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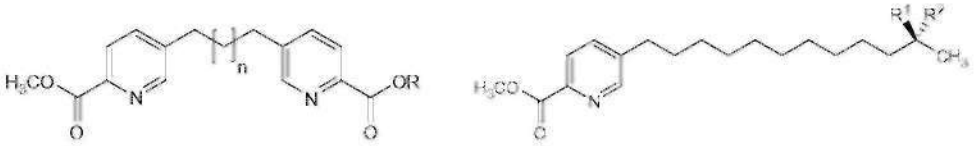
Chakapong Intaraudom, Nattawut Boonyuen, Rapheephat Suvannakad, Pranee Rachtawee, Pattama Pittayakhajonwut*



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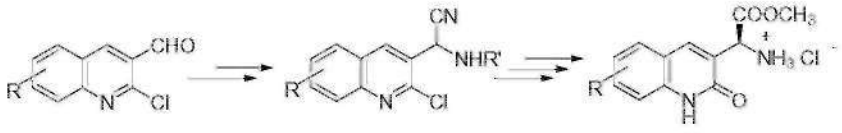

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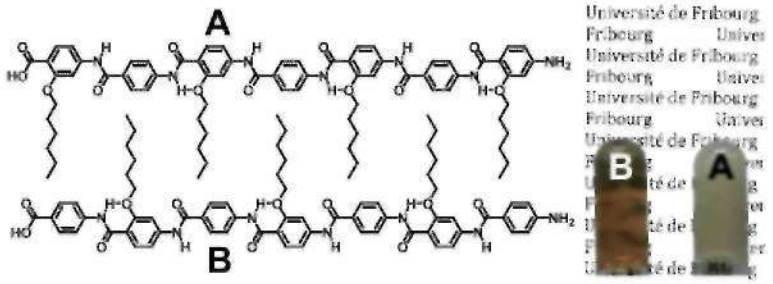
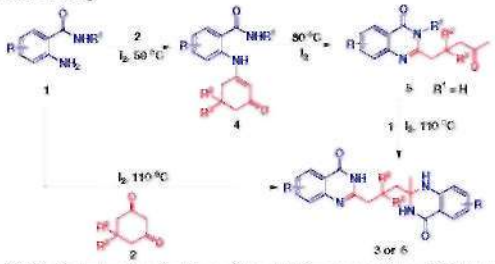
Helga Seyler, Andreas Kilbinger*



Structurally diversified products from the reactions of 2-aminobenzamides with 1,3-cyclohexanediones catalyzed by iodine pp 757–760
Lian Lu, Mei-Mei Zhang, Hong Jiang, Xiang-Shan Wang*



Controlling the reaction temperature at 50 °C, 80 °C, and 110 °C, respectively, the iodine-catalyzed reaction of 2-aminobenzamides with 1,3-cyclohexanediones gave structurally diversified products. In the latter, it gave bis-quinazolin-4(3*H*)-ones unexpectedly, with 1,3-cyclohexanediones ring-opening.

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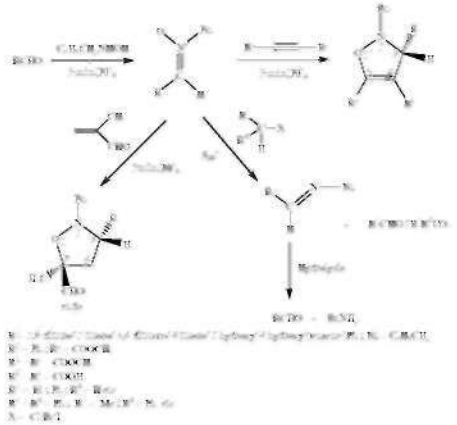

1-Butyl-3-methylimidazolium based ionic liquids are found to accelerate significantly the intermolecular 1,3-dipolar cycloaddition of *N*-benzyl fluoro nitrones derived in situ from aldehydes and benzylhydroxylamine, with electron deficient alkynes to afford enhanced rates and improved yields of isoxazolines while with enals exclusively *endo* isoxazolidines are obtained with high selectivity. Synthetic potentiality of the novel isoxazolines and nitrones has also been tested successfully.

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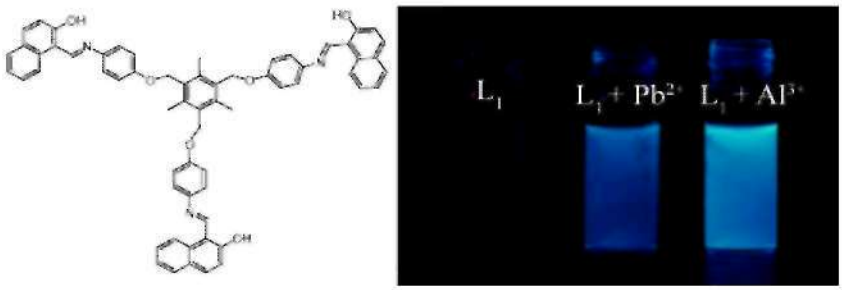
Barun Kumar Datta, Chirantan Kar, Arghya Basu, Gopal Das*



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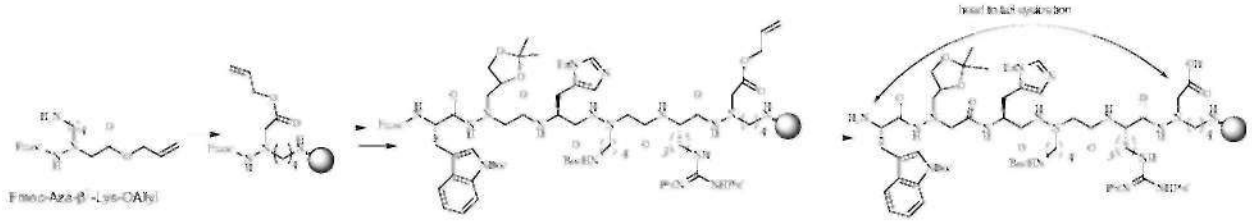

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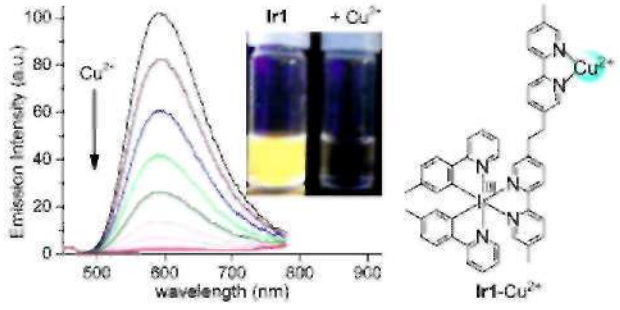

