Regenerating reptile retinas: a comparative approach to restoring retinal ganglion cell function

DL Williams

Abstract

Transection or damage to the mammalian optic nerve generally results in loss of retinal ganglion cells by apoptosis. This cell death is seen less in fish or amphibians where retinal ganglion cell survival and axon regeneration leads to recovery of sight. Reptiles lie somewhere in the middle of this spectrum of nerve regeneration, and different species have been reported to have a significant variation in their retinal ganglion cell regenerative capacity. The ornate dragon lizard *Ctenophoris ornatus* exhibits a profound capacity for regeneration, whereas the Tenerife wall lizard *Gallotia galloti* has a more variable response to optic nerve damage. Some individuals regain visual activity such as the pupillomotor responses, whereas in others axons fail to regenerate sufficiently. Even in *Ctenophoris*, although the retinal ganglion cell axons regenerate adequately enough to synapse in the tectum, they do not make long-term topographic connections allowing recovery of complex visually motivated behaviour. The question then centres on where these intraspecies differences originate. Is it variation in the innate ability of retinal ganglion cells from different species to regenerate with functional validity? Or is it variances between different species in the substrate within which the nerves regenerate, the extracellular environment of the damaged nerve or the supporting cells surrounding the regenerating axons? Investigations of retinal ganglion cell regeneration between different species of lower vertebrates *in vivo* may shed light on these questions. Or perhaps more interesting are *in vitro* studies comparing axon regeneration of retinal ganglion cells from various species placed on differing substrates.

Eye (2017) 31, 167–172; doi:10.1038/eye.2016.224;
published online 11 November 2016

Introduction

Have you ever wondered why the rod of Asclepius, that ancient sign of medical healing,¹ is characterised by a snake entwined around a staff (Figure 1)? The healing temples of Asclepius were apparently inhabited by snakes, which crawled freely around the floors where the patients seeking healing spent the night. The snake was a sign of the duality of life and death. Often venomous to be sure, but also a sign of life renewed. The regular ecdysis or sloughing of the entire skin occurring regularly as the snake grew through life was central to this sign. As Comutus, a first century CE philosopher notes ‘Asclepius derived his name from healing soothingly and from deferring the withering that comes with death. For this reason, therefore, they give him a serpent as an attribute, indicating that those who avail themselves of medical science undergo a process similar to the serpent in that they, as it were, grow young again after illnesses and slough off old age.’ Pliny the Elder (23–79 CE) wrote ‘The snake, when the membrane covering its body has been contracted by the cold of winter, throws it off in the spring and thus becomes sleek and youthful in appearance. The same animal, too, on finding its sight weakened during its winter retreat, anoints and refreshes its eyes by rubbing itself on the fennel plant.’² The ophidian eye is indeed covered by fused eyelids, the spectacle, which becomes opaque before it is shed with the rest of the skin as the reptile increases in size through life. Very interesting, you might say, but what has this to do with the retinal ganglion cell?

Department of Veterinary Medicine, University of Cambridge, Queen's Veterinary School Hospital, Cambridge, UK

Correspondence:
DL Williams, Department of Veterinary Medicine, University of Cambridge, Queen's Veterinary School Hospital, Madingley Road, Cambridge, CB3 0ES, UK
Tel: +44 (0)1223 337621;
Fax: +44 (0)1223 232977.
E-mail: dlw33@cam.ac.uk

Received: 21 September 2016
Accepted: 22 September 2016
Published online:
11 November 2016

Figure 1 Asclepius and his rod with obligatory snake coiled around it.

Tissue regeneration in lower vertebrates

Lower vertebrates, that is to say fish, amphibians, and reptiles, generally grow throughout life, not reaching a maximal adult size as do the high vertebrate birds and mammals. And generally in association with this, they have also maintained the ability to regenerate,

Figure 2 (a) The *Gallotia galloti* the Tenerife wall lizard. (b) *Ctenophorus ornatus* the ornate dragon lizard.

be it the regrowth of the lizard tail after autotomy,^{3,4} limb regeneration in amphibia,⁵ cardiac regrowth in fish⁶ and, of more interest in this paper, lens and retinal replication in newts⁷ and fish.⁸ In truth, the involvement of reptiles in the alliterative title of this paper does leave us with somewhat of a problem, since the retinal ganglion cell (henceforth RGC) regeneration involved in neural regrowth after optic nerve injury (ONI) as either a crush episode or frank axonotomy is not as complete as in the piscine or amphibian examples we will cover later. Nevertheless, the variation in RGC regeneration between reptile species may give opportunities to study what molecular and cellular mechanisms are at work in RGC development and redevelopment.

