

First record of leucism in a primary burrowing crayfish: *Distocambarus crockeri* Hobbs & Carlson, 1983 (Decapoda: Astacidea: Cambaridae) from South Carolina, USA

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ABSTRACT

The piedmont prairie burrowing crayfish, *Distocambarus crockeri* Hobbs & Carlson, 1983, is a primary burrowing crayfish endemic to South Carolina, USA. The species is the most widely distributed and studied of the five species of *Distocambarus* Hobbs, 1981. Surveys have clarified the species distribution and habitat, but, like other burrowing crayfishes, our understanding of its ecology is lacking. We report the discovery of two leucistic specimens of *D. crockeri* collected in an ephemeral pool in a roadside ditch on 14 March 2022 in Edgefield County, South Carolina. To the best of our knowledge, our discovery represents the first documented case of leucism in a primary burrowing crayfish. We discuss the significance of this observation in relation to other recorded observations of color anomalies throughout crayfishes as well as potential environmental causes of leucism.

KEYWORDS: albinism, color abnormalities, color morphs, crayfish color, Crustacea, color mutations, pigments

Freshwater crayfishes are recognized as ecosystem engineers and keystone species (Crocker & Barr, 1968; Momot et al., 1978; Creed, 1994; Momot, 1995; Usio, 2000), mostly because of their burrowing ability. Hobbs (1981) assigned crayfishes into three burrowing categories based on behavior: tertiary, secondary, and primary. Tertiary and secondary burrowing crayfishes construct relatively simple burrows with a single vertical tunnel that remains connected to perennial lentic or lotic surface waters. Tertiary and secondary burrowing crayfishes often only burrow when sheltering themselves from predators, avoiding desiccation, or when reproducing. Conversely, primary burrowing crayfishes spend the majority of their lives within and around their terrestrial burrows and do not require a direct connection to perennial surface waters. Primary burrowing crayfishes construct complex burrows which often exhibit multiple portals (surface openings for burrows), tunnels, and chambers that can occur significant distances from permanent bodies of water (Hobbs, 1981). Unsurprisingly, with over 150 described species of primary burrowing crayfishes, their secluded underground lifestyle has thus far resulted in a poor understanding of their biology. According to the International Union for Conservation

of Nature (IUCN), 19.5% of primary burrowing crayfishes are categorized as being data deficient, but 12.9% of primary burrowers have not been evaluated (Bloomer *et al.*, 2021; IUCN, 2022).

One unanswered question relating to primary burrowing crayfishes relates to their coloration and color patterns (Schuster, 2020). Despite spending a majority of their lives underground, primary burrowing crayfishes can exhibit vibrant colors, with many species displaying conspicuous red, orange, and blue colors and distinct color patterns (Schuster, 2020). The function of such vibrant colors and conspicuous patterns in primary burrowing crayfishes remains unknown, but it has been speculated to be important in intra/interspecific competition, communication, camouflage, and species identification (Schuster, 2020). Although some primary burrowing crayfishes may possess conspicuous coloration, other primary burrowing crayfishes exhibit inconspicuous, drab colorations, like many tertiary and secondary burrowing crayfishes. Tertiary and secondary burrowing crayfishes often display coloration that provides camouflage and advantages for life in open-water stream habitats, where they are presumably exposed to increased predation pressures compared

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to primary burrowers (Kent, 1901; Thacker *et al.*, 1993; Füreder & Machino, 2002; Sacchi *et al.*, 2021).

Atypical colors phenotypes and color patterns have been documented in several species of crayfishes, although almost entirely within tertiary and secondary burrowers. For instance, an individual of Cambarus bartonii bartonii Fabricius, 1798 from eastern Pennsylvania was discovered with a bilaterally partitioned color phenotype with the left half of the body dark olive green and the right half light amber (Hartzell, 2017). A similar observation of color bilateralism was recorded in Faxonius immunis Hagen, 1870 (Dowell & Winier, 1969). Blue-color phenotypes appear to be common in crayfishes, with documented instances of atypical blue coloration appearing in three of four families: Cambaridae (Procambarus clarkii Girard, 1852, P. hagenianus Faxon, 1884, P. acutus Girard, 1852, Cambarus carolinus Erichson, 1846, and Creaserinus hedgpethi Hobbs, 1948), Parastacidae (Cherax destructor Clark, 1936), and Astacidae (Pontastacus leptodactylus Eschscholtz, 1823) (Hayes & Reimer, 1975; Walker et al., 2000; Schuster, 2020; Kale et al., 2021). Atypical color phenotypes are also popular in the pet trade, with orange phenotypes of Cambarellus patzcuarensis Villalobos, 1943 (Weiperth et al., 2017) and blue phenotypes of Procambarus alleni Faxon, 1884 being sold worldwide.

