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**Education:** BS (1974): Pharmaceutical Institute, Tohoku University  
(with Prof. Seiichi Takano)

MS (1976): Graduate Course of Pharmaceutical Institute, Tohoku University  
(with Prof. Seiichi Takano)

PhD (1979): Graduate Course of Pharmaceutical Institute, Tohoku University  
(with Prof. Seiichi Takano)

**Experience:** Postdoctoral Fellow (1979, April ~ 1981, May)  
Chemistry Department, Harvard University  
(with Prof. Yoshito Kishi)

Assistant Professor (1981, June ~ 1993, January)  
Pharmaceutical Institute, Tohoku University  
(with Prof. Seiichi Takano)

Associate Professor (1993, February ~ 1994, March)  
Faculty of Pharmaceutical Sciences, Tokushima Bunri University  
(with Prof. Mugio Nishizawa)

Associate Professor (1994, April ~ 1997, March)  
School of Pharmaceutical Sciences, Nagasaki University  
(with Prof. Hiroshi Irie)

Professor (1997, April ~ 2002, March)  
School of Pharmaceutical Sciences, Nagasaki University

Professor (2002, April ~)

Graduate School of Biomedical Sciences, Nagasaki University

**Award:** 1987: The Pharmaceutical Society of Japan Award of Tohoku Branch  
for Young Scientists

1994: The Pharmaceutical Society of Japan Award for Young Scientists

2000: The Tokyo Biomedical Research Foundation Award

2001: Nagase Science and Technology Foundation Award

2014: The Pharmaceutical Society of Japan Award

**Research Interest:** Synthetic Methodology, Asymmetric Synthesis, Natural Product Synthesis

1. Synthetic studies directed toward kaitocephalin: a highly stereocontrolled route to the right-hand pyrrolidine core.  
Keisuke Takahashi, Natsumi Haraguchi, Jun Ishihara, and Susumi Hatakeyama  
*Synlett*, **2008**, 671-674.
2. Asymmetric Total Synthesis of (+)-Phoslactomycin B  
Setsuya Shibahara, Masataka Fujino, Masashi Tashiro, Keisuke Takahashi, Jun Ishihara, and Susumi Hatakeyama  
*Org. Lett.* **10**, 2139-2142 (2008).
3. Entry to Heterocycles Based on Indium-Catalyzed Conia-Ene Reactions: Asymmetric Synthesis of (–)-Salinosporamide A  
Keisuke Takahashi, Michiko Midori, Kei Kawano, Jun Ishihara, and Susumi Hatakeyama  
*Angew. Chem. Int. Ed.*, **47**, 6244-6246 (2008).
4. Asymmetric synthesis of (+)-trachyspic acid  
Kenji Morokuma, Yuko Taira, Yumiko Uehara, Setsuya Shibahara, Keisuke Takahashi, Jun Ishihara and Susumi Hatakeyama  
*Tetrahedron Lett.*, **49**, 6043-6045 (2008).
5. Enantioselective Route to Aryl(1,3-butadien-2-yl)methanols: Formal Synthesis of (–)-Sporochnol A  
Daisuke Yanagimoto, Kazuyuki Kawano, Keisuke Takahashi, Jun Ishihara, and Susumi Hatakeyama  
*Heterocycles*, **77**, 249-253 (2009).
6. Indium-catalyzed Conia-ene reaction for alkaloid synthesis  
Susumi Hatakeyama  
*Pure Appl. Chem.*, **81**, 217-226 (2009).
7. Synthetic studies of vitamin D analogs. Part 35. An improved synthesis of 1-epi-ED-71, a biologically interesting diastereomer of 1a,25-dihydroxy-2b -(3-hydroxypropoxy)vitamin D<sub>3</sub> (ED-71).  
Kohei Eto, Ayako Fujiyama, Mai Kaneko, Keisuke Takahashi, Jun Ishihara, and Susumi Hatakeyama, Yoshiyuki Ono, Noboru Kubodera  
*Heterocycles*, **77**, 323-331 (2009).
8. Synthesis of 1 $\alpha$ ,25-Dihydroxy-2 $\beta$ -(3-hydroxypropoxy)vitamin D<sub>3</sub> (Eldecalcitol) and Related Compounds by the Trost Convergent Methodology.  
Noboru Kubodera and Susumi Hatakeyama  
*Heterocycles*, **79**, 145-162 (2009).
9. A New Variant of Reformatsky–Claisen Rearrangement Mediated by Indium Chloride

