
Review: Angew. Chem. Int. Ed. 2000

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Group seminar Kalesse & Gaich

2. 11. 2011



Leibniz
Universität
Hannover

Publication Record

- 4617 pages
- 1086 papers in total
- 33 „Total Synthesis“ as research topic

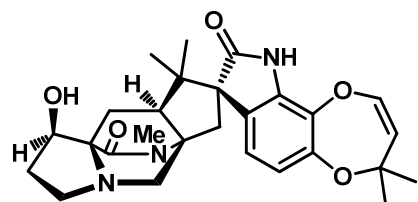
Notable events

- Wladimir Putin is elected President of Russia
- First year of the FPÖ-ÖVP Government in Austria
- American Beauty wins Academy Award for Best Picture
- Maschen-Draht-Zaun by Stefan Raab is No. 1 for 14 weeks
- First permanent resident aboard the ISS
- EXPO 2000 in Hannover

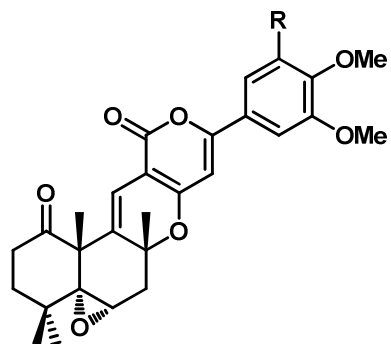
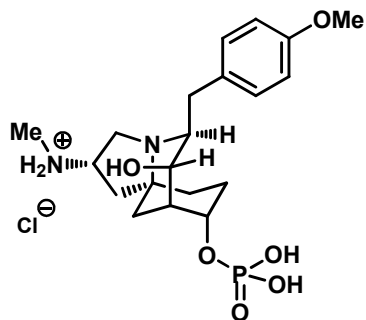
Most cited papers

- 1) (2729) **Ionic Liquids – new „Solution“ for Transition Metal Catalysis;**
Wasserscheid, P.; Keim, W. *Angew. Chem. Int. Ed.* **2000**, *39(21)*, 3772 – 3789;
- 2) (1488) **Olefin Metathesis and Beyond;**
Fürstner, A. *Angew. Chem. Int. Ed.* **2000**, *39(17)*, 3012 – 3043;
- 3) (1363) **Multicomponent Reactions with Isocyanides;**
Domling, A.; Ugi, I. *Angew. Chem. Int. Ed.* **2000**, *39(18)*, 3168 – 3210;

