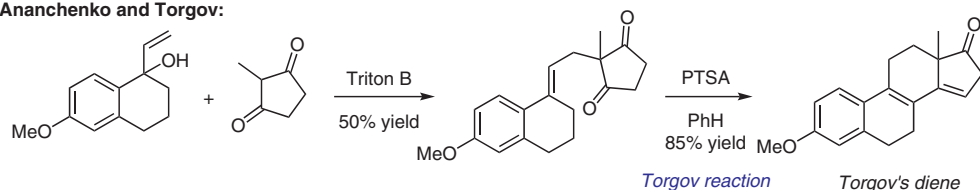


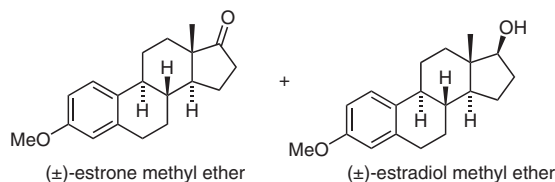
S. N. ANANCHENKO, I. V. TORGOV\* (USSR ACADEMY OF SCIENCE, MOSCOW, USSR)  
New Syntheses of Estrone, d,l-8-Iso-oestrone and d,l-19-Nortestosterone  
*Tetrahedron Lett.* **1963**, 4, 1553–1558.

## Torgov Cyclization

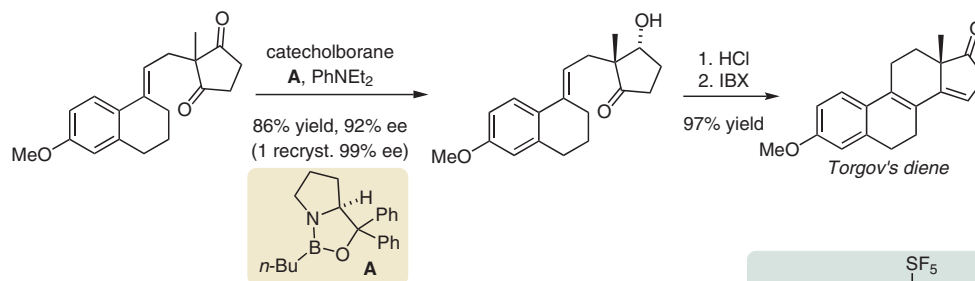
### Ananchenko and Torgov:



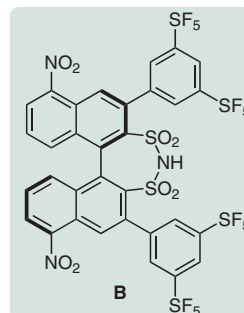
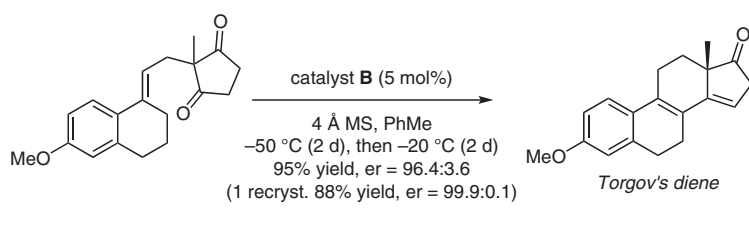
1. Pd/CaCO<sub>3</sub>, H<sub>2</sub>  
2. K, NH<sub>3</sub>  
1:1 mixture of products



### Corey and co-workers:



### List and co-workers:



**Significance:** In 1963, Ananchenko and Torgov described the carbocyclization of an achiral diketone to give a tetracyclic intermediate (Torgov's diene) that provided an early pathway for the total synthesis of steroid hormones. This method has been used for the industrial-scale production of certain steroids. However, an enantioselective variant of the Torgov reaction has long remained elusive.

**Comment:** Corey and co-workers reported an enantioselective and diastereoselective reduction of the achiral diketone by using an oxazaborolidine catalyst A (*J. Am. Chem. Soc.* **2007**, 129, 10346). Treatment of the mono-ketone with methanolic HCl followed by IBX oxidation gave Torgov's diene in high optical purity. List and co-workers developed a catalytic asymmetric Torgov cyclization using a novel dinitro-substituted disulfonimide catalyst B (*Angew. Chem Int. Ed.* **2014**, 53, 8770).