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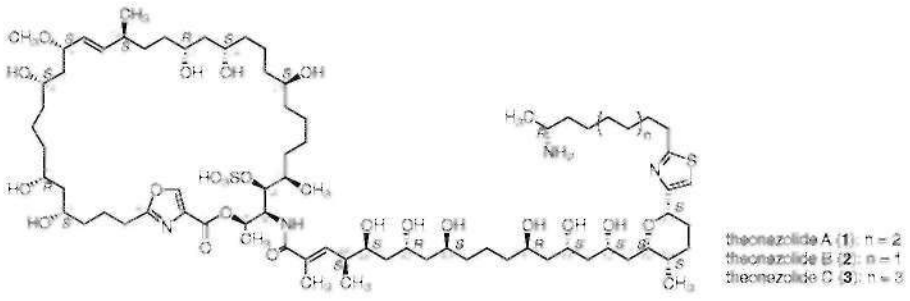

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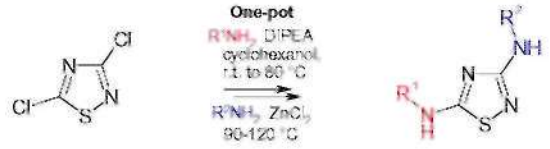
Howard A. Beeley, Sébastien Degorce, Craig S. Harris*, Jonathan Lecoq, Rémy Morgentin, David Perkins



The synthesis of bis(amino)-1,2,4-thiadiazoles usually relies on a four step sequence. Herein, we communicate to our knowledge the first one-pot synthesis of bis(amino)-1,2,4-thiadiazoles via a double S_NAr approach.

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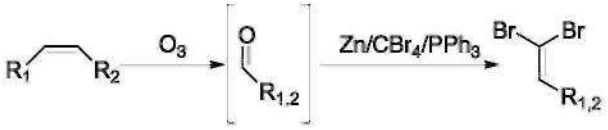
Brenna Arlyce Brown*, Jonathan G. C. Veinot



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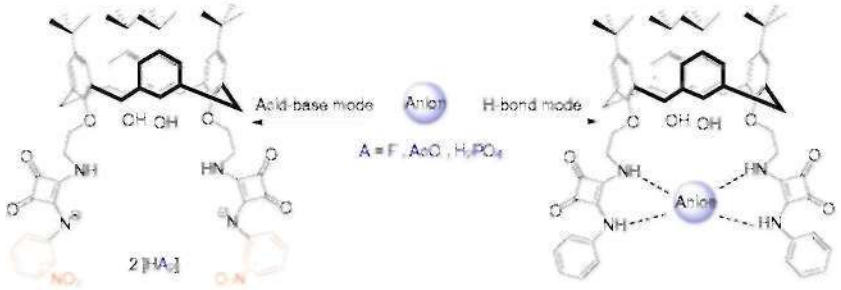

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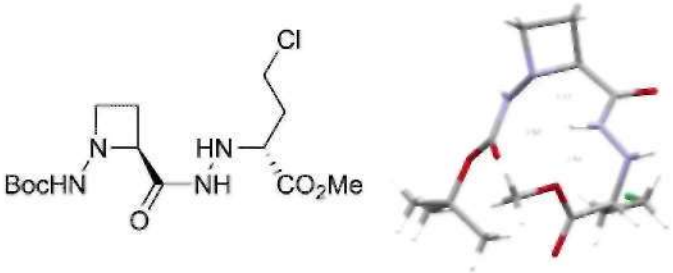

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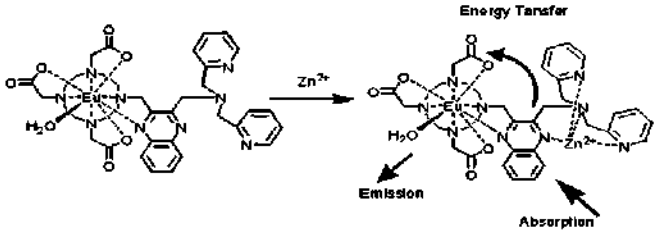

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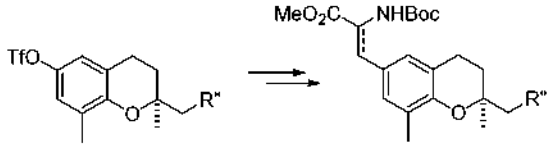

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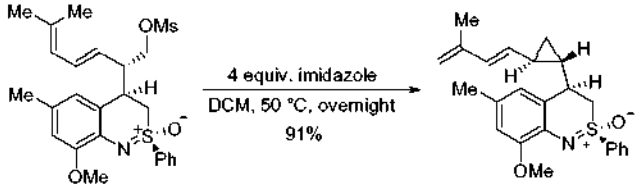

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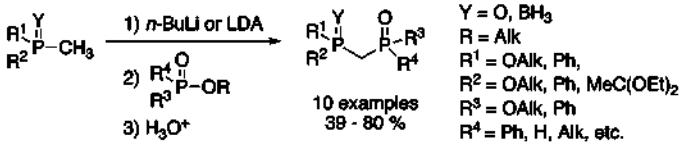
Laurent Gavara, Fabien Gelat, Jean-Luc Montchamp*



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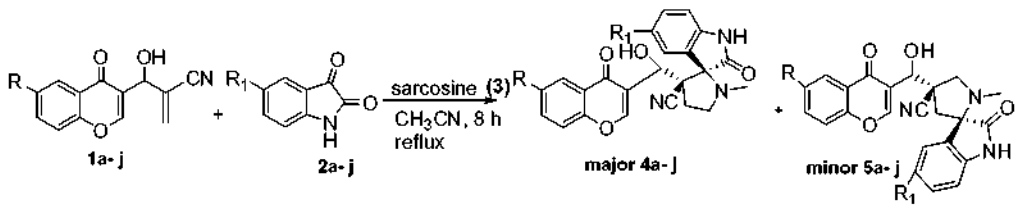
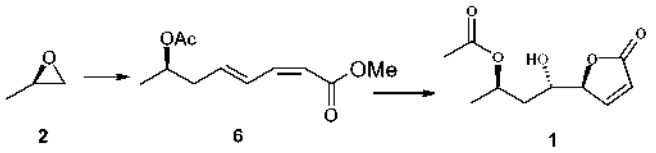
Panneerselvam Yuvaraj, Boreddy S. R. Reddy*



3-Spiropyrrolidine-oxindole unit is a privileged heterocyclic motif forming the core of a large family of alkaloid natural products with strong bioactivity profile and interesting structural properties. A novel regioselective synthesis of functionalized 3-spiropyrrolidine-3-spirooxindoles from 4-oxo-4*H*-chromone derivatives was accomplished by the [3+2] cycloaddition of azomethineylides with Baylis–Hillman adducts.

