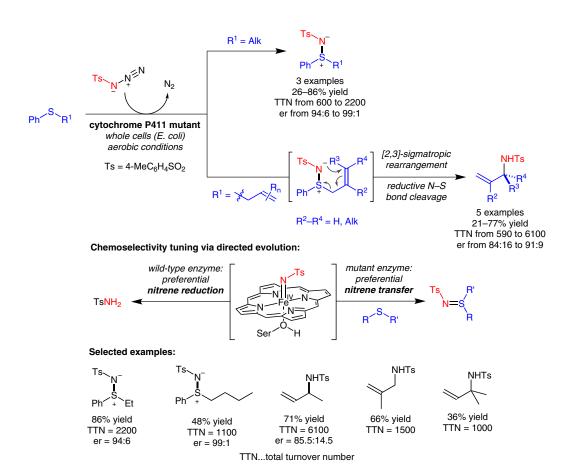
C. K. PRIER, T. K. HYSTER, C. C. FARWELL, A. HUANG, F. H. ARNOLD\* (CALIFORNIA INSTITUTE OF TECHNOLOGY, PASADENA, USA)

Asymmetric Enzymatic Synthesis of Allylic Amines: A Sigmatropic Rearrangement Strategy *Angew. Chem. Int. Ed.* **2016**, *55*, 4711–4715.

## **Enzymatic Nitrene Transfer/Sigmatropic Rearrangement to Access Allylic Amines**



**Significance:** Arnold and co-workers report an enzymatic synthesis of allylic amines through a sulfimidation/[2,3]-sigmatropic rearrangement of phenyl allyl sulfides with tosyl azide. A mutant variant of cytochrome P411 from *Bacillus megaterium* efficiently catalyzes a highly enantioselective nitrene transfer to the sulfides, and permits a subsequent rearrangement with partial retention of the stereochemical information. In a scale-up experiment with reduced catalyst loading, 0.1 mmol of substrate was converted into the corresponding allylic amine in 71% yield and a remarkable total turnover number of 6100.

**SYNFACTS Contributors:** Benjamin List, Lucas Schreyer Synfacts 2016, 12(06), 0635 Published online: 17.05.2016 **DOI:** 10.1055/s-0035-1562137; **Reg-No.:** B02216SF

**Comment:** The authors have successfully employed directed evolution to achieve a chemoselective nitrene transfer over the competing reduction. Furthermore, they achieved a sigmatropic rearrangement of the intermediate allylic sulfimides, a process unknown in wild-type biological systems.

## Category

Organo- and Biocatalysis

## Key words

sigmatropic rearrangement

nitrene

directed evolution

allylic amines

enzymes