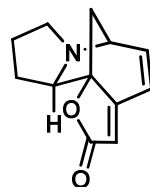
Detailed Syntheses



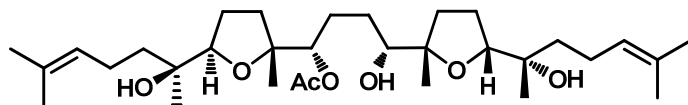
Paraherquamide A

Arisugacin A R = H
Territrems B R = OMe

FR901483

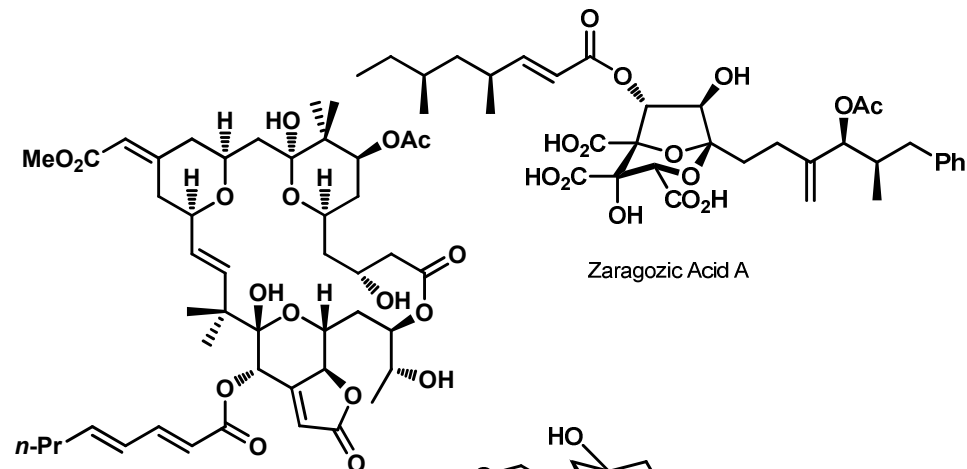


(-)-Norsecurinine



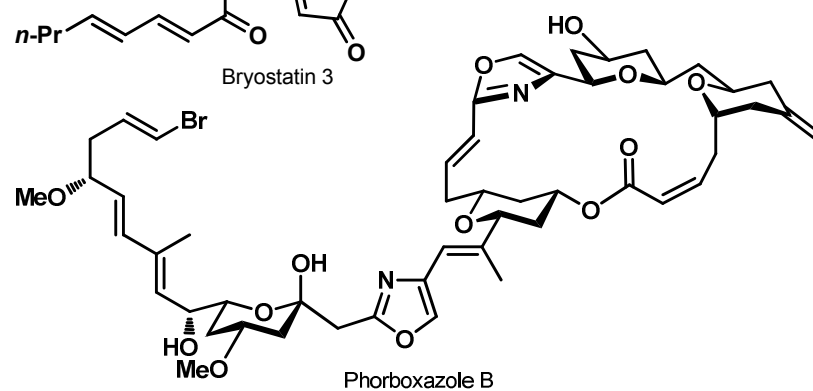
(+) -14-deacetyl eurylene

Concise Syntheses



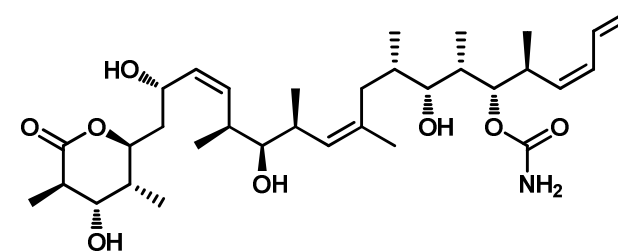
Zaragozic Acid A

Bryostatin 3



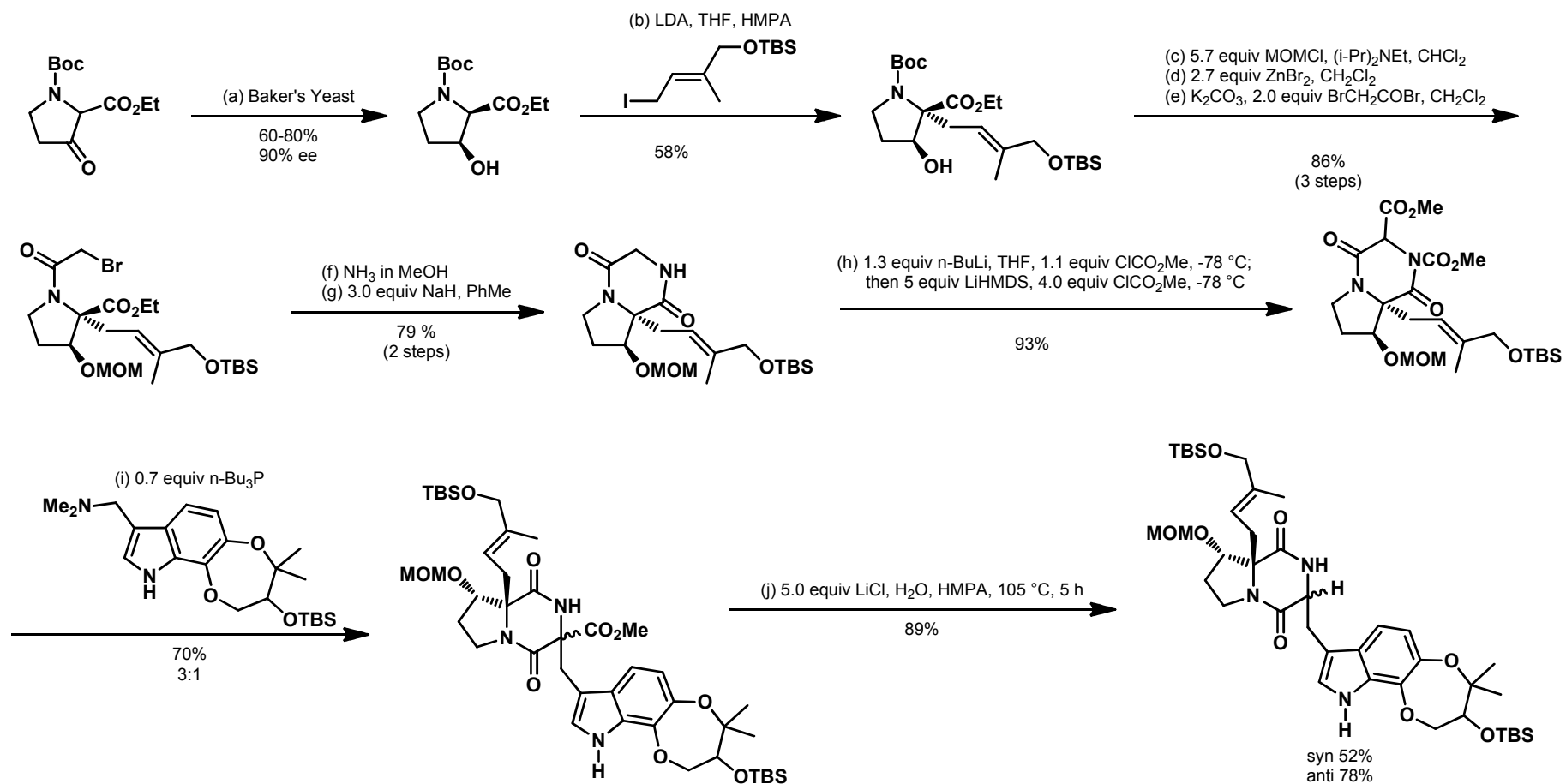
Phorboxazole B

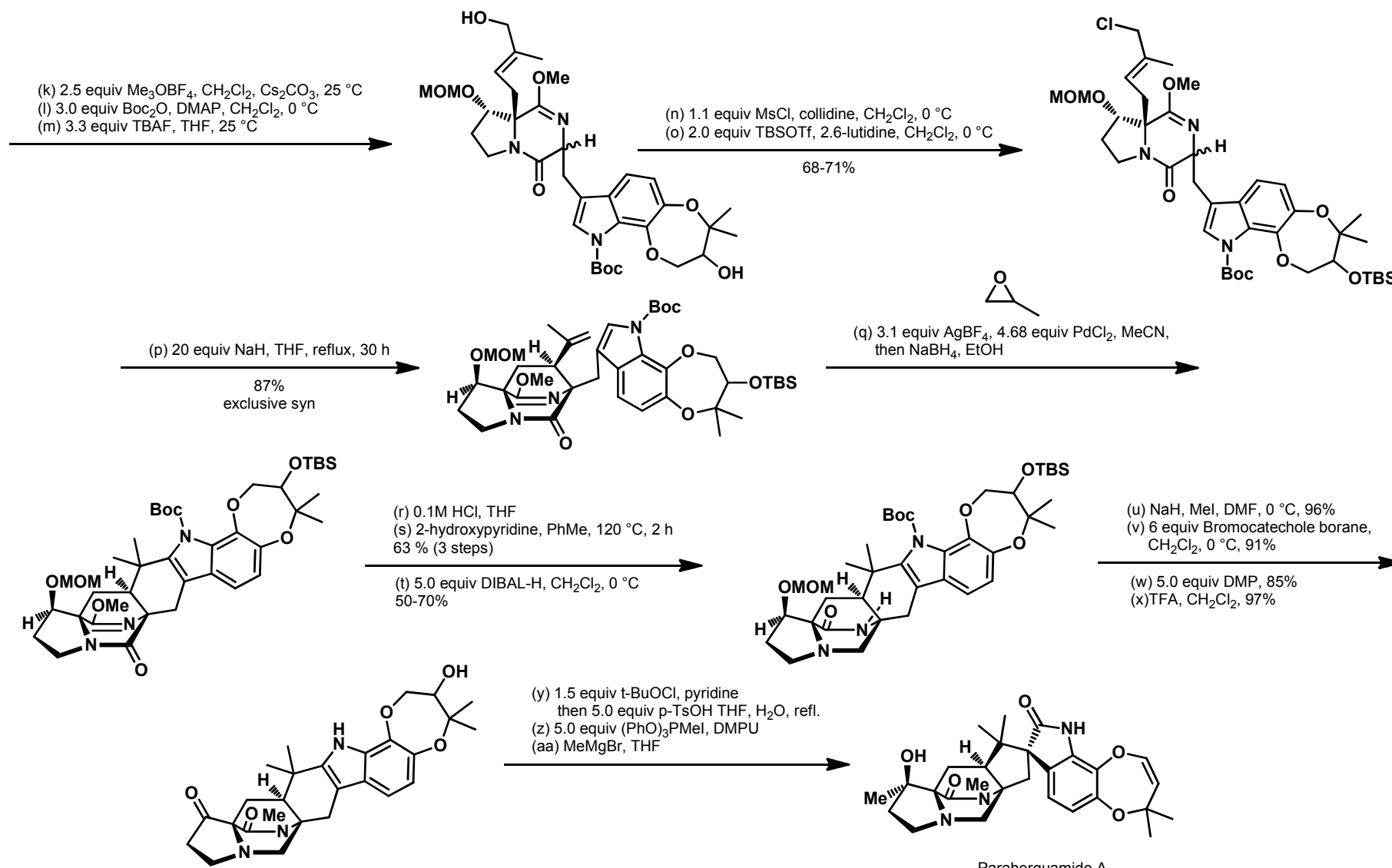
(-)-Spirotyprostatin B



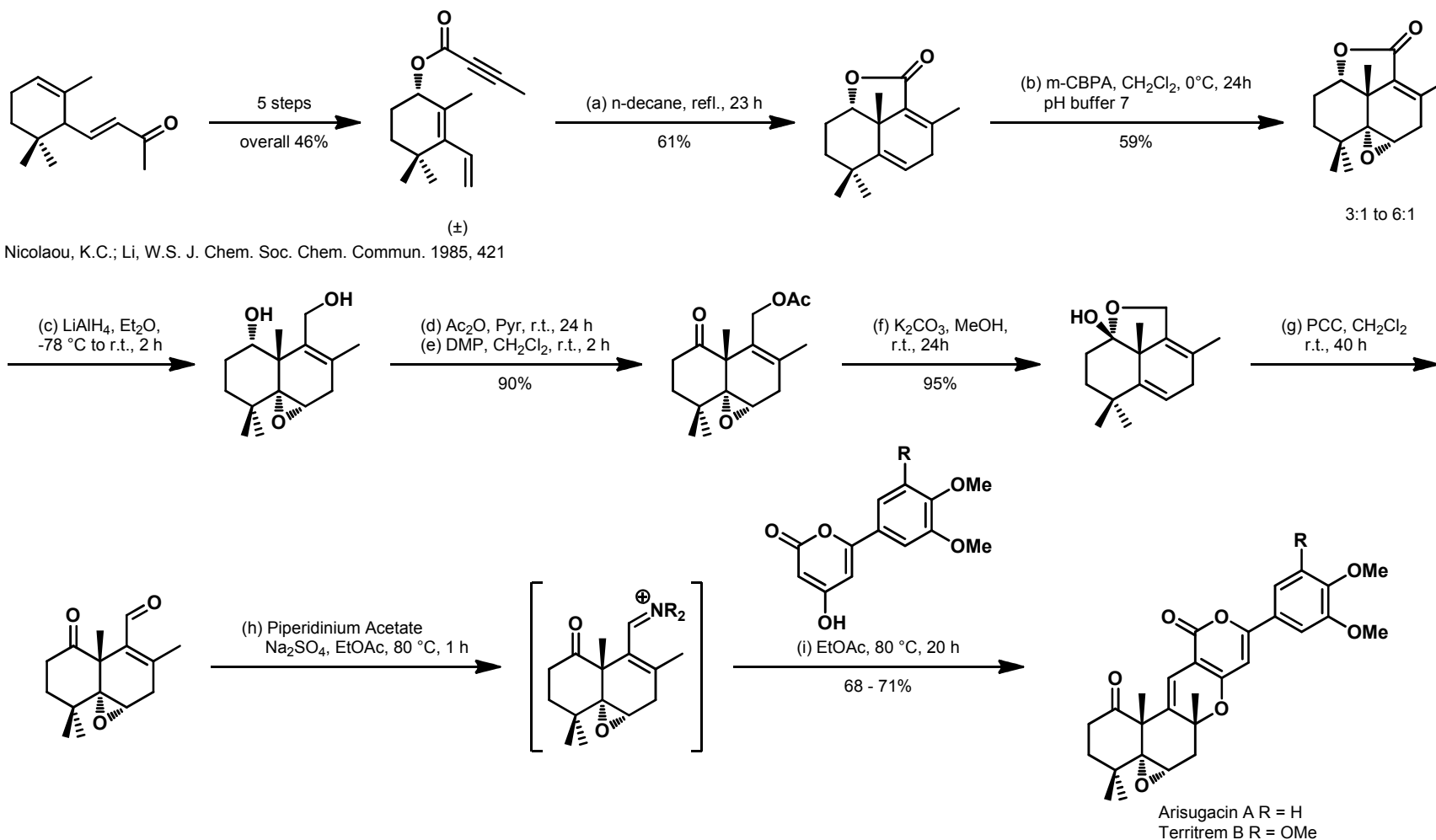
(+) -Discodermolide

Asymmetric, Stereocontrolled Total Synthesis of Paraherquamide A;
Williams, R. M.; Cao, J.; Tsujishima, H. *Angew. Chem. Int. Ed.* 2000, 39(14), 2540 - 2544