Concise and protecting group-free synthesis of botryolide-E pp 828–829
D. Chandra Rao, D. Kumar Reddy, V. Shekhar, Y. Venkateswarlu*



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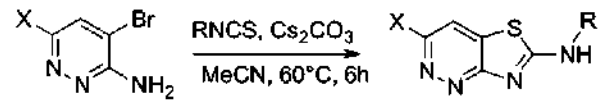
Alessandro Stella, Steven De Jonghe, Kenneth Segers, Piet Herdewijn*



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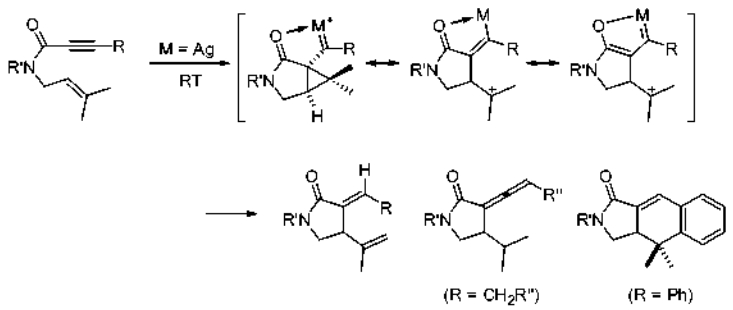

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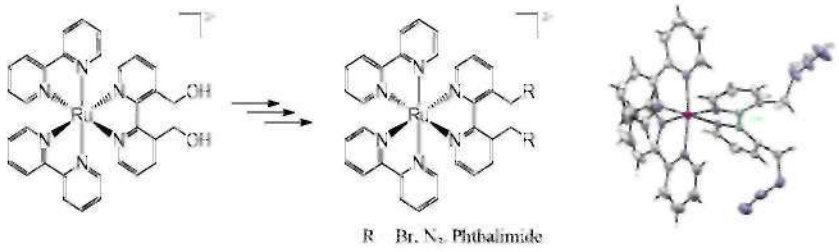

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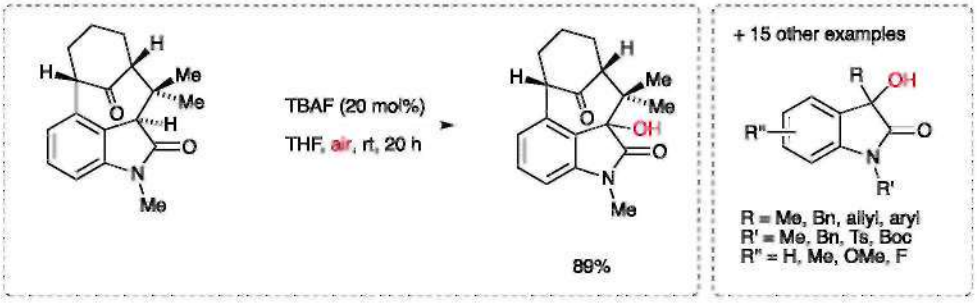
Benjamin R. Buckley*, Beatriz Fernández D.-R.



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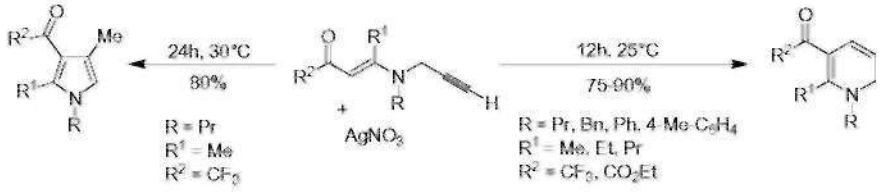

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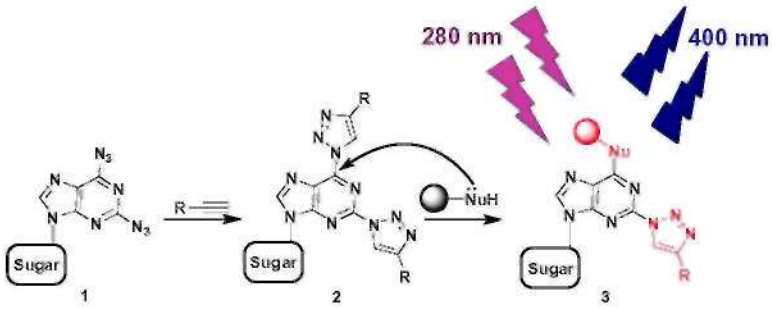

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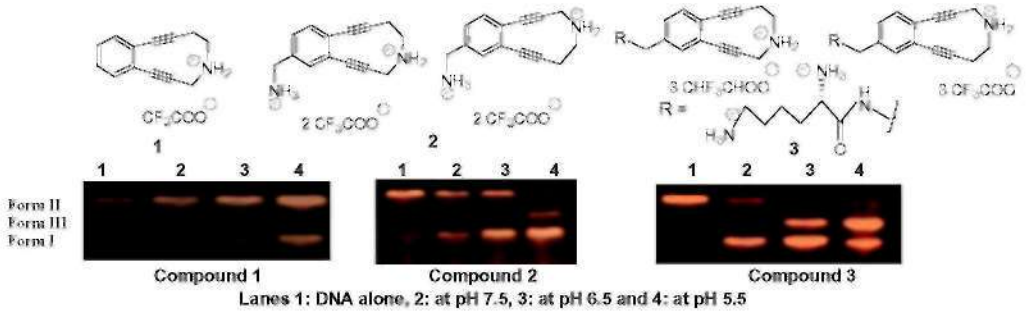
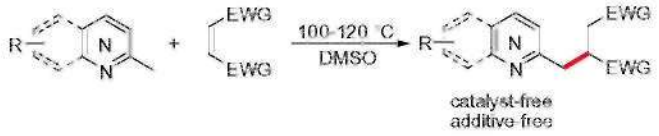
Ishita Hatial, Partha S. Addy, Ananta K. Ghosh, Amit Basak*



