

Spotlight on Angewandte's Sister Journals

12396 – 12399



"If I were not a scientist, I would be a fisherman or a farmer."

What I look for first in a publication is its graphical abstract ..."

This and more about Shu-Hong Yu can be found on page 12402.

Service

Author Profile

Shu-Hong Yu _____ 12402



G. Mugesh

G. J. Sanjayan

I. S. Lee

D.-H. Min

C. R. Bertozzi

News

Shanti Swarup Bhatnagar Prize:
G. Mugesh and G. J. Sanjayan _____ 12403

KCS–Wiley Young Chemist Award:
I. S. Lee and D.-H. Min _____ 12403

Heinrich Wieland Prize:
C. R. Bertozzi _____ 12403



Heinz A. Staab, who was for many years Director at the Max Planck Institute (MPI) for Medical Research in Heidelberg, and previously President of the Max Planck Society (MPG) and the Gesellschaft Deutscher Chemiker (GDCh, German Chemical Society), passed away on July 29, 2012 at the age of 86 in Berlin after a long illness.

Obituaries

Heinz A. Staab (1926–2012)

M. W. Haenel* _____ 12404 – 12405

The Pauson–Khand Reaction

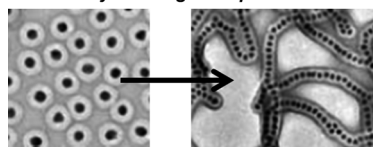
Ramon Rios Torres

Books

reviewed by N. Kann _____ 12406

No, it's not frogspawn! Polymer-coated gold nanoparticles can be assembled into extended mesoscopic chains with precise dimensional control. Here, the conditions can be adjusted to promote the fusion of polymeric ligands into cylindrical micellar aggregates. This type of colloidal polymerization offers a new and versatile route to a variety of mesoscopic assemblies of nanoparticles.

Polymerizing Nanoparticles



Highlights

Nanoparticle Chains

J. Pyun* _____ 12408 – 12409

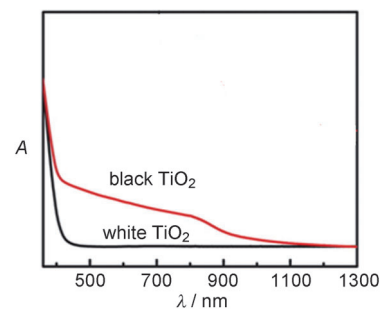
Self-Assembly and Colloidal
Polymerization of Polymer–Nanoparticle
Hybrids into Mesoscopic Chains

Photocatalysis

Y. H. Hu* ————— **12410–12412**

A Highly Efficient Photocatalyst—
Hydrogenated Black TiO₂ for the
Photocatalytic Splitting of Water

In black and white: The hydrogenation of TiO₂ can extend its optical absorption into the visible and infrared region and change its color from white to black. Furthermore, the hydrogenated black TiO₂ exhibits excellent photocatalytic activity for the splitting of water to yield H₂.

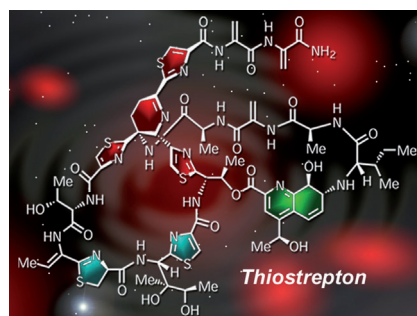


Reviews

Total Synthesis

K. C. Nicolaou* ————— **12414–12436**

How Thiostrepton Was Made in the
Laboratory



An adventurous undertaking: The synthetic conquest of thiostrepton was achieved in 2004. In this vivid account the author describes the laboratory odyssey with its many intriguing twists and turns that led to this memorable total synthesis.

Communications

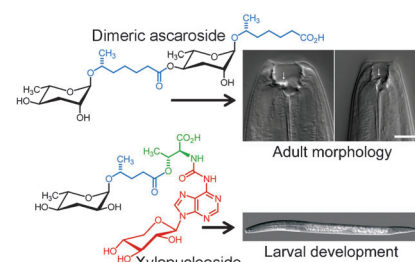
Metabolomics

N. Bose, A. Ogawa, S. H. von Reuss,
J. J. Yim, E. J. Ragsdale, R. J. Sommer,*
F. C. Schroeder* ————— **12438–12443**



Complex Small-Molecule Architectures
Regulate Phenotypic Plasticity in
a Nematode

Chemistry the worm's way: The nematode *Pristionchus pacificus* constructs elaborate small molecules from modified building blocks of primary metabolism, including an unusual xylopyranose-based nucleoside (see scheme). These compounds act as signaling molecules to control adult phenotypic plasticity and dauer development and provide examples of modular generation of structural diversity in metazoans.

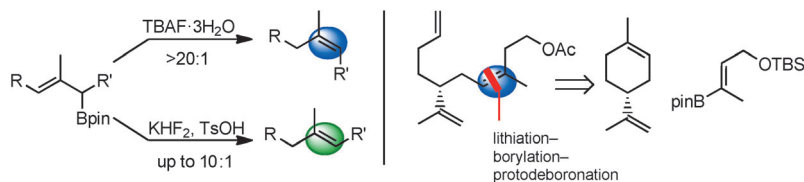


Frontispiece

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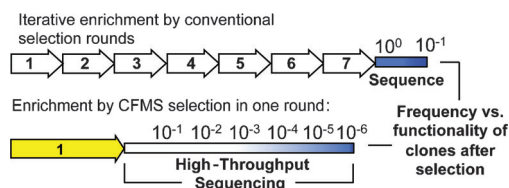
E-allylic boronic esters undergo a highly diastereoselective protodeboronation with TBAF·3 H₂O to give Z-trisubstituted alkenes. The selectivity can be switched to give predominantly the E-alkene instead

by using KHF₂/TsOH (see scheme). The utility of the methodology has been illustrated in a short synthesis of a component of the sex pheromone of the Californian red scale beetle.

