



The following Communications have been judged by at least two referees to be “very important papers” and will be published online at www.angewandte.org soon:

Y. Fu, Q. Dai, W. Zhang, J. Ren, T. Pan,* C. He*

AlkB Domain of Mammalian ABH8 Catalyzes Hydroxylation of 5-Methoxycarbonylmethyluridine at the Wobble Position of tRNA

H. Zheng, J. Gao*

Highly Specific Heterodimerization Mediated by Quadrupole Interactions

M. Willis, M. Götz, A. K. Kandalam, G. F. Ganteför,* P. Jena*

Hyperhalogens: A New Class of Highly Electronegative Species

V. Mazumder, M. Chi, K. L. More, S. Sun*

Synthesis and Characterization of Multimetallic Pd/Au and Pd/Au/FePt Core/Shell Nanoparticles

S.-Y. Moon, J.-S. Bae, E. Jeon, J.-W. Park*

Organic Sol-Gel Synthesis: Solution-Processable Microporous Organic Networks

D. V. Esposito, S. T. Hunt, A. L. Stottlemeyer, K. D. Dobson, B. E. McCandless, R. W. Birkmire, J. G. Chen*

Low-Cost Hydrogen-Evolution Catalysts Based on Monolayer Platinum on Tungsten Monocarbide (WC) Substrates

K. Sasaki, H. Naohara, Y. Cai, Y. M. C. Liu, M. B. Vukmirovic, J. X. Wang, R. R. Adzic

Platinum-Monolayer-Protected Cores: High-Stability Electrocatalysts for Fuel-Cell Cathodes

R. Matsui, K. Seto, K. Fujita, T. Suzuki, A. Nakazaki, S. Kobayashi
Unusually E-Selective Ring-Closing Metathesis to Form Eight-Membered Rings

J. S. Chen, T. Zhu, C. M. Li, X. W. Lou*

Building Hematite Nanostructures Using Oriented Attachment

S. Sun, G. Zhang, D. Geng, Y. Chen, R. Li, M. Cai, X. Sun*

A New Highly Durable Platinum Nanocatalyst for PEM Fuel Cells: Multiarmed Starlike Nanowire Single Crystals

T. J. Hebden, A. J. S. John, D. G. Gusev, W. Kaminsky, K. I. Goldberg, D. M. Heinekey*

Preparation of a Dihydrogen Complex of Cobalt

Y. Zhang, G. M. Miyake, E. Y.-X. Chen*

Alane-Based Classical and Frustrated Lewis Pairs in Polymer Synthesis: Rapid Polymerization of Methyl Methacrylate and Naturally Renewable Methylene Butyrolactones to High-Molecular-Weight Polymers



Richard F. Heck



Ei-ichi Negishi



Akira Suzuki

News

Nobel Prizes 2010:

Richard F. Heck _____ 8300

Ei-ichi Negishi _____ 8300

Akira Suzuki _____ 8300



“The part of my job which I enjoy the most is working with motivated and talented students.

I am waiting for the day when someone will discover a cure for major diseases such as Alzheimer's disease and cancer ...”

This and more about Milko E. van der Boom can be found on page 8301.

Author Profile

Milko E. van der Boom _____ 8301

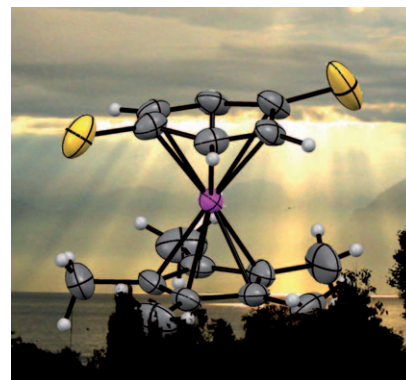
Highlights

Reactive Species

C. G. Hartinger* — 8304–8305

Trapping Unstable Benzoquinone Analogues by Coordination to a $[(\eta^5\text{-C}_5\text{Me}_5)\text{Ir}]$ Fragment and the Anticancer Activity of the Resulting Complexes

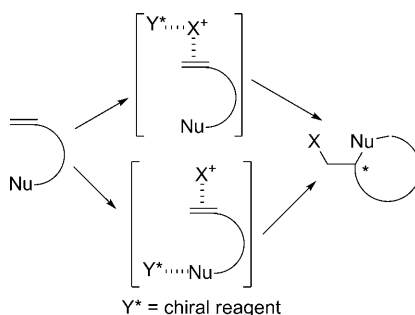
Gotcha! Unstable dithio- and diseleno-benzoquinones can be trapped by coordination to an $[(\eta^5\text{-C}_5\text{Me}_5)\text{Ir}]$ moiety (see structure; gray C, white H, pink Ir, yellow Se). In vitro investigations on the anticancer activity of the resulting complexes showed them to be comparable to *cis*-platin. These organometallic species could form the basis for new anticancer agents.



Asymmetric Synthesis

G. Chen, S. Ma* — 8306–8308

Enantioselective Halocyclization Reactions for the Synthesis of Chiral Cyclic Compounds



Ground-breaking progress has been made in the synthesis of chiral cyclic compounds from nonchiral unsaturated substrates with a nucleophilic functionality. A number of highly enantioselective electrophilic halocyclizations based on either the interaction of a chiral Lewis acid with an unsaturated substrate or the generation of a chiral electrophilic intermediate in situ from an electrophile and a chiral reagent (see scheme) were developed.

Essays

History in the Making

R. R. Ernst* — 8310–8315

Zurich's Contributions to 50 Years Development of Bruker

Humble beginnings: It is 50 years since the foundation of Bruker Physik AG in Karlsruhe. In this Essay, Prof. Richard R. Ernst looks at the early years of the company, and how researchers in Zurich contributed to its success in the field of NMR instrumentation.



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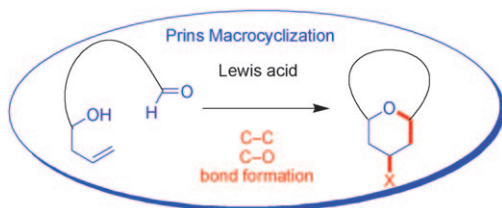
individuals who are personal members of a national chemical society prices are available on request. Postage and handling charges included. All prices are subject to local VAT/sales tax.

