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Animal Pigment Bilirubin Discovered in Plants

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Cyclic tetrapyrroles occur throughout the plant kingdom and include vital biosynthetic products such as chlorophyll and heme. In plants, oxidative degradation of heme forms first biliverdin-IXα and subsequently phytochromobilin, the precursor of the phytochrome chromophore, an essential light sensing molecule.[1] In animals, oxidative degradation of heme also leads to the formation of biliverdin-IXα, but it is transformed into the yellow-orange pigment bilirubin-IXα. Here, we present spectroscopic and chromatographic evidence that bilirubin (Figure 1) is the major pigment of the orange aril of *Strelitzia nicolai* Regel & Koern. (Strelitziaceae, order Zingiberales), the white bird of paradise tree.

This is the first example of bilirubin in a plant,[2] a finding which likely necessitates the revision of the plant tetrapyrrole pathway since there is currently no known mechanism of bilirubin production in the plant kingdom.

S. nicolai is native to South Africa and widely cultivated in the tropics. It produces woody capsular fruits which contain orange arillate seeds. Analytical high-performance liquid chromatography (HPLC) of the aril extract[3] revealed one major peak, which had a UV−visible spectrum with a maximum absorbance at 444 nm (Figure S1). After purification using preparative scale HPLC, the isolated pigment was analyzed by ^{1}H NMR, ^{13}C NMR (Brucker, 400 MHz, $(CD_3)_2S{=}O$), and liquid chromatography−positive ion electrospray mass spectrometry (LC−ESI) (Thermo-Finnigan LCQ).

The ^{1}H NMR and ^{13}C NMR spectra of the isolated pigment matched published values of authentic bilirubin[4,5,10] (Tables S1 and S2). Identification was further supported by ^{1}H NMR analysis of bilirubin standard (Aldrich). This yielded a spectrum which matched that of the *S. nicolai* pigment. Both the positive ion ES mass spectrum and the product ion spectrum matched those of authentic bilirubin standard and previous published data[6] (molecular ion, *m*/*z* 585 $(M + H)^+$, product ion *m*/*z* 299).

Given the unexpected discovery of bilirubin in plants, it was essential to confirm the identity of the pigment as bilirubin-IXα, and not other isomers. Previous chromatographic studies have demonstrated that the ability of bilirubin-IXα to undergo intramolecular hydrogen bonding makes it significantly less polar than bilirubin-IX β, γ, or δ[7,8] (Figure 1).

In our HPLC analyses, a single peak was observed when bilirubin-IXα standard was coinjected with the isolated pigment, thereby eliminating the possibility that the pigment was bilirubin-IX β, γ, or δ. Furthermore, the visible spectrum of bilirubin-IXα has an intense peak at 458 nm in dimethylsulfoxide (DMSO), which is approximately 50 nm longer than bilirubin-IX β, γ, or δ.[7] Other bilirubin isomers, including bilirubin-IIIα and bilirubin-XIIIα, were eliminated because their ^{1}H NMR spectra are substantially different.[9]

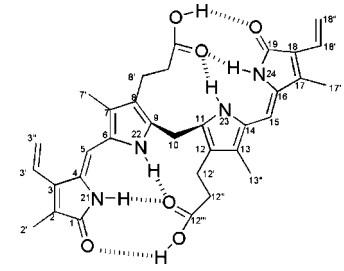

Figure 1. Bilirubin-IXα demonstrating intramolecular H-bonding.

The occurrence of bilirubin is not restricted to *S. nicolai*. Two other species in the Strelitziaceae, *Phenakospermum guyanense* Endl., and *S. reginae* Aiton, the bird of paradise, contain aril pigments which coeluted with authentic bilirubin in HPLC and had similar UV−visible spectra. We are currently examining species in related families. This information, in combination with studies on the synthesis of bilirubin-IXα in *S. nicolai*, will provide the basis for a more thorough understanding of the evolutionary origin of this pigment in plants.

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Supporting Information Available: Methods, UV−vis spectrum, and ^{1}H and ^{13}C NMR data. This material is available free of charge via the Internet at http://pubs.acs.org.

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