An addition of benzylic sp3 C–H to electron-deficient olefins pp 858–860
Hong-Ying Li, Li-Juan Xing, Tong Xu, Peng Wang, Rui-Hua Liu*, Bin Wang*



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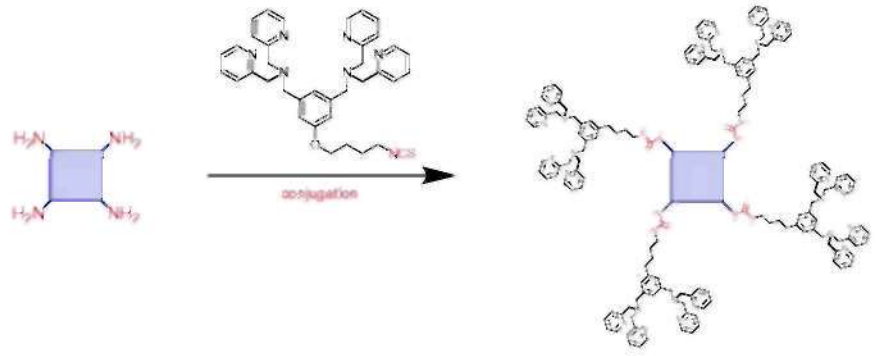
Shuzhang Xiao, Serhan Turkyilmaz, Bradley D. Smith*



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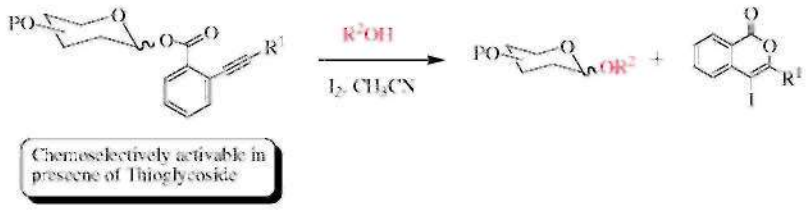
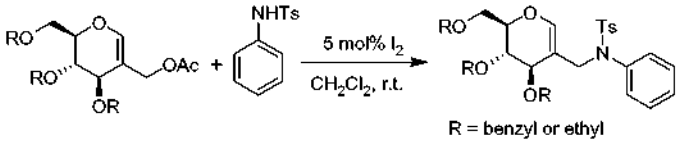
Samrat Dutta, Swarbhanu Sarkar, Shyam Ji Gupta, Asish Kumar Sen*



S_N2 substitution reaction of 2-*C*-acetoxymethyl glycals catalyzed by iodine: a novel synthesis of 2-*C*-*N*-arylamidomethyl glycals **pp 871–873**
J. S. Yadav, G. Narasimhulu, N. Umadevi, Y. Vikram Reddy, B. V. Subba Reddy*



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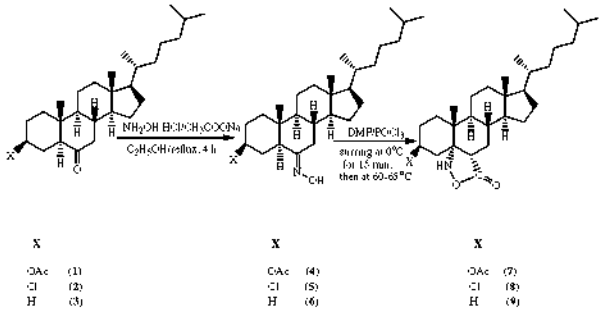
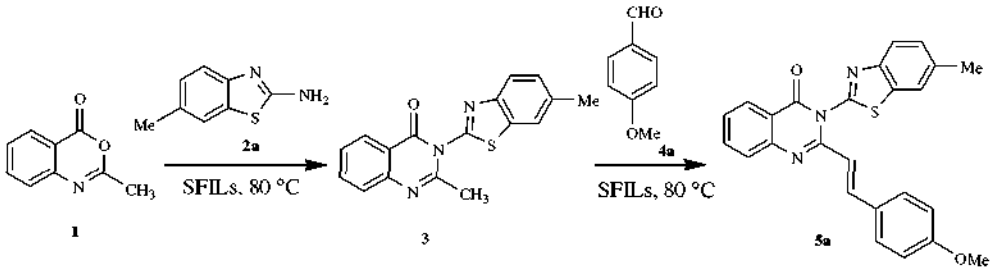
Shamsuzzaman*, Hena Khanam, Ashraf Mashrai, Nazish Siddiqui

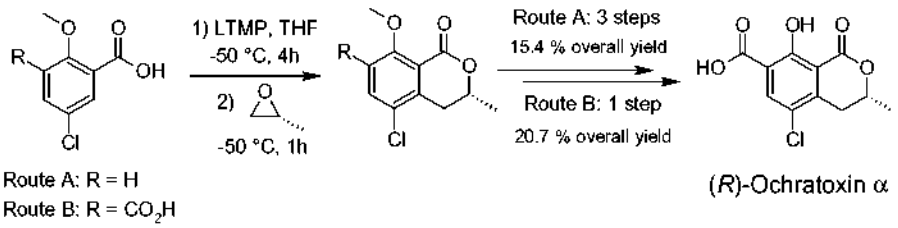


An efficient, ionic liquid mediated one-pot, three component sequential synthesis of 3-benzothiazolyl-2-styrylquinazolin-4(3*H*)-ones **pp 878–882**
Mahendra Kumar*, Kailash Sharma, Dinesh Kumar Sharma, Anand Kumar Arya



An efficient and diversity oriented one-pot three component sequential synthetic method has been presented for the synthesis of 3-benzothiazolyl-2-styrylquinazolin-4(3*H*)-ones. The synthetic method involves the reaction of 3,1-benzoxazinone with 2-aminobenzothiazole and subsequently with aromatic aldehyde using SO_3H-functionalized ionic liquids (SFILs) as solvent/catalyst.

