

SPOTLIGHTS ...

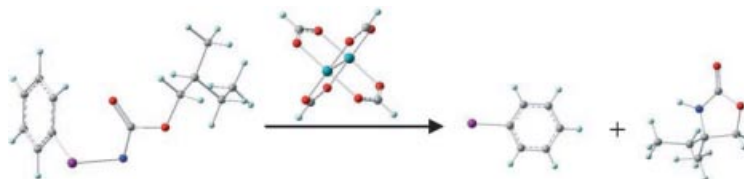
Reaction Mechanisms

X. Lin, C. Zhao, C.-M. Che,* Z. Ke, D. L. Phillips

A DFT Study on the Mechanism of $\text{Rh}_2^{\text{II,II}}$ -Catalyzed Intramolecular Amidation of Carbamates

Chem. Asian J.

DOI: 10.1002/asia.200700068



Alone yet together: DFT calculations elucidate the catalytic process of a dirhodium tetracarboxylate catalyzed intramolecular C–H bond-amidation reaction. Out of six reaction pathways

that involve a metal–nitrene complex, the singlet concerted pathway that produces the product with retention of chirality at C is predicted to be the predominant one.

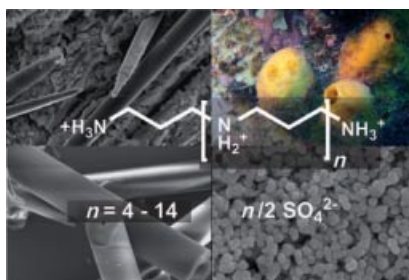
Biomineralization

S. Matsunaga, R. Sakai,* M. Jimbo, H. Kamiya

Long-Chain Polyamines (LCPAs) from Marine Sponge: Possible Implication in Spicule Formation

ChemBioChem

DOI: 10.1002/cbic.200700305



LCPAs in sponge spicules relate two distinct marine organisms, diatoms and sponges, by “silica deposition”. LCPA-directed silica deposition has been proposed for the formation of silica walls in diatoms, while other chemical factors have been suggested for sponges. Here we show that LCPAs can be an additional factor involved in spiculogenesis in the sponge.

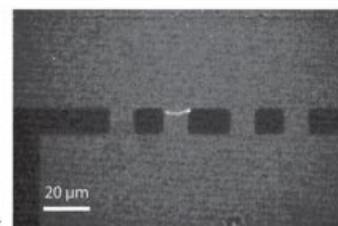
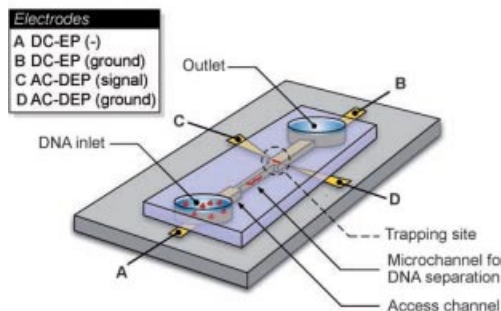
Dielectrophoretic Trapping

M. Kumemura,* D. Collard,* C. Yamahata, N. Sakaki, G. Hashiguchi, H. Fujita

Single DNA Molecule Isolation and Trapping in a Microfluidic Device

ChemPhysChem

DOI: 10.1002/cphc.200700268



DNA bridges: A double stranded λ -DNA molecule is isolated by dc electrophoresis and trapped by ac dielectrophoresis between aluminium electrodes in a microfluidic chip [see device (left) in figure and DNA bridge

between electrodes (right)]. This single-molecule trapping technique is highly effective, allowing long DNA fragments to be instantly captured in stretched formation.

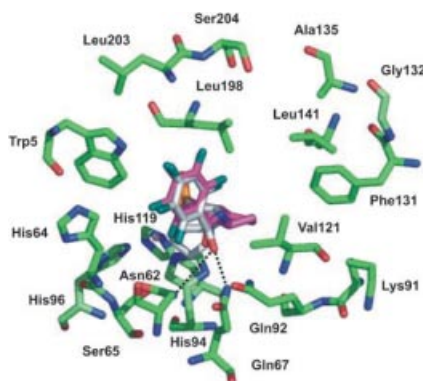
Inhibitor Design

A. Thiry,* B. Masereel, J.-M. Dogné, C. T. Supuran, J. Wouters, C. Michaux

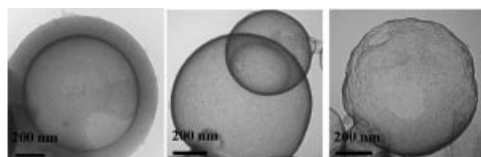
Exploration of the Binding Mode of Indanesulfonamides as Selective Inhibitors of Human Carbonic Anhydrase Type VII by Targeting Lys91

