# Spotlights ...

## Liquid Crystals

B.-H. Tan, M. Yoshio, T. Kato\*

Induction of Columnar and Smectic Phases for Spiropyran Derivatives: Effects of Acidichromism and Photochromism

Chem. Asian J.

DOI: 10.1002/asia.200700225

R<sup>2</sup>
R<sup>3</sup>
CH<sub>3</sub>SO<sub>3</sub>H
CH<sub>3</sub>SO<sub>5</sub>CH<sub>3</sub>
Nonmesomorphic

Nonmesomorphic

Nonmesomorphic

Nonmesomorphic

By means of chemical or light? Liquidcrystalline phases of spiropyran derivatives are formed by acid-induced spiroprotonated-merocyanine isomerization. On the contrary, photoirradiation of the spiropyran compounds does not lead to the formation of a mesophase.

### **Natural Products**

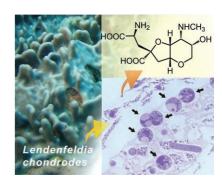
R. Sakai,\* K. Yoshida, A. Kimura, K. Koike, M. Jimbo, K. Koike, A. Kobiyama, H. Kamiya

Cellular Origin of Dysiherbaine, an Excitatory Amino Acid Derived from a Marine Sponge

ChemBioChem

DOI: 10.1002/cbic.200700498

Pinpointing responsibility: The marine-sponge toxin dysiherbaine was shown by immunohistochemical methods to be localized in the cells of the endosymbiotic cyanobacteria Synechosystis harbored in the host sponge Lendenfeldia chondrodes (see light micrograph of cells). Chemical analysis indicated the presence of two chemotypes of the cyanobacterium, only one of which appears to produce the toxin.



### **Electronic Engineering**

N. Armaroli\*

Electronic Excited-State Engineering

Chem Phys Chem

DOI: 10.1002/cphc.200700794

Long-living complexes: A Cu<sup>1</sup>- bisphenanthroline complex exhibits a 15-fold prolongation of its excited state lifetime due to the planned intervention of an appended anthracene fragment (see picture). This elegant example of electronic excited-state engineering extends the range of possibilities for improving the photophysical properties of Cu<sup>1</sup> complexes.



# Drug Discovery

J. Doyon,\* E. Coesemans, S. Boeckx, M. Buntinx, B. Hermans, J. P. Van Wauwe, R. A. H. J. Gilissen, A. H. J. De Groot, D. Corens, G. Van Lommen\*

Discovery of Potent, Orally Bioavailable Small-Molecule Inhibitors of the Human CCR2 Receptor

ChemMedChem

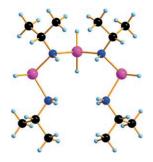
DOI: 10.1002/cmdc.200700276

The synthesis of small-molecule CCR2 receptor inhibitors is presented. The initial lead, although quite potent, was metabolically unstable both in vitro and in vivo. Isosteric replacement of one of

the ester functions with small heterocycles led to highly potent, orally bioavailable and metabolically stable compounds.

# ... on our Sister Journals





LiGaH<sub>4</sub> can be made to react with a 50% molar excess of the amine hydrochloride [RNH<sub>3</sub>]Cl to afford the cationic gallane derivative [(RH<sub>2</sub>N)<sub>2</sub>GaH<sub>2</sub>]<sup>+</sup>Cl<sup>-</sup>, with R = Me or *i*Pr, in 45–65% yield. A significant secondary product for R = *i*Pr is the trigallium compound [{(*i*PrH<sub>2</sub>N)GaH<sub>2</sub>NH-*i*Pr}<sub>2</sub>GaH<sub>2</sub>]<sup>+</sup>Cl<sup>-</sup>. The structures and other properties of such compounds give evidence of their mediating the formation of neutral amidogallanes, as well as having wider possible implications.

#### Cationic Gallane Derivatives

C. Y. Tang, A. R. Cowley, A. J. Downs,\* S. Marchant, S. Parsons

Formation and Characterization of the Cationic Gallane Derivatives  $[(RH_2N)_2GaH_2]Cl$  (R = Me or *i*Pr) and  $[\{(iPrH_2N)GaH_2NHiPr\}_2GaH_2]Cl$ 

Eur. J. Inorg. Chem.

DOI: 10.1002/ejic.200701120



Does the mechanistic pathway of the title reaction pass through an enamino-lactone intermediate? Both the experimental and theoretical data have allowed us to suggest a plausible bifunctional catalytic role for proline in the Baylis—Hillman reaction between MVK or EVK and aryl aldehydes using hydrogen carbonate as a co-catalyst.

### **Organocatalysis**

M. Gruttadauria,\* F. Giacalone,
P. Lo Meo, A. Mossuto Marculescu,
S. Piala, P. Noto

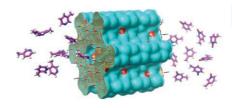
S. Riela, R. Noto

First Evidence of Proline Acting as a Bifunctional Catalyst in the Baylis-Hillman Reaction Between Alkyl Vinyl Ketones and Aryl Aldehydes

Eur. J. Org. Chem.

DOI: 10.1002/ejoc.200701112

Its green and selective! Single-site heterogeneous catalysts (shown here) can be judiciously combined with a solid source of active oxygen for the single-step, solvent-free, and environmentally benign production of Niacin (used in the preparation of Vitamin B<sub>3</sub>) and other fine-chemical, pharmaceutical, and agrochemical intermediates.



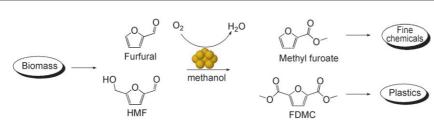
### Heterogeneous Catalysis

R. Raja,\* J. M. Thomas,\* M. Greenhill-Hooper, S. V. Ley, F. A. Almeida Paz

Facile, One-Step Production of Niacin (Vitamin B<sub>3</sub>) and Other Nitrogen-Containing Pharmaceutical Chemicals with a Single-Site Heterogeneous Catalyst

Chem. Eur. J.

DOI: 10.1002/chem.200701679



Aerobic exercise: The biomass-derived platform chemicals furfural and hydroxymethylfurfural (HMF) are readily oxidized in methanol in the presence of oxygen and a supported gold nanoparticle catalyst to afford the corresponding

methyl esters (see scheme). Thus, furfural was oxidized to methyl furoate under very mild conditions, and HMF was converted into furan-2,5-dimethylcarboxylate (FDMC), a potential polymer building block, with high yields.

### Heterogeneous Catalysis

E. Taarning, I. S. Nielsen, K. Egeblad, R. Madsen, C. H. Christensen\*

Chemicals from Renewables: Aerobic Oxidation of Furfural and Hydroxymethylfurfural over Gold Catalysts

ChemSusChem

DOI: 10.1002/cssc.200700033