

贾彦兴



个人信息:

生于 1975年5月1日, 河北满城
地址 北京大学药学院、天然药物及仿生药物国家重点实验室
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学习经历:

1993-1997 理学学士, 兰州大学化学系应用化学专业
1997-2002 理学博士, 兰州大学化学系有机化学专业 (导师: 涂永强 院士)
2002-2007 博士后, 法国国家科学研究中心, 天然产物化学研究所 (导师: 祝介平)

任职:

2007- 副教授(课题组长), 北京大学

研究方向与兴趣:

1. 具有重要生物活性天然产物的全合成及仿生合成;
2. 生物活性天然产物的结构改造以及构效关系研究;
3. 有机合成新方法、新策略的研究。

获奖情况:

2010 Asian Core Program Lectureship Award
2009 新世纪优秀人才支持计划
2009 北京大学医学部优秀青年学者
2008-2009 教学优秀奖
2008 第十一届中国药学会-施维雅青年药物化学奖
2002 甘肃省科技进步奖, 一等奖(排名第五)
1997 兰州大学优秀毕业生
1997 张淑明奖学金

承担项目:

1. 2010.1-2012.12: 国家自然科学基金委面上项目, “Celogentins 的合成及构效关系研究”。(20972007)
2. 2010.1-2013.12: 国家自然科学基金委重点项目, “海绵 baculiferin 型新生物碱探针对 HIV-1 Vif 和 APOBEC3G 的靶向调控及作用机理研究”。(30930109/C180202)
3. 2010.1-2014.8: 科技部 973, “具有重要生物活性的天然产物的化学合成”。(2010CB833200)
4. 2010.1-2012.12: 教育部创新团队发展计划, “基于内源性物质的先导物发

- 现”。(IRT0902)
- 2009.1-2011.12: 国家自然科学基金委青年基金, “几种吲哚类生物碱的全合成研究”。(20802005)
 - 2009.1-2011.12: 教育部博士点新教师基金, “Celogentin C 的合成研究”。(200800011055)
 - 2009.9-2010.09: 中国科学院上海药物研究所新药研究国家重点实验室开放研究课题, “Celogentins 的全合成及构效关系研究”。
 - 2009.1-2009.12: 兰州大学功能有机分子化学国家重点实验室开放课题, “Celogentins 的全合成研究”。
 - 教育部留学回国人员科研启动基金, “杀鱼菌素类天然产物的全合成研究”。
 - 2008.1-2008.12: 兰州大学功能有机分子化学国家重点实验室开放课题, “几种 3, 4-二取代吲哚类生物碱的全合成研究”。
 - 2007.9-2008.8: 国家自然科学基金委主任基金, “杀鱼菌素类天然产物的全合成研究”。(20842004)
 - 2007.11-2010.10: 北京大学医学部、天然药物及仿生药物国家重点实验室(引进人才基金), “几种具有重要生物活性的吲哚类生物碱的全合成研究”。

论文及专著:

迄今为止已在国内外重要学术刊物上发表研究论文 40 篇。

独立课题组后:

40. “Total Synthesis of (±)-Decursivine and (±)-Serotobenine: A Cascade Witkop Photocyclization/Elimination/O-Michael Addition Approach”
Hua Qin, Zhengren Xu, Yuxin Cui, Yanxing Jia*
Angew. Chem. Int. Ed. **2011**, *50*, (online, DOI: 10.1002/anie.201100495).
39. “Total Synthesis of (-)-Indolactam V”
Zhengren Xu, Fengying Zhang, Lihe Zhang, Yanxing Jia*
Org. Biomol. Chem. **2011**, *9*, 2512–2517.
38. “Total Synthesis of Lamellarins D, H, and R and Ningalin B”
Qingjiang Li, Jingqian Jiang, Aili Fan, Yuxin Cui, Yanxing Jia*
Org. Lett. **2011**, *13*, 312–315.
37. “Total Synthesis of Clavicipitic Acid and Aurantioclavine: Stereochemistry of Clavicipitic Acid Revisited”
Zhengren Xu, Weimin Hu, Qiang Liu, Lihe Zhang, Yanxing Jia*
J. Org. Chem. **2010**, *75*, 7626–7635.
36. “One-Pot AgOAc-Mediated Synthesis of Polysubstituted Pyrroles from Primary Amines and Aldehydes: Application to the Total Synthesis of Purpurone”

Qingjiang Li, Aili Fan, Zhiyao Lu, Yuxin Cui, Wenhan Lin, Yanxing Jia*
Org. Lett. **2010**, *12*, 4066–4069.

