their boomerang-like cross section. This allows each barbule to work as three coloured mirrors: a yellow-orange reflector in the plane of the feather, and two symmetrically positioned bluish reflectors at respective angles of about 30°. Movement during the parotia’s courtship displays thereby achieves much larger and more abrupt colour changes than is possible with ordinary iridescent plumage. To our knowledge, this is the first example of multiple thin film or multi-layer reflectors incorporated in a single structure (engineered or biological). It nicely illustrates how subtle modification of the basic feather structure can achieve novel visual effects. The fact that the parotia’s breast feathers seem to be specifically adapted to give much stronger colour changes than normal structural coloration implies that colour change is important in their courtship display.

Keywords: Lawes’ parotia; iridescence; feather reflectance; multi-layers; thin film; scattering

1. INTRODUCTION

Bird feathers have two main mechanisms of structural coloration: either a spongy keratin matrix with air spaces or multi-layers created by regularly arranged melanin rodlets in a keratin matrix [1–3]. As air, keratin and melanin have different refractive indexes, light is reflected at the interfaces [4]. With keratin sponges, the spacing of the air–keratin interfaces is more or less independent of orientation, so that the reflection is diffuse, rather than directional [5–7]. This diffuse reflection means that the visual appearance of keratin sponges tends to resemble ordinary pigmented materials, but birds use them to create blues and greens that are difficult to make with plumage pigments [8]. Feathers with a thin-film cortex and/or regularly arranged melanin rodlets give directional (mirror-like) reflections, which are described by multi-layer theory [9–12]. Appropriate spacing and orientation of the elements produce intensely saturated colours and directional effects, which are not achieved by ordinary pigments [5,10]. Multi-layer reflectors are inherently iridescent—that is, the wavelength (approximately hue) of the reflected light varies with the viewing geometry [10].

The male bird of paradise Lawes’ parotia (Parotia lawesii; figure 1) displays brilliant iridescent breast feathers to attract females [13–15]. Iridescence emerges from the barbules, which, in common many other feathers, contain orderly arrays of melanin rodlets embedded in a keratin matrix [1,16]. However, whereas in cross-section barbules generally have a more or less flattened oval shape, sometimes with slight bends and turns (e.g. [1,17]), the parotia barbules have a unique boomerang-like shape [1,16]. Here, we show that the parotia breast feathers are (as far as we know) uniquely modified to greatly enhance the colour changes when the males perform their exhilarating ballerina dance on the forest floor [15]. Each feather barbule appears to incorporate three separate coloured mirrors that reflect light in different directions. This allows abrupt switches between yellow, blue and black. These observations exemplify how a modification of the basic feather anatomy can produce a new visual effect, and give insight into the evolution and function of structural coloration in avian displays.

2. MATERIAL AND METHODS

We photographed a male Lawes’ parotia from labelled skins in the collection of the Queensland Museum, Brisbane (figure 2a,b). The same museum provided four breast feathers for study with conventional light microscopy (epi-illumination as well as transmitted light) and with transmission electron microscopy. We examined the feathers with an imaging scatterometer ([18]; see also [19]), which yields images of the light reflected into a 180° hemisphere.
by locally illuminated feather barbules. Furthermore, we measured the feather reflectance spectra from different directions in a plane perpendicular to the barbule axis with a bifurcated fibre-optic probe. The instrument comprised six light guides, delivering light from a halogen-deuterium lamp to the feather. These illuminating light guides collectively surrounded a central fibre that acted as a collector of scattered light and which delivered it to a photodiode-array-spectrometer (Avantes, AvaSpec-2048-2). A white diffusing reflectance standard (Avantes WS-2) served as the reference. In addition, we measured the feather reflectance with an angular reflectance measurement set-up consisting of two fibres rotating, independently from each other, around the same axis in a plane containing the barbule axis [5]. One fibre acted as the light source, delivering light from a xenon lamp, and illuminated an area with a diameter approximately 4 mm; the other fibre captured the reflected light and delivered it to the spectrometer. The two fibres were symmetrically rotated with respect to the normal to the feather plane.

3. RESULTS

The breast-plate feathers of male Lawes’ parotia have a conventional structure; barbs emanate from a central rachis, and barbules emanate from the barbs. Each feather has a central region of black barbs, which is surrounded by a broad margin of brilliant coloured barbs (figure 2a,b). Viewed from above, the coloured barbules appear as a row of cushion-shaped segments (approx. 20 x 25 μm²; figure 2c–f). With the feather normal to the line of sight and illuminated with a co-axial beam (i.e. as if viewed face-on with the sun behind the viewer) the segments are coloured yellow-orange (figure 2d,e).