Minireviews

Macrocyclization Strategies

E. A. Crane, K. A. Scheidt* — 8316–8326

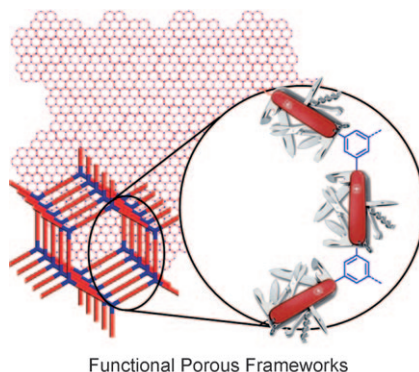
Prins-Type Macrocyclizations as an Efficient Ring-Closing Strategy in Natural Product Synthesis



A “Prins” among reactions: The first report of a Prins macrocyclization appeared in 1979, but only since late 2008 has this strategy gained momentum in natural product synthesis. Convergent,

selective, and high yielding, the Prins macrocyclization forms tetrahydropyran-containing macrocycles in a merged C–O and C–C bond-forming event (see scheme).

Pores for thought: This Review describes porous functional materials in which the organic groups act as a part of the pore wall. The focus is on functional mesoporous organosilicas as well as meso- and microporous polymers. The large number of functional groups in such porous materials allows, for example, a new approach to bridging the gap between homogeneous and heterogeneous catalysis.



Functional Porous Frameworks

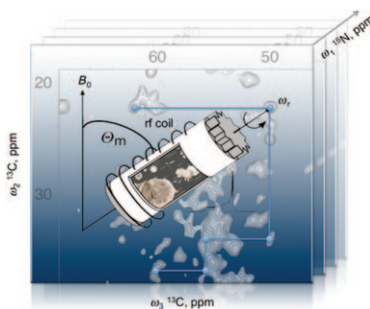
Reviews

Porous Materials

A. Thomas* — 8328–8344

Functional Materials: From Hard to Soft Porous Frameworks

The bigger they are, the harder they fall: Recent advancements in NMR spectroscopy, biophysics, and related research areas have resulted in solid-state NMR (ssNMR) spectroscopy having the potential to study complex biomolecular systems at atomic resolution in situ. Methodological challenges of ssNMR spectroscopy for studying complex biomolecules are described as well as recent and possible applications.



Spectroscopic Methods

M. Renault, A. Cukkemane, M. Baldus* — 8346–8357

Solid-State NMR Spectroscopy on Complex Biomolecules

Communications



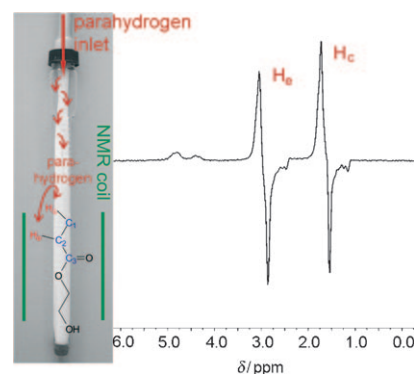
NMR Signal Enhancement

M. Roth, P. Kindervater, H.-P. Raich,
J. Bargon, H. W. Spiess,*
K. Münnemann* 8358–8362



Continuous ^1H and ^{13}C Signal
Enhancement in NMR Spectroscopy and
MRI Using Parahydrogen and Hollow-
Fiber Membranes

Parahydrogen-induced polarization is used for the continuous generation of hyperpolarized molecules. Constant ^1H and ^{13}C NMR signal enhancement is demonstrated using a continuous delivery of parahydrogen by hollow fiber membranes (see picture, left), which allows the acquisition of two-dimensional NMR signals requiring multiple acquisitions with constant initial polarization.



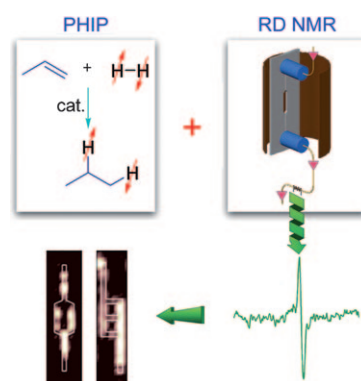
Imaging in Microfluidics

V.-V. Telkki,* V. V. Zhivonitko, S. Ahola,
K. V. Kovtunov, J. Jokisaari,
I. V. Koptiyug 8363–8366



Microfluidic Gas-Flow Imaging Utilizing
Parahydrogen-Induced Polarization and
Remote-Detection NMR

Sensitivity enhancement of several orders of magnitude provided by combining parahydrogen-induced polarization (PHIP) and remote-detection (RD) NMR techniques enables gas-flow visualization in microfluidic devices (see scheme). A 7700-fold reduction in the imaging time compared to the current state-of-the-art is achieved, and it can also be used to significantly improve the spatial resolution in such experiments.



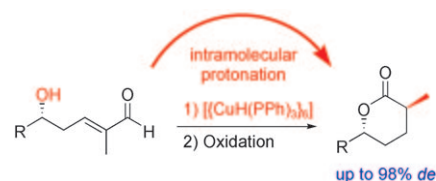
Stereoselective Protonation

A. Kena Diba, C. Noll, M. Richter,
M. T. Gieseler, M. Kalesse* 8367–8369



Intramolecular Stereoselective
Protonation of Aldehyde-Derived Enolates

Picking sides: Asymmetric protonation of the titled compounds poses a most significant challenge and has been addressed by taking advantage of internal protonation and subsequent hemiacetal formation to avoid epimerization (see scheme). The substrates employed in these transformations can be easily accessed through a sequence of vinylogous aldol reactions with subsequent conjugate reductions.



Enantioselective Catalysis

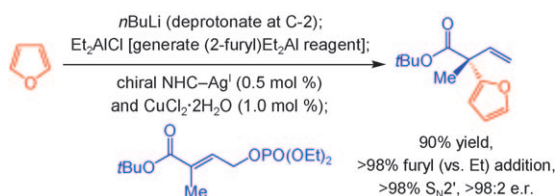
F. Gao, Y. Lee, K. Mandai,
A. H. Hoveyda* 8370–8374



Quaternary Carbon Stereogenic Centers
through Copper-Catalyzed Enantio-
selective Allylic Substitutions with Readily
Accessible Aryl- or Heteroarylolithium
Reagents and Aluminum Chlorides

The case of the notorious aryls is solved: The first efficient catalytic and enantioselective method for allylic substitutions that furnish quaternary carbon stereogenic centers by additions of aryl- or

heteroarylmetals is reported (see scheme). Highly site- and enantioselective processes begin with readily available organolithium reagents.