- Jun Ishihara, Noriko Koyama, Yukihiro Nishino, Keisuke Takahashi, and Susumi Hatakeyama  
*Synlett*, **2009**, 2351-2355.
10. Total Synthesis of (+)-Fostriecin and (+)-Phoslactomycin B  
Setsuya Shibahara, Masataka Fujino, Yasumasa Tashiro, Nanako Okamoto, Tomoyuki Esumi, Keisuke Takahashi, Jun Ishihara, and Susumi Hatakeyama  
*Synthesis*, **2009**, 2935-2953.
11. Enantio- and stereoselective route to the phoslactomycin family of antibiotics: formal synthesis of (+)-fostriecin and (+)-phoslactomycin B  
Shaheen M. Sarkar, Everlyne N. Wanzala, Setsuya Shibahara, Keisuke Takahashi, Jun Ishihara, and Susumi Hatakeyama  
*Chem. Commun.*, **2009**, 5907-5909.
12. Synthetic Studies on Vitamin D Analogs. 37. Synthesis of 20-Epi-1a,25-dihydroxy-2b-(3-hydroxypropoxy)vitamin D<sub>3</sub>: 20-Epi-ED-71  
Madoka Yoshino, Kohei Eto, Keisuke Takahashi, Jun Ishihara, Susumi Hatakeyama, Yoshiyuki Ono, Hitoshi Saito, and Noboru Kubodera  
*Heterocycles* **81**, 381-394 (2010).
13. Synthetic Study on Clutilolide Based on a Remote Chelation Controlled Ireland-Claisen Rearrangement  
Jun Ishihara, Okihisa Tokuda, Kazunori Shiraishi, Yukihiro Nishino, Keisuke Takahashi, and Susumi Hatakeyama  
*Heterocycles*, **80**, 1067-1079 (2010).
14. 145. Synthesis and preliminary biological evaluation of 20-epi-eldecalcitol [20-epi-1a,25-dihydroxy-2b-(3-hydroxypropoxy)vitamin D<sub>3</sub>: 20-epi-ED-71]  
Susumi Hatakeyama, Madoka Yoshino, Kohei Eto, Keisuke Takahashi, Jun Ishihara, Yoshiyuki Ono, Hitoshi Saito, and Noboru Kubodera  
*J. Steroid. Biochem.*, **121**, 25-28 (2010).
15. Indium-catalyzed Conia-ene Reaction and Total Syntheses of Biologically Active Alkaloids  
Keisuke Takahashi and Susumi Hatakeyama  
*J. Synth. Org. Chem. Jpn.*, **68**, 951-961 (2010).
16. Organocatalytic asymmetric synthesis of quinine and quinidine  
Shaheen M. Sarkar, Yuko Taira, Ayako Nakano, Keisuke Takahashi, Jun Ishihara, and Susumi Hatakeyama  
*Tetrahedron Lett.*, **52**, 923-927 (2011).
17. Indium-mediated Reformatsky-Claisen rearrangement  
Jun Ishihara, Yuki Watanabe, Noriko Koyama, Yukihiro Nishino, Keisuke Takahashi, and

Susumi Hatakeyama

- Tetrahedron*, **67**, 3659-3667 (2011).
18. Synthesis of 1-Deoxyeldecalcitol, a Biologically Interesting Analog of 1a,25-Dihydroxy-2b-(3-hydroxypropoxy)vitamin D<sub>3</sub> (Eldecalcitol)  
Hirotaka Sasaki, Kohei Eto, Keisuke Takahashi, Jun Ishihara, Susumi Hatakeyama, and Noboru Kubodera  
*Heterocycles*, **83**, 1385-1394 (2011).
19. Total Synthesis of NW-G01, a Cyclic Hexapeptide Antibiotic, and 34-epi-NW-G01  
Setsuya Shibahara, Takaaki Matsubara, Keisuke Takahashi, Jun Ishihara, and Susumi Hatakeyama  
*Org. Lett.*, **13** (17), 4700–4703 (2011).
20. Total Synthesis of Oxazolomycin A  
Kohei Eto, Madoka Yoshino, Keisuke Takahashi, Jun Ishihara, and Susumi Hatakeyama  
*Org. Lett.*, **13** (19), 5398–5401 (2011).
21. Process Development for the Practical Production of Eldecalcitol by Linear, Convergent and Biomimetic Syntheses  
Noboru Kubodera and Susumi Hatakeyama  
*Anticancer Res.*, **32**, 303-310 (2012)
22. Total Synthesis of (−)-Kaitocephalin Based on a Rh-Catalyzed C–H Amination  
Keisuke Takahashi, Daisuke Yamaguchi, Jun Ishihara, and Susumi Hatakeyama  
*Org. Lett.*, **14**, 1644–1647 (2012).
23. Stereocontrolled Total Synthesis of (−)-Englerin A  
Keisuke Takahashi, Keita Komine, Yuichi Yokoi, Jun Ishihara, and Susumi Hatakeyama  
*J. Org. Chem.*, **77**, 7364–7370 (2012).
24. Organocatalytic asymmetric syntheses of inthomycins A, B and C  
Madoka Yoshino, Kohei Eto, Keisuke Takahashi, Jun Ishihara, and Susumi Hatakeyama  
*Org. Biomol. Chem.*, **10**, 8164-8174 (2012).
25. A New Method for the Protection of Carboxylic Acids with a Triisopropylsiloxyethyl Group  
Hikaru Yoshimura, Kohei Eto, Keisuke Takahashi, Jun Ishihara, and Susumi Hatakeyama  
*Chem. Pharm. Bull.*, **60**, 1334-1339 (2012).
26. Recent Developments in the Reformatsky-Claisen Rearrangement  
Jun Ishihara and Susumi Hatakeyama  
*Molecules*, **17**, 14249-14259 (2012).
27. Stereoselective Synthesis of the Fused γ-Lactone/δ-Lactone Core of Ophiodilactones

Takaaki Matsubara, Keisuke Takahashi, Jun Ishihara, and Susumi Hatakeyama  
*Heterocycles*, **86**, 155-158 (2012).