E/Z-Alkene Synthesis

M. J. Hesse, C. P. Butts, C. L. Willis,*
V. K. Aggarwal* — 12444 – 12448

Diastereodivergent Synthesis of Trisubstituted Alkenes through Protodeboronation of Allylic Boronic Esters: Application to the Synthesis of the Californian Red Scale Beetle Pheromone



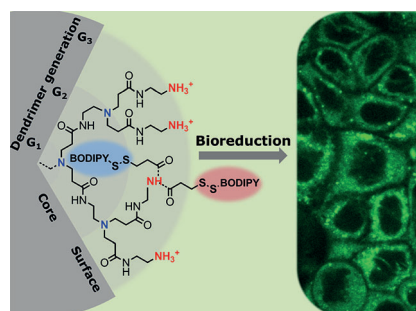
In a single round: By combining the high-efficiency enrichment through the continuous-flow magnetic separation (CFMS) technique with the analytical power of next-generation sequencing, the generation of antibody mimetics with a single

round of mRNA display is made possible. This approach eliminates iterative selection cycles and provides a path to fully automated ligand generation (see picture).

Ligand Design

C. A. Olson, J. Nie, J. Diep, I. Al-Shyoukh,
T. T. Takahashi, L. Q. Al-Mawsawi,
J. M. Bolin, A. L. Elwell, S. Swanson,
R. Stewart, J. A. Thomson, H. T. Soh,
R. W. Roberts, R. Sun* — 12449 – 12453

Single-Round, Multiplexed Antibody Mimetic Design through mRNA Display

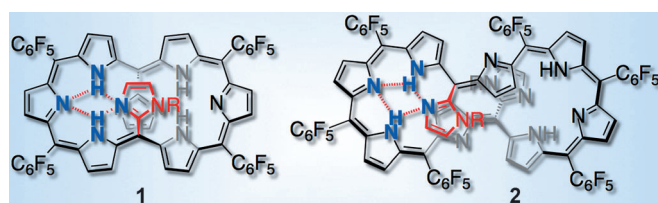


Disulfides enhance the transfection efficacy and reduce the toxicity of cationic gene delivery polymers. A quantitative analysis is provided of the bioreduction of a dynamic bioreducible dendritic polycationic probe in four cell lines. Such knowledge is indispensable for understanding and optimizing bioreducible drug and gene delivery systems.

Bioreduction

L. Brülisauer, N. Kathriner, M. Prenrecaj,
M. A. Gauthier,
J.-C. Leroux* — 12454 – 12458

Tracking the Bioreduction of Disulfide-Containing Cationic Dendrimers



Stable antiaromatic expanded porphyrins were designed by the judicious implementation of *meso*-imidazolyl groups, which cause stabilization through the creation of a hydrogen-bonding network (see structures) that overcomes antiaro-

matic electronic destabilization. Both the [28]hexaphyrin **1** and the [36]octaphyrin **2**, which contain imidazolyl groups at two opposite *meso* positions, are shown to be stable Hückel antiaromatic species.

Porphyrinoids

H. Mori, Y. M. Sung, B. S. Lee, D. Kim,*
A. Osuka* — 12459 – 12463

Antiaromatic Hexaphyrins and Octaphyrins Stabilized by the Hydrogen-Bonding Interactions of *meso*-Imidazolyl Groups



The German Chemical Society (GDCh) invites you to:



Angewandte Anniversary Symposium

GDCh
Eine Zeitschrift der Gesellschaft Deutscher Chemiker

Tuesday, March 12, 2013

Henry Ford Building / FU Berlin

Speakers



Carolyn R.
Bertozzi



François
Diederich



Alois
Fürstner



Roald Hoffmann
(Nobel Prize 1981)



Susumu
Kitagawa



Jean-Marie Lehn
(Nobel Prize 1987)



E.W. "Bert"
Meijer



Frank
Schirrmacher
(Publisher, FAZ)



Robert
Schlögl



George M.
Whitesides



Ahmed Zewail
(Nobel Prize 1999)

More information:



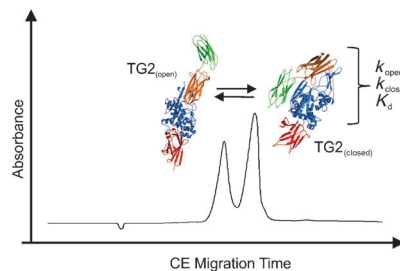
angewandte.org/symposium



 **WILEY-VCH**


GESELLSCHAFT
DEUTSCHER CHEMIKER

Conformational analysis: Capillary electrophoresis (CE) allows for the rapid separation of slowly interconverting protein conformers. Kinetic analysis (k_{open} , k_{closed} , and K_d) of electropherograms in the presence and absence of effector ligands allows the measurement of kinetic and thermodynamic constants associated with conformational changes and ligand binding.



Protein Conformations

C. M. Clouthier, G. G. Mironov,
V. Okhonin, M. V. Berezovski,
J. W. Keillor* ————— 12464–12468

Real-Time Monitoring of Protein
Conformational Dynamics in Solution
Using Kinetic Capillary Electrophoresis



Rise or fall: Complex-structured free-standing polymer films with molecular order in three dimensions were prepared through photoalignment of polymerizable

liquid crystals. The resulting films deform into cone and saddle shapes upon heating.

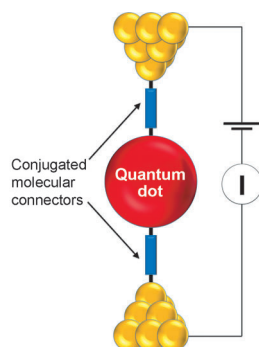
Smart Materials

L. T. de Haan, C. Sánchez-Somolinos,
C. M. W. Bastiaansen,
A. P. H. J. Schenning,*
D. J. Broer* ————— 12469–12472

Engineering of Complex Order and the
Macroscopic Deformation of Liquid
Crystal Polymer Networks



Making contact to a quantum dot: Single quantum-dot electronic circuits are fabricated by wiring atomically precise metal chalcogenide clusters with conjugated molecular connectors. These wired clusters can couple electronically to nanoscale electrodes and be tuned to control the charge-transfer characteristics (see picture).



