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Publication details, including instructions for authors and subscription information: <a href="http://www.informaworld.com/smpp/title~content=t713618290">http://www.informaworld.com/smpp/title~content=t713618290</a>

## The Novel Phosphadiazacalixcrown Compounds

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**To cite this Article** Antipin, Igor S., Stoikov, Ivan I., Nikonov, Grigory N. and Konovalov, Alexander I.(1999) 'The Novel Phosphadiazacalixcrown Compounds', Phosphorus, Sulfur, and Silicon and the Related Elements, 147: 1, 9

To link to this Article: DOI: 10.1080/10426509908053485 URL: http://dx.doi.org/10.1080/10426509908053485

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# The Novel Phosphadiazacalixcrown Compounds

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A new series of calixcrown compounds (II) containing some VA group elements (N, P) was synthesized. 1,3-bis-(aminoethoxy) calix[4]arene (I) was used as starting platform for the preparation of calixcrown compounds. Reaction of (I) with bis( $\alpha$ -hydroxyalkyl)phenylphosphines in toluene lead to phosphadiazacalixcrown compounds (II). All obtained receptors are hydrolytic stable and are not oxidized by air. A more convenient synthesis of (II) involves treatment of (I) with 2,5-diphenyl-1,3,2,5-dioxaboraphosphorinanes (III) which are stable on air unlike bis-( $\alpha$ -hydroxyalkyl) phenylphosphines. Structure of obtained macrocycles was established by <sup>1</sup>H and <sup>31</sup>P NMR spectroscopy.

To enchance the binding ability of the novel receptors to cations and pronodonor molecules the macrocycles (II) were oxidized with the near-quantitative yields to corresponding phosphine oxides (III) by a hydrogen peroxide in acetone or phosphine thiooxides by a sulfur addition.

Phosphine oxides (III) possess an array of potential binding sites for interaction with  $\alpha$ -amino and hydroxy acids. Unsubstituted macrocycle (III) (R=H) will be examine as earrier for membrane transport of the zwitterionic form of aromatic amino acids.

This investigation was supported by Russian Foundation for Basic Research (grant N 98-03-33051).