Retinal ganglion cell regeneration in reptiles

The wall lizard *Gallotia galloti* (Figure 2a) is found on the Canary Islands and the striking colour of the males suggests the importance of vision for mate choice in this species. In fact it is more likely to be the markings, which reflect electromagnetic radiation in the ultraviolet spectrum that are most important in sexual selection.⁹ Indeed across reptile genera, as with avian vision photoreceptor sensitivity in the ultraviolet is very important.^{10,11} Never let us think that as trichomats we are at some evolutionary pinnacle in visual development! Although we might admire the wide variation in

colouration of the ornate crevice dragon lizard *Ctenophorus ornatus*¹² vision in the ultraviolet spectrum is again crucial to this species too.¹³

But where the *Gallotia* and *Ctenophorus* lizards may be similar in their use of ultraviolet signalling, their response to optic nerve axotomy is surprisingly different. Perhaps the first thing to say is that both lizards, as with all reptiles, show a considerable survival of RGC after ONI. In birds and mammals, degeneration of RGC after axonal injury is widespread as first recognised by Santiago Ramon y Cajal as long ago as 1928.¹⁴ Axons start to sprout at the site of the injury in these higher vertebrates, but degenerate within a week and are lost with apoptosis of the RGC cell body. After a standard optic nerve crush injury in the *Gallotia* and *Ctenophorus* lizards, however, axon regeneration occurs even in the face of a glial scar and the absence of proliferating cells in the retina itself. But although the *Gallotia* lizard has an average RGC loss of around 30% and slow regeneration over 6 months or so with 60% of neurectomised animals regaining a pupillary light reflex,¹⁵ *Ctenophorus* has a significantly better axonal regeneration with an accelerated time course compared with *Gallotia*. Its axons reach the visual centres of the brain within a month. Although this sounds impressive, functionality depends on correct topographic localisation of migrating neurons and here even *Ctenophorus* falls short of our expectations. Although all seems well to start with post injury and pupillary light reflexes are restored at 2–3 months post injury, the topographic arrangement of the retinal projections is not maintained in the regenerating optic nerve. Vision allowing such essential behaviours as accurate apprehension of prey items is not regained after ONI, nor are the retinotectal projections in these lizards stable. Regenerating axons seem to be continually searching, as it were, for the correct projection rather than forming robust synaptic connections as shown anatomically^{16,17} or electrophysiologically.¹⁸ And so perhaps now is the time to stray a bit further back in evolution and consider the anamniotes, amphibians, and fish and their responses to optic nerve damage as opposed to the poorer regenerative capacity of the amniotes, those vertebrates developing in an egg with an amnion.

Different animals, different questions?

Maybe as well as looking at the wrong genera, reptiles rather than amphibians and fish, perhaps we are asking the wrong question in trying to define what causes optic nerve regeneration in animals where it occurs. Maybe we should be asking what prevents RGC regeneration in higher species. Is it the nerve itself unable to survive and experience renewal or is it the environment of the

damaged nerve, which is key in preventing nerve regeneration? Do amphibia and fish have RGC, which themselves are intrinsically more able to regenerate or is it the environment of the ONI scar, which is the key variable? At a more fundamental level, is it merely RGC survival that explains the restoration of visual processes after ONI or is optic nerve regeneration more related to *de novo* RGC neurogenesis? It used to be thought that the retinae of anamniote species continually produced new neurons and that this accounted for restoration of the optic nerve. In fish, retinal stem cells do continually give rise to new RGC. However, studies on adult *Xenopus*, the African clawed toad, showed that in these animals neurogenesis in the retina ceased once adulthood has been reached.¹⁹ Even so, RGC axons regenerate in adult *Xenopus*.²⁰ In fish, on the other hand, retinal neurogenesis occurs throughout life with stem cells present and continually proliferating.²¹ Before we get too excited about this as a critical difference between fish and mammals, it has to be remembered that the central nervous system of every genera seems to have latent stem cells,^{22,23} which might be able to give rise to new nerve cells in the right circumstances, though much more promisingly in lower vertebrates.