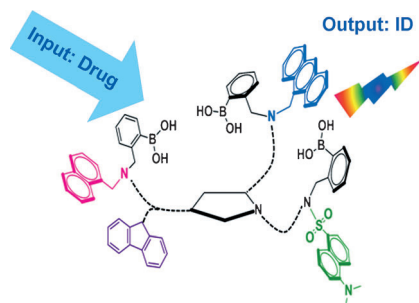
Molecular Electronics

X. Roy, C. L. Schenck, S. Ahn,
R. A. Lalancette, L. Venkataraman,*
C. Nuckolls,*
M. L. Steigerwald* ————— 12473–12476

Quantum Soldering of Individual
Quantum Dots



Inside Back Cover



Working together to uncover the truth: A molecule-sized diagnostic system combining several recognition elements and four fluorescence-emission channels enabled the identification of a wide range of pharmaceuticals on the basis of distinct photophysical processes. The molecular sensor (see simplified representation; ID = identification) was also used to analyze drug concentrations and combinations in urine samples in a high-throughput manner.

Molecular-Scale Diagnostics

B. Rout, L. Unger, G. Armony, M. A. Iron,
D. Margulies* ————— 12477–12481

Medication Detection by a Combinatorial
Fluorescent Molecular Sensor



Back Cover

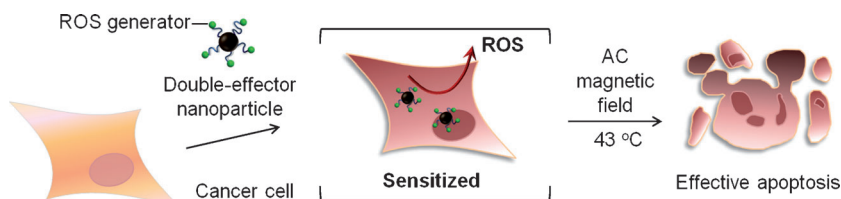


Synergistic Apoptosis

D. Yoo, H. Jeong, C. Preihs, J.-s. Choi,
T.-H. Shin, J. L. Sessler,*
J. Cheon* — 12482 – 12485



Double-Effector Nanoparticles:
A Synergistic Approach to Apoptotic
Hyperthermia



Highly efficient apoptotic hyperthermia is achieved using a double-effector nanoparticle that can generate reactive oxygen species (ROS) and heat. ROS render cancer cells more susceptible to subsequent heat treatment, which remarkably

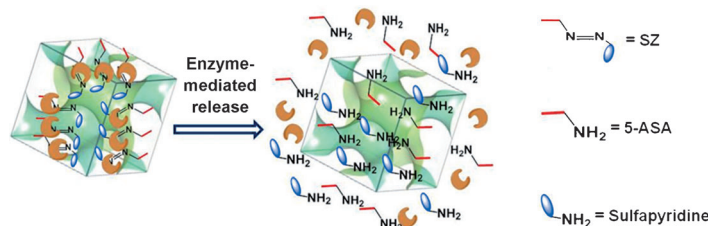
increases the degree of apoptotic cell death. Xenograft tumors (100 mm³) in mice are completely eliminated within 8 days after a single mild magnetic hyperthermia treatment at 43 °C for 30 min.

Targeted Drug Delivery

A. Popat, B. P. Ross, J. Liu,
S. Jambhrunkar, F. Kleitz,*
S. Z. Qiao* — 12486 – 12489



Enzyme-Responsive Controlled Release of
Covalently Bound Prodrug from
Functional Mesoporous Silica
Nanospheres



I want to break free: Mesoporous silica nanoparticles are functionalized with sulfasalazine (SZ; see scheme), a prodrug of 5-aminosalicylic acid (5-ASA) and sulfapyridine, to generate enzyme-responsive

nanocarriers. In the presence of the colon-specific enzyme azo-reductase (orange), 5-ASA and sulfapyridine are efficiently released.

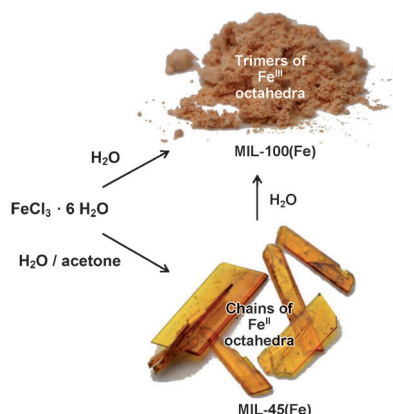
Metal–Organic Frameworks

T. Birsa Čelič, M. Rangus, K. Lázár,
V. Kaučič,
N. Zabukovec Logar* — 12490 – 12494



Spectroscopic Evidence for the Structure
Directing Role of the Solvent in the
Synthesis of Two Iron Carboxylates

Crystal engineering: The synthesis of the known compounds MIL-100(Fe) and MIL-45(Fe) is characterized by spectroscopy. The products are obtained under identical conditions by varying the solvent from pure water to a mixture of water and acetone. The starting solution, the gel, and the final reaction product were characterized by X-ray absorption spectroscopy (see picture).

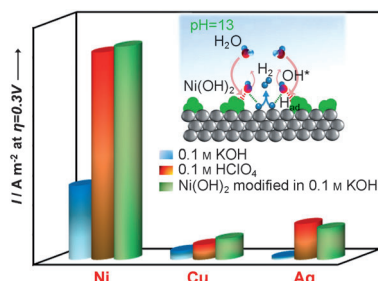


Electrochemistry

N. Danilovic, R. Subbaraman,
D. Strmcnik, K.-C. Chang, A. P. Paulikas,
V. R. Stamenkovic,
N. M. Markovic* — 12495 – 12498

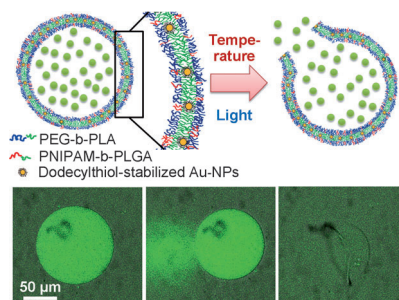


Enhancing the Alkaline Hydrogen
Evolution Reaction Activity through the
Bifunctionality of Ni(OH)₂/Metal
Catalysts



Active in alkaline environment: The activity of nickel, silver, and copper catalysts for the electrochemical transformation of water to molecular hydrogen in alkaline solutions was enhanced by modification of the metal surfaces by Ni(OH)₂ (see picture; *I* = current density and *η* = overpotential). The hydrogen evolution reaction rate on a Ni electrode modified by Ni(OH)₂ nanoclusters is about four times higher than on a bare Ni surface.