In transmitted light microscopy, the barbule segments are brown, as is typical of melanin pigmentation [5], and feature a prominent axial line (figure 2c), which is also seen in epi-illumination (figure 2d). High magnification (figure 2e) shows a longitudinal grating of dark lines separated by 0.5–0.6 μm. There are 12–15 lines over the 8 μm horizontal section arrowed in figure 2e. When the barbule is rotated, a bluish colour appears on one or other side of the barbule segments (figure 2f).

For further study of the optical properties of the feather, we used transmission electron microscopy. An approximately longitudinal section (figure 3a) shows a characteristic barbule, about 5 μm thick. The barbule has a clear (presumably) keratin cortex about 250–300 nm thick, which surrounds approximately 25 layers of pigmented rodlets, with a diameter approximately 60–90 nm, set in a keratin matrix. The pigment layers are separated by approximately 200 nm and are approximately perpendicular to the symmetry plane, but slightly curved. In the cross section, the barbule has a distinctive, boomerang-like shape: the upper surface consists of two flattish planes, width approximately 8 μm, separated by an angle of about 120°. About 15 rodlet planes terminate on the upper surface of each side of the barbule (figure 3b).

The ‘roof-line’ along the upper surface of the barbule segments (figure 3b) immediately explains the axial line seen on the barbules with light microscopy (figure 2c,d). Also, the number of the melanin layers abutting the sides of the upper surface in figure 3b corresponds with the number of the dark lines seen in the barbule segments with the light microscope (figure 2e). The implication is that the entire barbule surface appears yellow-orange owing to optical interference in the pigment layers within the barbule segments as pictured in the diagram of figure 3c. A light ray parallel to the symmetry plane incident at the 30° slanted top surfaces is refracted with an angle 18.7°, assuming that the dominant medium of the barbule segment is keratin with an effective refractive index of 1.56 (e.g. [11]). The refracted ray will hit the multi-layer inside the barbule normally when the multi-layer is tilted over an angle of 11.3°. The tilted multi-layers as seen in figure 3b may, therefore, function to counteract angular spreading of the reflected light.

In addition to the refraction, the incident light will be partially reflected at the surface. Apparently, the layering of the barbule cortex selectively favours the reflection of blue light, as was observed when rotating the barbules (figure 2f). With incident light in the symmetry plane, the direction of the reflected light beams from the two surface sides will then be −60° and +60° (figure 3c).

To investigate the feather’s optical properties further, we placed an isolated barb, glued to a pipette, in our imaging scatterometer [18] and illuminated an approximately 200 μm diameter spot (figure 4a) with a beam of white light normal to the feather surface (figure 4b). The diagram of figure 4b illustrates how the barb reflects light from the normal incident beam into a hemisphere (the red circles indicate reflection angles of 5°, 30°, 60° and 90°).

The distribution of scattered light projected into a polar coordinate system yields the far-field scattering pattern (figure 4c,d); the red circles in the polar plot represent the same reflection angles as the circles of figure 4b. With illumination normal to the barb ‘face’ (figure 4c), the barbules reflect yellow-orange light approximately along the direction of the illumination (as for a fronto-parallel reflector), while blue-green light is reflected in two directions about 60° to the normal direction (cf. figures 3c and 4b). Rotating the barb by about 10° results in an angular shift of the scattering pattern (figure 4d), and the colours of the two reflected side beams change in opposite spectral directions; that is, one side beam becomes greener, the other more violet. Similarly, if less obviously, the colour of the central, yellow-orange beam also changes, with long-wavelength (redder) or short-wavelength (greener) yellow coloration according to the direction of reflection (figure 4d). These observations are entirely consistent with the barb behaving as a set of three multi-layer mirrors (figure 4e,f; [5,10]), where a decrease (increase) in the angle of incidence shifts the reflection to longer (shorter) wavelengths.

The light reflected by the different mirrors is coloured quite differently. To investigate the spectra of the light reflected by the breast feathers of the parotia, we applied two different spectrophotometric methods. Firstly, we used a bifurcated fibre-optic probe, which measures a small area (diameter 1–2 mm). It gave a narrow-band spectrum with a peak at about 600 nm when the probe was normal to the feather surface (figure 5a). The peak moved to shorter wavelengths upon rotation of the feather around the barbule axis. A bluish reflectance, peaking at about 500 nm, emerged at an angle of about 30° (figure 5a). Secondly, we used a pair of optical fibres that could be rotated independently around the same
axis, one acting as the light source, the other as the light collector. The reflectance of a feather with the barbule symmetry axis perpendicular to the rotation axis was measured as a function of the angle of light incidence. The fibre delivering the light was rotated in steps of 10°. To obtain the maximal reflectance, the fibre capturing the reflected light had to be rotated over the same angle but in the opposite direction. The peak of the resulting spectra shifted progressively towards shorter wavelengths and the reflectance amplitude decreased with larger angles of incidence (figure 5b).