Celebrating 50 Years of Innovation

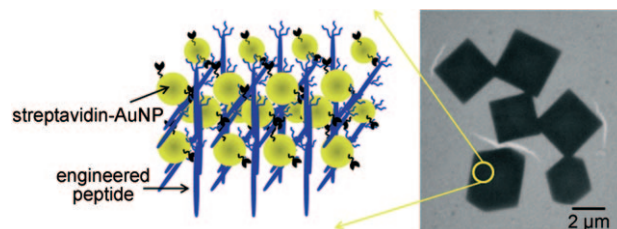
Celebrating 50 years of innovative excellence, technical leadership, integrity and long term commitment to quality. With thanks to our customers for their numerous collaborations that set the ground for the inventive approaches, that has enabled us to deliver the most innovative solutions. We look forward to continuing cooperation in an ever-challenging future.

think forward

Analytical Solutions

Bionanotechnology

P. Kaur, Y. Maeda, A. C. Mutter,
T. Matsunaga, Y. Xu,
H. Matsui* — 8375–8378



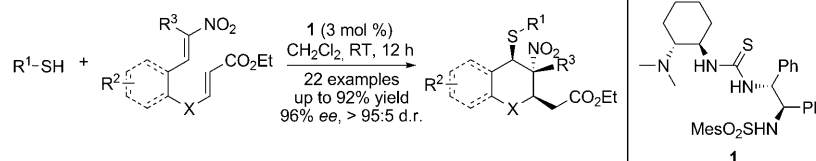
Three-Dimensional Directed Self-Assembly of Peptide Nanowires into Micrometer-Sized Crystalline Cubes with Nanoparticle Joints

Micro construction sites: Engineered peptides and ligand-functionalized gold nanoparticles have been assembled into micrometer-scale 3D cube-shaped crystals (see picture), creating a physical

framework for a biomimetic assembly strategy. The NPs join the peptide nanowires through streptavidin–biotin interactions to create cubic unit cells.

Asymmetric Synthesis

X.-F. Wang, Q.-L. Hua, Y. Cheng, X.-L. An,
Q.-Q. Yang, J.-R. Chen,*
W.-J. Xiao* — 8379–8383



Organocatalytic Asymmetric Sulfamichael/Michael Addition Reactions: A Strategy for the Synthesis of Highly Substituted Chromans with a Quaternary Stereocenter

Simply complex: Diverse and structurally complex chroman derivatives with a quaternary stereocenter have been obtained through the titled reaction of thiols with nitroolefin enoates using the bifunctional

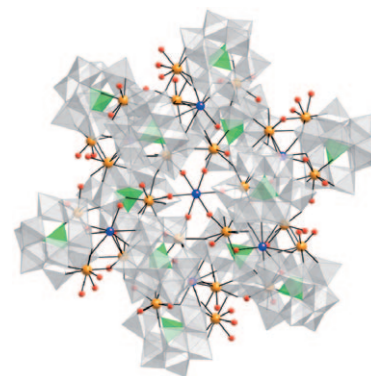
catalyst **1**. The reaction features simple experimental procedures, high yield, enantiomeric excess, and excellent diastereoselectivity.

Polyoxometalates

S. Reinoso,* M. Giménez-Marqués,
J. R. Galán-Mascarós, P. Vitoria,
J. M. Gutiérrez-Zorrilla — 8384–8388

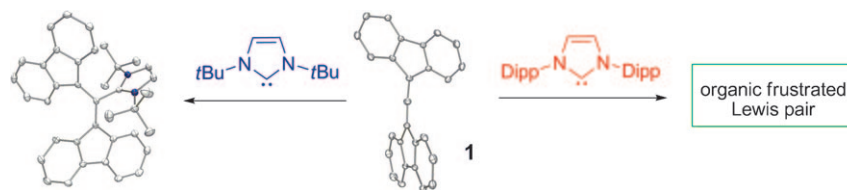
Giant Crown-Shaped Polytungstate Formed by Self-Assembly of Ce^{III}-Stabilized Dilacunary Keggin Fragments

The largest tungstogermanate and third-largest polytungstate (see picture) was obtained by K⁺-directed self-assembly of three unprecedented types of [GeW₁₀O₃₈]¹²⁻ fragments stabilized by coordination of Ce ions on the vacancies. In its Ce₆O₄₂ central ring, which displays antiferromagnetic interactions, a K⁺ cation is captured by internal water molecules. Gray octahedra WO₆, green tetrahedra GeO₄; Ce orange, K purple, O red.



Organic Frustrated Lewis Pairs

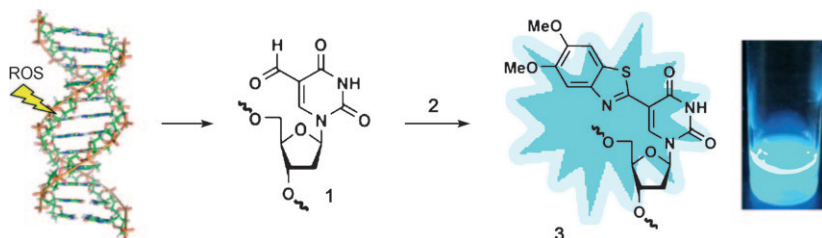
B. Inés, S. Holle, R. Goddard,
M. Alcarazo* — 8389–8391



Heterolytic S–S Bond Cleavage by a Purely Carbogenic Frustrated Lewis Pair

Boranes were getting borin': Owing to the Lewis acidic character of the central carbon atom, a bisfluorenyl-substituted allene **1** could be used instead of B(C₆F₅)₃ for the generation of frustrated Lewis

pairs. Mixtures of the allene and the bulky N-heterocyclic carbene shown in red (Dipp = diisopropylphenyl) caused heterolytic cleavage of the S–S bond in disulfides.



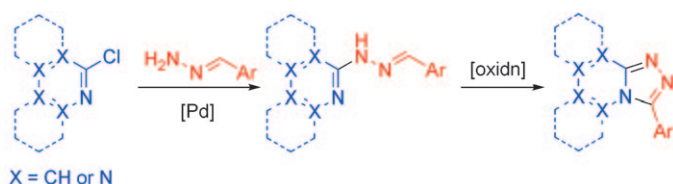
Chemoselectivity in DNA: The thymidine lesion 5-formyl-2'-deoxyuridine (**1**) induces the mutation of DNA. Its derivatization with the specific fluorogenic reagent 2-amino-4,5-dimethoxythiophenol (**2**)

gives **3**, which shows strong fluorescence. This fast method for the detection of **1** requires no enzymatic digestion, HPLC separation, or MS analysis. ROS = reactive oxygen species.