Microfluidics: Thermo- and photoresponsive polymersomes are assembled using capillary microfluidic devices. Encapsulants can be selectively released from the thermoresponsive polymersomes if they are incubated at and above temperatures of 40 °C, whereas the photoresponsive polymersomes selectively release encapsulants if illuminated with laser light (see picture; NP = nanoparticle).



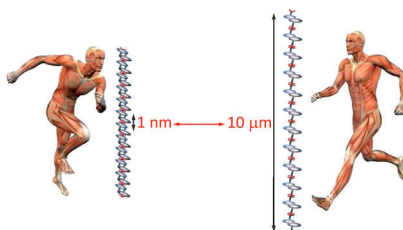
Soft Matter

E. Amstad, S.-H. Kim,
D. A. Weitz* 12499–12503

Photo- and Thermoresponsive
Polymersomes for Triggered Release



Pumping iron: Double-threaded rotaxanes can be linked to coordination units and polymerized in the presence of iron or zinc ions. pH modulation triggers cooperative contractions (or extensions) of the individual rotaxanes, thus resulting in an amplified motion of the muscle-like supramolecular chains with changes of their contour lengths of several micrometers (see picture).



Nanotechnology

G. Du, E. Moulin, N. Jouault, E. Buhler,*
N. Giuseppone* 12504–12508

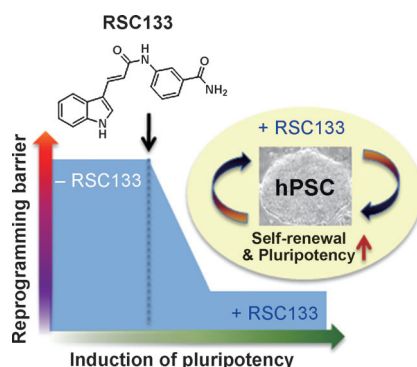
Muscle-like Supramolecular Polymers:
Integrated Motion from Thousands of
Molecular Machines



Front Cover



Booster of pluripotency: RSC133, a new synthetic derivative of indoleacrylic acid/indolepropionic acid, exhibits dual activity by inhibiting histone deacetylase and DNA methyltransferase. Furthermore it potentially improves the reprogramming of human somatic cells into a pluripotent state and aids the growth and maintenance of human pluripotent stem cells (hPSCs).



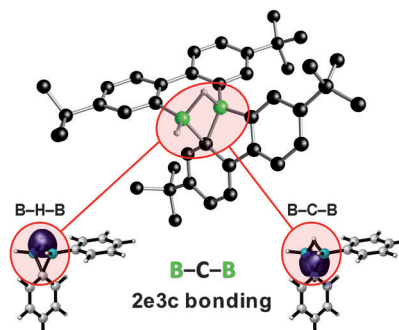
Pluripotency

J. Lee, Y. Xia, M.-Y. Son, G. Jin, B. Seol,
M.-J. Kim, M. J. Son, M. Do, M. Lee,
D. Kim, K. Lee,*
Y. S. Cho* 12509–12513

A Novel Small Molecule Facilitates the
Reprogramming of Human Somatic Cells
into a Pluripotent State and Supports the
Maintenance of an Undifferentiated State
of Human Pluripotent Stem Cells



Finally, boron did it too: The first example of a dimeric organyl(hydro)borane with a B-B-bridging aryl ring has been elucidated (see picture; B green/blue, C black/gray). It features a B–C–B two-electron–three-center bond and a largely unperturbed aromatic π -electron system.



Bond Theory

A. Hübner, M. Diefenbach, M. Bolte,
H.-W. Lerner, M. C. Holthausen,*
M. Wagner* 12514–12518

Confirmation of an Early Postulate:
B–C–B Two-Electron–Three-Center
Bonding in Organo(hydro)boranes

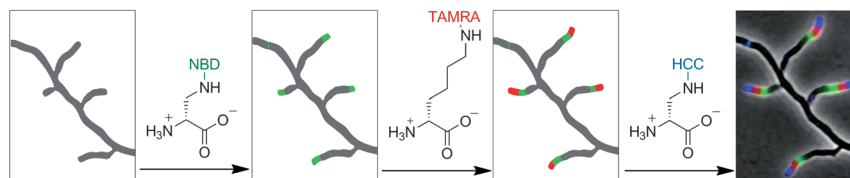


Fluorescence Imaging

E. Kuru, H. V. Hughes, P. J. Brown, E. Hall,
S. Tekkam, F. Cava, M. A. de Pedro,
Y. V. Brun,*
M. S. VanNieuwenhze* — 12519–12523



In Situ Probing of Newly Synthesized
Peptidoglycan in Live Bacteria with
Fluorescent D-Amino Acids



Tracking a bug's life: Peptidoglycan (PG) of diverse bacteria is labeled by exploiting the tolerance of cells for incorporating different non-natural D-amino acids. These nontoxic D-amino acids preferably label the sites of active PG synthesis,

thereby enabling fine spatiotemporal tracking of cell-wall dynamics in phylogenetically and morphologically diverse bacteria. HCC = 7-hydroxycoumarin, NBD = 7-nitrobenzofurazan, TAMRA = carboxytetramethylrhodamine.

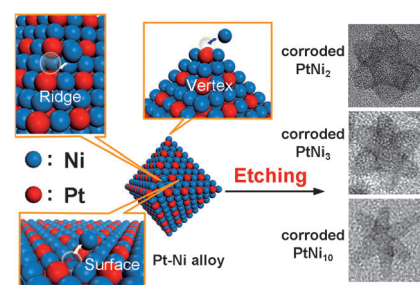
Alloys

Y. E. Wu, D. S. Wang, Z. Q. Niu,
P. C. Chen, G. Zhou,
Y. D. Li* — 12524–12528



A Strategy for Designing a Concave Pt–Ni Alloy through Controllable Chemical Etching

This corrosion: Octahedral Pt–Ni alloy nanoparticles (NPs) are converted into concave Pt₃Ni NPs by a coordination-assisted chemical-etching process (see scheme). The corroded concave Pt–Ni NPs have a higher density of low-coordinate atoms in steps sites, a decisive property in heterogeneous catalysis.