Efficient synthesis of (*R*)-ochratoxin alpha, the key precursor to the mycotoxin ochratoxin A **pp 883–886**
Cesar Antonio Lenz, Michael Rychlik*



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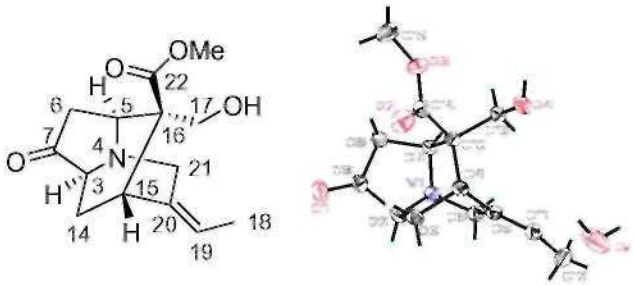
Shuang Liang*, Chun-Yong He, László F. Szabó, Yi Feng*, Xiao Lin, Yuan Wang



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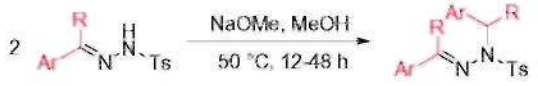
Jin-Biao Liu, Hui Yan, Gui Lu*



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Kamalesh Debnath, Sudipta Pathak, Animesh Pramanik*

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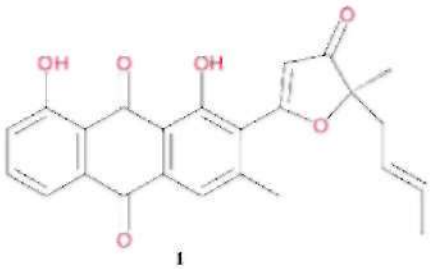
Ritesh Raju, Oleksandr Gromyko, Viktor Fedorenko, Jennifer Herrmann, Andriy Luzhetskyy, Rolf Müller*



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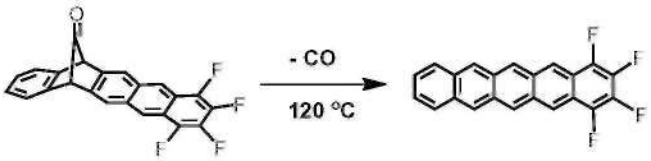

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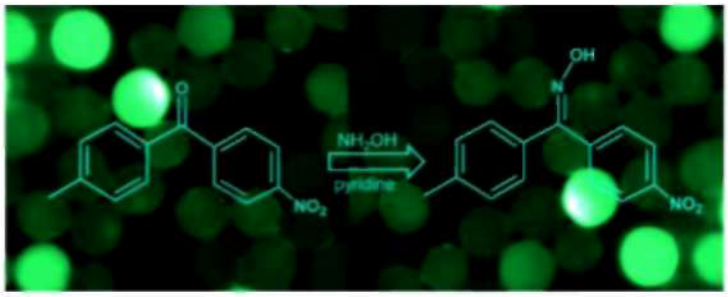
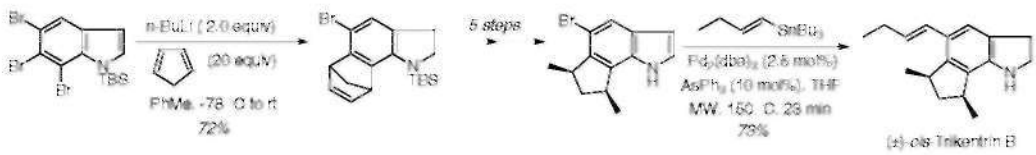
Sebastian Lüttenberg, Frank Sondermann, Jürgen Scherkenbeck*



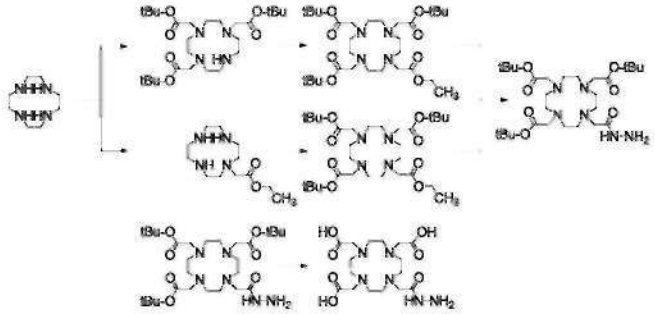
A red fluorescent 'turn-on' chemosensor for Hg^{2+} based on triphenylamine–triazines derivatives with aggregation-induced emission characteristics **pp 909–912**
Hao Zhang, Yi Qu, Yuting Gao, Jianli Hua*, Jing Li, Bo Li

A new sensitive and selective red fluorescence 'turn on' chemosensor **1** for Hg^{2+} was developed by taking advantage of AIE feature of triphenylamine–triazines motif and the specific binding of thymine with Hg^{2+}. Moreover, chemosensor **1** exhibited large two-photon absorption cross-section (3328 GM).

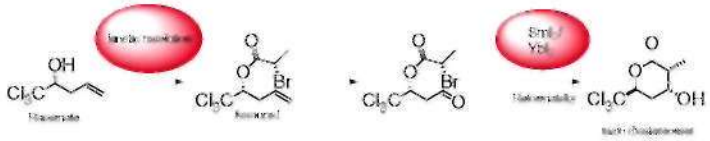
Total synthesis of (±)-*cis*-trikentrin B via intermolecular 6,7-indole aryne cycloaddition and Stille cross-coupling **pp 913–917**
Nalin Chandrasoma, Neil Brown, Allen Brassfield, Alok Nerurkar, Susana Suarez, Keith R. Buszek*



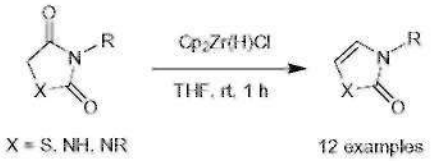
Comparison and systematic optimization of synthetic protocols for DOTA–hydrazide generation **pp 918–920**
Felix Fuge, Marek Weiler, Jessica Gätjens, Twan Lammers, Fabian Kiessling*



Synthesis of β-hydroxy-δ-trichloromethyl-δ-valerolactones by intramolecular samarium/ytterbium diiodide-mediated Reformatsky reaction **pp 921–924**
Thies M. Schulze, Jörg Grunenberg, Stefan Schulz*