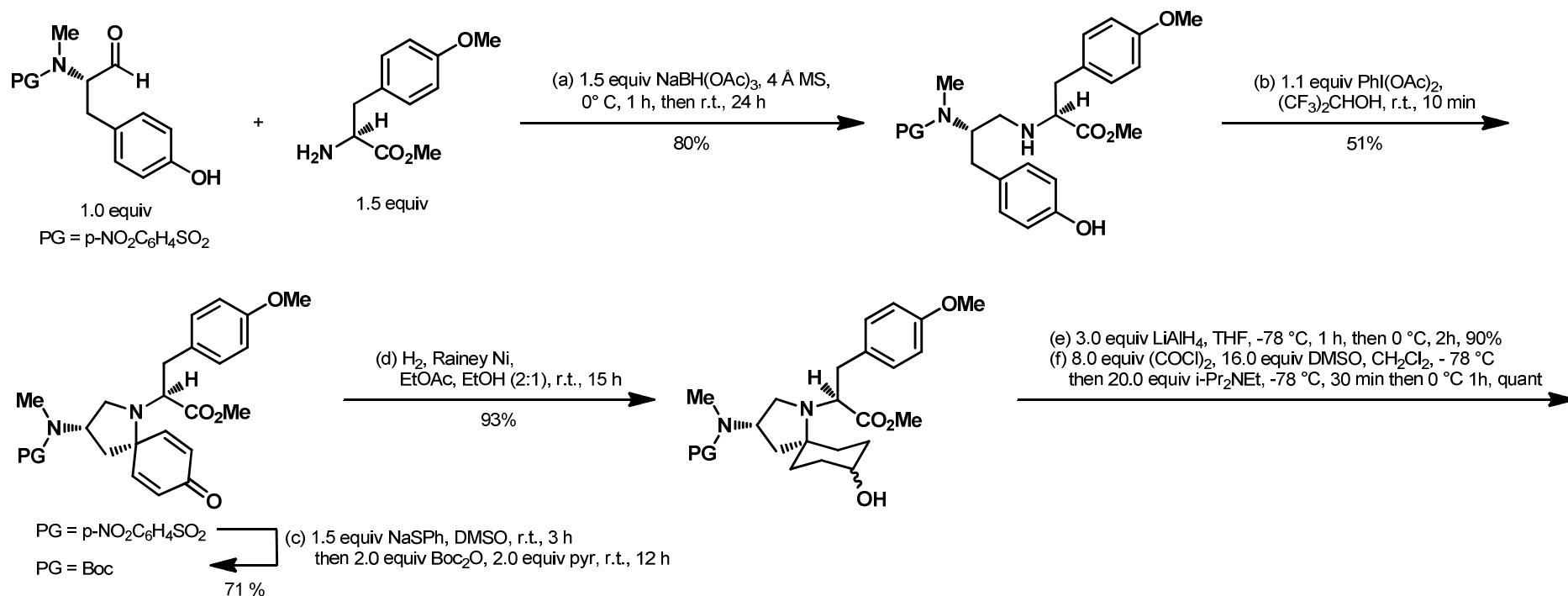


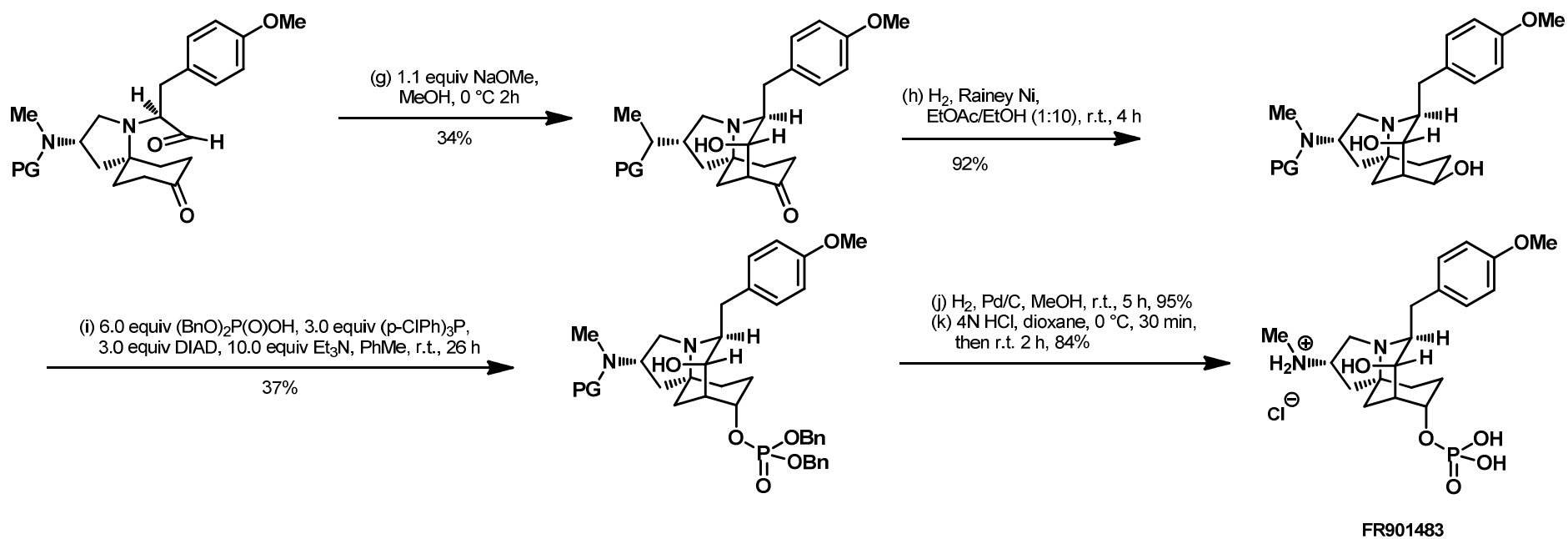


A Concise Stereoselective Route to the Pentacyclic Framework in Arisugacin A and Territrem B;
Hsung, R. P.; Zehnder, L. R.; Wang, J.; Golding, G. M. *Angew. Chem. Int. Ed.* 2000, 39(21), 3876 - 3879



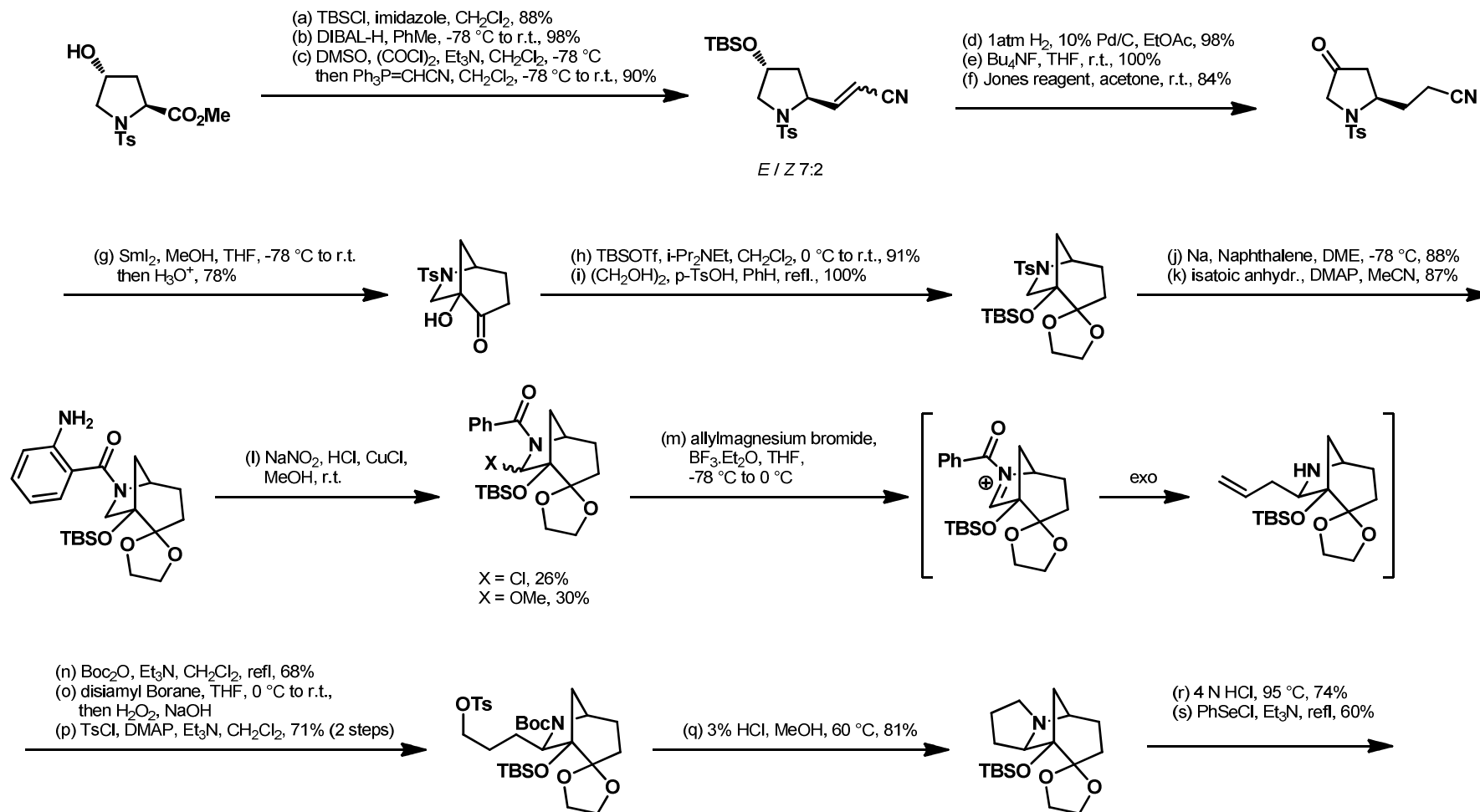
An Enantiospecific Synthesis of the Potent Immunosuppressant FR901483;
Sorensen, E. J.; Scheffler, G.; Seike, H. *Angew. Chem. Int. Ed.* 2000, 39(24), 4593 - 4596