35. “Stereocontrolled and Efficient Total Synthesis of (–)-Stephanotic Acid Methyl Ester and (–)-Celogentin C”
Weimin Hu, Fengying Zhang, Zhengren Xu, Qiang Liu, Yuxin Cui, Yanxing Jia*
Org. Lett. **2010**, *12*, 956–959.
34. “Water-Controlled Regioselectivity of Pd-Catalyzed Domino Reaction Involving a C–H Activation Process: Rapid Synthesis of Diverse Carbo- and Heterocyclic Skeletons”
Zhiyao Lu, Chunmei Hu, Jiajie Guo, Jing Li, Yuxin Cui, Yanxing Jia*
Org. Lett. **2010**, *12*, 480–483.
33. “Artificial Transcription Factors which Mediate Double-Strand DNA Cleavage”
Chao Li, Chao Du, Hua Tian, Chao Jiang, Min Du, Yan Liu, Renzhong Qiao*, Yanxing Jia*, and Yufen Zhao*
Chem. Eur. J. **2010**, *16*, 12935–12940.
32. “Formal synthesis of semiaquilegin A”
Jing Li, Yong Jiang, Qingjiang Li, Qiang Xiao, Yanxing Jia*, Pengfei Tu*
Tetrahedron Lett. **2010**, *51*, 1121–1123.
31. “Palladium-Catalyzed Synthesis of Tryptamines and Tryptamine Homologues: Synthesis of Psilocin”
Chunmei Hu, Hua Qin, Yuxin Cui, Yanxing Jia*
Tetrahedron **2009**, *65*, 9075–9080.
30. “Efficient Total Synthesis of (–)-*cis*-Clavicipitic Acid”
Zhengren Xu, Qingjiang Li, Lihe Zhang, Yanxing Jia*
J. Org. Chem. **2009**, *74*, 6859–6862.
29. “Total Synthesis of (–)-Incarvilline and (–)-Incarvillateine”
Fengying Zhang, Yanxing Jia*
Tetrahedron **2009**, *65*, 6840–6843.
28. “Palladium-Catalyzed Indole and Azaindole synthesis by Direct Annulation of Electron-Poor *o*-Chloroanilines and Chloroaminopyridines with Aldehydes”
Zhengren Xu, Weimin Hu, Fengying Zhang, Qingjiang Li, Zhiyao Lü, Lihe Zhang, Yanxing Jia*
Synthesis **2008**, 3981–3987.