DNA-Damage Detection

W. Hirose, K. Sato,
A. Matsuda* 8392–8394

Selective Detection of 5-Formyl-2'-deoxyuridine, an Oxidative Lesion of Thymidine, in DNA by a Fluorogenic Reagent



The palladium-catalyzed intermolecular coupling of aldehyde-derived hydrazones with chloroazines, followed by oxidative cyclization under mild conditions afforded

access to a broad variety of bicyclic heterocyclic scaffolds (see scheme) that have potential for use in drug discovery.

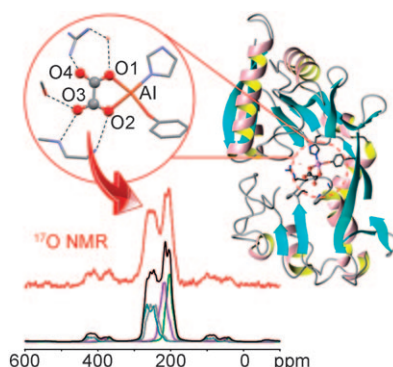
Synthetic Methods

O. R. Thiel,* M. M. Achmatowicz,*
A. Reichelt, R. D. Larsen 8395–8398

Palladium-Catalyzed Coupling of Aldehyde Hydrazones: Practical Synthesis of Triazolopyridines and Related Heterocycles



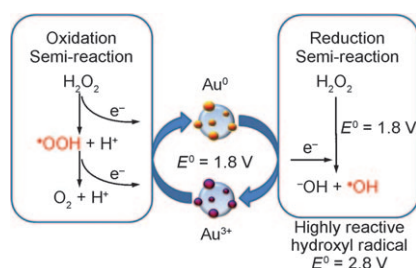
Oxygen, oxygen, everywhere! Poor sensitivity has hindered the development of solid-state ^{17}O NMR spectroscopy as a practical technique for the structural elucidation of protein complexes. However, this has now changed and it has been demonstrated that multinuclear ^{17}O , ^{27}Al , ^{13}C NMR parameters can be used to aid structural refinement for a protein-bound ligand molecule (see picture).



NMR Spectroscopy

J. Zhu, E. Ye, V. Tersikh,
G. Wu* 8399–8402

Solid-State ^{17}O NMR Spectroscopy of Large Protein–Ligand Complexes



Nanojewels as catalysts: Degradation of phenol with a small excess of H_2O_2 can be achieved by using gold nanoparticles supported on Fenton-treated diamond nanoparticles as a highly selective (at least 79%) and efficient (turnover number: 321 000) catalyst for the Fenton reaction at room temperature. The proposed mechanism is shown schematically in the picture.

Catalytic Fenton Reaction

S. Navalon, R. Martin, M. Alvaro,
H. Garcia* 8403–8407

Gold on Diamond Nanoparticles as a Highly Efficient Fenton Catalyst



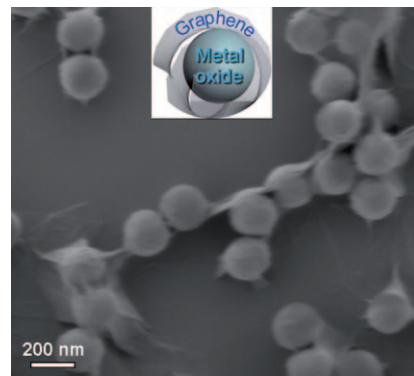
Electrochemistry

S. Yang, X. Feng,* S. Ivanovici,
K. Müllen* 8408–8411



Fabrication of Graphene-Encapsulated
Oxide Nanoparticles: Towards High-
Performance Anode Materials for Lithium
Storage

Storage shells: The assembly of negatively charged graphene oxide and positively charged oxide nanoparticles by electrostatic interactions, and subsequent chemical reduction, leads to metal oxides encapsulated in flexible and ultrathin graphene shells (see picture). These electrochemically active nanoparticles show a remarkable lithium storage capacity, with an excellent cycle performance.

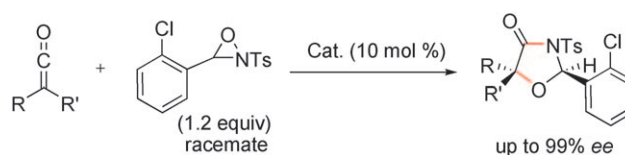


Asymmetric Catalysis

P.-L. Shao, X.-Y. Chen,
S. Ye* 8412–8416



Formal [3+2] Cycloaddition of Ketenes
and Oxaziridines Catalyzed by Chiral
Lewis Bases: Enantioselective Synthesis
of Oxazolin-4-ones



Choose the right cat.: A highly enantioselective synthesis of oxazolin-4-ones by the formal [3+2] cycloaddition of ketenes and a racemic oxaziridines has been

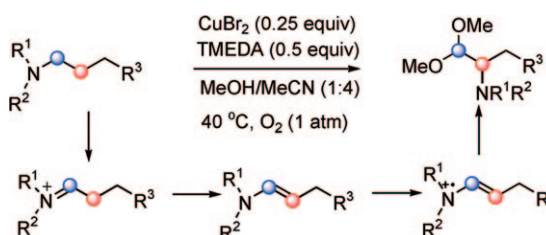
developed (see scheme; cat. = N-heterocyclic carbenes for disubstituted ketenes or cinchona alkaloids for monosubstituted ketenes, Ts = 4-toluenesulfonyl).

Oxidative Rearrangement

J.-S. Tian, T.-P. Loh* 8417–8420



Copper-Catalyzed Rearrangement of
Tertiary Amines through Oxidation of
Aliphatic C–H Bonds in Air or Oxygen:
Direct Synthesis of α -Amino Acetals



A surprising turn of events: Mechanistic studies, including trapping, control, and isotope-labeling experiments, led to the proposal of a rearrangement mechanism

involving oxidation of aliphatic C–H bonds (see scheme; TMEDA = tetraethylethylenediamine).

Isopeptide Bonds

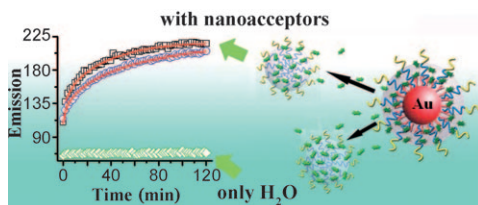
R. M. Hagan, R. Björnsson,
S. A. McMahon, B. Schomburg,
V. Braithwaite, M. Bühl, J. H. Naismith,
U. Schwarz-Linek* 8421–8425



NMR Spectroscopic and Theoretical
Analysis of a Spontaneously Formed
Lys–Asp Isopeptide Bond



One bond makes all the difference: Three suitably positioned amino acid side chains (see picture) and a hydrophobic environment are all that is required for an amidation reaction with remarkable consequences. An emerging central building block of bacterial surface proteins owes its stability to a spontaneously formed isopeptide bond. The impact of this bond on protein structure and dynamics and the mechanism of its formation are scrutinized in detail.