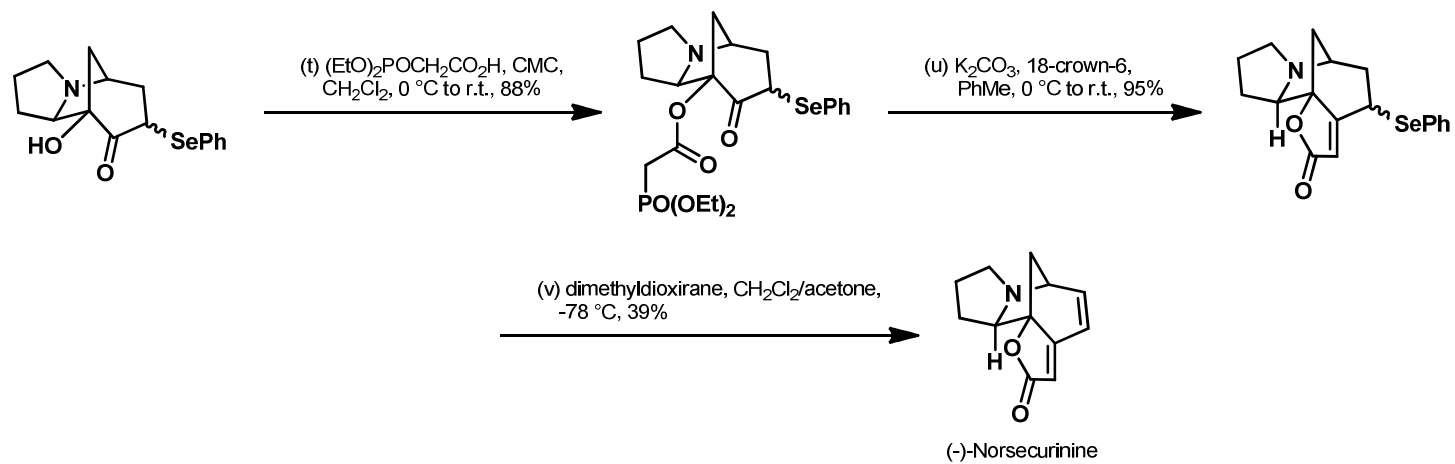




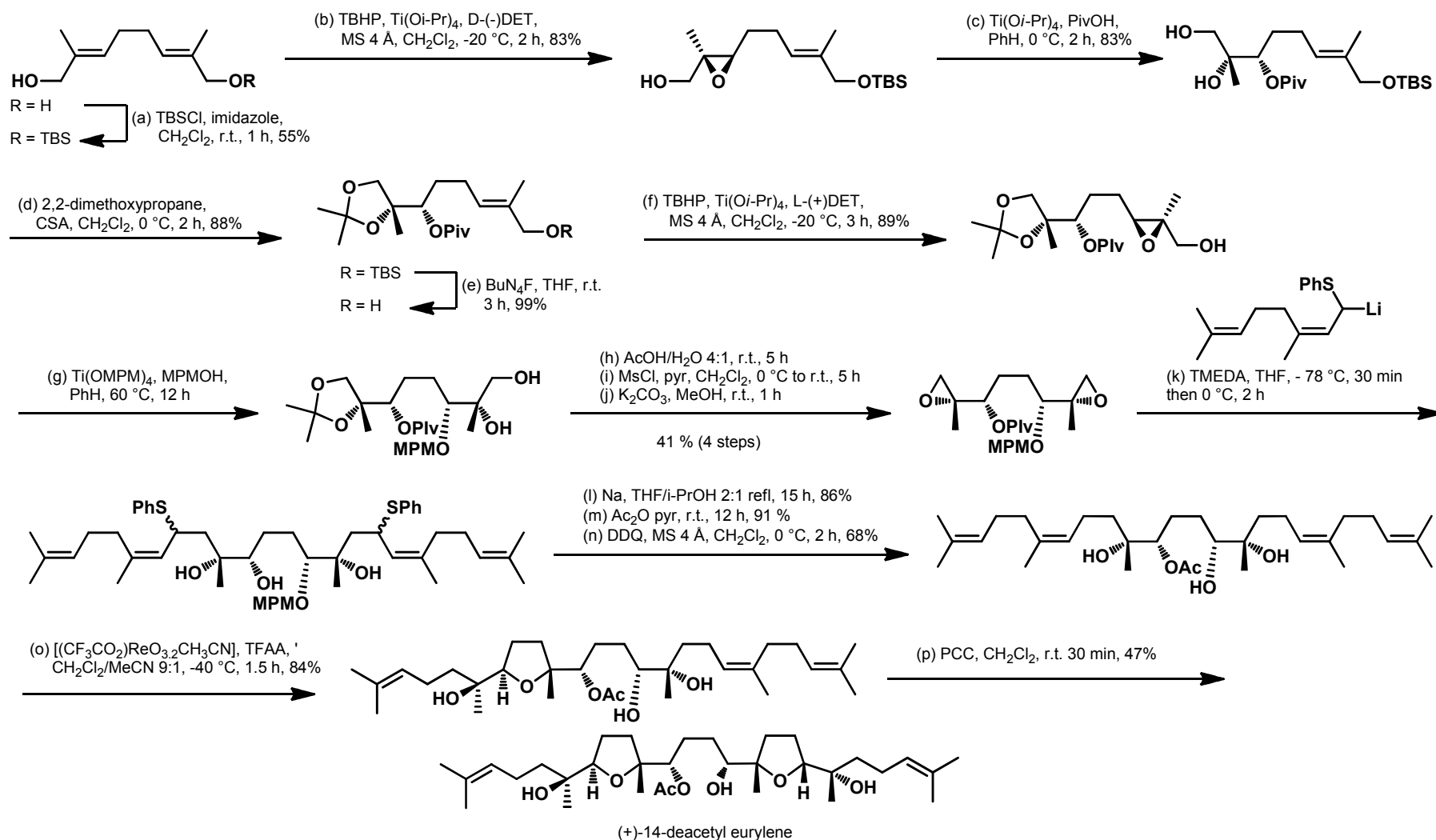
A New Enantioselective Approach to Total Synthesis of the Securinega Alkaloids:
Application to (-)-Norsecurinine and Phyllantine;

Weinreb, S.M.; Han, G.; LaPorte, M. G.; Folmer, J. J.; Werner, K. M.; *Angew. Chem. Int. Ed.* 2000, 39(1), 237 - 240