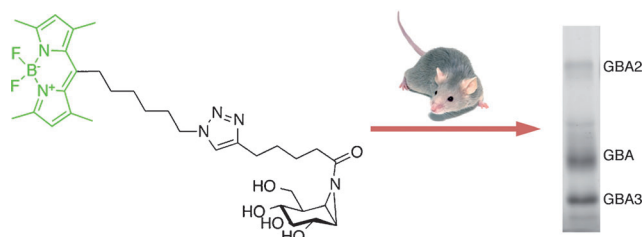


Enzyme Labeling

W. W. Kallemeijn, K.-Y. Li, M. D. Witte,
A. R. A. Marques, J. Aten, S. Scheij,
J. Jiang, L. I. Willems,
T. M. Voorn-Brouwer,
C. P. A. A. van Roomen, R. Ottenhoff,
R. G. Boot, H. van den Elst,
M. T. C. Walvoort, B. I. Florea,
J. D. C. Codée, G. A. van der Marel,
J. M. F. G. Aerts,*
H. S. Overkleeft* — 12529–12533



Novel Activity-Based Probes for Broad-Spectrum Profiling of Retaining β -Exoglucosidases In Situ and In Vivo



A high-end label: Cyclophellitol aziridine-type activity-based probes allow for ultra-sensitive visualization of mammalian β -glucosidases (GBA1, GBA2, GBA3, and LPH) as well as several non-mammalian β -glucosidases (see picture). These

probes offer new ways to study β -exoglucosidases, and configurational isomers of the cyclophellitol aziridine core may give activity-based probes targeting other retaining glycosidase families.

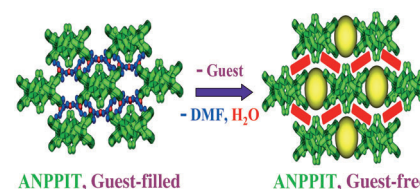
Multifunctional Organic Materials

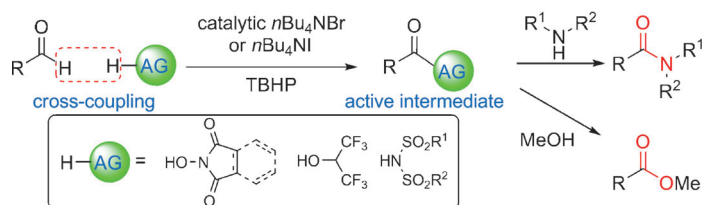
S. Dalapati, R. Saha, S. Jana, A. K. Patra,
A. Bhaumik, S. Kumar,
N. Guchhait* — 12534–12537



A Multifunctional Porous Organic Schottky Barrier Diode

Mesoporous materials: A multifunctional porous organic material (ANPPIT; see picture) has been synthesized and characterized. Multifunctionality of the compound has been determined from nitrogen adsorption, guest-dependent luminescence, and electrical conductivity measurements.





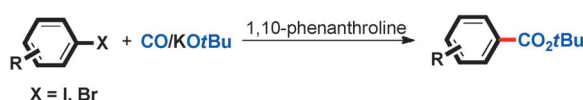
Formation on demand: An organocatalytic cross-coupling reaction of aldehydes with *N*-hydroxyimides, hexafluoroisopropyl alcohol, and sulfonimides has been developed. The resulting active intermediates can be directly converted into

amides or esters in one pot. This simple method makes use of readily available starting materials, and the newly discovered activating reagents should find broad application in the synthesis of amides and esters.

Cross-Coupling Reactions

B. Tan, N. Toda,
C. F. Barbas III* 12538–12541

Organocatalytic Amidation and Esterification of Aldehydes with Activating Reagents by a Cross-Coupling Strategy



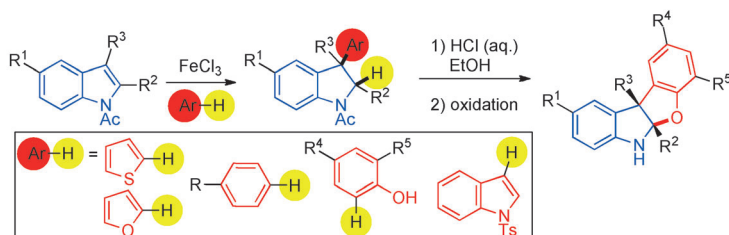
Transitions: The title reaction has been developed for the synthesis of a variety of *tert*-butyl benzoates by employing 1,10-phenanthroline as an additive. Various functional groups were tolerated and

heteroaryl iodides were also suitable substrates. Preliminary mechanism studies were conducted and indicate the participation of radical intermediates.

Radical Reactions

H. Zhang, R. Shi, A. Ding, L. Lu, B. Chen,
A. Lei* 12542–12545

Transition-Metal-Free Alkoxy carbonylation of Aryl Halides



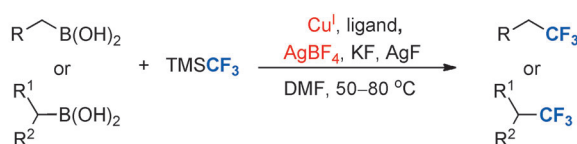
IRONic electrophilic indoles! The C3-regioselective hydroarylation of *N*-acetyl indoles with aromatic nucleophiles mediated by FeCl₃ features a rare example of the electrophilic reactivity of the indole core in a Friedel–Crafts reaction. This

indole umpolung allows us straightforward access to the tetracyclic benzofuroindoline motif found in the natural product diazonamide A, which is a potent antitumor agent.

Indole Chemistry

R. Beaud, R. Guillot, C. Kouklovsky,
G. Vincent* 12546–12550

FeCl₃-Mediated Friedel–Crafts Hydroarylation with Electrophilic *N*-Acetyl Indoles for the Synthesis of Benzofuroindolines



New couple: The Cu-promoted trifluoromethylation of primary and secondary alkylboronic acids with TMS-CF₃ extends the scope of transition-metal-catalyzed trifluoromethylation reactions to sp³-hy-

bridized carbon centers. It also represents one of the first examples for Cu-catalyzed C–C cross-coupling reactions of alkylboronic acid derivatives.

