

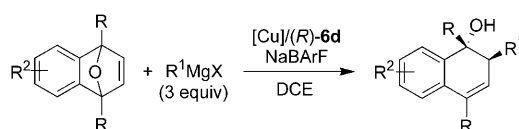
## Cross-coupling

W. Zhang, S.-F. Zhu, X.-C. Qiao,  
Q.-L. Zhou\*

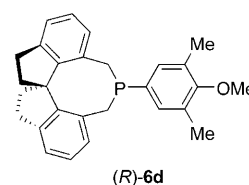
Highly Enantioselective Copper-Catalyzed  
Ring Opening of Oxabicyclic Alkenes  
with Grignard Reagents

*Chem. Asian J.*

DOI: 10.1002/asia.200800159



34 examples, 71–95% yields  
*trans/cis*: >99/1, *ee*: up to  
99.6%  
TON: up to 9000



**Open sesame:** A highly efficient enantioselective desymmetrization of oxabicyclic alkenes is established with chiral copper complexes of spiro phosphines as cata-

lysts. Excellent enantioselectivities (up to 99.6% *ee*) and high catalytic activity (TON = 9000) are obtained.

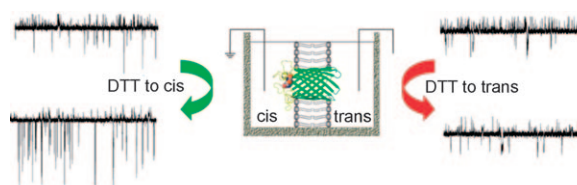
## Biosensors

M. Chen, Q.-H. Li, H. Bayley\*

Orientation of the Monomeric Porin  
OmpG in Planar Lipid Bilayers

*ChemBioChem*

DOI: 10.1002/cbic.200800444



**The orientations of single OmpG pores in planar lipid bilayers** were determined from the sidedness of the response of an extracellular disulfide bond to dithiothreitol (DTT). With this knowledge, the binding of a cyclodextrin adapter presented

to OmpG from the extracellular or periplasmic side was investigated. The information on the interaction between the cyclodextrin and OmpG serves to advance the use of OmpG as a biosensor.

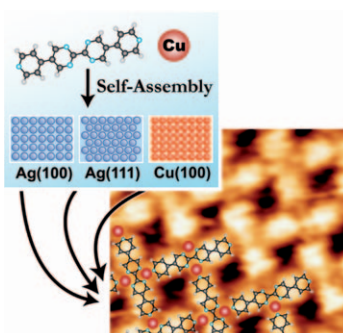
## Self-Assembly

S. L. Tait,\* A. Langner, N. Lin,  
R. Chandrasekar, O. Fuhr, M. Ruben,\*  
K. Kern

Assembling Isostructural Metal–Organic  
Coordination Architectures on Cu(100),  
Ag(100) and Ag(111) Substrates

*ChemPhysChem*

DOI: 10.1002/cphc.200800575



**Isostructural coordination architectures** in two dimensions on different substrates require sufficient metal–organic bonding strength to overcome templating effects from the surface. The network structure in this STM image was grown on Cu(100) and was also produced on Ag(111) and Ag(100) surfaces, due to robust three-fold N–Cu coordination interactions stabilizing the network.

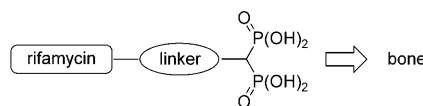
## Prodrugs

R. Reddy, E. Dietrich, Y. Lafontaine,  
T. J. Houghton, O. Belanger, A. Dubois,  
F. F. Arhin, I. Sarmiento, I. Fadhil,  
K. Laquerre, V. Ostiguy, D. Lehoux,  
G. Moeck, T. R. Parr, Jr., A. Rafai Far\*

Bisphosphonated Benzoxazinorifamycin  
Prodrugs for the Prevention and  
Treatment of Osteomyelitis

*ChemMedChem*

DOI: 10.1002/cmdc.200800255



**Benzoxazinorifamycins are potent anti-bacterial agents** currently in development. Tethering these antibiotics to a bisphosphonate functional group by a cleavable linker allows the delivery of these agents to osseous tissues, where they can be released over time to treat bone infections. Various linker strategies are presented herein to develop osteotropic prodrugs, the activities of which are examined *in vitro* and *in vivo*.