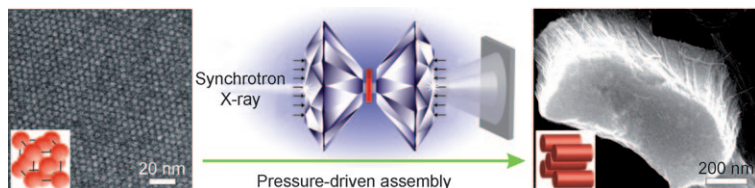
Quick release: Polymer-coated gold nanoparticles (AuNPs) are used as nano-carriers and nanoscale acceptors are used to represent proteins and lipid membranes in a model drug-delivery system.

The fluorescence of a drug is quenched in the vicinity of the AuNPs but reemerges upon its release (see picture). The critical role of nanoacceptors in facilitating drug release is demonstrated.

Drug Delivery

H. Wang, J. Xu, J. Wang, T. Chen, Y. Wang, Y. W. Tan, H. Su, K. L. Chan, H. Chen* — 8426–8430

Probing the Kinetics of Short-Distance Drug Release from Nanocarriers to Nanoacceptors



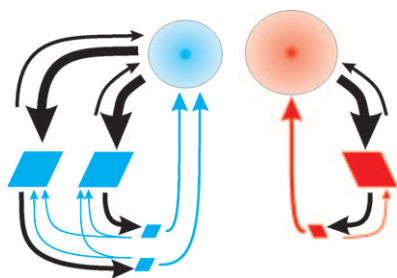
Forced into line: External pressure was used to engineer nanoparticle assembly. Reversible manipulation of the unit-cell dimensions of a 3D ordered nanoparticle array under a hydrostatic pressure field enabled the fine-tuning of the interparticle

distance. Under a uniaxial pressure field, nanoparticles were forced to contact and coalesce into nanorods or nanowires and ordered ultrahigh-density arrays (see picture; small arrows denote pressure).

High-Pressure Chemistry

H. Wu, F. Bai, Z. Sun, R. E. Haddad, D. M. Boye, Z. Wang, H. Fan* — 8431–8434

Pressure-Driven Assembly of Spherical Nanoparticles and Formation of 1D-Nanostructure Arrays

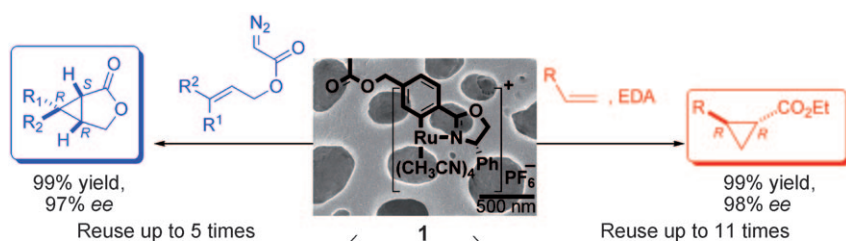


Change of heart: A solution-phase enantiomeric excess that has the handedness of the minor population of the solid phase is observed during grinding of a slurry of racemic conglomerate crystals. This excess is the driving force for a net flux of molecules from crystals of the minor handedness to crystals of the major handedness, thus explaining the complete deracemization of the solid phase (see picture: blue: *S* form, red: *R* form).

Chiral Resolution

W. L. Noorduin, W. J. P. van Enckevort, H. Meekes, B. Kaptein, R. M. Kellogg, J. C. Tully, J. M. McBride, E. Vlieg* — 8435–8438

The Driving Mechanism Behind Attrition-Enhanced Deracemization



Along came poly: A novel macroporous polymer-supported chiral Ru^{II}/pheox catalyst (1, see SEM image) afforded excellent reactivity and enantioselectivity in inter- and intramolecular cyclopropana-

tion reactions with a broad range of substrates. The catalyst showed no leaching and could be reused up to 11 times, even after storage.

Macroporous Catalysts

A. Abu-Elfotoh, K. Phomkeona, K. Shibatomi, S. Iwasa* — 8439–8443

Asymmetric Inter- and Intramolecular Cyclopropanation Reactions Catalyzed by a Reusable Macroporous-Polymer-Supported Chiral Ruthenium(II)/Phenylloxazoline Complex



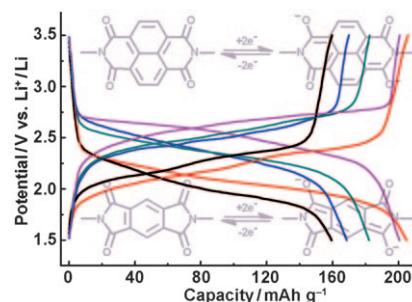
Lithium Batteries

Z. Song, H. Zhan,* Y. Zhou . 8444–8448



Polyimides: Promising Energy-Storage Materials

Plastic batteries: Polyimides are proposed as cathode materials for rechargeable lithium batteries. Although they are regarded as insulators, five polyimides with different structures all show good electrochemical activity and some of them show promising performance, which could allow their use in a new generation of “green battery” applications.



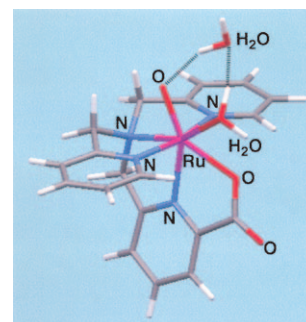
Oxo Complexes

T. Kojima,* Y. Hirai, T. Ishizuka, Y. Shiota, K. Yoshizawa, K. Ikemura, T. Ogura, S. Fukuzumi* 8449–8453



A Low-Spin Ruthenium(IV)–Oxo Complex: Does the Spin State Have an Impact on the Reactivity?

Spin doesn't matter: A ruthenium(II)–aqua complex bearing a pentadentate pyridylamine with a carboxylate group as a ligand affords a seven-coordinate low-spin ($S=0$) ruthenium(IV)–oxo complex (see structure) by oxidation through proton-coupled electron transfer. Comparison of the reactivity of the low-spin and an intermediate-spin ($S=1$) Ru^{IV} –oxo complexes revealed that the spin state does not affect the reactivity of catalytic oxidation of organic compounds.