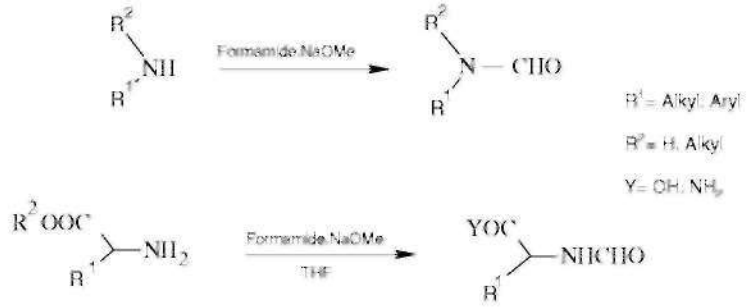


Schwartz reagent mediated synthesis of thiazolones and imidazolones from thiazolidine-2,4-diones and imidazolidine-2,4-diones **pp 925–928**
Srinivasa Reddy Dandepally, Radouane Elgoummadi, Alfred L. Williams*

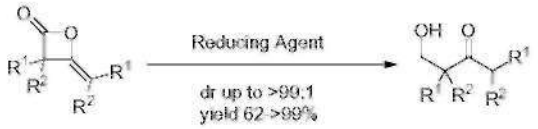


A novel reduction/elimination method of thiazolidine-2,4-dione and imidazolidine-2,4-dione derivatives using Schwartz reagent to synthesize numerous thiazolones and imidazolones in a single step is reported.

A convenient procedure for N-formylation of amines **pp 929–931**
Sony Joseph, Prasenjit Das, Bindu Srivastava, Hashim Nizar*, Mohan Prasad



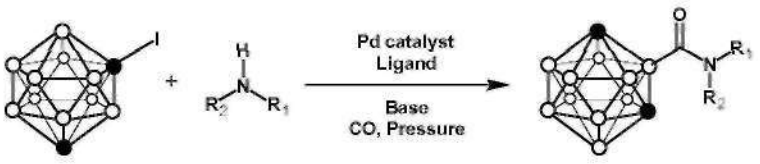
Diastereoselective formation of β-hydroxyketones by the reduction of Ketene dimers **pp 932–935**
Pei-Hsun Wei, Melanie A. Gary, Divya Nalla, Gero D. Harzmann, Ahmad A. Ibrahim, Kyle R. Dayak, Nessan J. Kerrigan*



A general method for the diastereoselective formation of β-hydroxyketones by the reduction of ketene dimers was developed. Methylphenylketene dimer was reduced with optimal diastereoselectivity (dr up to 6:1) using $LiBH_4$. However, more generally LiAlH4 was found to be the most effective reducing system with respect to diastereoselectivity (dr up to >99:1) and yield (62–>99% for 10 examples).

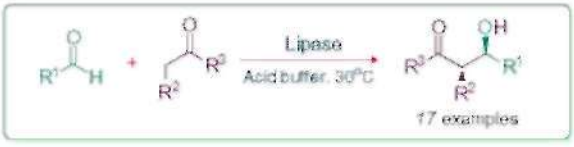
The use of aqueous potassium dichloroiodate for the synthesis of ureas **pp 936–940**
Gil Mendes Viana, Lúcia Cruz de Sequeira Aguiar*, Jonas de Araújo Ferrão, Alessandro Bolis Costa Simas, Marcela Guariento Vasconcelos

Synthesis of *m*-carboranyl amides via palladium-catalyzed carbonylation **pp 941–944**
Kiran Babu Gona, Vanessa Gómez-Vallejo, Jordi Llop*

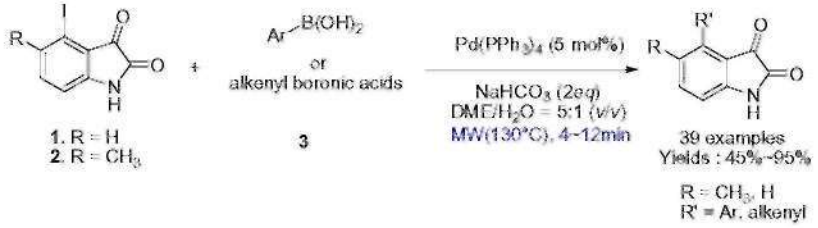


One-pot one-step reaction for the synthesis of secondary and tertiary *m*-carboranyl amides via palladium catalyzed carbonylation.

Biocatalytic asymmetric aldol reaction in buffer solution **pp 945–948**
Zong-Bo Xie, Na Wang*, Guo-Fang Jiang, Xiao-Qi Yu*

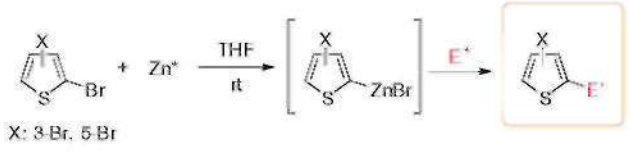


Efficient synthesis of bulky 4-substituted-isatins via microwave-promoted Suzuki cross-coupling reaction pp 949–955
Yu-Chao Liu, Chen-Jin Ye, Qiong Chen*, Guang-Fu Yang*



Mechanistic aspect of ring transformations in the reaction of 5-nitro-4-pyrimidinone with acetophenone derivatives and cycloalkanones depending on the electron density/ring size of the ketone pp 956–959
Nagatoshi Nishiwaki*, Ryuichi Sugimoto, Kazuhiko Saigo, Kazuya Kobiro

Site-selective mono-oxidative addition of active zinc into carbon–bromine bond of dibrominated-thiophenes: preparation of thienylzinc reagents and their applications pp 960–964
Hye-Soo Jung, Hyun-Hee Cho, Seung-Hoi Kim*