There are also several reports of individuals with white color phenotypes (see below), which may be a result from genetic conditions known as leucism and albinism. Albinism results from a genetic mutation in which there is a total absence of the pigment melanin (Lutz, 2008), resulting in an entirely white phenotype. By contrast, leucism, sometimes referred to as partial albinism, occurs when the majority, but not all, of the body exhibits a lack of melanin due to genetic mutation, resulting in a white phenotype with the possibility of pigmented features, such as the eyes (Sage, 1962; Jehl, 1985; Fleck *et al.*, 2016). Although there is substantial confusion regarding the nomenclature of white-color aberrations (see van Grouw (2021) for a discission of these issues in birds), we discuss leucism following the definition provided above, in which a majority of the animal in question is white, but some regions may remain pigmented. Nevertheless, it is worth highlighting that crustacean coloration is believed to be primarily driven by carotenoid-based pigments, and not melanin-based pigments (Tlusty & Hyland, 2005; Wade et al., 2005). Still, we use the word leucism to describe these color phenotype aberrations in alignment with previous literature. Within natural populations of crayfishes, there are reports of leucism in P. leptodactylus (Kale et al., 2020), P. clarkii (Nakatani, 1999), Faxonius propinguus Girard, 1852 (Dunham et al., 1979) and Faxonius rusticus Girard, 1852 (Glon, 2016). Further, leucism is believed to result from a genetic mutation within a single locus (Owen & Skimmings, 1992; Fleck et al., 2016), which allows leucistic crayfish to be bred in the pet trade. For example, within the pet trade, leucistic or "white" Procambarus clarkii are widely available (Faulkes, 2015a, b; Putra et al., 2018). Unsurprisingly, within primary burrowing crayfishes, few observations of atypical coloration have been reported in the literature. While describing two new species of Lacunicambarus, Glon et al. (2020) commented on abnormalities in the base color for two banded mudbugs, Lacunicambarus freudensteini Glon et al., 2020: one specimen exhibited an abnormal powder blue coloration, whereas a second one exhibit pink coloration. Additionally, there are reports of a population of prairie crayfish, Procambarus gracilis Bundy in Forbes, 1876 in which many individuals possess blue coloration (Secker, 2013).

We report the discovery of two leucistic individuals of the piedmont prairie burrowing crayfish, *Distocambarus crockeri* Hobbs & Carlson, 1983, a primary burrowing crayfish endemic to South Carolina, USA. Individuals of this species are typically inconspicuous, being dark brown to grayish brown with stripes or patterning sometimes present. As one of the five species of *Distocambarus*, *D. crockeri* has the widest distribution, spanning



Figure 1. The roadside ditch where we collected two leucistic individuals of Distocambarus crockeri (A, B).

Edgefield, McCormick, Greenwood, and Saluda counties in South Carolina (Hobbs & Carlson, 1983; unpublished data).

Both leucistic individuals were collected in Edgefield County on March 2022, (33.921730, -82.126683) at a roadside ditch (RSD) site *D. crockeri* historically inhabited (Welch & Eversole, 2006). Intense anthropogenic manipulation from logging occurred at the site at the time of sampling (Fig. 1). Pine-tree management frequently occurs in Edgefield and surrounding counties, with clear-cutting as a common practice to initiate new plots for the timber industry. At the collection site, a culvert-fed pool approximately 25 m long × 1.5 m wide filled the RSD with approximately 12 cm of turbid water containing a large amount of trash. The entire RSD habitat had an open canopy and surrounding vegetation composed of grasses, sedges, and rushes, with evidence of mowing.

Because no visible surface burrows were detected, we collected individuals by dip-netting the entire length of the RSD six times to disturb any crayfish in the water column and increase our chances of collection. The first attempt resulted in the collection of two apparent leucistic individuals of *D. crockeri* (Fig. 2) in addition to many other typically colored individuals. *Distocambarus crockeri* was the only species collected at the site (N = 68). We presumed that both individuals were leucistic based on their clear-white unpigmented carapaces with the only pigmented