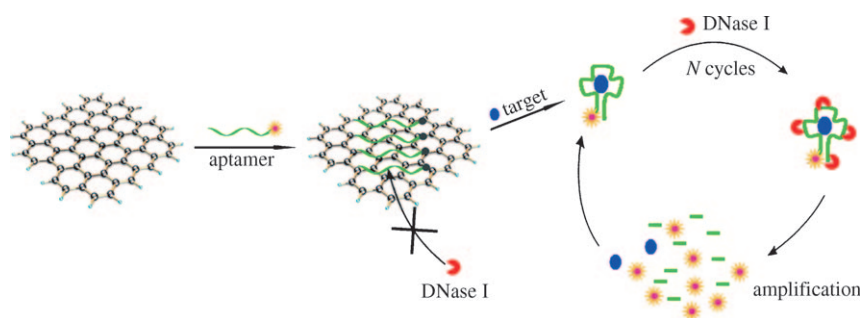


Biological Assays

C.-H. Lu, J. Li, M.-H. Lin, Y.-W. Wang, H.-H. Yang,* X. Chen, G.-N. Chen 8454–8457

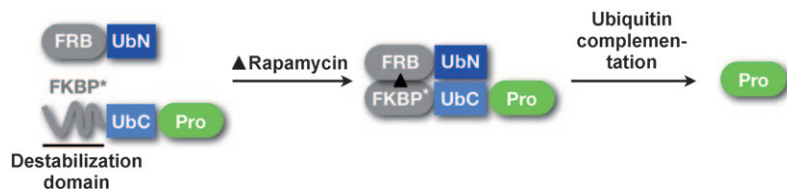


Amplified Aptamer-Based Assay through Catalytic Recycling of the Analyte



Turn up the volume: In a simple and highly sensitive aptamer-based assay that relies on the ability of nanomaterials to protect DNA from nuclease cleavage, the aptamer is released from the nanomaterial when challenged with a target. The

nuclease then cleaves the free aptamer and releases the target, which can initiate a new cycle. Recycling of the target in this way leads to significant signal amplification (see diagram).



Rapamycin to the rescue: Native protein concentration was precisely controlled in living cells by using the small molecule rapamycin. The addition of rapamycin “shielded” a destabilization domain from degradation and induced complementa-

tion of split ubiquitin (UbN and UbC); as a result, proteins (Pro) of interest were released. This technology enables the investigation of native protein biology over a broad concentration range.

Protein Biology

H. D. Lau, J. Yaegashi, B. W. Zaro, M. R. Pratt* 8458–8461

Precise Control of Protein Concentration in Living Cells



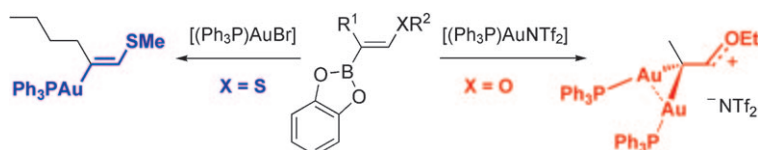
When structural flexibility is a plus: A link between structural flexibility of biomimetic antimicrobials and their ability to penetrate into the hydrophobic core and disrupt the integrity of bacterial lipid model membranes has been established

using liquid surface X-ray scattering techniques. Results indicate that the modes of interaction of flexible and conformationally restrained antimicrobials with the bacterial membranes are different (see picture).

Antimicrobial Peptides

A. Ivankin, L. Livne, A. Mor, G. A. Caputo, W. F. DeGrado, M. Meron, B. Lin, D. Gidalevitz* 8462–8465

Role of the Conformational Rigidity in the Design of Biomimetic Antimicrobial Compounds



Disturbing neighbors: Alkenylgold species with a heteroatom substituent are thought to be key intermediates in gold-catalyzed *trans* additions of protic nucleophiles to alkynes. One reason for the scarcity of such compounds lies in the non-inno-

cence of the neighboring heteroatom, which may enforce the uptake of a second gold fragment with formation of surprisingly robust species with a *gem*-digold unit adjacent to a largely cationic center (see scheme).

Homogeneous Catalysis

G. Seidel, C. W. Lehmann, A. Fürstner* 8466–8470

Elementary Steps in Gold Catalysis: The Significance of *gem*-Diauration



Donuts anyone? Molecules based on a *meta*-linked aromatic segment self-assemble into hexameric macrocycles that, in turn, stack on top of each other to

from elongated tubular helical suprastructures. The formed tubules dissociate into discrete stacks of toroids in response to the addition of silver salt (see picture).

Self-Assembly

H.-J. Kim, S.-K. Kang, Y.-K. Lee, C. Seok, J.-K. Lee, W.-C. Zin, M. Lee* 8471–8475

Self-Dissociating Tubules from Helical Stacking of Noncovalent Macrocyces

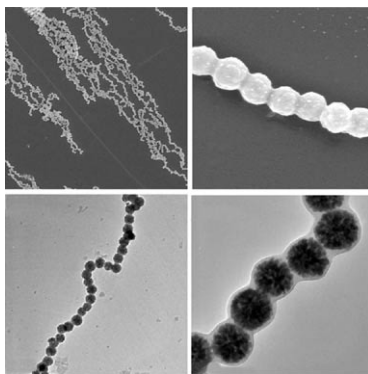


Imaging Agents

J. Zhou, L. Meng,* X. Feng, X. Zhang,
Q. Lu* 8476–8479



One-Pot Synthesis of Highly Magnetically Sensitive Nanochains Coated with a Highly Cross-Linked and Biocompatible Polymer



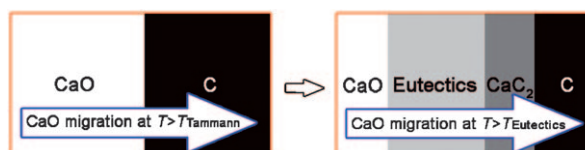
Neptune's necklace: Fe_3O_4 colloidal nanocrystal clusters (ca. 200 nm in diameter) were organized into one-dimensional nanochains and coated with a highly cross-linked polymer in a facile one-pot synthesis (see picture). The nanochains have enhanced hydrogen T_2 relaxation rates, facile magnetic manipulation, good water dispersivity, and excellent biocompatibility.