Cross-Coupling Reactions

J. Xu, B. Xiao, C.-Q. Xie, D.-F. Luo, L. Liu,
Y. Fu* 12551–12554

Copper-Promoted Trifluoromethylation of Primary and Secondary Alkylboronic Acids

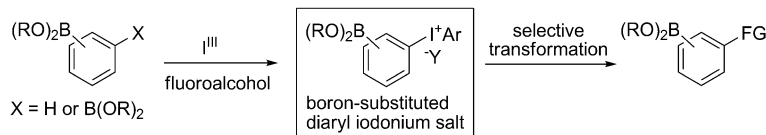


Boron Compounds

M. Ito, I. Itani, Y. Toyoda, K. Morimoto,
T. Dohi, Y. Kita* — 12555 – 12558



Synthesis of Boron-Substituted
Diaryliodonium Salts and Selective
Transformation into Functionalized Aryl
Boronates



Dormant boron awaits its true destiny in diaryliodonium salts synthesized from aryl boronate derivatives according to two alternative general methods with hyper-valent iodine(III) reagents and fluoroalcohol solvents: transformation of an aryl

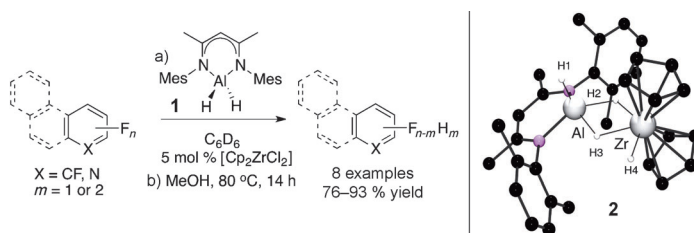
C–H bond and boron–iodine(III) exchange (see scheme; FG = functional group). The salts could be functionalized by both catalyst-free and metal-catalyzed reactions without loss of the boron functionality.

Hydrides

S. Yow, S. J. Gates, A. J. P. White,
M. R. Crimmin* — 12559 – 12563



Zirconocene Dichloride Catalyzed
Hydrodefluorination of C_{sp^2} –F bonds



A two-metal job: Four-coordinate aluminum dihydrides such as **1** are reported as terminal reductants for the selective title

reaction. The heterobimetallic complex **2** has been isolated and shown to be catalytically competent.

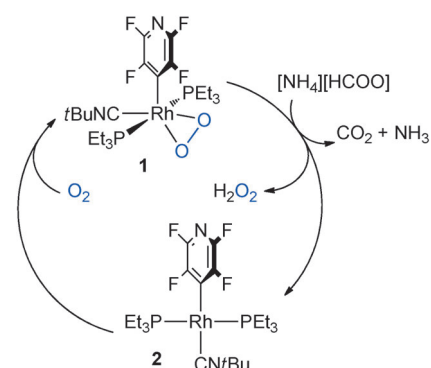
Oxygenation Chemistry

G. Meier, T. Braun* — 12564 – 12569



Hydrogenation of a Rhodium Peroxido
Complex by Formate Derivatives:
Mechanistic Studies and the Catalytic
Formation of H_2O_2 from O_2

Hydrogenation of dioxygen: The rhodium peroxido complex **1**, which can be prepared from **2** and dioxygen, can be reduced with dihydrogen sources to yield hydrogen peroxide. In a catalytic experiment, hydrogen peroxide is produced from dioxygen and ammonium formate under ambient conditions in the presence of **1** (see scheme).

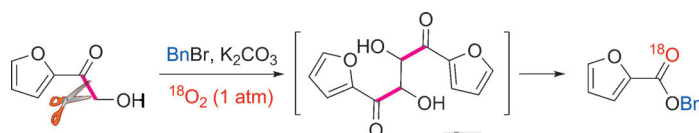


Synthetic Methods

H. Liu, C. Dong, Z. Zhang, P. Wu,
X. Jiang* — 12570 – 12574

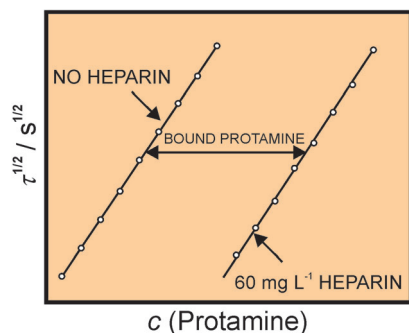


Transition-Metal-Free Aerobic Oxidative
Cleavage of C–C Bonds in α -Hydroxy
Ketones and Mechanistic Insight to the
Reaction Pathway



Clear cut: For the title reaction, O_2 , the ideal oxidant, was used as the only oxidizing reagent. The dimer intermediate

(see scheme) and isotopic labeling control experiments with $^{18}O_2$ partially disclosed the reaction mechanism.



A permselective membrane electrode allows the rapid and operationally reversible detection of the polycationic polypeptide protamine in physiological samples. Anticoagulant levels of heparin can be measured in undiluted whole blood by adding a known excess of its antidote protamine to discrete blood samples.

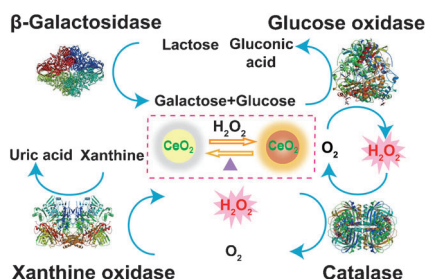
Membranes

G. A. Crespo, M. G. Afshar, E. Bakker* 12575 – 12578

Reversible Sensing of the Anticoagulant Heparin with Protamine Permselective Membranes



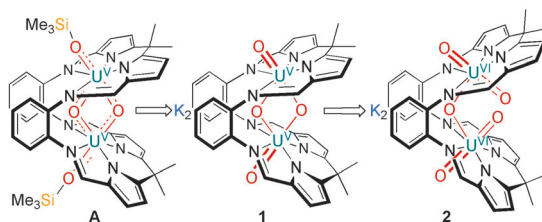
A label-free, resettable, and colorimetric logic network has been realized by utilizing thermally regenerable cerium oxide nanoparticles and biocatalytic reactions. Coupling switchable CeO_2 nanoparticles with biocomputing would convert molecular recognition events into colorimetric outputs and make logic gates feasible to reset.