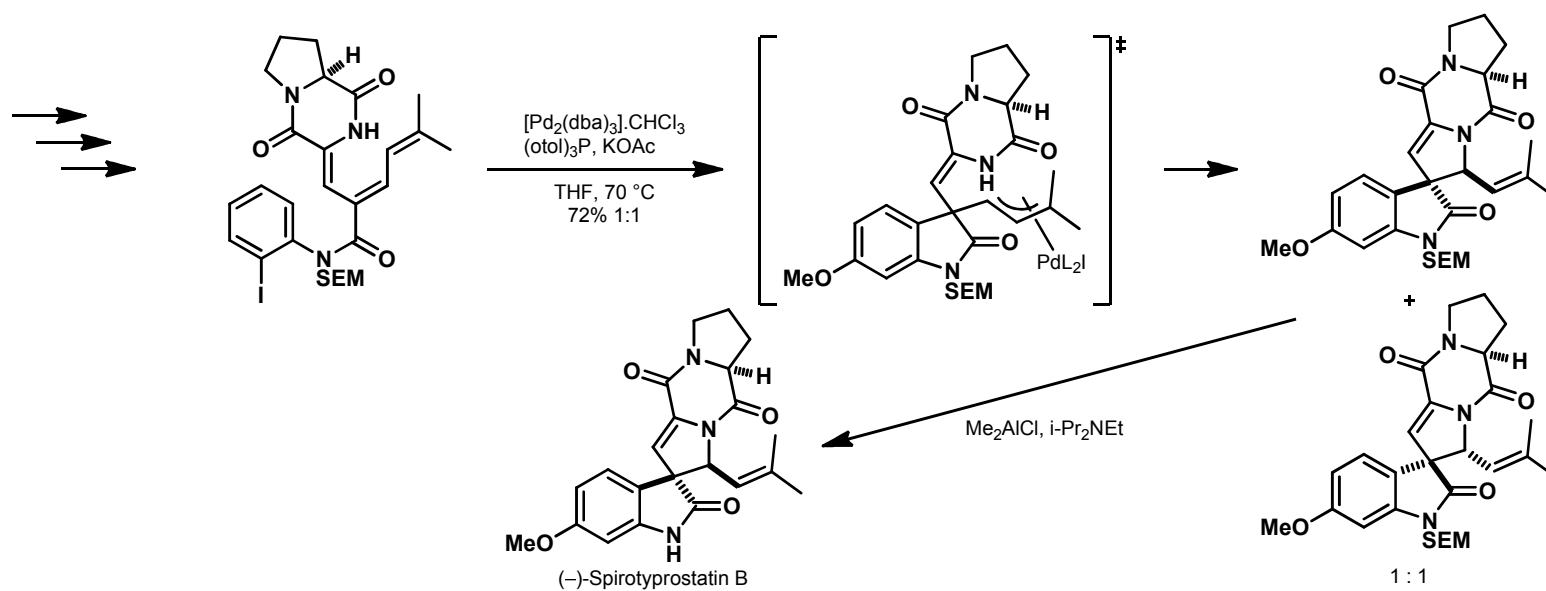




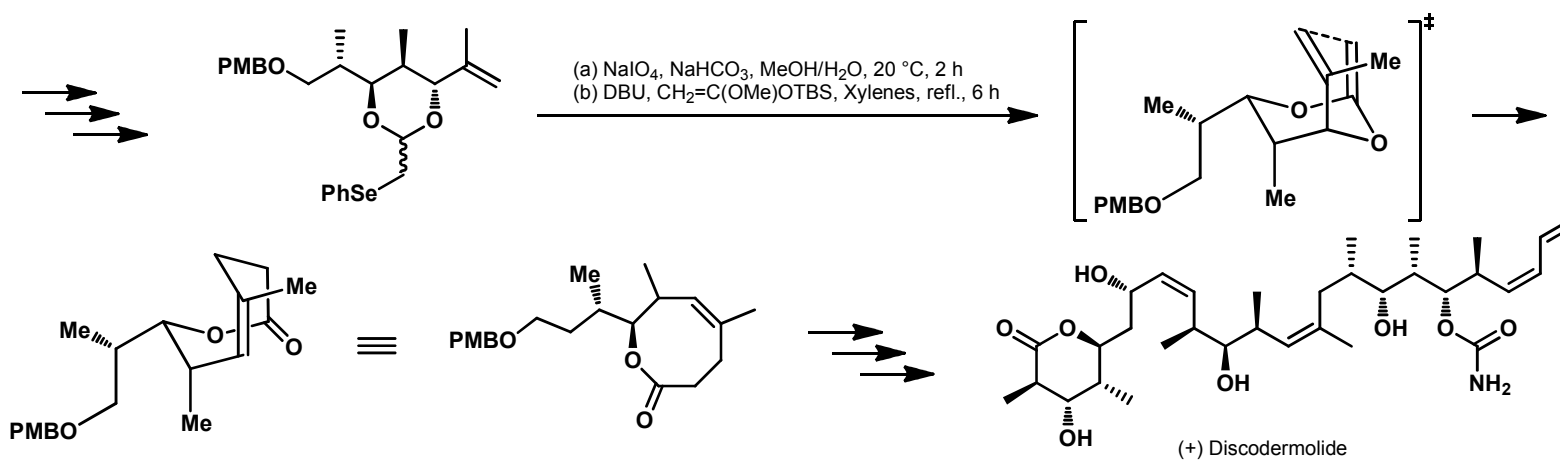
Total Synthesis of (+)-Eurylene and (+)-14-Deacetyleurylene;

Morimoto, Y.; Muragaki, K.; Iwai, T.; Morishita, Y.; Kinoshita, T. *Angew. Chem. Int. Ed.* 2000, 39(22), 4082 - 4084

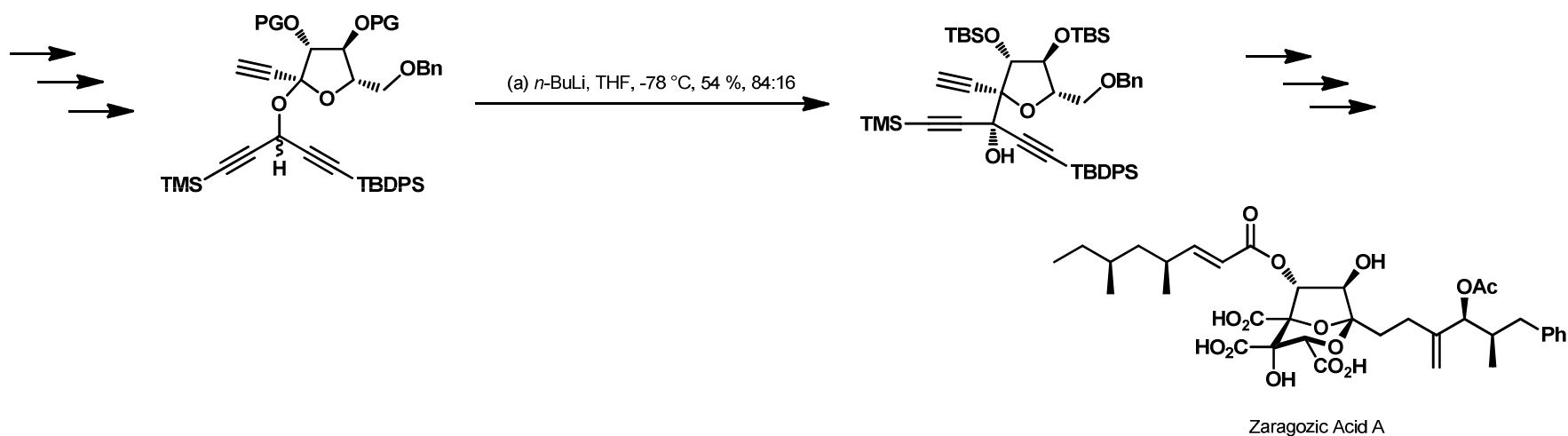
Total Synthesis of (–)-Spirotryprostatin B and Three Stereoisomers;
Overman, L. E.; Rosen, D. M. *Angew. Chem. Int. Ed.* 2000, 39(24), 4596 – 4599



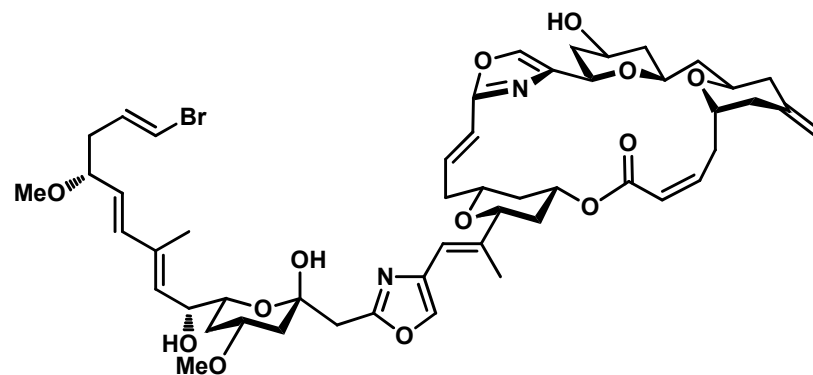
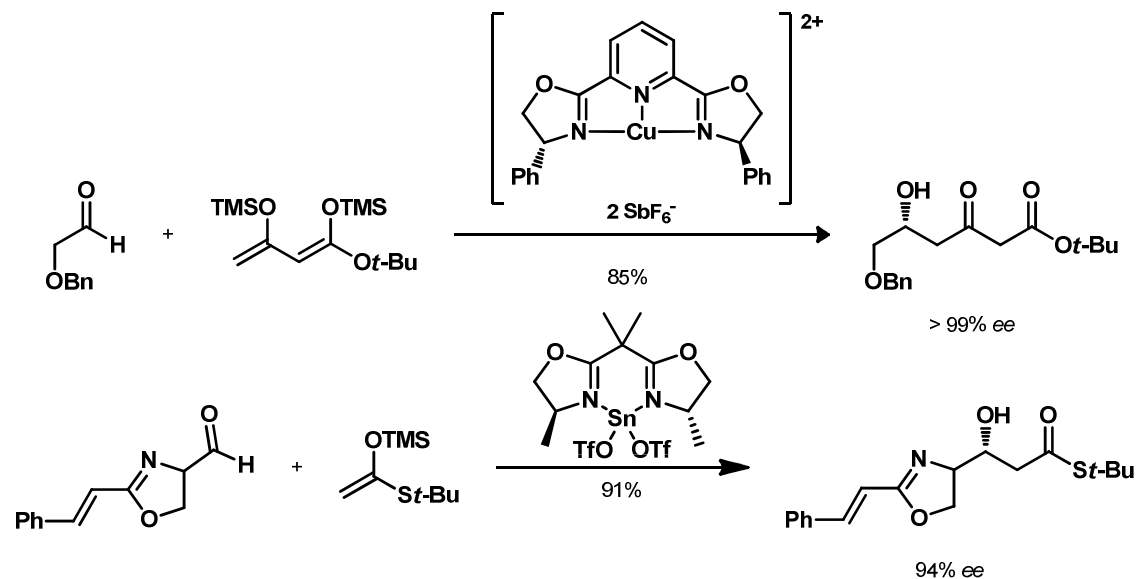
Total Synthesis of the Antimicrotubule Agent (+)-Discodermolide Using Boron-Mediated Aldol Reactions;
 Paterson, I.; Florence, G. J.; Gerlach, K.; Scott, J. P. *Angew. Chem. Int. Ed.* 2000, 39(2), 377- 380