博士、博士后期间：

27. “Total Synthesis of Complestatin (Chloropeptin II)”
Zihui Wang, Michèle Bois-Choussy, **Yanxing Jia**, Jieping Zhu*
Angew. Chem. Int. Ed. **2010**, *49*, 2018–2022.
26. “Synthesis of Diastereomers of Complestatin, and Chloropeptin, High Substrate-dependent Atropstereoselectivity of Intramolecular Suzuki-Miyaura Reaction”
Yanxing Jia, Michèle Bois-Choussy, Jieping Zhu*
Angew. Chem. Int. Ed. **2008**, *47*, 4167–4172.
25. “Synthesis of DEFG Ring of Complestatin and Chloropeptin I: Highly Atropdiastereoselective Macrocyclization by Intramolecular Suzuki-Miyaura Reaction”
Yanxing Jia, Michèle Bois-Choussy, Jieping Zhu*
Org. Lett. **2007**, *9*, 2401–2404.
24. “Palladium-Catalyzed Enantioselective Domino Heck-Cyanation Sequence: Development and Application to the Total Synthesis of Physostigmine”
Artur Pinto, **Yanxing Jia**, Luc Neuville, Jieping Zhu*
Chem. -A Eur. J. **2007**, *13*, 961–967.
23. “Importance of the Structure of Vancomycin Binding Pocket in Designing Compounds Active Against Vancomycin-resistant Enterococci (VRE)”
Yanxing Jia, Michèle Bois-Choussy, Adriano Malabarba, Cristina Brunati, Jieping Zhu*
J. Antibiotics. **2006**, *59*, 543–552.
22. “Palladium-Catalyzed, Modular Synthesis of Highly Functionalized Indoles and Tryptophans by Direct Annulation of Substituted *o*-Haloanilines and Aldehydes”
Yanxing Jia, Jieping Zhu*
J. Org. Chem. **2006**, *71*, 7826–7834.
21. “A Novel Strategy for the Key Fully Substituted Cyclopentenedione Moiety of Madindolines via AlEt₃-Promoted Tandem Reductive Rearrangement of α -Hydroxy Epoxides”
Shuanhu Gao, **Yanxing Jia**, Xuezhi Zhao, Yongqiang Tu*
Chin. J. Chem. **2006**, *24*, 595–597.
20. “Design and Synthesis of Simple Macrocycles Active Against Vancomycin-resistant *Enterococci* (VRE)”
Yanxing Jia, Nianchun Ma, Zuosheng Liu, Michèle Bois-Choussy, Eduardo Gonzalez-Zamora, Adriano Malabarba, Cristina Brunati, Jieping Zhu*
Chem. Eur. J. **2006**, *12*, 5334–5351.

19. "Synthesis of Ring-A-Substituted Tryptophan by a Palladium-Catalyzed Heteroannulation Reaction"
Yanxing Jia, Jieping Zhu*
Synlett **2005**, 2469–2472.
18. "Identification of Synthetic Compounds Active Against VRE: The Role of the Lipidated Aminoglucose and the Structure of Glycopeptide Binding Pocket"
Yanxing Jia, Eduarda Gonzalez-Zamora, Nianchun Ma, Zuosheng Liu, Michèle Bois-Choussy, Adriano Malabarba, Cristina Brunati, Jieping Zhu*
Bioorg. Med. Chem. Lett. **2005**, *15*, 4594–4599.
17. "Synthesis of β -Amino- α -Hydroxy Esters and β -Amino- α -Azido Ester by Sharpless Asymmetric Aminohydroxylation, By-Products Analysis"
Zuosheng Liu, Nianchun Ma, **Yanxing Jia**, Michèle Bois-Choussy, Adriano Malabarba, Jieping Zhu*
J. Org. Chem. **2005**, *70*, 2847–2850.
16. "Design and synthesis of macrocycles active against vancomycin-resistant enterococci (VRE): the interplay between D-Ala-D-Lac binding and hydrophobic effect"
Nianchun Ma, **Yanxing Jia**, Zuosheng Liu, Eduarda Gonzalez-Zamora, Michèle Bois-Choussy, Adriano Malabarba, Cristina Brunati, Jieping Zhu*
Bioorg. Med. Chem. Lett. **2005**, *15*, 743–746.
15. "One or More C-H Bond(s) Formed by Substitution: Reduction of Carbon-Nitrogen, -Phosphorus, -Arsenic, -Antimony, -Bismuth, -Carbon, -Silicon, -Germanium, -Boron, and -Metal Bonds" in "Comprehensive Organic Functional Group Transformations II", Eds. Katritzky, A. R., Taylor, R. J. K.; Volume Ed. Cossy, J. Elsevier, **2005**, *1*, 31-78
Jérôme Blanchet, **Yanxing Jia**, Jieping Zhu*
14. "4-Nitrophenyltriflate" in *Electronic Encyclopedia of Reagents for Organic Synthesis (e-EROS)* Ed; L. A. Paquette, Wiley-VCH, Weinheim. **2004**
Yanxing Jia, Jieping Zhu*
13. "Synthetic studies of the HIV-1 protease inhibitive didemnaketals: stereocontrolled synthesis of an ester side chain"
Xuezhi Zhao, Yongqiang Tu*, Lei Peng, Xueqiang Li, **Yanxing Jia**
Tetrahedron Lett. **2004**, *45*, 3713–3716.
12. "An Interesting AlEt₃-promoted Stereoselective Tandem Rearrangement/Reduction of α -Hydroxy (or Amino) Heterocyclopropane"
Xin Li, Bin Wu, Xuezhi Zhao, **Yanxing Jia**, Yongqiang Tu*, Derun Li
Synlett **2003**, 623–626.