ChemMedChem

DOI: 10.1002/cmdc.200700057



The binding mode of (*R*)- (magenta) and (*S*)-1-pentafluorophenylamido-5-sulfonamide (light gray) inside the human carbonic anhydrase VII active site is shown. A shared hydrogen bond between the carbonyl group of each compound and Gln67 and Asn62 is observed. To increase hCA VII selectivity, structural modifications of these compounds are proposed in order to design new anticonvulsant agents.



Mesoporous Carbon

Silica

Metal Oxide hollow spheres

PF-PEO polymer blend can be used as a novel organic template to synthesize mesoporous carbon or silica hollow spheres. The mesoporous carbon

hollow spheres can act as a hard template for the preparation of mesoporous oxide hollow spheres.

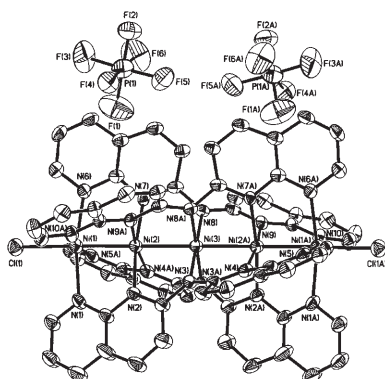
Hollow Mesoporous Spheres

C.-Y. Chang-Chien, C.-H. Hsu,
T.-Y. Lee, C.-W. Liu, S.-H. Wu,
H.-P. Lin,* C.-Y. Tang, C.-Y. Lin

Synthesis of Carbon and Silica Hollow Spheres with Mesoporous Shells using Polyethylene Oxide/Phenol Formaldehyde Polymer Blend

Eur. J. Inorg. Chem.

DOI: 10.1002/ejic.200700210



Metal string! The first linear nickel framework in which the usual sequence of Ni^{II} atoms has been reduced by two electrons is presented. The electronic structure of the metal framework appears intermediate between a localized picture corresponding to $\text{Ni}^{\text{II}}\text{-Ni}^{\text{I}}\text{-Ni}^{\text{II}}\text{-Ni}^{\text{I}}\text{-Ni}^{\text{II}}$ and a fully delocalized model represented as $(\text{Ni}_2)^{3+}\text{-Ni}^{\text{II}}\text{-(Ni}_2)^{3+}$.

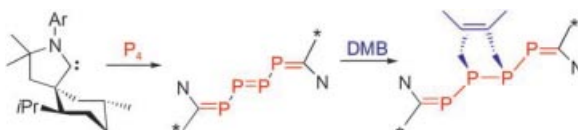
Mixed-Valent Compounds

I. P.-C. Liu, M. Bénard,* H. Hasanov,
I.-W. P. Chen, W.-H. Tseng, M.-D. Fu,
M.-M. Rohmer, C.-h. Chen,
G.-H. Lee, S.-M. Peng*

A New Generation of Metal String Complexes: Structure, Magnetism, Spectroscopy, Theoretical Analysis, and Single Molecular Conductance of an Unusual Mixed-Valence Linear $[\text{Ni}_5]^{8+}$ Complex

Chem. Eur. J.

DOI: 10.1002/chem.200700750



Broken down and added onto: An enantiomerically pure cyclic (alkyl)-(amino)carbene cleanly activates P_4 , affording highly reactive products that can be further used for the diastereoselective construction of P_4 -containing

molecules that feature phosphorus-carbon bonds (see scheme; DMB = 2,3-dimethylbutadiene; portions of the products have been omitted, * denotes stereogenic carbon center).

P_4 Activation

J. D. Masuda, W. W. Schoeller,
B. Donnadieu, G. Bertrand*

Carbene Activation of P_4 and Subsequent Derivatization

Angew. Chem. Int. Ed.

DOI: 10.1002/anie.200703055



On these pages, we feature a selection of the excellent work that has recently been published in our sister journals. If you are reading these pages on a computer, click on any of

the items to read the full article. Otherwise please see the DOIs for easy online access through Wiley InterScience.