A Stereoselective Total Synthesis of Zaragozic Acid A based on an Acetal [1,2] Wittig Rearrangement;
Tomooka, K.; Kikuchi, M.; Igawa, K.; Suzuki, M.; Keong, P.-H.; Nakai, T. *Angew. Chem. Int. Ed.* 2000, 39(24), 4502 - 4505

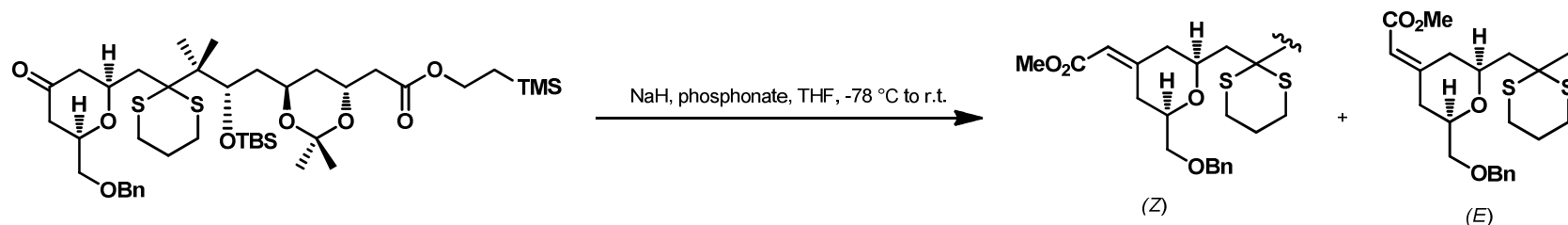


Asymmetric Synthesis of Phorboxazole B – Part I: Synthesis of the C₂₀-C₃₈ and C₃₉-C₄₆ Subunits;
 Evans, D. A.; Cee, V. J.; Smith, T. E.; Fitch, D. M., Cho, P. S. *Angew. Chem. Int. Ed.* 2000, 39(14), 2533 - 2536

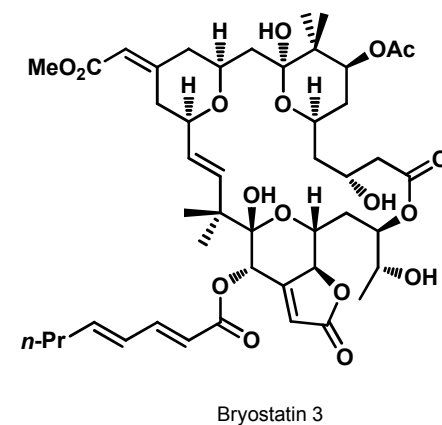


Phorboxazole B

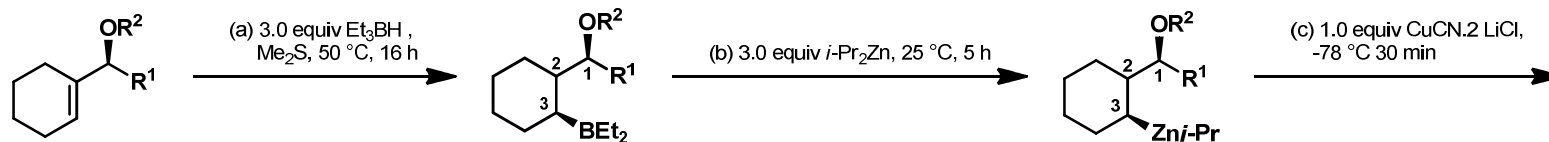
Total Synthesis of Bryostatin 3;

Nishiyama, S.; Yamamura, S.; Ohmori, K.; Ogawa, Y.; Obitsu, T.; Ishikawa, Y. *Angew. Chem. Int. Ed.* 2000, 39(13), 2290 - 2294

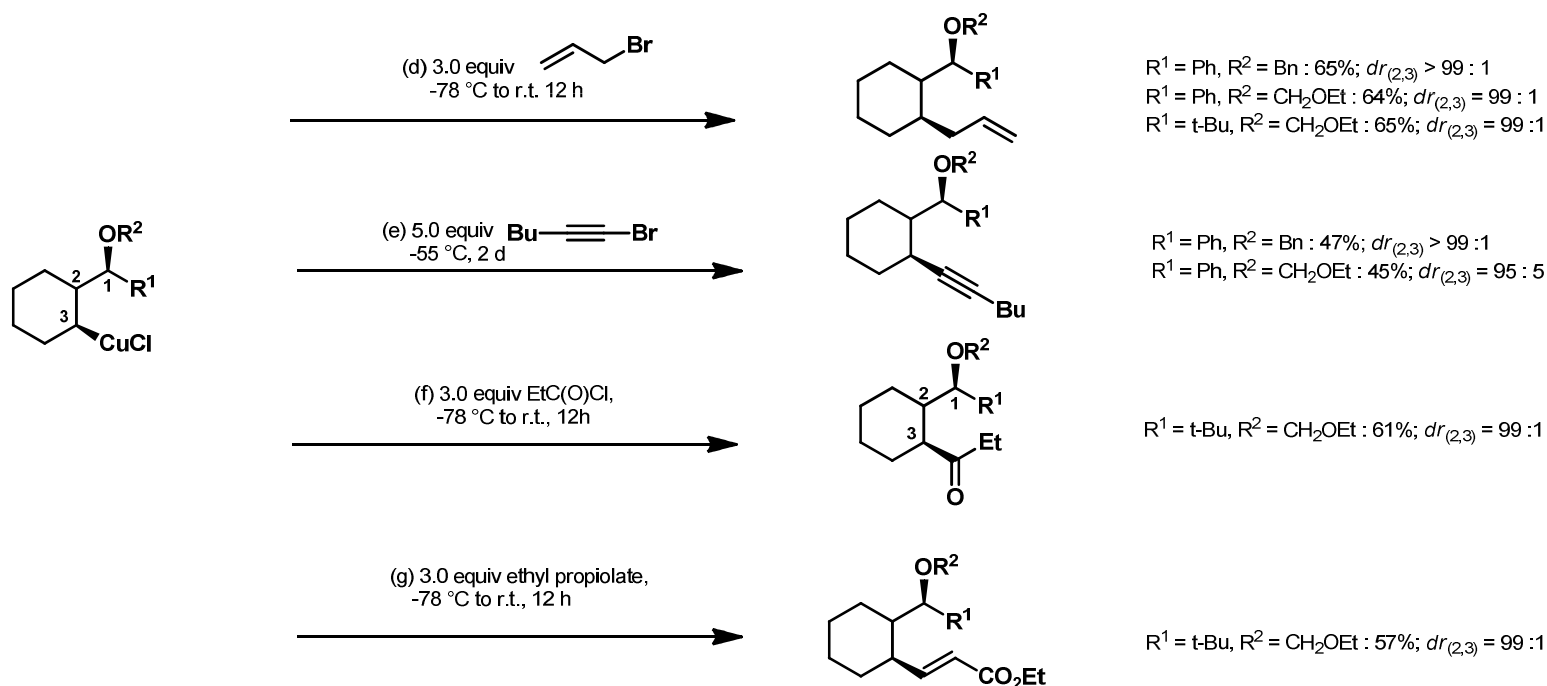
Entry	Phosphonate	Yield	Z : E
1		94	1.6 : 1
2		90	2.0 : 1
3		92	4.0 : 1
4		85	2.3 : 1

