

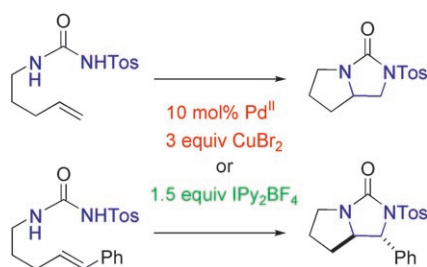
### Amination

K. Muñoz,\* C. H. Hövelmann,  
E. Campos-Gómez, J. Barluenga,\*  
J. M. González, J. Streuff, M. Nieger

Intramolecular Diamination of Alkenes  
with Palladium(II)/Copper(II) Bromide  
and  $\text{IPy}_2\text{BF}_4$ : The Role of Halogenated  
Intermediates

*Chem. Asian J.*

DOI: 10.1002/asia.200700373



**Complementary procedures** for the diamination of terminal and internal alkenes with tethered ureas and related groups as the nitrogen source provide access to cyclic urea, sulfamide, and guanidine derivatives. Opposite stereochemical pathways were identified for the diamination of terminal alkenes with  $\text{IPy}_2\text{BF}_4$  and under palladium catalysis with the reoxidant  $\text{CuBr}_2$ , but identical ones were found for internal alkenes.

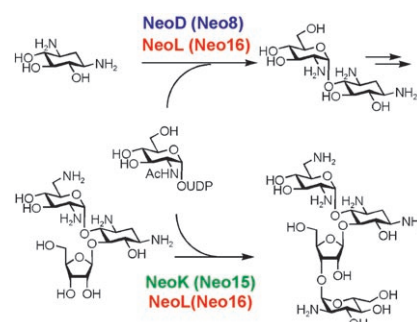
### Biosynthesis

K. Yokoyama, Y. Yamamoto, F. Kudo,  
T. Eguchi\*

Involvement of Two Distinct  
*N*-Acetylglucosaminyltransferases and a  
Dual-Function Deacetylase in Neomycin  
Biosynthesis

*ChemBioChem*

DOI: 10.1002/cbic.200700717



**A new enzyme family:** The glycosylation steps in the biosynthesis of neomycin were clarified through recombinant enzyme assays. NeoD catalyzes the first transglucosamylation, and NeoK catalyzes the second, while NeoL catalyzes both deacetylations. We found that NeoK and its homologous proteins constitute a novel glycosyltransferase family.

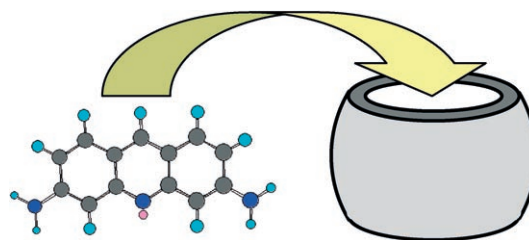
### Host–Guest Complexes

P. Montes-Navajas, A. Corma,  
H. Garcia\*

Complexation and Fluorescence of  
Tricyclic Basic Dyes Encapsulated in  
Cucurbiturils

*ChemPhysChem*

DOI: 10.1002/cphc.200700735



**Encapsulating cucurbiturils:** Tricyclic dyes can be inserted into cucurbiturils to form host–guest complexes (see figure). These complexes vary in stoichiometry,

depending on the cucurbituril involved. The stoichiometry of the complex plays a major role in the photophysical behaviour of the dyes.

### Receptor Antagonist

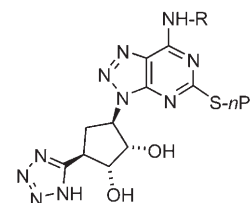
H. Ye, C. Chen, H.-C. Zhang,\*  
B. Haertlein, T. J. Parry, B. P. Damiano,  
B. E. Maryanoff\*

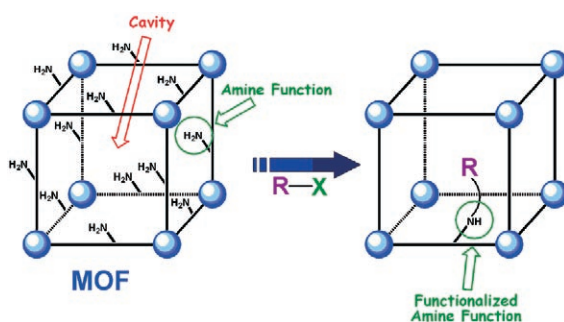
Carba-nucleosides as Potent Antagonists  
of the Adenosine 5'-Diphosphate (ADP)  
Purinerger Receptor ( $\text{P2Y}_{12}$ ) on Human  
Platelets

*ChemMedChem*

DOI: 10.1002/cmdc.200700310

**Antagonizing a key platelet purinerger receptor.** The wide clinical use of clopidogrel has highlighted the importance of platelet ADP receptor ( $\text{P2Y}_{12}$ ) antagonists for preventing adverse cardiovascular events. We synthesized a series of novel carba-nucleosides and examined their usefulness as  $\text{P2Y}_{12}$  antagonists. Some tetrazole derivatives were high-affinity receptor antagonists and potent inhibitors of human platelet aggregation.