Biomass

G. Li, Q. Liu, Z. Liu,* Z. C. Zhang,* C. Li,
W. Wu 8480–8483



Production of Calcium Carbide from Fine Biochars



Rising from the ashes: Fine chars produced from biomass can be used as a feedstock for the production of CaC_2 in an autothermal process that operates at a

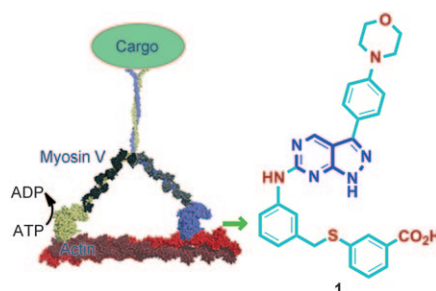
temperature 500°C lower than the traditional electric arc process and with greatly reduced reaction time (see scheme for the reaction of CaO and C forming CaC_2).

Enzyme Inhibition

K. Islam, H. F. Chin, A. O. Olivares,
L. P. Saunders, E. M. De La Cruz,*
T. M. Kapoor* 8484–8488



A Myosin V Inhibitor Based on Privileged Chemical Scaffolds



Switching the motor off: Privileged chemical scaffolds were used as a starting point for the development of a specific chemical inhibitor **1** of myosin V, a key motor protein required for intracellular transport (see picture; ADP=adenosine diphosphate, ATP=adenosine triphosphate). The potency of **1**, which does not compete directly with nucleotide binding, is comparable to that of other known motor-protein inhibitors used as probes in chemical biology.

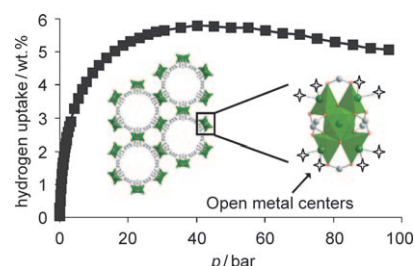
Porous Materials

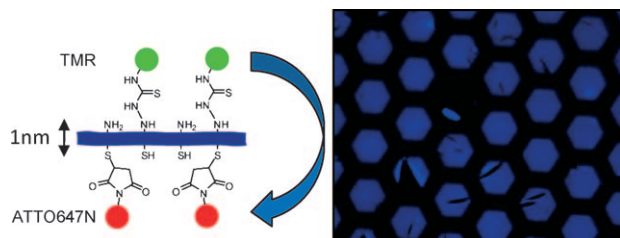
K. Gedrich, I. Senkovska, N. Klein,
U. Stoeck, A. Henschel, M. R. Lohe,
I. A. Baburin, U. Mueller,
S. Kaskel* 8489–8492



A Highly Porous Metal–Organic Framework with Open Nickel Sites

A healthy appetite: A new metal–organic framework (DUT-9; see picture) has an unprecedented topology that contains $\{\text{Ni}_5(\mu_3\text{-O})_2(\text{CO}_2)_6\}$ clusters (green polyhedra; C gray, O red). DUT-9 has very high storage capacities for methane, hydrogen, and carbon dioxide because of the presence of open metal sites (marked by stars).





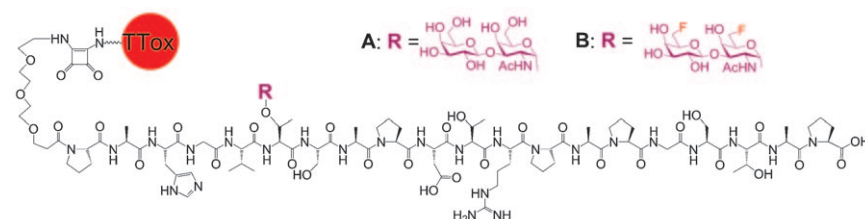
Buttered side up: A one nanometer thick membrane with different functional groups on each side was fabricated from self-assembled monolayers of aromatic molecules. The amino side of this Janus membrane was modified with tetrame-

thylrhodamine (TMR) fluorescent dye, and the thiol side with ATTO647N (see picture). The functionalization and functionality of the nanomembrane were demonstrated by X-ray photoelectron spectroscopy and fluorescence microscopy.

Nanosheets

Z. Zheng, C. T. Nottbohm, A. Turchanin, H. Muzik, A. Beyer, M. Heilemann, M. Sauer, A. Götzhäuser* — 8493–8497

Janus Nanomembranes: A Generic Platform for Chemistry in Two Dimensions



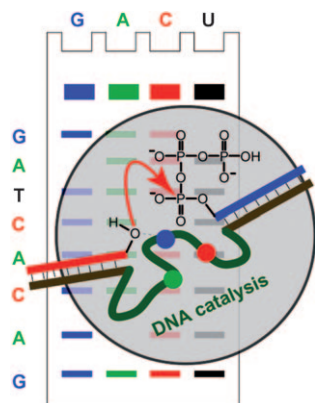
A “shot in the arm” for cancer therapy: Coupling of synthetic glycopeptide tandem-repeat sequences of the epithelial mucin MUC1 with the Thomsen–Friedenreich (T) antigen (A) or a difluoro analogue (B) to tetanus toxoid (TTTox) affords synthetic vaccines, which induce very

strong immune responses in mice overriding the natural tolerance of the immune system. The induced antibodies are selectively directed against the tumor-associated MUC1 structures and strongly bind to breast cancer cells of the MCF-7 cell line.

Antitumor Vaccines

A. Hoffmann-Röder,* A. Kaiser, S. Wagner, N. Gaidzik, D. Kowalczyk, U. Westerlind, B. Gerlitzki, E. Schmitt, H. Kunz* — 8498–8503

Synthetic Antitumor Vaccines from Tetanus Toxoid Conjugates of MUC1 Glycopeptides with the Thomsen–Friedenreich Antigen and a Fluorine-Substituted Analogue

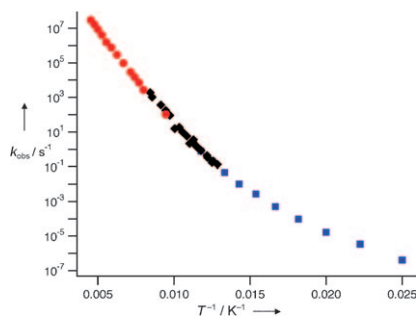


Complete analysis! Combinatorial mutation interference analysis (CoMA) is a highly efficient method to identify catalytically essential nucleotides in deoxyribozymes by the simultaneous assessment of all possible mutations in the active site of the catalyst. The application of CoMA for two different deoxyribozymes revealed indispensable guanosine nucleotides for DNA-catalyzed RNA ligation.