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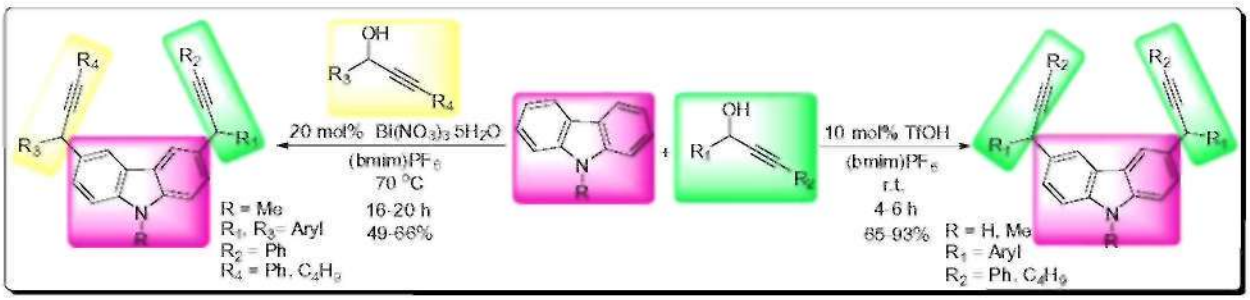
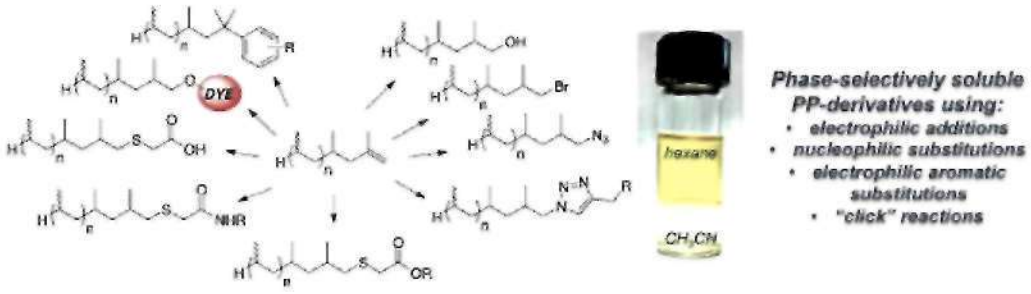
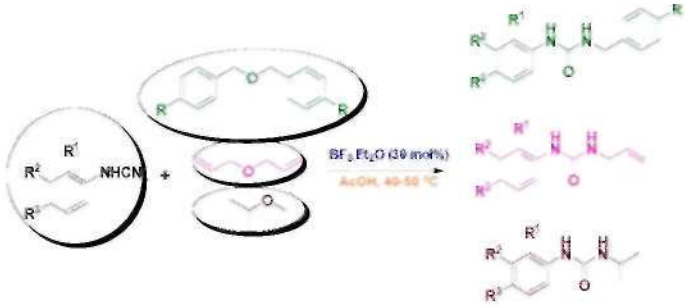
G. G. K. S. Narayana Kumar, Kenneth K. Laali*



Terminal functionalization of atactic polypropylene: a new soluble polymer support pp 970–974
Binhong Lin, Devin Lawler, Gregory P. McGovern, Christopher A. Bradley, Christopher E. Hobbs*



An efficient transformation of ethers to *N,N*-disubstituted ureas in a Ritter type reaction pp 975–979
Veladi Panduranga, Basavaprabhu, Vommina V. Sureshbabu*



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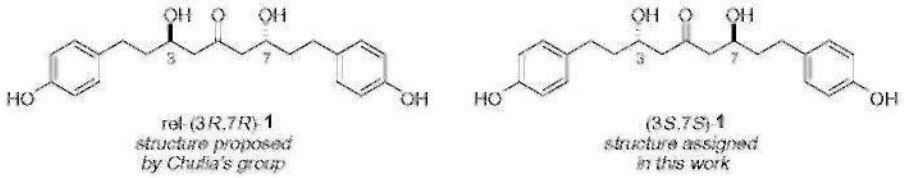
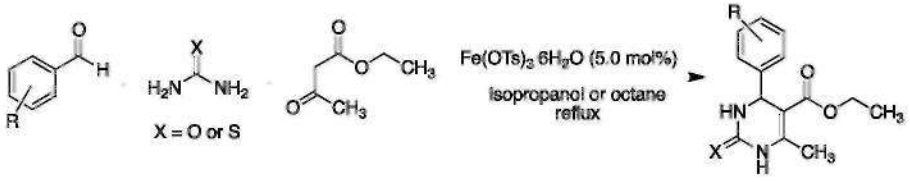
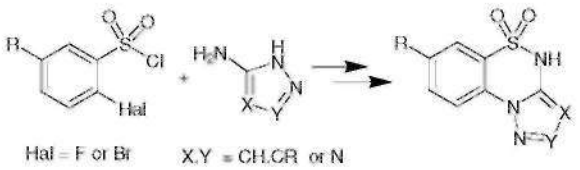
Luiz C. Dias*, Paula K. Kuroishi, Ellen C. Polo, Emílio C. de Lucca Jr.



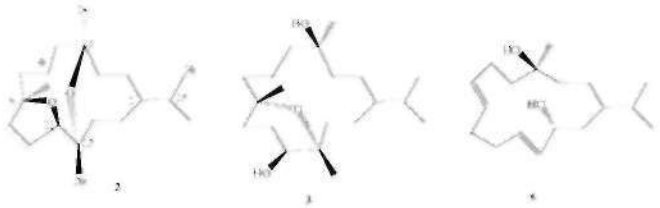
Iron(III) tosylate catalyzed synthesis of 3,4-dihydropyrimidin-2(1*H*)-ones/thiones via the Biginelli reaction pp 983–985
Jacob T. Starcevich, Thomas J. Laughlin, Ram S. Mohan*



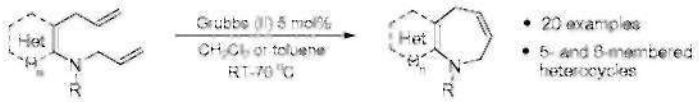
A facile synthesis of annulated azolo[*c*][1,2,4]thiadiazine *S,S*-dioxides **pp 986–988**
Artem Cherepakha*, Vladimir O. Kovtunenko, Andrey Tolmachev



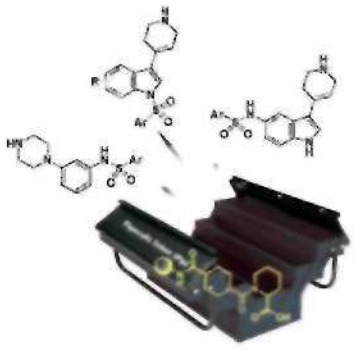
Bioactive cembranoids from the Red Sea soft coral *Sarcophyton glaucum* **pp 989–992**
Rania F. Abou El-Ezz, Safwat A. Ahmed, Mohamed M. Radwan, Nahla A. Ayoub, Manal S. Afifi, Samir A. Ross, Pawel T. Szymanski, Hesham Fahmy, Sherief I. Khalifa*