It was back in 1927 that Matthey reported optic nerve regeneration in amphibia²⁴ but not until more than 20 years later that Sperry documented the same results in fish.²⁵ With regard to RGC survival, we know, as noted above, that after ONI in the rat the RGC undergo apoptosis within 3–5 days post injury. Terminal deoxynucleotidyl transferase-mediated dUTP nick end labelling staining demonstrates apoptotic signals in the rat retina 7 days after ONI,²⁶ whereas in the fish no such signals are seen at any time point after axotomy.²⁷ What accounts for this RGC survival? Presence of pro- and anti-apoptotic factors differ substantially between piscine and mammalian RGC post ONI. Phospho-Akt and phospho-Bad are anti-apoptotic molecules from the phosphatidylinositol-3-kinase system, which increase after ONI in the fish²⁸ but decrease in the rat.²⁹ Insulin-like growth factor-1 and brain-derived neurotrophic factor are both key activators in the phosphatidylinositol-3-kinase system and levels of IGF-12 increase in the goldfish after ONI but rapidly decrease in the rat.²⁷ Heat shock proteins (HSP) also seem key in RGC survival. HSP are chaperon molecules protecting cells from a wide variety of environmental and physiological insults. HSP70 mRNA increases over threefold in the first few hours after ONI in the zebra fish but not after mammalian ONI.³⁰ Here inhibition of HSP activity blocked expression of the anti-apoptotic protein Bcl-2 and increased levels of the apoptotic protein Bax.

Is there a central trigger factor behind these various changes? Purpurin, a retinol binding protein, increases

markedly but transiently in the fish retina after ONI suggesting that retinoid activity may have an important role in recovery after ONI, acting as a molecular signalling cascade.³¹ Is the same true in other species? Retinoic acid signalling is also upregulated in the frog after ONI³² and while there is normally no regeneration after ONI in mammals, a model using the neuroprotective herbal iridoid genipin does allow RGC regeneration. Inhibition of retinoic acid receptor beta expression by use of siRNA inhibits the neuritogenic actions of IPRG001, a genipin derivative.³³

Ganglion cell growth or scar permissivity?

We have said that on the one hand, there are intrinsic factors in the anamniote RGC, which explain their survival and regeneration but on the other, the extracellular environment after the ONI is critical too. The glial scar resulting from the ONI in mammals is inhibitory to nerve regeneration. Oligodendrocytes and myelin are potent inhibitors of axonal regrowth in the mammalian central nervous system,³⁴ but in the fish and amphibian this regrowth is not inhibited in the same manner with oligodendrocytes successfully remyelinating regenerating optic nerve axons,³⁵ which can then reform synapses in the optic tectum.³⁶ The area of ONI in fish is characterised by increased extracellular matrix molecules, such as tenascin, chondroitin sulphate, and laminin.^{37,38} If we return to the lizards we began with, straddling the gap between the axonal restoration of the fish and the failure of regeneration in mammals, RGC axons in *Gallotia galloti* regenerate successfully even in the presence of inhibitory myelin and oligodendrocytes. Their sensitivity seems to be less than neurons from higher vertebrates. Intricate work from Lang and Stuermer's group culturing *Gallotia* retinal explants on lizard or rat glial cell cultures or rat dorsal root ganglion explants on lizard optic nerve explants showed that lizard RGC growth cones traversed rat oligodendrocyte cultures, whereas rat neurons are inhibited by lizard oligodendrocytes.³⁹ It seems that it may be the lizard RGC which has the ability to regenerate in whatever environment it is placed. Perhaps it is its response to signals that go unnoticed by mammalian RGC is the key factor in its survival and regeneration. Hypertrophic gliosis in the region of ONI (Figure 3) is mediated by axons in the locality with the presence of growing optic nerve fibres being essential to oligodendrocyte and type II astrocyte differentiation.⁴⁰

Mammalian relevance?

Although the work on fish, amphibians, and reptile ONI is fascinating, its true relevance must be in what it tells us about repair of mammalian optic nerves and, though it

Figure 3 GFAP staining shows astrocytosis in injured optic nerve (right).