A new metal–organic framework with amino groups oriented inside the pores has been synthesized. The post-synthetic modification of the cavities in this MOF with two different functionalities is for

the first time clearly evidenced by X-ray crystallography. The cavities of the MOF can be transformed without modifying the original 3D structure of the MOF.

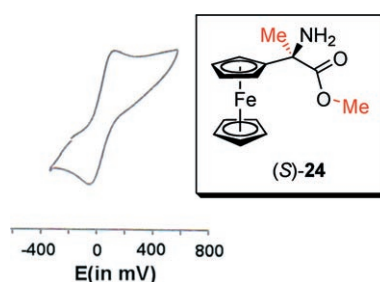
#### Post-Synthetically Modified MOFs

J. S. Costa, P. Gamez,\* C. A. Black, O. Roubeau, S. J. Teat, J. Reedijk

Chemical Modification of a Bridging Ligand Inside a Metal–Organic Framework while Maintaining the 3D Structure

*Eur. J. Inorg. Chem.*  
DOI: 10.1002/ejic.200800002

Are  $\alpha$ -ferrocenyl- $\alpha$ -amino esters stable? The answer is yes provided that the  $\alpha$ -carbon is tetrasubstituted and that very mild reaction conditions are involved in their generation. Thus, the (*S*) enantiomer of methyl  $\alpha$ -ferrocenylalaninate **24**, prepared in around 90% *ee* from 2-ferrocenylpropene, is a stable compound that can be stored for prolonged periods of time without appreciable decomposition.

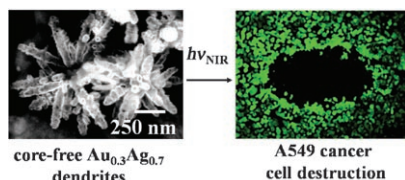


#### Ferrocenyl Amino Acids

R. M. Moreno, M. Catasús, C. López, A. Moyano\*

Enantiocontrolled Preparation of the First Stable  $\alpha$ -Ferrocenylalanine Derivatives

*Eur. J. Org. Chem.*  
DOI: 10.1002/ejoc.200800029



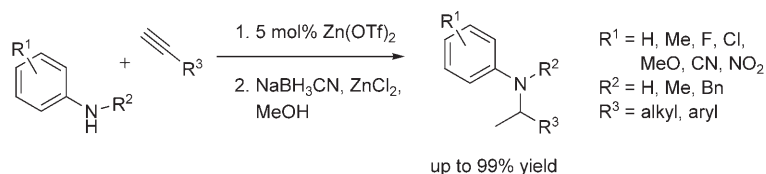
**Gold–silver therapy:** Core-free  $\text{Au}_x\text{Ag}_{1-x}$  nanostructured dendrites have been developed that show absorption in the NIR region, exhibiting potential as photothermal therapeutic agents. An effective photothermal capability was found in hollow  $\text{Au}_{0.3}\text{Ag}_{0.7}$  nanostructured dendrites treated with A549 lung cancer cells.

#### Gold–Silver Nanostructures

K.-W. Hu, C.-C. Huang, J.-R. Hwu, W.-C. Su, D.-B. Shieh, C.-S. Yeh\*

A New Photothermal Therapeutic Agent: Core-Free Nanostructured  $\text{Au}_x\text{Ag}_{1-x}$  Dendrites

*Chem. Eur. J.*  
DOI: 10.1002/chem.200800114



**Zinc green:** Easily available zinc salts are active and practical catalysts for the intermolecular hydroamination of terminal alkynes with anilines. The reactions proceed in the presence of  $\text{Zn}(\text{OTf})_2$  with

excellent regioselectivity (> 99%) and with high yields. Moreover, difficult functional groups such as nitro and cyano substituents are tolerated by the catalyst.

#### Homogeneous Catalysis

K. Alex, A. Tillack, N. Schwarz, M. Beller\*

General Zinc-Catalyzed Intermolecular Hydroamination of Terminal Alkynes

*ChemSusChem*  
DOI: 10.1002/cssc.200700160