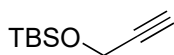


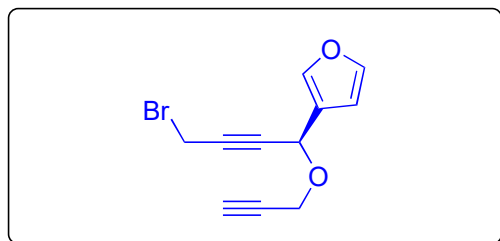
Enantioselective Total Synthesis of (+)-Salvileucalin B

Sergiy Levin, Roger R. Nani, and Sarah E. Reisman.

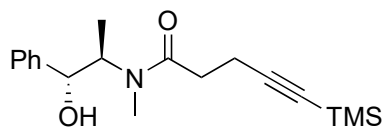
J. Am. Chem. Soc. **2011**, 133, 774-776.



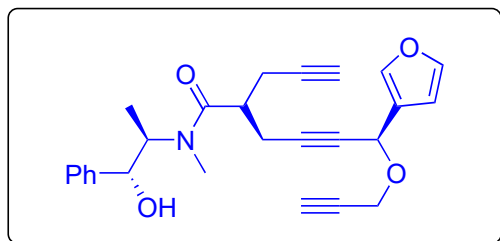
1 - 4



A

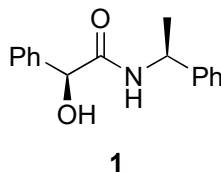


5 - 6

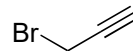


B

- 1) Me_2Zn , **1**, PhMe, 70 °C then 3-furaldehyde, 0 to 22 °C
- 2) NaH, **2**, DMF
- 3) 1M HCl, MeOH
- 4) MsCl, NEt_3 , THF then LiBr



1

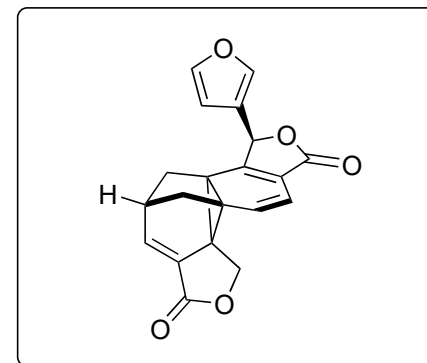


2

- 5) LiHMDS, LiCl, THF, -78 to 22 °C then **A**, -78 °C
- 6) TBAF, CH_2Cl_2

- 1) Name of chiral ligand **1**?
[Mandelamide.](#)

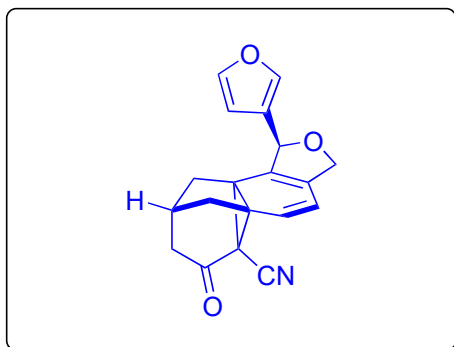
- 5) Name of the auxiliary on the starting material? Who developed this chemistry?
[Pseudoephedrine.](#) [Myers.](#)



(+)-Salvileucalin B

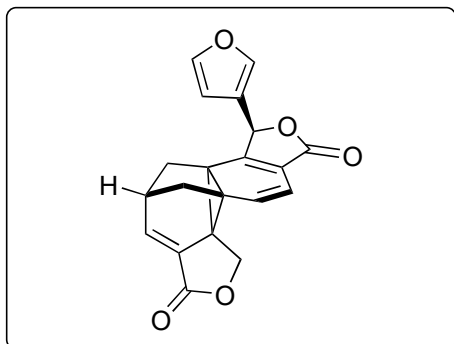
B

7-13



C

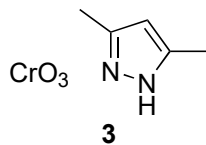
14-18



(+)-Salvileucalin B

- 7) $\text{RuCp}^*(\text{cod})\text{Cl}$ (8 mol%), CH_2Cl_2 , 45 °C
- 8) $n\text{-Bu}_4\text{NOH}$, $t\text{-BuOH}/\text{H}_2\text{O}$, 90 °C
- 9) $(\text{COCl})_2$, DMF (cat.) then CH_2N_2 , THF
- 10) AgTFA , MeOH, NEt_3 , THF, -30 to 22 °C
- 11) NaCH_2CN , THF, -78 to 22 °C
- 12) $(\text{imid})\text{SO}_2\text{N}_3$, pyridine
- 13) $\text{Cu}(\text{hfacac})_2$ (10 mol%), CH_2Cl_2 , 120 °C, microwave

- 14) NaHMDS , -78 °C then Tf_2NPh
- 15) DIBAL , CH_2Cl_2 , -40 °C then AcOH aq.
- 16) DIBAL , CH_2Cl_2 , -40 °C then AcOH aq.
- 17) $\text{Pd}_2(\text{dba})_3$ (5 mol%), dppf (10 mol%), CO, DIPEA, THF
- 18) **3**, CH_2Cl_2 , -35 °C



7) *Hint*: 3 rings are formed.

10) Name of the reaction?
[Arndt-Eistert Homologation.](#)

15) Name of the reaction?
[Retro-Claisen rearrangement.](#)