Viscosity-Controlled Stereoselective Inversion during the Photochemical Denitrogenation of a Stereolabeled Diazabicyclo[2.1.1]heptene(DBH)-Type Azoalkane

Waldemar Adam,\*†Vicente Martí,†Coskun Sahin† and Alexei V. Trofimov†,‡

<sup>†</sup>Institute of Organic Chemistry, University of Würzburg

D-97074 Würzburg, Germany

E-mail: adam@chemie.uni-wuerzburg.de, Fax: +49 (931) 8884756,

Internet: http://www-organik.chemie.uni-wuerzburg.de

<sup>‡</sup>Institute of Biochemical Physics, Russian Academy of Sciences

117977 Moscow, Russia

Synthesis of Starting materials: The azoalkane 1 and housanes 2(inv) and 2(ret) were synthesized and characterized as previously described.

Photolysis and Product Studies: A solution of 0.07 mmol of the azoalkane 1 in the particular solvent (distilled prior to use) was transferred to an NMR tube, deaerated with a stream of argon gas for 10 min, and irradiated with the 333-, 353- and 364-nm laser lines of the argonion laser (INNOVA 100, Coherent Company) at 20 °C and 760 torr. The conversions, mass balances, and product distributions (error ca. 5% of the stated value) were determined by quantitative capillary GC analysis and corrected for the response factors of the starting material and the products. These analyses were performed on a GC 6000 Vega Series 2 equipped with a FID and a Shimadzu C-R 6 A electronic integrator. A 30-m RTX 1 wide-bore capillary column with an integral diameter of 0.53 mm and a film thickness of 1.5 μm was used.