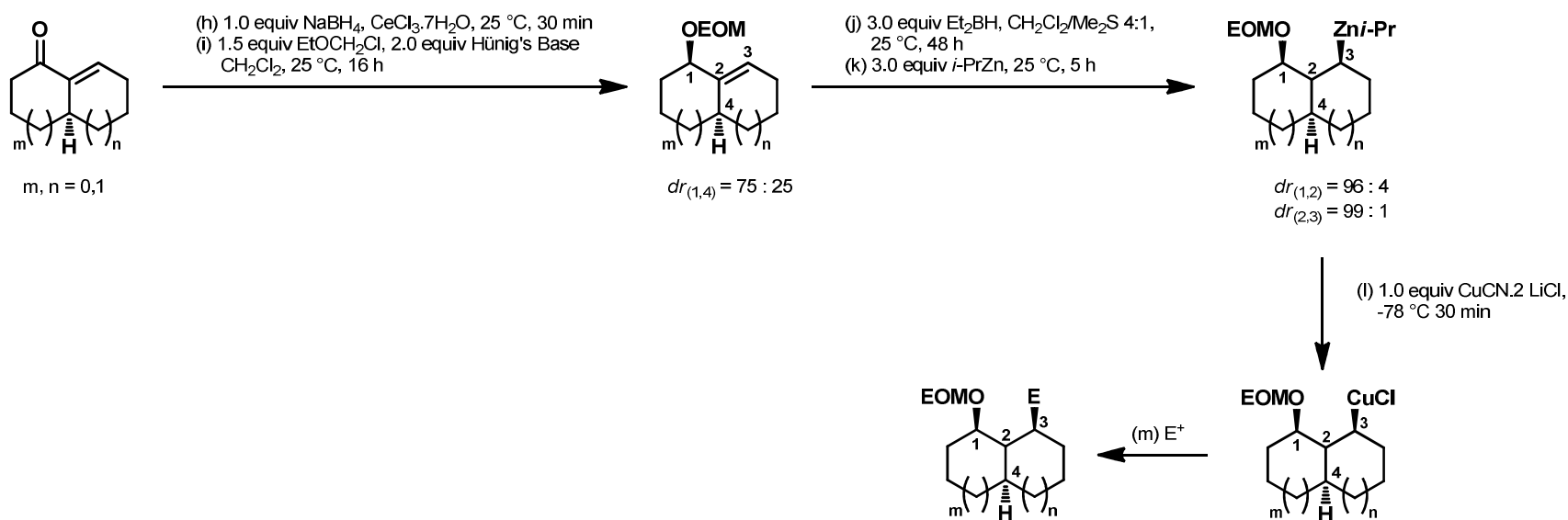


Highly diastereoselective Synthesis of Monocyclic and Bicyclic Secondary Diorganozinc Reagents with defined Configuration;
Knochel, P.; Boudier, E.; Hupe, E. *Angew. Chem. Int. Ed.* 2000, 39(13), 2294 - 2997

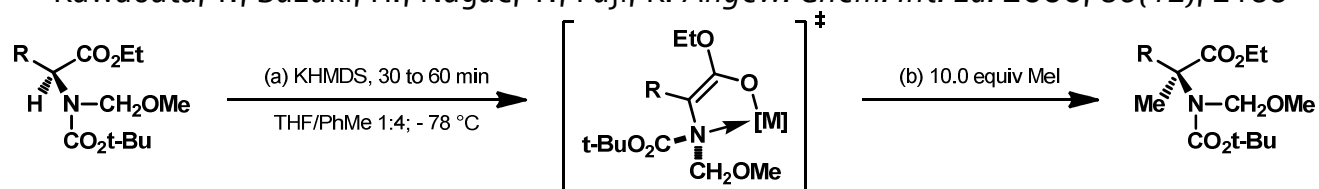


$\text{R}^1 = \text{Ph}$, $\text{R}^2 = \text{Bn}$; $dr_{(1,2)} = 93 : 7$
 $\text{R}^1 = \text{Ph}$, $\text{R}^2 = \text{CH}_2\text{OEt}$; $dr_{(1,2)} = 99 : 1$
 $\text{R}^1 = t\text{-Bu}$, $\text{R}^2 = \text{CH}_2\text{OEt}$; $dr_{(1,2)} = 99 : 1$





A Chiral, Nonracemic Enolate with Dynamic Axial Chirality: Direct Asymmetric α -Methylation of α -Amino Acid Derivatives;
Kawabata, T.; Suzuki, H.; Nagae, Y.; Fuji, K. *Angew. Chem. Int. Ed.* 2000, 39(12), 2155 - 2157

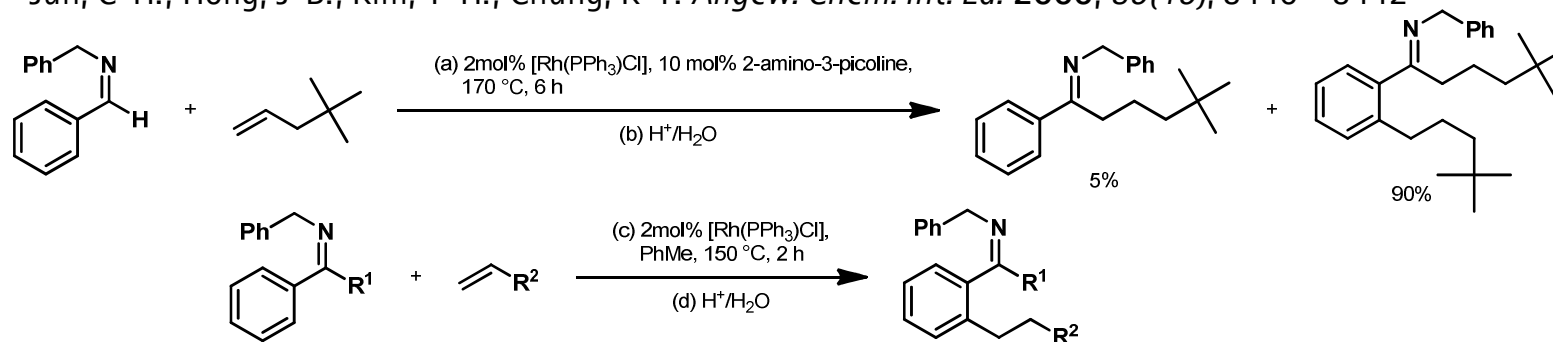


Entry	R ^a	Yield [%]	ee [%] ^b
1		96	81
2		83	93
3		94	79
4		95	80
5		88	76
6		81	87
7		78	78

^aee value for each substrate was >99 %; ^bdetermined by HPLC using columns with chiral stationary phase

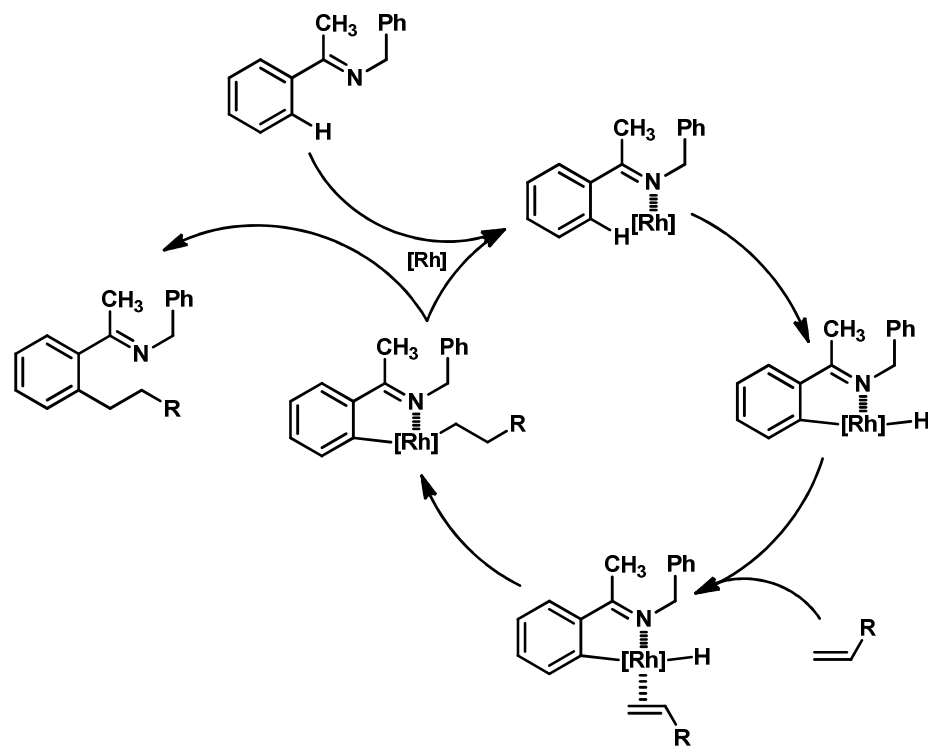
The catalytic Alkylation of Aromatic Imines by Wilkinson's Complex: The Domino Reaction of Hydroacylation and *ortho*-Alkylation

