REPLACEMENT OF THE ACINITRO GROUP BY AN ARYLAZO GROUP IN 1,4-DINITROCYCLOOCTATETRAENE AND 1,8-DINITRONAPHTHALENE DIANIONS

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UDC 542.91:541.6:547.556.33

The reaction of 1,4-dinitrocyclooctatetraene and 1,8-dinitronaphthalene with the dipotassium salt of the cyclooctatetraene dianion ( $C_8H_8K_2$ ) leads to the formation of dianions (I) and (II), respectively. Dianions (I) and (II) in THF at -10 to -50°C react with anyldiazonium tetrafluoroborides p-RC<sub>6</sub>H<sub>4</sub>N<sub>2</sub>BF<sub>4</sub> [R = NO<sub>2</sub>, SCN, OCH<sub>3</sub>, and N(CH<sub>3</sub>)<sub>2</sub>] to form the products of the substitution of the acinitro group by the arylazo group in yields up to 90%. The substitution in dianion (I) affects one or both nitro groups with the formation of the corresponding mono or bisazo derivatives of cyclooctatetraene [only the monoazo derivative was obtained for R = N(CH<sub>3</sub>)<sub>2</sub>].

In all cases, only one nitro group in diamion (II) is substituted, leading to 1-nitro-8-arylazonaphthalenes.

The synthesis of the dipotassium salts of diamions (I) and (II) and monitoring of their formation were carried by analogy to our previous procedure [1]. The structure of the azo dyes obtained was established by PMR, IR, and mass spectroscopy, elemental and x-ray diffraction structural analysis.

## LITERATURE CITED

1. Z. V. Todres, Izv. Akad. Nauk SSSR, Ser. Khim., 1749 (1970).

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