pains me as a veterinary surgeon to have to say this, its impact really lies on how it can help us in understanding restoration of human optic nerve structure and function not regaining lizard or fish vision! As we move to recovery from ONI in fish to mammals perhaps we can move to an animal which, while clearly a mammal, has optic nerve regeneration much more similar to that in fish and amphibians. This is the naked mole rat (*Heterocephalus glaber*).⁴¹ Little bigger than a standard laboratory mouse (*Mus musculus*), the naked mole rat has a lifespan not of 2–3 years as with *Mus* but rather of 30 years. Living underground with low-oxygen levels in large eusocial colonies, *Heterocephalus* is nyctophilous, living with its body temperature matching the ambient temperature. These animals not only have an exceptionally long lifespan but little in the way of senescence and rare reports of neoplasia. Its cells have high levels of telomerase, which may explain the substantial longevity but also ironically high levels of oxidative stress.⁴² So with all these unusual characteristics perhaps it is not surprising that the naked mole rat's response to ONI is very different from those in more conventional mammals. Rather than the low percentage of RGCs that survive ONI in other mammals, in *Heterocephalus*, 70% of RGC survive ONI and three times more optic nerve axons show regeneration after ONI. So what differs between conventional laboratory rodents and *Heterocephalus* with regard to the damaged optic nerve? The JAK/STAT pathway seems to play an important part in optic nerve regeneration in fish⁴³ and, while there is nearly no optic nerve p-STAT3 immunoreactivity in *Mus*, many RGC in *Heterocephalus* express nuclear p-STAT3 after ONI. Not only are there intrinsic differences in *Heterocephalus* RGC but extrinsic

Figure 4 Boa constrictor with an apparently buphthalmic glaucomatous globe but actually with a bullous spectaculopathy.

variations may be critical, particularly those in astrocytes. Immunohistochemistry for GFAP, a marker for reactive astrocytes shows little difference between *Heterocephalus* and *Mus* optic nerves 3 days after ONI, whereas at 14 days, there is little GFAP immunoreactivity in *Mus* but intense staining in *Heterocephalus*.⁴⁴ In most mammals, the fibrotic scars after ONI tend to preclude the formation of glial bridges across the lesion, whereas in fish, amphibians, and naked mole rats astrocyte involvement in the regenerating optic nerve seems to be key. Astrocytes have key roles in remodelling the optic nerve scar, but their activity and neuronal responses to it differ substantially between species.

Conclusion

The reason for being concerned about RGC regeneration after ONI is of course to prevent the sight threatening damage of glaucoma. So perhaps rather than looking at experimental optic nerve crush models, we should be searching for spontaneously glaucomatous reptiles. Cases such as that in Figure 4 appear exactly what we are looking for—an animal with an enlarged glaucomatous globe. But here we need to go back to Pliny the Elder with whom we started. The apparent rejuvenation experienced by the snakes he saw was in fact shedding of the skin and transparent spectacle. And the swelling seen in Figure 4 is not an enlarged globe but rather a swollen spectacle in an animal with an occluded nasolacrimal duct, in what we term a bullous spectaculopathy.⁴⁵ All of which goes to show how difficult it can be to extrapolate from these findings in species very different from the experimental laboratory mammals we are more used to investigating. Part of the fascination of investigating responses to ONI in these lower vertebrates is quite how different they are, how ready they are to regenerate. They are so different, however, in so many varying ways that it can be very difficult to determine what it is, RGC or scar biology or

both, which accounts for this dramatic variation in RGC regeneration in these lower vertebrates compared with the vast majority of mammals.

Conflict of interest

The author declares no conflict of interest.