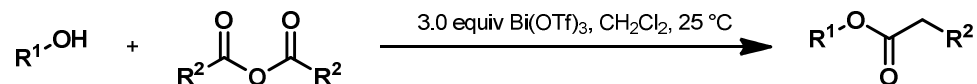
Jun, C-H.; Hong, J-B.; Kim, Y-H.; Chung, K-Y. *Angew. Chem. Int. Ed.* 2000, 39(19), 3440 - 3442

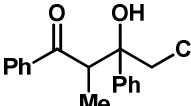
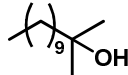
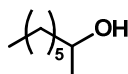
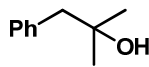
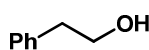
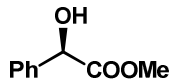


Entry	R ¹	R ²	Yield [%] ^a
1	Me	<i>t</i> -C ₄ H ₉	97 (100)
2	Me	C ₆ F ₅	91 (100)
3	Me	Cy	65 (68)
4	Me	<i>n</i> -C ₄ H ₉	94 (97) ^b
7	Me	TMS	92 (96)
9	Me		42 (47) ^c
10	Me		35
11	Et	TMS	93 (100) ^d
12	<i>n</i> -C ₅ H ₁₁	TMS	73(80) ^e

^a GC yields are given in parentheses; ^b 5.0 equiv of olefin were used; ^c 3.0 equiv of olefin were used; ^d 12% of di-*ortho* alkylation included; ^e 16% of di-*ortho* alkylation included

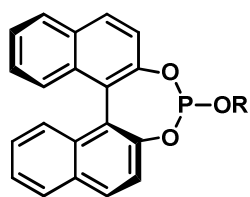
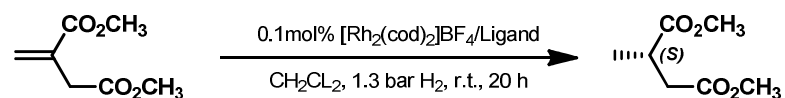


Highly Efficient and Versatile Acylation of Alcohols and $\text{Bi}(\text{OTf})_3$ Otera, J.; Orita, A.; Tanahashi, C.; Kakuda, A. *Angew. Chem. Int. Ed.* 2000, 39(16), 2877 - 2879

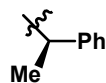
Entry	Alcohol	Anhydride / equiv	Mol% cat.	Time [h]	Yield [%]
1		$\text{Ac}_2\text{O} / 10$	3.0	1	96 ^a
2		$\text{Ac}_2\text{O} / 10$	0.5	7	95 ^b
3		$(\text{PhCO})_2\text{O} / 0.5$ $(t\text{-BuCO})_2\text{O} / 3.0$	3.0 3.0	8 4	2 ^c 99
4		$(\text{PhCO})_2\text{O} / 5.0$	3.0	48	0 ^c
5		$(\text{PhCO})_2\text{O} / 1.5$	3.0	24	90 ^c
6	1-adamantol	$(t\text{-BuCO})_2\text{O} / 1.5$	3.0	4	94
7		$(t\text{-BuCO})_2\text{O} / 1.5$	3.0	4	97

^a MeCN used as solvent; ^b THF used as solvent; ^c determined by GC

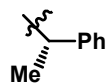
Highly enantioselective Rh–Catalyzed Hydrogenation Reactions based on Chiral Monophosphite Ligands

Reetz, M.T.; Mehler, G. *Angew. Chem. Int. Ed.* 2000, 39(21), 3889 – 3890

Ligand



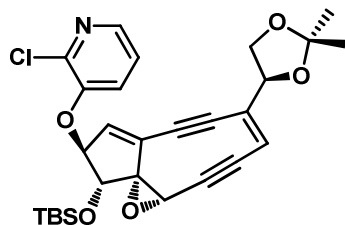
(R)-2-(1-phenylethyl)



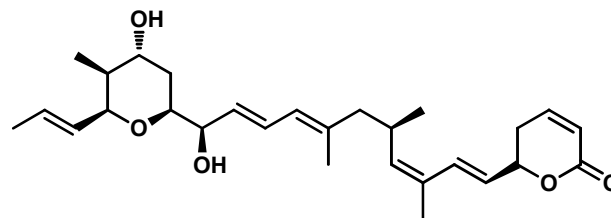
(S)-2-(1-phenylethyl)

Entry	R	Rh : Ligand	Mol% Rh	Conversion [%]	ee [%]
1	Me	1 : 1	0.1	100	89.2
2	<i>i</i> -Pr	1 : 1	0.1	100	97.6
3	Ph	1 : 1	0.1	100	96.6
4	<i>o</i> -Br-Ph	1 : 1	0.1	100	89.8
5	2,6-(Me)2-Ph	1 : 1	0.1	78	39.2
6	2,6-(Ph)2-Ph	1 : 1	0.1	8	28.6
7	(<i>R</i>)-2-(1-phenylethyl)	1 : 1	0.1	100	99.2
8	(<i>S</i>)-2-(1-phenylethyl)	1 : 1	0.1	100	98.2
9	(<i>R</i>)-2-(1-phenylethyl)	1 : 1	0.04	100	99.4
10	(<i>R</i>)-2-(1-phenylethyl)	1 : 1	0.02	100	99.4
11	(<i>R</i>)-2-(1-phenylethyl)	1 : 1	0.01	49	96.2
12	(<i>R</i>)-2-(1-phenylethyl)	1 : 2	0.1	100	99.6
13	(<i>R</i>)-2-(1-phenylethyl)	1 : 4	0.1	100	99.5
14	(<i>R/S</i>)-2-(1-phenylethyl)	1 : 1	0.1	100	98.8

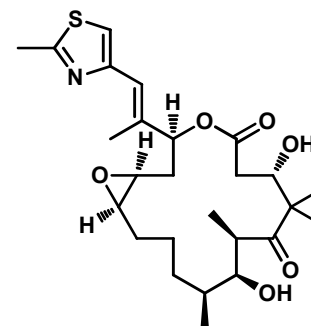
Syntheses not covered

**Kredaricin Core**

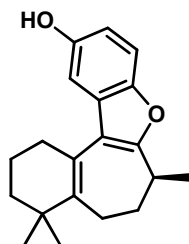
Myers, A. G.; Goldberg, S. D.
Angew. Chem. Int. Ed. **2000**, 39(15), 2732 - 2735

**(+)-Ratjadone**

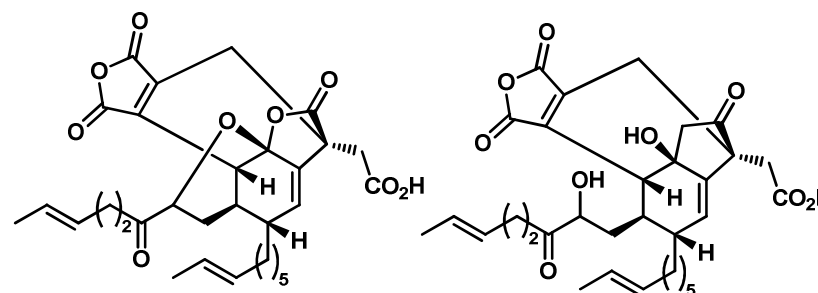
Kalesse, M.; Christmann, M.; Bhatt, U.; Quitschalle, E. C.;
Angew. Chem. Int. Ed. **2000**, 39(23), 4364 - 4366

**Epithilone B**

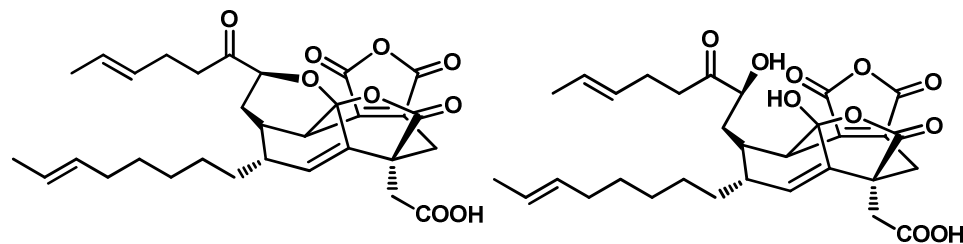
Shibasaki, M.; Sawada, D.;
Angew. Chem. Int. Ed. **2000**, 39(1), 209 - 213

**Frondosin B**

Danishefsky, S. J.; Inoue, M.; Frontier, A. J.
Angew. Chem. Int. Ed. **2000**, 39(4), 761 - 764

**CP-263,114 and CP-225,917**

Danishefsky, S. J.; Tan, Q.;
Angew. Chem. Int. Ed. **2000**, 39(24), 4509 - 4511

**(-)-CP-263,114 (Phorimide B) and (+)-CP-225,917 (Phorimide A)**

Nicolau, K. C.; Jung, J.-K.; Yoon, W. H.; He, Y.; Zong, Y.-L.; Baran, P. S.
Angew. Chem. Int. Ed. **2000**, 39(10), 1829 - 1832