Boolean Logic

Y. Lin, C. Xu, J. Ren,* X. Qu* 12579 – 12583

Using Thermally Regenerable Cerium Oxide Nanoparticles in Biocomputing to Perform Label-free, Resettable, and Colorimetric Logic Operations



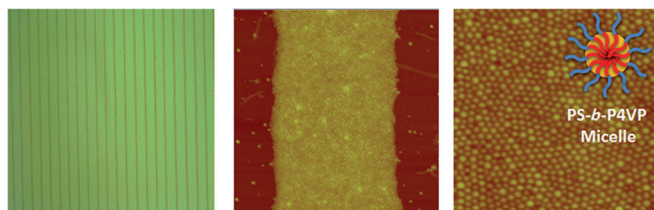
Switching on uranium(V) reactivity: The silylated uranium(V) dioxo complex $[(\text{Me}_3\text{SiOUO})_2(\text{L})_2]$ (**A**) is inert to oxidation, but after two-electron reduction to $[(\text{Me}_3\text{SiOUO})_2(\text{L})]^{2-}$ (**1**), it can be desilylated to form $[\text{OU}(\mu\text{-O})_2\text{UO}(\text{L})_2]^{2-}$ (**2**) with

reinstated uranyl character. Removal of the silyl group uncovers new redox and oxo rearrangement chemistry for uranium, thus reforming the uranyl motif and involving the $\text{U}^{\text{VI}}/\text{V}$ couple in dioxygen reduction.

Uranyl Complexes

G. M. Jones, P. L. Arnold,* J. B. Love* 12584 – 12587

Controlled Deprotection and Reorganization of Uranyl Oxo Groups in a Binuclear Macrocyclic Environment



Earning their stripes: A hierarchical assembly of micelles composed of an amphiphilic diblock copolymer, poly(styrene)-*block*-poly(4-vinylpyridine) (PS-*b*-P4VP), were made by combining controlled evaporative self-assembly of the

confined PS-*b*-P4VP toluene solution in a cylinder-on-Si geometry with spontaneous self-assembly of micelles. This method gave microscopic stripes of nanometer-sized PS-*b*-P4VP micelles within the stripes (see pictures).

Nanoparticle Gradient

W. Han, M. Byun, B. Li, X. Pang, Z. Lin* 12588 – 12592

A Simple Route to Hierarchically Assembled Micelles and Inorganic Nanoparticles



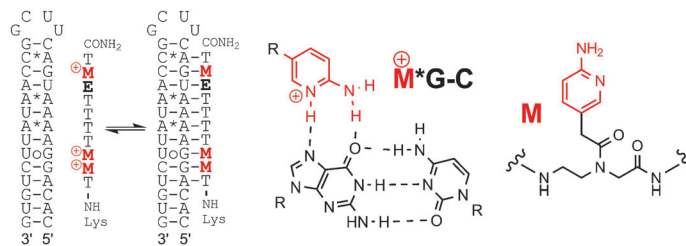


RNA Recognition

T. Zengeya, P. Gupta,
E. Rozners* — 12593 – 12596



Triple-Helical Recognition of RNA Using
2-Aminopyridine-Modified PNA at
Physiologically Relevant Conditions



Peptide nucleic acids containing thymidine and 2-aminopyridine (**M**) nucleobases form stable and sequence-selective triple helices with double-stranded RNA at physiologically relevant conditions. The

M-modified PNA showed unique RNA selectivity by having two orders of magnitude higher affinity for the double-stranded RNAs than for the same DNA sequences.

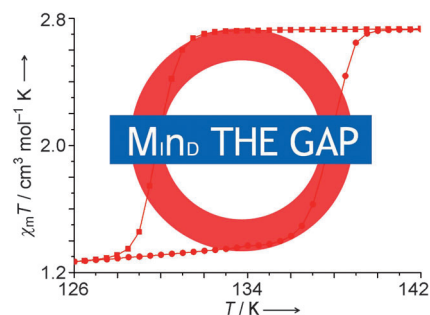
Spin Transitions

P. N. Martinho, B. Gildea, M. M. Harris,
T. Lemma, A. D. Naik, H. Müller-Bunz,
T. E. Keyes, Y. Garcia,
G. G. Morgan* — 12597 – 12601



Cooperative Spin Transition in
a Mononuclear Manganese(III) Complex

Mind the gap: A complete, cooperative spin transition for a mononuclear Mn^{III} complex is reported with an 8 K hysteresis window. Raman spectra collected at a single temperature in warming and cooling modes confirm the electronic bistability within the hysteresis loop. The source of the cooperativity is a disconnection in the hydrogen-bonded 1D chains that connect adjacent cations owing to an order–disorder transition in the PF_6^- counterion.

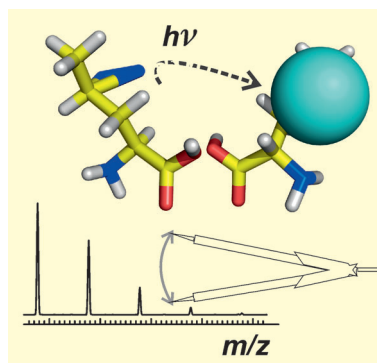


Peptide Structures

K. Kölbel,* C. H. Ihling,
A. Sinz* — 12602 – 12605



Analysis of Peptide Secondary Structures
by Photoactivatable Amino Acid
Analogues



Photochemical cross-linking was applied to trap intramolecular interactions in peptides. The incorporation of diazirine-labeled amino acid analogues in combination with high-resolution mass spectrometry made it possible to catch reverse-turn conformations within peptides, exactly map their self-interacting surfaces, and discriminate between stable and transient interactions.

