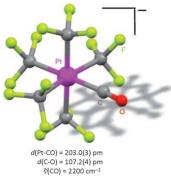




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 $V(CO) = 2200 \text{ cm}^{-1}$  $\delta_C = 158.1 \text{ ppm}$ 

Chem. Eur. J.

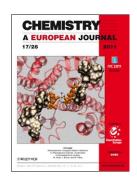
DOI: 10.1002/chem.201100626

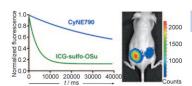
## Organoplatinum(IV) Compounds

S. Martínez-Salvador, J. Forniés,\* A. Martín, B. Menjón

Highly Trifluoromethylated Platinum Compounds

Rarest of the rare: A rare  $Pt^{IV}$  carbonyl derivative with unexpected thermal stability has been prepared and structurally characterized (see picture). Structural and spectroscopic properties point to the CO molecule acting as a mainly  $\sigma$ -donor ligand. Various other organoplatinum(IV) compounds that contain the " $Pt(CF_3)_5$ " unit are also reported.





Chem. Asian J.

DOI: 10.1002/asia.201100041

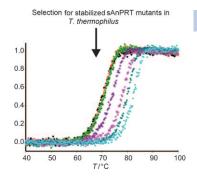
## **Imaging Agents**

Enzymes

A. Samanta, M. Vendrell, S.-W. Yun, Z. Guan, Q.-H. Xu, Y.-T. Chang\*
A Photostable Near-Infrared Protein Labeling Dye for In Vivo Imaging

A NIR death experience: A highly photostable NIR protein-labeling dye (CyNE790) that maintained excellent fluorescent properties after protein conjugation has been synthesized. Furthermore, the adequacy of the dye for NIR imaging was confirmed by showing that the recognition properties of a labeled anti-EGFR antibody remained unaffected.





ChemBioChem

DOI: 10.1002/cbic.201000770

#### T. Schwab, R. Sterner\*

Stabilization of a Metabolic Enzyme by Library Selection in *Thermus thermophilus* 

If you can stand the heat: A metabolic enzyme was stabilized by random mutagenesis and in vivo selection performed at around  $80\,^{\circ}\text{C}$  by using an extremely thermophilic host bacterium that has a natural competence for DNA uptake. The beneficial mutations identified by this approach led to an increase in the thermal stability of the purified protein of  $13\,^{\circ}\text{C}$ .







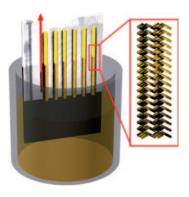


## Self-Assembled Fibers

S. Wang, M. Kivala, I. Lieberwirth, K. Kirchhoff, X. Feng, W. Pisula,\* K. Müllen\*

Dip-Coating-Induced Fiber Growth of a Soluble Heterotriangulene

Long-range well-aligned fibers are fabricated from a soluble heterotriangulene derivative via simple dip coating by controlling the concentration of the compound. This procedure is an alternative route for the generation of unique microstructures of organic semiconductors—both for both fundamental studies and for practical device applications.



ChemPhysChem

DOI: 10.1002/cphc.201100199

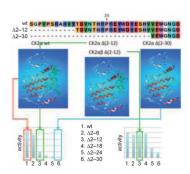


### **Computational Medicinal Chemistry**

A. Cristiani,\* G. Costa, G. Cozza, F. Meggio, L. Scapozza, S. Moro

The Role of the N-Terminal Domain in the Regulation of the "Constitutively Active" Conformation of Protein Kinase  $CK2\alpha$ : Insight from a Molecular Dynamics Investigation

**Always on**: Protein kinase CK2 is a constitutively active enzyme. Mutation studies show that CK2 activity is regulated by the interaction between the N-terminal region and the kinase domain; progressive deletions of the N-terminal tail elicit decreased kinase activity. MD simulations were carried out on wild-type,  $\Delta 2$ –12, and  $\Delta 2$ –30 deletion mutants of CK2 $\alpha$  to explore how the N-terminal tail affects CK2 conformational behavior.



ChemMedChem

DOI: 10.1002/cmdc.201100046

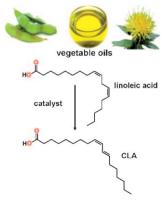


## Renewable Resources

A. Philippaerts, S. Goossens, P. A. Jacobs, B. F. Sels\*

Catalytic Production of Conjugated Fatty Acids and Oils

Conjugated vegetable oils are attractive bioresources for use in paints, inks, coatings, and plastics as well as ingredients for functional foods. Technologies for the conjugation of linoleic acid and vegetable oils using (bio)catalysts (enzymes, homogeneous, and heterogeneous catalysts) are discussed in this Review. Special attention is given to the conjugated linoleic acids productivity and their isomer distribution.

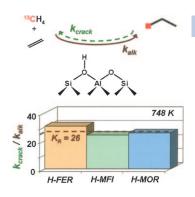


ChemSusChem

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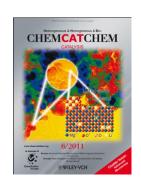
ChemCatChem DOI: 10.1002/cctc.201100051

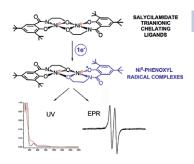
### Zeolites

R. Gounder, E. Iglesia\*

Catalytic Alkylation Routes via Carbonium-Ion-Like Transition States on Acidic Zeolites

Zeolites going crackers: Brønsted acid sites in zeolites (H-FER, H-MFI, H-MOR) catalyze CH<sub>4</sub>-alkene reactions at high temperatures (> 700 K) via carbonium-ion-like transition states. The ratio of rate constants for forward and reverse reactions (alkane alkylation-cracking, alkene dimerization-cracking) equal their respective equilibrium constants ( $K_R$ ). In contrast, relative rates of CH<sub>4</sub> and alkene reactions with a given alkoxide are influenced by the local environment around acid sites.





Eur. J. Inorg. Chem. DOI: 10.1002/ejic.201100232

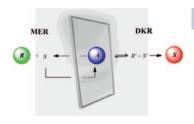
### **Radical Complexes**

L. Benisvy,\* R. Wanke, M. F. C. Guedes da Silva, A. J. L. Pombeiro\*

A Dianionic Dinickel(II) Complex and Its Oxidised Phenoxyl Radical

A dianionic dinickel(II) complex bearing new trianionic NO<sub>2</sub> ligands is reported. It displays two one-electron oxidations resulting in monoand diphenoxyl radical complexes, as indicated by UV/Vis/NIR and EPR spectroscopic data of the electrochemically oxidised species.





Eur. J. Org. Chem. DOI: 10.1002/ejoc.201100467

## Asymmetric Catalysis

A. Laurell, C. Moberg\*

Opposite Enantiomers from Minor Enantiomer Recycling and Dynamic Kinetic Resolution Using a Single Biocatalyst

The (R) enantiomer of the O-acetylated cyanohydrin from (E)-2-butenal was obtained by minor enantiomer recycling, whereas the (S) enantiomer of the same product was obtained by dynamic kinetic resolution with the use of the same catalyst.



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