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Verdazyl [2,4-diphenyl-6-aryl-3,4-dihydro-sym-tetrazin-1(2H)-yl] radicals I and verdazylium salts II are readily interconvertible; in studies of the reactions of these compounds they
are often simultaneously present in solution [1]. We have found that I and II in relatively
concentrated solutions react with one another to give 1,5-diphenyl-3-aryl-6-(2,4-diphenyl-6aryl-1,2,3,4-tetrahydro-sym-tetrazin-1-yl)-5,6-dihydro-sym-tetrazinium salts III and leucoverdazyls:

III a  $\operatorname{Ar}^1 = \operatorname{Ar}^2 = \operatorname{C}_6 \operatorname{H}_5$ ; b  $\operatorname{Ar}^1 = \operatorname{C}_6 \operatorname{H}_5$ ,  $\operatorname{Ar}^2 = \operatorname{C}_6 \operatorname{H}_4 \operatorname{OCH}_3 - p$ ; c  $\operatorname{Ar}^1 = \operatorname{C}_6 \operatorname{H}_4 \operatorname{OCH}_3 - p$ ,  $\operatorname{Ar}^2 = \operatorname{C}_6 \operatorname{H}_5$ ;

Salt II evidently acts as a CH acid in this reaction.

A solution of verdazyl radical I ( $\sim$ 0.2 mole) and salt II ( $\sim$ 0.1 mole) in dry acetonitrile was refluxed for  $\sim$ 20 min, after which the mixture was diluted with absolute ether, and salt III precipitated in the form of fine brown crystals in  $\sim$ 85% yield. Salt IIIa (X = Br) had mp 220-222°C (dec.). IR spectrum (KBr): 1600 (C=N) and 2900 cm<sup>-1</sup> (CH<sub>2</sub>). Electronic spectrum (CH<sub>3</sub>CN):  $\lambda_{max}$  490 nm (log  $\epsilon$  4.37). PMR spectrum (CF<sub>3</sub>COOH): 6.6-8.0 (m, 30 aromatic H + 6-H) and 5.2 ppm (s, 2H, CH<sub>2</sub>).

Leucoverdazyls IV were isolated in  $\sim 90\%$  yields in the form of hydrochlorides [2] by the addition of concentrated hydrochloric acid to ether solutions.

Salts III were stable in the solid state. In aqueous acetonitrile they are rapidly hydrolyzed and oxidized, and bright-blue salts precipitate. A salt with the composition  $C_{40}H_{33}BrN_8O$  was obtained from IIIa (X = Br). Electronic spectrum (CH<sub>3</sub>CN):  $\lambda_{max}$  655 nm (log  $\epsilon$  4.11). Treatment of a solution of this salt (or IIIa) in CH<sub>3</sub>CN with aqueous alkali gave velvet-green crystals of the  $C_{40}H_{32}N_8O$  base. Electronic spectrum (CH<sub>3</sub>CN),  $\lambda_{max}$  (log  $\epsilon$ ): 585 (4.08), 845 (4.14), and 925 nm (4.12). Molecular weight (M) (by the Rast method) 645; the calculated value was 640.5.

Verdazyl radicals and verdazylium salts undergo virtually no reaction with one another in solutions at low concentrations ( $\sim 10^{-4}$  mole/liter).

## LITERATURE CITED

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