11. “An efficient synthetic approach to the aromatic sesquiterpenoids via a SmI₂-promoted construction of quaternary centre”
Xuezhi Zhao, **Yanxing Jia**, Yongqiang Tu*
J. Chem. Res(S). **2003**, 27, 54–55.
10. “Convergent synthesis of the key core spiroketals of the HIV-1 protease inhibitors didemnaketals”
Yanxing Jia, Xin Li, Pingzhen Wang, Bin Wu, Xuezhi Zhao, Yongqiang Tu*
J. Chem. Soc., Perkin trans. 1 **2002**, 565–570.
9. “A convergent synthesis of the spiroketal moiety of the HIV-1 protease inhibitors didemnaketals”
Yanxing Jia, Xin Li, Bin Wu, Xuezhi Zhao, Yongqiang Tu*
Tetrahedron **2002**, 58, 1697–1708.
8. “Synthetic Studies of the HIV-1 Protease Inhibitive Didemnaketals: Stereocontrolled Synthetic Approach to the Key Mother Spiroketal”
Yanxing Jia, Bin Wu, Xin Li, Shikuo Ren, Yongqiang Tu*, Albert S. C. Chan, William Kitching
Org. Lett. **2001**, 3, 847–849.
7. “A New One-Pot Synthesis of 2-Quaternary 1,3-Diketones”
Shikuo Ren, Fei Wang, Huani Dou, Chunan Fan, Lan He, Zhenlei Song, Wujiong Xia, Derun Li, **Yanxing Jia**, Xin Li, Yong qiang Tu*
Synthesis **2001**, 16, 2384–2388.
6. “Diastereo-recognizable reaction between Cr(VI) reagents and tertiary α -hydroxy epoxide”
Yongqiang Tu*, Shikuo Ren, **Yanxing Jia**, Baomin Wang, Albert S. C. Chan, Michael C. K. Choi
Tetrahedron Lett. **2001**, 42, 2141–2144.
5. “手性 3, 7-二甲基-5, 6-二羟基-辛-7-烯醛中间体的合成”
贾彦兴, 吴滨, 涂永强*
高等学校化学学报. **2001**, 22, 584–586.
4. “Synthetic Studies of Didemnaketals Analogue-Construction of the Intermediate (3*S*,5*S*,6*R*)-3,7-Dimethyl-5,6,7-trihydroxy-octanal”
Yanxing Jia, Bin Wu, Pingzhen Wang, Yongqiang Tu*
Chin. Chem. Lett. **2000**, 11, 509–510.
3. “Didemnaketals A 类似物的合成研究- C1-C7 中间体的合成”
贾彦兴, 王平珍, 吴滨, 涂永强*

高等学校化学学报。2000, 21, 729-730.

2. “Synthetic Studies of an Analogue of HIV-1 Protease Inhibitors of Didemnaketals: Construction of the C1-C8 Intermediate”

Pingzhen Wang, **Yanxing Jia**, Yongqiang Tu*, Bin Wu
Chin. Chem. Lett. **1999**, 10, 749-750.

1. “Didemnaketals A 类似物的合成研究- C8-C15 中间体的合成”

涂永强, * 刘增路, 王平珍, **贾彦兴**
高等学校化学学报. 1999, 20, 81-83.

发明专利:

2. 李清江, **贾彦兴**; 1, 3, 4-三取代或 3, 4 二取代的吡咯环化合物的制备方法。
申请号: 201010177801.7

1. 姜勇, 李菁, **贾彦兴**, 屠鹏飞; 一种制备长栲利素 A 的方法。申请号:
200910236246.8