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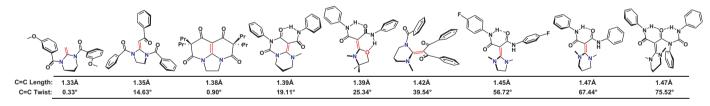


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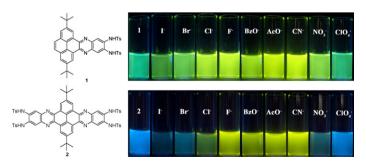


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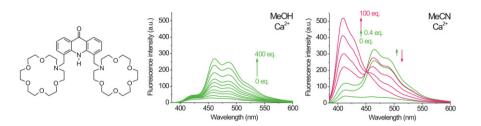
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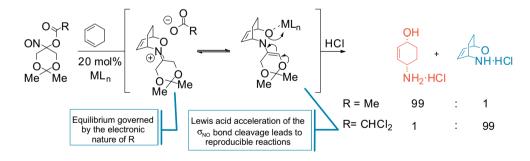
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A piperidinium triflate catalyzed Biginelli reaction

Chennan Ramalingan, Su-Jung Park, In-Sook Lee, Young-Woo Kwak*

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$$\begin{array}{c|c} & & & & \\ & & & \\ & & & \\ &$$

$Fe(ClO_4)_3 \cdot \times H_2O$ -Catalyzed direct C–C bond forming reactions between secondary benzylic alcohols with different types of nucleophiles

pp 2995-3003

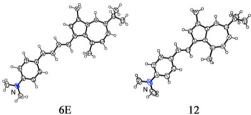
Ponnaboina Thirupathi, Sung Soo Kim*

A mild and efficient $Fe(ClO_4)_3 \cdot \times H_2O$ -catalyzed direct C-C bond coupling reactions of 1,3-dicarbonyl compounds, electron-rich heteroarenes and arenes, and 4-hydroxycoumarin with secondary benzylic alcohols have been described.

Preparation, crystal structure, and spectroscopic, chemical, and electrochemical properties of (2E,4E)-4-[4-(dimethylamino)phenyl]-1-(3-guaiazulenyl)-1,3-butadiene compared with those of (E)-2-[4-(dimethylamino)phenyl]-1-(3-guaiazulenyl)ethylene

pp 3004-3015

Shin-ichi Takekuma*, Hiroto Matsuoka, Toshie Minematsu, Hideko Takekuma



Wittig reaction of 3-[4-(dimethylamino)phenyl]propanal with (3-guaiazulenylmethyl)triphenylphosphonium bromide in ethanol containing NaOEt at 25 °C for 24 h under argon gives the title compound **6E** in 19% isolated yield. Preparation and crystal structure as well as spectroscopic, chemical, and electrochemical properties of **6E** compared with those of **12** are reported in detail.



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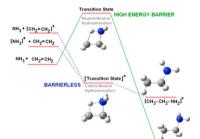
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Exploring the versatility of the Johnson–Claisen rearrangement: access to functionally versatile δ -ethoxycarbonyl- α,β -unsaturated nitriles

pp 3050-3057

Kelly L. Cosgrove, Ross P. McGeary*

Subjecting cyanohydrins derived from enals to standard Johnson–Claisen conditions does not lead to the desired products. We have now developed methodology for accomplishing this reaction, giving δ -ethoxycarbonyl- α , β -unsaturated nitriles in good yields.

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Stéphane Gérard*, Andrea Renzetti, Bérangère Lefevre, Antonella Fontana, Paolo de Maria, Janos Sapi

Inverse electron demand asymmetric cycloadditions of cyclic carbonyl ylides catalyzed by chiral Lewis acids—scope and limitations of diazo and olefinic substrates

pp 3070-3089

Hiroyuki Suga*, Satoshi Higuchi, Motoo Ohtsuka, Daisuke Ishimoto, Tadashi Arikawa, Yuta Hashimoto, Shunta Misawa, Teruko Tsuchida, Akikazu Kakehi, Toshihide Baba

$$\begin{array}{c|cccc} O & Rh_2(OAc)_4 & R^2 \\ \hline R^2 & Chiral \\ \hline N_2 & Lewis acid \\ \hline OR & R^1 & OR \\ \end{array}$$

Benzo, R^1 = H, R^2 = OMe, R = c-Hex: 96% ee (endo) Benzo, R^1 = Acyl, R^2 = OMe, R = c-Hex: 73 - 97% ee (endo) R^1 = H, R^2 = Alkyl and Ph, R = Bu: 75 - 84% ee (exo)



*Corresponding author

(1)+ Supplementary data available via ScienceDirect

COVER

 α -Acyloxynitroso derivatives are a class of heterodienophiles leading to valuable 1,4-syn-aminoalcohols in good yields starting from 1,3-dienes. The discovery that a α -oxygenated moiety led to a domino [4+2] cycloaddition/ σ (N-O) bond cleavage in the presence of a catalytic amount of Lewis acid was investigated in detail, through kinetic profiling of the reaction.

Details can be found in Tetrahedron, 2010, 66, 2969–2980.

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