

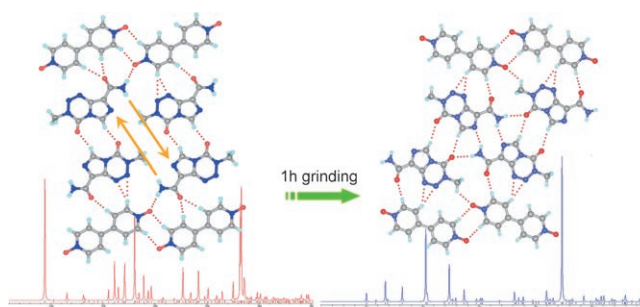
Crystal Polymorphism

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A. Nangia*

Polymorphs and Polymorphic Cocrystals
of Temozolomide

Chem. Asian J.

DOI: 10.1002/asia.200800070



How to morph: Unused hydrogen-bond donors/acceptors in 2:1 temozolomide-4,4'-bipyridine-*N,N'*-dioxide cocrystals (orange arrows) form intermolecular

N—H...N bonds in the stable polymorph (right). The structural origins of synthon polymorphism in cocrystals are remarkably similar to those of polymorphs.

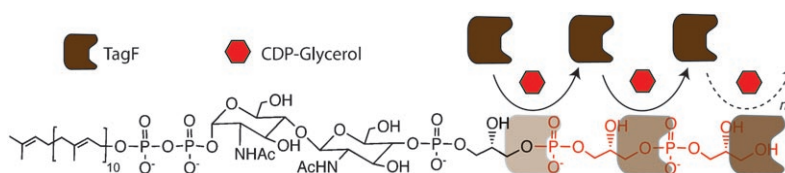
Teichoic Acid

M. P. Pereira, J. W. Schertzer,
M. A. D'Elia, K. P. Koteva,
D. W. Hughes, G. D. Wright,
E. D. Brown*

The Wall Teichoic Acid Polymerase TagF
Efficiently Synthesizes Poly(glycerol
phosphate) on the TagB Product Lipid III

ChemBioChem

DOI: 10.1002/cbic.200800026



One piece at a time: Glycerol phosphate polymerization by the TagF enzyme is investigated through the use of a synthetic analogue of a teichoic acid intermediate. Analysis of polymerization

products and reaction kinetics of the enzyme suggest a distributive assembly of the teichoic acid polymer on the TagB product lipid III.

Molecular Machines

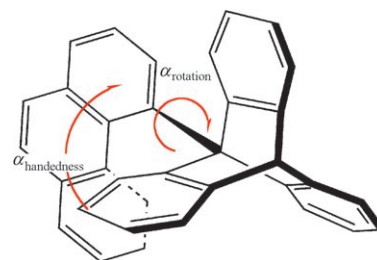
M. Llunell, P. Alemany,* J. M. Bofill*

Conformational Analysis of Molecular
Machines: Internal Rotation and
Enantiomerization in Triptycyl[3]helicene

ChemPhysChem

DOI: 10.1002/cphc.200800052

No independent rotation: Detailed analysis of the PES for triptycyl[3]helicene shows that rotation around the helicene–triptycyl bond cannot be considered independently from other degrees of freedom when analyzing its stereodynamic behavior. The possibility of enantiomerization of the helicene pawl results in more complex dynamics than previously expected.



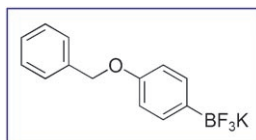
Drug Design

N. Lecat-Guillet, Y. Ambroise*

Discovery of Aryltrifluoroborates as
Potent Sodium/Iodide Symporter (NIS)
Inhibitors

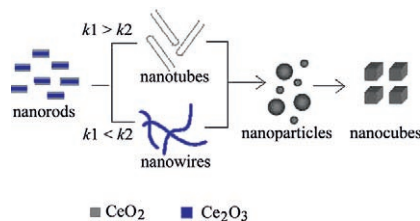
ChemMedChem

DOI: 10.1002/cmdc.200800049



The structure-based design of sodium/iodide symporter (NIS) inhibitors identified new active compounds. The organotrifluoroborate shown was found to inhibit iodide uptake with an IC₅₀ value of 0.4 μM on rat-derived thyroid cells. The biological activity is rationalized by the presence of the BF₃[−] ion as a minimal binding motif for substrate recognition at the iodide binding site.