DNA Enzymes

F. Wachowius, F. Javadi-Zarnaghi, C. Höbartner* — 8504–8508

Combinatorial Mutation Interference Analysis Reveals Functional Nucleotides Required for DNA Catalysis



Relax! In the dilute mixed-crystal system $[\text{Zn}_{1-x}\text{Fe}_x(\text{bbtr})_3](\text{ClO}_4)_2$, $x = 2\%$ (bbtr = 1,4-di(1,2,3-triazol-1-yl)butane), the iron(II) centers are predominantly in the high-spin state. The low-spin state can be populated as a metastable state by irradiation with near-IR light; the rate constant of the low-spin \rightarrow high-spin relaxation spans 14 orders of magnitude between 40 and 220 K (see picture).

Spin Crossover

I. Krivokapic, P. Chakraborty, R. Bronisz, C. Enachescu, A. Hauser* — 8509–8512

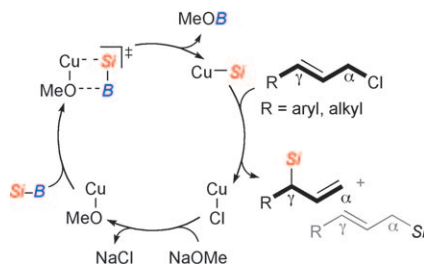
Significant Variation of the Singlet–Quintet Intersystem Crossing Rate Constant in an Iron(II) High-Spin Complex as a Function of Temperature

Allylic Substitution

D. J. Vyas, M. Oestreich* — 8513–8515



Copper-Catalyzed Si–B Bond Activation in Branched-Selective Allylic Substitution of Linear Allylic Chlorides



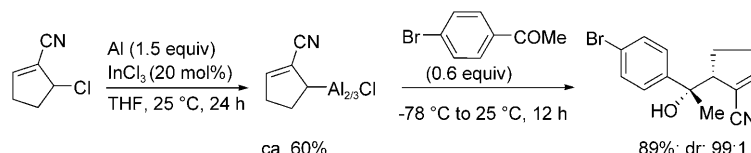
The harder they come, the better they leave! Chloride is superior to all common leaving groups in the γ -selective allylic substitution of linear precursors. This approach involves a novel copper-catalyzed Si–B bond activation (see scheme; $\gamma/\alpha \geq 98:2$, 7 examples; Si = SiMe₂Ph and B = Bpin with pin = pinacolato).

Aluminum Reagents

Z. Peng, T. D. Blümke, P. Mayer, P. Knochel* — 8516–8519



Diastereoselective Synthesis of Homoallylic Alcohols with Adjacent Tertiary and Quaternary Centers by Using Functionalized Allylic Aluminum Reagents



Al be there for you: Sensitive functional groups, including ester and cyano groups, can be present in allylic aluminum reagents prepared by aluminum insertion in the presence of InCl₃. These aluminum organometallic compounds undergo

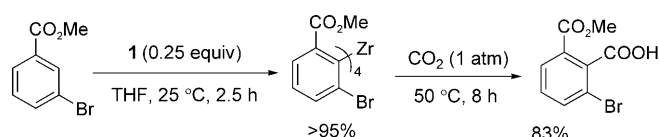
addition reactions to various functionalized aldehydes or ketones with remarkable diastereoselectivities allowing the construction of two adjacent quaternary and tertiary centers (see scheme).

Directed Zirconation

M. Jeganmohan, P. Knochel* — 8520–8524



tmp₄Zr: An Atom-Economical Base for the Metalation of Functionalized Arenes and Heteroarenes



Zirconium bases reach lots of places: Only 25% of zirconium metal is required for the metalation of various functionalized aromatic compounds and heterocycles. All four 2,2,6,6-tetramethylpiperidyl (tmp) groups of the new base

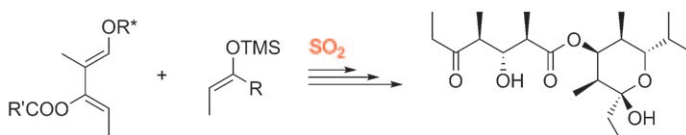
tmp₄Zr·4 MgCl₂·6 LiCl (**1**) are used up in the metalation process. The resulting zirconium species smoothly undergo palladium-catalyzed cross-couplings and addition to CO₂ and epoxides (see scheme).

Natural Product Synthesis

S. Laclef, M. Turks, P. Vogel* — 8525–8527



Total Synthesis and Determination of the Absolute Configuration of (–)-Dolabriferol



A reaction cascade combining sulfur dioxide with a 1-oxy-1,3-diene and (E)-silyl enol ether formed the basis of a short total synthesis of (–)-dolabriferol (see scheme; TMS = trimethylsilyl). The absolute con-

figuration of this natural product, which was first extracted from a gastropod mollusc in 1996, was established unequivocally.



Supporting information is available on www.angewandte.org (see article for access details).



A video clip is available as Supporting Information on www.angewandte.org (see article for access details).

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Corrigendum

Throughout this Communication (DOI: 10.1002/anie.200907175), the "pyrene" dye molecule should be correctly referred to as "perylene". The authors apologize for this mistake.

Inversion of Particle-Stabilized Emulsions to Form High-Internal-Phase Emulsions

G. Sun, Z. Li, T. Ngai* _____ 2163–2166

Angew. Chem. Int. Ed. **2010**, 49

DOI 10.1002/anie.200907175

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The Faculty of Chemistry wishes to appoint a

Professor (W3) of Functional Supramolecular Chemistry

to commence as soon as possible.

The professor is expected to perform research and teaching in inorganic chemistry and suitably complement the research interests of the Institute of Inorganic Chemistry. Applicants should be recognised for exceptional achievements in a modern area of inorganic chemistry, preferably in the area of „Functional Supramolecular Chemistry“. Participation in existing and planned research collaborations of the faculty (such as the PhD program CaSuS or the SFB 803) would be desirable. The teaching duties include inorganic chemistry courses for chemistry majors and other students.

The requirements for an appointment as a professor are specified by § 25 of the Niedersächsischen Hochschulgesetzes of February 26th, 2007 (Nds. GVBl. 2007 S. 69). The Stiftungsuniversität Göttingen has the right to make this appointment. Further details are available on request.

Applications from scientists working abroad are particularly welcome. A part time appointment is in principle possible. Appropriately qualified severely handicapped applicants will be given preference. The University would like to increase the percentage of women in areas in which they are underrepresented and thus encourages suitably qualified women to apply.

Online applications with CV, publications list and details of teaching and research experience should be received not later than **December 12th, 2010**. Applications by regular mail are accepted.

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