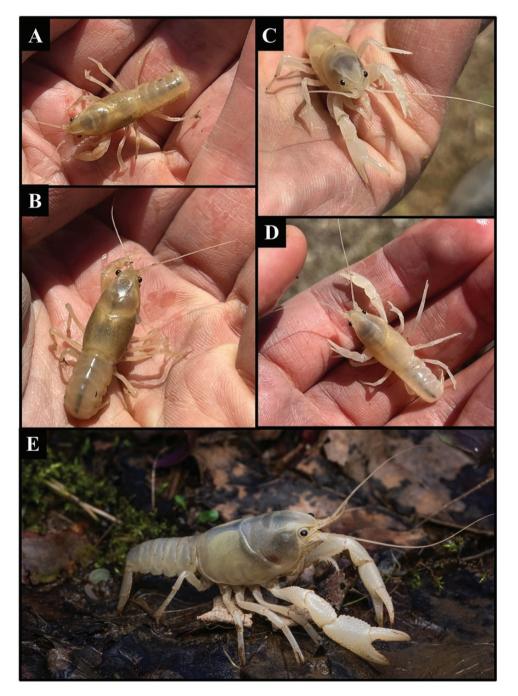


Figure 2. Female leucistic *Distocambarus crockeri* in situ (A, B); male leucistic individual in situ (C, D); male individual after a fourth molt in captivity at West Liberty University; photo by ZAG (E).

region of the body being their eyes. Both crayfishes appeared to have recently molted based on their exoskeleton pliability. We sexed each individual and measured their carapace length with digital calipers. One specimen was a form II male with a carapace length of 17.5 mm and the second a female with a carapace length of 16.33 mm and a regenerated right claw. Based on previously published data, both specimens were subadults (i.e., too small to reproduce; Eversole & Welch, 2013). The leucistic female did not survive beyond collection. The male, however, was taken to West Liberty University alive, where it has since molted five times.

While in captivity, the male *D. crockeri* was fed leaf-litter *ad libidum* in a $15 \times 15 \times 5.5$ cm container. During this time, the crayfish's white exoskeleton slowly developed darker colors, but the dominant coloration remained white. After molting, the crayfish returned to the bright white coloration with pigmented eyes (Fig. 2E), which confirms leucism (as a genetic phenotype), ruling out the lack of dietary pigments as a cause of coloration. We acknowledge, however, that leaf litter alone may not have supplied the complete range of pigments available in the natural diet(s) of primary burrowing crayfishes. This discovery is the first reported case of leucism in a primary burrowing crayfish.

Although color abnormalities such as leucism are rarely reported in crayfishes, one population of leucistic *F. propinguus* is known to have co-occurred with typical colored individuals of the same species over 40 years in Lake Simcoe, Ontario, Canada (Dunham et al., 1979; Jordan et al., 1986; Hamr et al., 2019). Since this leucistic F. propinquus population was first discovered, rusty crayfish F. rusticus have invaded Lake Simcoe and compete sympatrically (Bowles, 2009). Following the invasion of F. rusticus, the abundance of leucistic F. propinguus has dramatically decreased, with Hamr et al. (2019) recently collecting only six leucistic individuals, whereas Dunham et al. (1979) collected 119 in two days in 1977. Despite the recent decrease in population size, the Canadian population of leucistic F. propinguus demonstrates that leucism can occur and persist in natural populations for long periods despite the presumed disadvantage (such as increased vulnerability to predation or low genetic diversity) of leucistic phenotypes (Momot & Gall, 1971).

The occurrence of leucism in other taxa has been associated with mutations arising from radioactive pollution from a nuclear power plant (Ellegren et al., 1997; Møller & Mousseau, 2001) and inbreeding (Owen & Skimmings, 1992; Bensch et al., 2000), but the exact mechanism (i.e., physiological or genetic) causing such leucistic phenotypes is unknown. It is unclear if inbreeding is present in this population, or whether nuclear pollution is a likely cause of leucism in this instance. Nevertheless, it can be speculated that the recent impact of logging combined with substantial anthropogenic waste, such as plastic food bags and polystyrene foam cups and containers, in the particular habitat could have been the likely cause. We unfortunately did not collect water samples to determine water quality (e.g., dissolved oxygen, turbidity, nitrates, pH, and other potential factors). Because populations of primary burrowing crayfishes presumably have a reduced dispersal capacity and a reduced gene flow across populations compared to tertiary burrowing crayfishes (Hurt et al., 2019; Clay et al., 2020), the resulting atypical color phenotype from inbreeding may persist over time and eventually become common. It can also be

hypothesized that the appearance of abnormal colorations in primary burrowing crayfishes may be related to the unknown function of conspicuous coloration (orange, red, blues, etc.) exhibited by many primary burrowing crayfish species. Primary burrowing crayfishes presumably face reduced predation pressure due to their burrowing lifestyle compared to tertiary or secondary burrowing crayfishes inhabiting a permanent body of water. Since atypical color phenotypes like leucism presumably arise due to genetic mutations or inbreeding, there may be little to no selection against this trait once they emerge. These ideas, however, are entirely speculative, but are a potentially fertile area of research that could elucidate the genetic and ecological mechanisms relating to crayfish color phenotypes and color abnormalities.

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