References

- 1 Antoniou SA, Antoniou GA, Learney R, Granderath FA, Antoniou AI. The rod and the serpent: history's ultimate healing symbol. *World J Surg* 2011; **35**(1): 217–221.
- 2 Pliny the Elder. *Naturalis Historia* 8: 41. Available at: <http://www.perseus.tufts.edu/hopper/text?doc=Plin.+Nat.+toc>.
- 3 Alibardi L. Histochemical, biochemical and cell biological aspects of tail regeneration in lizard, an amniote model for studies on tissue regeneration. *Prog Histochem Cytochem* 2014; **48**(4): 143–244.
- 4 Gilbert EA, Payne SL, Vickaryous MK. The anatomy and histology of caudal autotomy and regeneration in lizards. *Physiol Biochem Zool* 2013; **86**(6): 631–644.
- 5 Kusumi K, Fisher RE. Studying mechanisms of regeneration in amphibian and reptilian vertebrate models. *Anat Rec (Hoboken)* 2012; **295**(10): 1529–1531.
- 6 Gamperl AK, Farrell AP. Cardiac plasticity in fishes: environmental influences and intraspecific differences. *J Exp Biol* 2004; **207**(Pt 15): 2539–2550.
- 7 Del Rio-Tsonis K, Tsonis PA. Eye regeneration at the molecular age. *Dev Dyn* 2003; **226**(2): 211–224.
- 8 Pushchina EV, Shukla S, Varaksin AA, Obukhov DK. Cell proliferation and apoptosis in optic nerve and brain integration centers of adult trout *Oncorhynchus mykiss* after optic nerve injury. *Neural Regen Res* 2016; **11**(4): 578–590.
- 9 Thorpe RS, Richard M Evidence that ultraviolet markings are associated with patterns of molecular gene flow. *Proc Natl Acad Sci USA* 2001; **98**(7): 3929–3934.
- 10 Pérez i de Lanuza G, Font E. Ultraviolet vision in lacertid lizards: evidence from retinal structure, eye transmittance, SWS1 visual pigment genes and behaviour. *J Exp Biol* 2014; **217**(Pt 16): 2899–2909.
- 11 Martin M, Le Galliard JF, Meylan S, Loew ER. The importance of ultraviolet and near-infrared sensitivity for visual discrimination in two species of lacertid lizards. *J Exp Biol*. 2015; **218**(Pt 3): 458–465.
- 12 Stuart-Fox DM, Moussalli A, Johnston GR, Owens IP. Evolution of color variation in dragon lizards: quantitative tests of the role of crypsis and local adaptation. *Evolution* 2004; **58**(7): 1549–1559.
- 13 Barbour HR, Archer MA, Hart NS, Thomas N, Dunlop SA, Beazley LD *et al*. Retinal characteristics of the ornate dragon lizard, *Ctenophorus ornatus*. *J Comp Neurol* 2002; **450**(4): 334–344.
- 14 Ramon y Cajal S. *Degeneration and Regeneration of the Nervous System*. Translated by RM May. Oxford Scholarship Online 2012; available at doi:10.1093/acprof:oso/9780195065169.001.0001.
- 15 Santos E, Romero-Alemán MM, Monzón-Mayor M, Yanes C. Variable functional recovery and minor cell loss in the ganglion cell layer of the lizard *Gallotia galloti* after optic nerve axotomy. *Exp Eye Res* 2014; **118**: 89–99.

