**SHORT COMMUNICATION**

Light sensitivity in a vertebrate mechanoreceptor?

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**ABSTRACT**
Using immunohistochemistry and western blot analysis, we demonstrate that melanopsin is localised in cells around the central pore of lateral line neuromasts in the African clawed frog, *Xenopus laevis*. Since melanopsin is a known photoreceptor pigment with diverse functions in vertebrates, we suggest that the lateral line of *Xenopus laevis*, which is primarily a mechanoreceptor, might also be light sensitive. Potential functions of such photosensitivity are discussed, including its role in mediating locomotor responses following dermal illumination.

**KEY WORDS:** Melanopsin, Lateral line, Mechanoreceptor, Photosensitivity, Multimodality, Phototaxis

**INTRODUCTION**
Although photoreceptors within the outer retina of vertebrate eyes are used by animals for image-forming light detection, extraretinal photoreceptors are widespread among non-mammalian vertebrates, occurring mainly in the brain, but also evident elsewhere in the body (Foster and Hankins, 2002). Such non-image-forming photoreceptors serve diverse functions, including: the regulation of circadian rhythms; mediating locomotor responses to dermal illumination; influencing pigment migration in chromatophores; and conferring direct light sensitivity to muscles within the iris.

Until relatively recently, it has been assumed that the only pigments capable of conferring photosensitivity to photoreceptors – even those located in structures outside the eye – use rod and cone opsins. However, in the past two decades, a number of opsins have been identified that are different enough to those of traditional photoreceptors to constitute separate gene families (Shand and Foster, 1999). One such photopigment opsin is melanopsin (OPN4). Initially shown to contribute to light-evoked pigment migration within dermal melanophores of *Xenopus laevis* (Provenzio et al., 1998), melanopsin has since been implicated in a number of roles, including conferring light-sensitivity to a subset of photoreceptive retinal ganglion cells (pRGCs) in mammals, which measure overall irradiance and underlie various non-imaging photoreceptive tasks (Hankins et al., 2008; Bailes and Lucas, 2010).

A chance observation during an investigation into iris photosensitivity suggested that the lateral line neuromasts of *Xenopus laevis* might contain melanopsin. Lateral line neuromasts are mechanoreceptors sensitive to water displacement, distributed across the body of many aquatic vertebrates (Dijkstra, 1963). In *Xenopus laevis*, they are grouped into raised ‘stitches’ arranged in characteristic patterns on the skin’s surface (Murray, 1955). The localisation of melanopsin within lateral line neuromasts suggests they may be sensitive to photic, as well as mechanical, stimuli. Here, we report on the presence and distribution of melanopsin within *Xenopus laevis* lateral lines and speculate on the functional significance of light sensitivity within this mechanoreceptor.

**RESULTS AND DISCUSSION**
Immunostaining using a polyclonal antibody (CERN972), raised against a *Xenopus laevis* melanopsin peptide, showed the majority of neuromasts on both dorsal and ventral surfaces of adult male and female pigmented and albino *Xenopus laevis* to be immunopositive (Fig. 1A). No differences in the distribution of melanopsin were observed between the different phenotypes.

Individual neuromasts showed dense immunopositive staining surrounding the central pore, with fine processes radiating outwards (Fig. 1B). In light-microscopic (Fig. 1C) and electron-microscopic (Fig. 1E) sections, dense immunopositive staining was located intracellularly in epidermal cells at the margins of the neuromast pore. As evident from wholemounts (Fig. 1B), immunostaining was not confined to the margin of the pore. In serial reconstructions of individual neuromasts, we also identified melanopsin in peripheral cells lying slightly deeper in the neuromast (Fig. 1D).