A template-free method is employed first to synthesize single-crystal CeO_2 nanorods at room temperature and pressure, and then the controlled conversion of nanorods into nanotubes, nanowires, and nanocubes is realized. Finally, CO oxidation properties of CeO_2 nanostructures were investigated systematically and CeO_2 nanorods were found to have excellent catalytic performance.



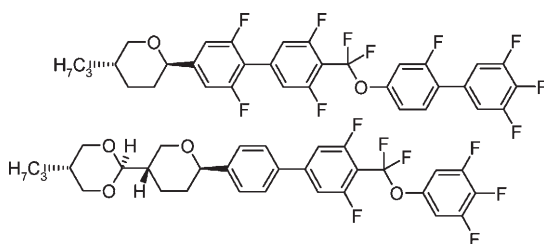
Controllable Ceria Nanostructures

C. Pan, D. Zhang,* L. Shi,* J. Fang

Template-Free Synthesis, Controlled Conversion, and CO Oxidation Properties of CeO_2 Nanorods, Nanotubes, Nanowires, and Nanocubes

Eur. J. Inorg. Chem.

DOI: 10.1002/ejic.200800047



Highly fluorinated liquid crystals with unprecedented polarity were obtained by introducing a tetrahydropyran moiety into their mesogenic core structure. The new materials exhibit an unusually

favourable combination of meso-phase properties, excellent solubility and low rotational viscosity, which renders them highly attractive for application in fast-switching active-matrix LCDs.

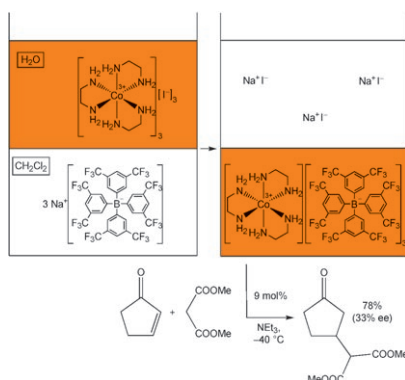
Fluorinated Liquid Crystals

P. Kirsch,* W. Binder, A. Hahn, K. Jähring, M. Lenges, L. Lietzau, D. Maillard, V. Meyer, E. Poetsch, A. Ruhl, G. Unger, R. Fröhlich

Super-Fluorinated Liquid Crystals: Towards the Limits of Polarity

Eur. J. Org. Chem.

DOI: 10.1002/ejoc.200800149



Werner complexes go organic! When orange aqueous solutions of $[\text{Co}(1,2\text{-diamine})_3]\text{X}_3$ ($\text{X} = \text{I}, \text{Cl}$) are treated with CH_2Cl_2 solutions of NaBAR_f (1:3 mol ratio), the aqueous phases decolorize (see scheme) and $[\text{Co}(1,2\text{-diamine})_3](\text{BAR}_f)_3 \cdot n\text{H}_2\text{O}$ can be isolated from the CH_2Cl_2 phase (78–89%). The ethylenediamine complex can catalyze enantioselective Michael additions, presumably by an outer coordination sphere mechanism involving hydrogen bonding.

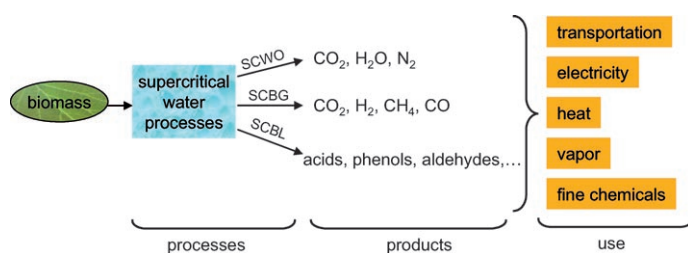
Enantioselective Catalysis

C. Ganzmann, J. A. Gladysz*

Phase Transfer of Enantiopure Werner Cations into Organic Solvents: An Overlooked Family of Chiral Hydrogen Bond Donors for Enantioselective Catalysis

Chem. Eur. J.

DOI: 10.1002/chem.200800226



Critical times, supercritical measures:

The supercritical water oxidation (SCWO) process has been studied intensively during the past 15 years and proved efficient in decomposing organic matter. Armed with this know-how,

supercritical water biomass valorization can now be developed for the production of gases and valuable chemicals (supercritical water biomass gasification (SCBG) or liquefaction (SCBL), respectively).

Supercritical Water

A. Loppinet-Serani,* C. Aymonier, F. Cansell

Current and Foreseeable Applications of Supercritical Water for Energy and the Environment

ChemSusChem

DOI: 10.1002/cssc.200700167