- 16 Dunlop SA, Tran N, Tee LB, Papadimitriou J, Beazley LD. Retinal projections throughout optic nerve regeneration in the ornate dragon lizard, *Ctenophorus ornatus*. *J Comp Neurol* 2000; **416**(2): 188–200.
- 17 Beazley LD, Sheard PW, Tennant M, Starac D, Dunlop SA. Optic nerve regenerates but does not restore topographic projections in the lizard *Ctenophorus ornatus*. *J Comp Neurol* 1997; **377**(1): 105–120.
- 18 Stirling RV, Dunlop SA, Beazley LD. Electrophysiological evidence for transient topographic organization of retinotectal projections during optic nerve regeneration in the lizard, *Ctenophorus ornatus*. *Vis Neurosci* 1999; **16**(4): 681–693.
- 19 Grant S, Keating MJ. Ocular migration and the metamorphic and postmetamorphic maturation of the retinotectal system in *Xenopus laevis*: an autoradiographic and morphometric study. *J Embryol Exp Morphol* 1986; **92**: 43–69.
- 20 Taylor JSH, Jack JL, Easter SS. Is the capacity for optic nerve regeneration related to continued retinal ganglion cell production in the frog? *Eur J Neurosci* 1989; **1**(6): 626–638.
- 21 Perron M, Harris WA. Retinal stem cells in vertebrates bioassays. *Bioessays* 2000; **22**(8): 685–688.
- 22 Alunni A, Bally-Cuif L. A comparative view of regenerative neurogenesis in vertebrates. *Development* 2016; **143**(5): 741–753.
- 23 Taupin P, Gage FH. Adult neurogenesis and neural stem cells of the central nervous system in mammals. *J Neurosci Res* 2002; **69**(6): 745–749.
- 24 Matthey R. Recupération de la vue après résection des nerfs optiques chez le triton. *CR Soc Biol* 1925; **93**: 904–906.
- 25 Sperry RW. Patterning of central synapses in regeneration of the optic nerve in teleosts. *Physiol Zool* 1948; **21**(4): 351–361.
- 26 Bien A, Seidenbecher CI, Böckers TM, Sabel BA, Kreutz MR. Apoptotic versus necrotic characteristics of retinal ganglion cell death after partial optic nerve injury. *J Neurotrauma* 1999; **16**(2): 153–163.
- 27 Koriyama Y, Homma K, Sugitani K, Higuchi Y, Matsukawa T, Murayama D *et al*. Upregulation of IGF-I in the goldfish retinal ganglion cells during the early stage of optic nerve regeneration. *Neurochem Int* 2007; **50**(5): 749–756.
- 28 Koriyama Y, Homma K, Kato S. Activation of cell survival signals in the goldfish retinal ganglion cells after optic nerve injury. *Adv Exp Med Biol* 2006; **572**: 333–337.
- 29 Homma K, Koriyama Y, Mawatari K, Higuchi Y, Kosaka J, Kato S. Early downregulation of IGF-I decides the fate of rat retinal ganglion cells after optic nerve injury. *Neurochem Int* 2007; **50**(5): 741–748.
- 30 Nagashima M, Fujikawa C, Mawatari K, Mori Y, Kato S. HSP70, the earliest-induced gene in the zebrafish retina during optic nerve regeneration: its role in cell survival. *Neurochem Int* 2011; **58**(8): 888–895.
- 31 Kato S, Matsukawa T, Koriyama Y, Sugitani K, Ogai K. A molecular mechanism of optic nerve regeneration in fish: the retinoid signaling pathway. *Prog Retin Eye Res* 2013; **37**: 13–30.
- 32 Duprey-Díaz MV, Blagburn JM, Blanco RE. Optic nerve injury upregulates retinoic acid signaling in the adult frog visual system. *J Chem Neuroanat* 2016; **77**: 80–92.
- 33 Koriyama Y, Takagi Y, Chiba K, Yamazaki M, Sugitani K, Arai K *et al*. Requirement of retinoic acid receptor β for genipin derivative-induced optic nerve regeneration in adult rat retina. *PLoS One* 2013; **8**(8): e71252.
- 34 Caroni P, Schwab ME. Oligodendrocyte- and myelin-associated inhibitors of neurite growth in the adult nervous system. *Adv Neurol* 1993; **61**: 175–179.
- 35 Wolburg H. Myelination and remyelination in the regenerating visual system of the goldfish. *Exp Brain Res* 1981; **43**(2): 199–206.
- 36 Stuermer CA, Easter SS Jr. A comparison of the normal and regenerated retinotectal pathways of goldfish. *J Comp Neurol* 1984; **223**(1): 57–76.
- 37 Becker CG, Schweitzer J, Feldner J, Schachner M, Becker T. Tenascin-R as a repellent guidance molecule for newly growing and regenerating optic axons in adult zebrafish. *Mol Cell Neurosci* 2004; **26**(3): 376–389.
- 38 Becker CG, Becker T. Repellent guidance of regenerating optic axons by chondroitin sulfate glycosaminoglycans in zebrafish. *J Neurosci* 2002; **22**(3): 842–853.
- 39 Lang DM, Monzón-Mayor M, Bandtlow CE, Stuermer CA. Retinal axon regeneration in the lizard *Gallotia galloti* in the presence of CNS myelin and oligodendrocytes. *Glia* 1998; **23**(1): 61–74.
- 40 Levine RL. Axon dependent glial changes during optic fiber regeneration in the goldfish. *J Comp Neurol* 1993; **333**(4): 543–553.
- 41 Buffenstein R. The naked mole-rat: a new long-living model for human aging research. *J Gerontol A Biol Sci Med Sci* 2005; **60**(11): 1369–1377.
- 42 Andziak B, O'Connor TP, Qi W, DeWaal EM, Pierce A, Chaudhuri AR *et al*. High oxidative damage levels in the longest-living rodent, the naked mole-rat. *Aging Cell* 2006; **5**(6): 463–471.
- 43 Elsaiedi F, Bemben MA, Zhao XF, Goldman D. Jak/Stat signaling stimulates zebrafish optic nerve regeneration and overcomes the inhibitory actions of Socs3 and Sfpq. *J Neurosci* 2014; **34**(7): 2632–2644.
- 44 Park KK, Luo X, Mooney SJ, Yungher BJ, Belin S, Wang C *et al*. Retinal ganglion cell survival and axon regeneration after optic nerve injury in naked mole-rats. *J Comp Neurol* 2016; epub ahead of print 28 June 2016; doi:10.1002/cne.24070.
- 45 Cullen CL, Wheler C, Grahn BH. Diagnostic ophthalmology. Bullous spectacleopathy in a king snake. *Can Vet J* 2000; **41**(4): 327–328.