4. DISCUSSION

By incorporating multiple tuned mirrors in a single barbule, Lawes’ parotia’s breast feathers produce strongly different colours. Melanin multi-layers within the barbules give yellow-orange reflections, while the cortex multi-layers cover the melanin multi-layer (figure 3), which distinctly from those expected for a classical multi-layer. The optical effect was overlooked. Somewhat similar dual coloration effects are achieved by the highly

a stack of layers with alternating low and high refractive index. Notably, increasing the angle of illumination shifts the reflectance spectrum to shorter wavelengths, together with a reflectance increase and a distinct polarization of the reflected light that is extreme at a (generalized) Brewster’s angle. These effects were experimentally demonstrated and quantitatively described for the elytra of the jewel beetle *Chrysochroa fulgidissima* [24]. Parotia breast feathers share some, but certainly not all, of these features. As expected for a multi-layer reflector, the peak wavelength of the barbule reflectance decreases with increasing angle of incidence (figure 5b). The interference condition predicts a reflectance peak for normal illumination at $\lambda_{\text{max}} = 2(n_l h + n_d h)$, where $n_l$ and $n_h$ are the low and high refractive index values, and $d_l$ and $d_h$ are the corresponding thicknesses of the alternating layers (e.g. [10, 20]). The distance of the layers inside the parotia barbules was found to be about 200 nm. With an effective layer thickness of $d_h = 50$ nm of the melanin layers and refractive indices $n_l = 1.56$ of the keratin [11] and $n_h = 1.69$ of the melanin [24], the expected peak wavelength is $\lambda_{\text{max}} = 637$ nm, somewhat higher than the measured 600 nm (figure 5a). More at variance with the expectations is the finding that the reflectance amplitude also decreases with increasing angle of incidence, which is opposite to the usual amplitude increase with decreasing angle. Strictly, the latter effect is true only for TE-polarized light, because the amplitude of TM-polarized light decreases with increasing angle of incidence up to the Brewster’s angle; only above that value does the amplitude increase [24]. However, for unpolarized light, which we applied in the experiment of figure 5b, the reflection is dominated by TE-light so that the reflectance of a classical multi-layer for unpolarized light increases with increasing angle of incidence. Furthermore, although preliminary experiments showed that the light reflected by the parotia feathers does indeed become polarized with increasing angle of incidence, the observed phenomena deviate distinctly from those expected for a classical multi-layer. The reason most probably is that a roof of thin-film keratin layers covers the melanin multi-layer (figure 3), which will strongly affect the reflectance. Further study of the intricate optics of the parotia barbules is needed for a detailed, quantitative understanding of the feather reflectance.

The fact that with approximately normal illumination two bluish side beams are reflected at angles of approximately $-60°$ and $+60°$ strongly suggests that they are produced by the sloping upper surfaces of the barbule, which are approximately $-30°$ and $+30°$ to the plane of the feather blade (figure 3b). The air/keratin/melanin interface of the barbule cortex presumably works as a thin-layer reflector—like oil on water. Many birds employ such thin-film reflectors to produce an iridescent ‘gloss’ (e.g. satin bowerbirds [25]; rock dove [26]; blue-back grassquit [11]).
curved multi-layers of the wing scales of the emerald swallowtail butterfly, *Papilio palinurus* [27] and the Madagascan sunset moth, *Chrysiridia rhipheus* [28], but there the structure produces polarization-sensitive colour mixing rather than abrupt hue-shifts.

(a) **The evolution of iridescence and role of colour change in avian visual displays**

Before mating, parotia females make multiple visits to courts, where the males use six main types of display, which include a range of movements described as ‘bobs’, ‘bows’, ‘dances’ and so-forth [14]. Frith & Beehler [29] comment that the courtship display of Lawes’ parotia is a ritualized set of danced steps and movements accompanied by intricate feather movements as complex (as that of) any bird; see [15], notably figs 14 and 15; and movie at http://macaulaylibrary.org/video/45933).

The parotia feather beautifully illustrates the versatility of feathers as optical devices [3], but what does it say about the evolution and function of iridescent coloration in displays? Multi-layer structural coloration is commonly associated with display plumage [30], the peacock being the best-known case [12,31], and for human observers the iridescence is part of its beauty [32,33]. Nonetheless, in natural viewing conditions, ordinary multi-layer structures often give a rather small wavelength (or ‘hue’) shift.
in reflectance [5]. So far as we know there are no direct tests of how birds respond to different ‘components’ of the complex optical effects that are produced by structural colours [31], which include saturated, strongly directional and multi-peaked reflectance spectra, but it seems that the parotia’s breast plumage is specifically adapted to produce hue-shifts. It is difficult to be specific about the visual effects, but the fact that the feather seems to be engineered for changes in the angle between the feather and the viewer to give much larger chromatic variation in colour.

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