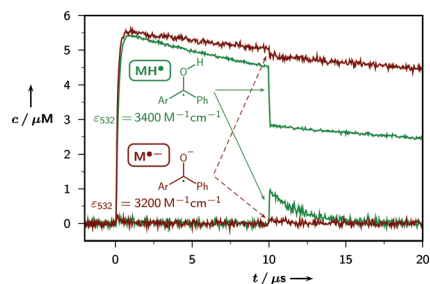
Photoionization

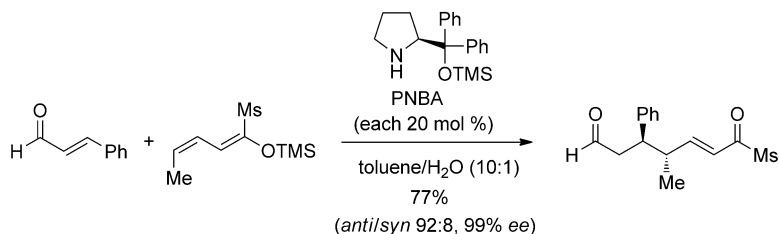
M. Goetz,* C. Kerzig — 12606 – 12608



Counterintuitive Influence of Protonation
on Radical-Anion Photoionization

Protonation of a radical anion $\text{M}^{\cdot-}$ greatly facilitates its green-light photoionization despite much less favorable energetics for the resulting ketyl radical MH^{\cdot} . The state symmetry of the radical species absorbing the ionizing photon might provide an explanation.





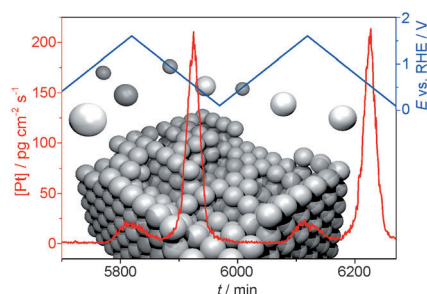
Now also acyclic: The first catalytic, enantioselective, vinylogous Michael reaction of linear, acyclic dienol silyl ethers was achieved. The reaction, based upon the principle of iminium ion catalysis, delivered 1,7-dioxo compounds in one

step with good yields, complete regio-, and excellent enantioselectivity. γ -Substituted dienol silyl ethers furnished products with two new stereogenic centers with good diastereoselectivity. Ms = mesityl, PNBA = *para*-nitrobenzoic acid.

Asymmetric Catalysis

V. Gupta, S. Sudhir V., T. Mandal, C. Schneider* 12609–12612

Organocatalytic, Highly Enantioselective Vinylogous Mukaiyama–Michael Reaction of Acyclic Dienol Silyl Ethers

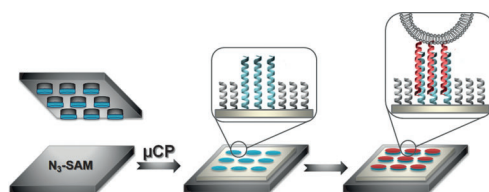


Platinum stability: Dissolution of Pt, which is one major degradation mechanism in, for example, hydrogen/air fuel cells, was monitored under potentiodynamic and potentiostatic conditions. The highly sensitive and time-resolving dissolution monitoring enables the distinction between anodic and cathodic dissolution processes during potential transient and chronoamperometric experiments, and the precise quantification of the amount of dissolved Pt.

Platinum Dissolution

A. A. Topalov,* I. Katsounaros, M. Auinger, S. Cherevko, J. C. Meier, S. O. Klemm, K. J. J. Mayrhofer* 12613–12615

Dissolution of Platinum: Limits for the Deployment of Electrochemical Energy Conversion?



Patchy surfaces: An azide-terminated self-assembled monolayer was patterned with the peptide sequence (EIAALEK)₃ by using microcontact printing. This sequence forms stable coiled-coil heterodimers with

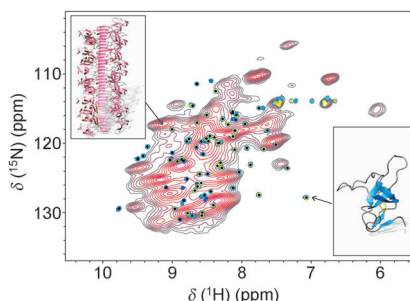
the complementary peptide (KIAALKE)₃. By introducing this peptide to the surface of phospholipid liposomes and cyclodextrin vesicles, liposomes and vesicles can be immobilized at the patterned surface.

Surface Patterning

J. Voskuhl, C. Wendeln, F. Versluis, E.-C. Fritz, O. Roling, H. Zope, C. Schulz, S. Rinnen, H. F. Arlinghaus, B. J. Ravoo,* A. Kros* 12616–12620

Immobilization of Liposomes and Vesicles on Patterned Surfaces by a Peptide Coiled-Coil Binding Motif

GrEASy fibrils: Hydrophobins are fungal proteins that assemble into an amphipathic fibrillar monolayer with amyloid properties and a hydrophobic face as water-resistant as Teflon. Solid-state NMR studies on EAS hydrophobin fibrils reveal direct evidence of a partial molecular rearrangement on assembly and an ordered β -sheet-rich core in the context of a whole protein in this functional amyloid.



Amyloids

V. K. Morris, R. Linser, K. L. Wilde, A. P. Duff, M. Sunde,* A. H. Kwan* 12621–12625

Solid-State NMR Spectroscopy of Functional Amyloid from a Fungal Hydrophobin: A Well-Ordered β -Sheet Core Amidst Structural Heterogeneity



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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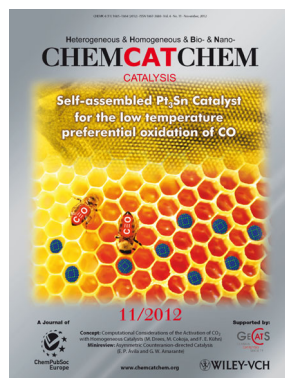


This article is accompanied by a cover picture (front or back cover, and inside or outside).

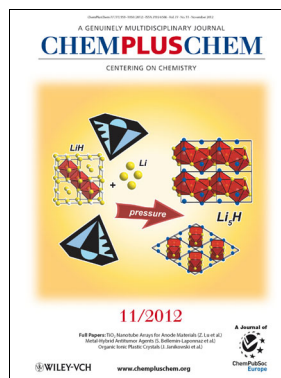
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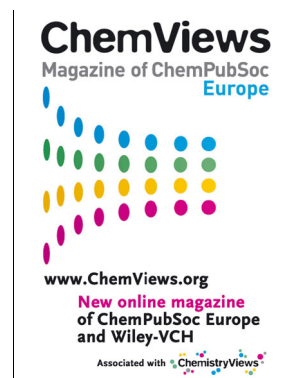
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