Immunoreactivity was also detected by the CERN972 antibody in western blot analysis of *Xenopus* brain and stitch samples at a mass consistent with melanopsin (Fig. 2). This is in agreement with previous identification of melanopsin expression in tadpole melanophores and brain and ocular structures of adult *Xenopus laevis* (Provenzio et al., 1998). Most samples present an upper immunoreactive band near 55 kDa and a lower band at 45–50 kDa. There are two isoforms of melanopsin in *Xenopus laevis* (OPN4x and OPN4m) (Bellingham et al., 2006), both of which would be detected by CERN972 and may be represented by the two bands in the stitch samples (Fig. 2). This would indicate that the two melanopsin orthologues most commonly found in non-mammalian vertebrates (Bellingham et al., 2006; Davies et al., 2011) are present in lateral line stitches of *Xenopus laevis*. The predicted mass for OPN4x is 60 kDa, but membrane proteins usually migrate with a somewhat lower apparent mass in SDS-PAGE. The full sequence of OPN4m is unknown, but comparison of OPN4x and OPN4m isoforms in other species suggests that they migrate with a similar apparent mass in SDS-PAGE (Davies et al., 2011; Bailes and Lucas, 2013). Bellingham et al. (2006) did not detect OPN4x mRNA in adult *Xenopus* skin tissue, which may be due to the low quantity of OPN4x mRNA, as it is only expressed in stitches, or it might indicate that the two strong bands we observe in stitch samples represent two splice variants of OPN4m. This phenomenon has already been observed in some mammalian species (Pires et al., 2009), such as the chicken (Torii et al., 2007), and in the elephant...
shark (Davies et al., 2012). The immunoreactivity at higher molecular masses is consistent with the formation of oligomeric complexes (dimers, trimers etc.), which is common under the conditions used for SDS-PAGE analysis.

Since melanopsin is a known photopigment, the presence of melanopsin immunoreactivity within *Xenopus laevis* lateral line neuromasts suggests that apart from being sensitive to mechanical stimuli, these sense organs may also be light sensitive. It is natural to speculate about the potential functional significance of such lateral line photosensitivity.

Many animals respond to dermal illumination with locomotor activity (Steven, 1963). Some previous evidence suggests the lateral line of larval lamprey mediates such dermal photosensitivity. Their lateral line nerves generate electrophysiological responses following illumination of the tail, and the lesioning of these nerves disrupts the behavioural response to such illumination (Deliagina et al., 1995; Ronan and Bodznick, 1991; Young, 1935). Our results suggest that the photosensitivity of the lateral line might be conferred by melanopsin. Interestingly, the light-driven electrophysiological responses of the lamprey lateral line nerves have a long latency, high threshold and do not adapt (Ronan and Bodznick, 1991), which are also characteristics of melanopsin-based retinal photoreceptors in mammals (Bailes and Lucas, 2010; Hughes et al., 2012).

A previous report suggests that adult *Xenopus laevis* are negatively phototactic (Denton and Pirenne, 1954). However, it is not known if they react to localised dermal illumination with locomotor activity. We confirmed the negative phototaxis of this species by observing their behaviour in an aquarium, only half of which was illuminated. In 89.8% trials (N=49) where the animals started in the lit half of the aquarium, they moved to the dark half of the tank within 3 min (mean latency 63 s). When they started in the dark half of the aquarium (N=39), by contrast, the frogs normally remained there for the duration of the experiment, spending on average 86.6% of their time in darkness and only rarely venturing into the light for brief periods of time.

We investigated whether focal illumination of the animal’s ventral surface, which could not be detected by their eyes, would induce a locomotor avoidance response. While they did appear to react to such stimuli, this was no more frequent than in control animals simply maintained in darkness. Thus, using focal ventral illumination, there was no evidence of dermally-induced locomotor activity in adult *Xenopus laevis*. It could be argued that ventral illumination is not the ideal stimulus, because in the wild, the underside of the animal will receive less illumination than other areas of the body. However, ventral neuromasts stained as heavily with melanopsin antibody as neuromasts elsewhere on the body. Furthermore, using focal ventral illumination was the only way to be certain that the illumination was not detected by the dorsally directed eyes of intact animals. Less systematic focal illumination of other areas of the body also failed to induce consistent locomotor responses.

Since focal illumination of the body surface did not induce a behavioural response, it seems likely that melanopsin in lateral line neuromasts of *Xenopus laevis* serves a function other than dermally-driven locomotor activity.

The activity of lateral line neuromasts is known to be modulated by the central nervous system using efferent neurons (Russell,
was immersed in phosphate buffered (pH 7.3) 4% paraformaldehyde at 4°C methanesulfonate (Sigma) followed by decapitation and pithing. The skin details.

smaller splice variants of OPN4m and/or OPN4x. See body of text for further represent full-length OPN4m (complete sequence is not known currently), or length OPN4m and/or OPN4x. The lower immunoreactive bands may either CERN972. The upper immunoreactive bands near 55 kDa represent full-

2828

Five X. laevis (Daudin 1802) were killed by overdose of tricaine methanesulfonate (Sigma) followed by decapitation and pithing. The skin was immersed in phosphate buffered (pH 7.3) 4% paraformaldehyde at 4°C for 3–4 h. Patches containing lateral line stitches were stored in phosphate buffered saline (PBS) until further processing or in 30% sucrose for cryosectioning.