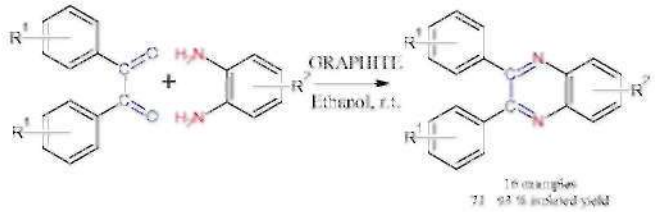
A ring-closing metathesis approach to heterocycle-fused azepines **pp 993–997**
Thomas A. Moss*



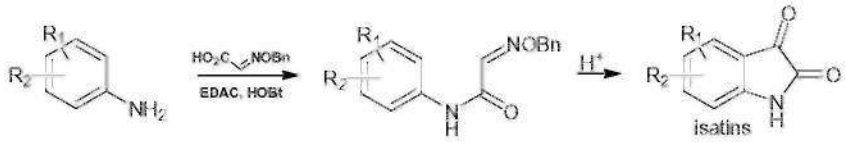
The pipecolic linker—an acid-labile handle for derivatization of secondary amines on a solid-support. Part 3 **pp 998–1002**
Paweł Zajdel*, Nicolas Masurier, Vittorio Canale, Pascal Verdie, Muriel Amblard, Maciej Pawłowski, Jean Martinez, Gilles Subra



Graphite catalyzed green synthesis of quinoxalines **pp 1003–1007**
Hari K. Kadam, Salman Khan, Rupesh A. Kunkalkar, Santosh G. Tilve*

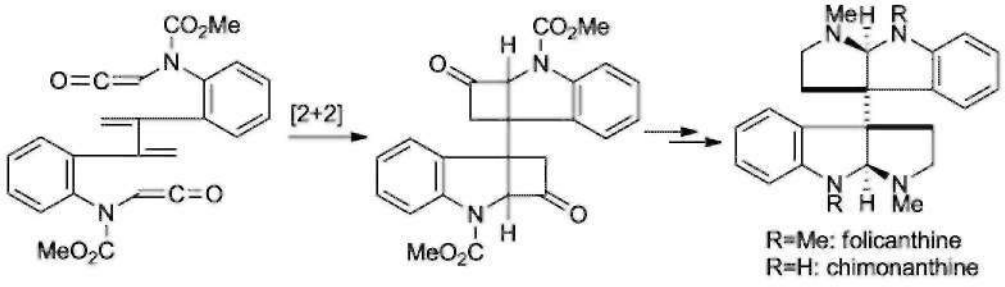


Synthesis of substituted isatins **pp 1008–1011**
Larry L. Klein*, Michael D. Tufano



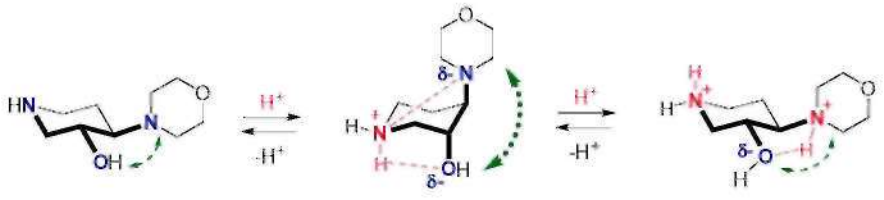
Isatins are valuable intermediates for heterocyclic chemistry. Most of the common methods for their production are less than adequate when the number and lipophilicity of substituents on the targeted isatin are increased. Our group desired such molecules and identified an alternative method for their production.

Total syntheses of (±)-folicanthine and (±)-chimonanthine via a double intramolecular carbamoylketene–alkene [2+2] cycloaddition **pp 1012–1014**
Takaaki Araki, Yuki Manabe, Kosuke Fujioka, Hiromasa Yokoe, Makoto Kanematsu, Masahiro Yoshida, Kozo Shishido*

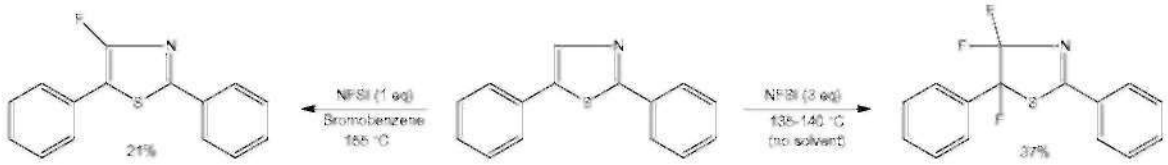


Specific naked eye sensing of cyanide by chromogenic host: studies on the effect of solvents **pp 1015–1019**
Jongmin Kang*, Eun Joo Song, Hyun Kim, Young-Hee Kim, Youngmee Kim, Sung Jin Kim, Cheal Kim*

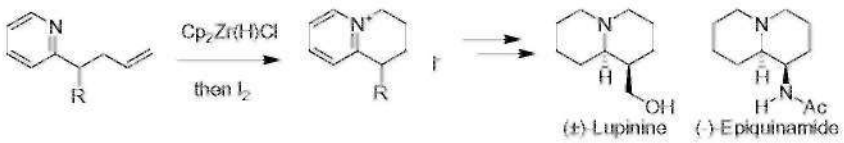
trans-3-Hydroxy-4-morpholinopiperidine—the pH-triggered conformational switch with a double flip **pp 1020–1024**
Andrey V. Samoshin, Hyun Joo, Andrei Ya Korneichuk, Ivan S. Veselov, Galina V. Grishina, Vyacheslav V. Samoshin*



Mono- and trifluorination of the thiazole ring of 2,5-diarylthiazoles using N-fluorobenzenesulfonimide (NFSI) **pp 1025–1028**
Julie M. Hatfield, Cheryl K. Eidell, Chad E. Stephens*



A hydrozirconation/iodination-mediated access to tetrahydroquinolizinium salts. Application to the synthesis of Lupinine and (−)-Epiquinamide **pp 1029–1031**
Majdi Hajri, Clément Blondelle, Agathe Martinez, Jean-Luc Vasse*, Jan Szymoniak



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*Corresponding author
ⓘ+ Supplementary data available via SciVerse ScienceDirect

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