For immunostaining, tissue was rinsed in PBS, immersed in 0.3% H2O2 in methanol for 30 min and rinsed again in PBS. Following immersion for 30 min in normal goat serum diluted in a solution of 1% Triton X-100 in PBS, tissue was incubated at 4°C for 24–48 h in the primary antibody diluted 1:2000 or 1:4000 in PBS (both dilutions produced identical staining patterns). The polyclonal antibody (CERN972) was raised against a 15-mer peptide covering residues 216–230 of X. laevis OPN4x (FLAIRSTGRNVQKLGR) (Provencio et al., 1998). The peptide was linked to rabbit serum albumin using SATA-MHS (Schieni et al., 1989). The resulting construct was injected in albino female New Zealand rabbits and processed as previously described (de Groot, 1985).

After primary antibody incubation, labelling was visualised using an avidin-biotinylated horseradish peroxidase second antibody procedure (Vector Elite ABC kit; Vector Laboratories, Peterborough, UK) applying diaminobenzidine as the chromogen (Sigma Fast; Sigma-Aldrich, Gillingham, UK).

Skin segments were viewed in wet mount to identify immunopositive regions. Some were prepared as wholemounts, while segments for fine structural observation were immersed in 2% aqueous osmium tetroxide for 1 h, before processing for araldite embedding. Semithin (1 µm) sections were cut (Ultracut E; Reichert-Jung, Depew, NY, USA) and counterstained with Toluidine Blue. Images were collected using an Olympus BH2 photomicroscope equipped with a Spot RT Color digital camera (Diagnostic Instruments Inc., Sterling Heights, MI, USA). For electron microscopy, no further enhancement to the contrast of the HRP label was required and sections were viewed on a LEO-EM912 electron microscope (Zeiss, Oberkochen, Germany) and recorded with a digital camera.

Molecular analysis

Samples of lateral line stitches, eye and various brain regions were removed from two animals killed as described above and frozen. All tissue was ground using a pre-chilled pestle and mortar prior to homogenisation in 2% (w/v) SDS, 50 mmol l−1 DTT with mini complete protease inhibitors (Roche). Samples were incubated at room temperature on a shaking platform for 2 h to improve solubilisation. The lysate was centrifuged at 23,000 g for 30 min at 20°C and the supernatant fraction used for SDS-PAGE and western blotting as described previously (Pires et al., 2009).

Every effort was made to avoid contamination of lateral line stitch samples with dermal melanoportes during dissection. If there was any minor contamination, this is unlikely to have been sufficient to produce the strong immunoreactive bands observed. We also found two clear bands in the stitch lanes on SDS-PAGE, while previous studies (Provencio et al., 1998; Bellingham et al., 2006) only detected one band in X. laevis melanophores. Hence, even if there was some contamination by melanophores, at least one of the observed bands is derived from lateral line stitches.

Phototaxis

Individual animals were removed from their home tank, during the light phase of their 12 h:12 h light:dark cycle, and placed in an experimental aquarium (20×30×20 cm). The sides of this aquarium were covered with black card and animals were observed from above. After 10 min acclimation in dim room light, the animal was placed in total darkness for 2 min, before one half of the aquarium was illuminated (3.41 W m−2) from below by a ‘light box’, consisting of two fluorescent tubes (Phillips 20W/47 Graphic A; Guildford, Surrey, UK) behind a white diffusing surface, for 3 min followed by 2 min of darkness. Animals were then exposed to light once more for another 3 min, for a maximum of 10 trials per animal. The half of the tank that was illuminated was varied randomly. Seven pigmented and two albino animals were tested.

We also investigated the ability of focal illumination to induce locomotor activity. The ventral surface of four animals was illuminated using the same protocol as above, but instead of illuminating half the aquarium the light source was covered except for a 1 cm round aperture that was positioned near the centre of the animal’s ventral surface when it was resting on the bottom of the tank. The time of any movement after the spot was turned on was noted (N=18). A similar number of control observations were made with
the stimulating spot in position but not switched on. Less systematically, we also tried directing light onto various parts of the body with both a narrow torch beam and low power lasers and observing any reaction.

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Competing interests
The authors declare no competing or financial interests.

Author contributions
G.E.B. and R.H.D. conceived the study. G.E.B. performed the immunohistochemistry, for which W.J.d.G. provided the antibody. H.-J.W. performed most of the microscopy. R.G.F. and M.T. carried out the western blot analysis and R.H.D. performed the phototactic experiments. All authors contributed to the interpretation of data. R.H.D. drafted the manuscript, which was edited by all authors, except G.E.B., prior to submission.

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