

General Information

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Contributions should be in English, but may be accompanied by an abstract in the language of the authors. The correspondence author will receive **page proofs** (in most cases as compressed PDF files). The correspondence author will also receive a complementary copy of the journal and a reprint PDF file, restricted to 25 printouts, free of charge. Manuscripts will be edited with a view to brevity and clarity. Queries regarding manuscripts should be sent to chemistry@wiley-vch.de.

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The authors must inform the editor of manuscripts submitted, soon to be submitted, or in press at other journals that have a bearing on the manuscript being submitted. *Chemistry—A European Journal* does not publish manuscripts that have already appeared in print or electronically.

The Notice to Authors as well as the Conditions for Publication are also available on the WWW on the *Chemistry* home page (<http://www.chemeurj.org/>).

Categories of Contributions

1. Concepts

The Concepts section contains short articles emphasizing the general concepts that have guided important developments in a specific area and their implications for future research. The reference sections should include the key papers that have contributed to conceptual advances in the field under review, rather than being fully comprehensive. The author should aim to

provide the nonspecialist reader with a useful conceptual guide and the expert with a new angle on a familiar problem. Although Concept articles are generally written upon invitation of the editor, unsolicited manuscripts are also welcome. All contributions are subject to peer review.

The Concept articles may be organized as the author wishes, but should include a very short Abstract (400 characters; an additional version in the authors' native language may also be included) that succinctly describes the concepts under discussion. A graphical abstract for the table of contents and keywords should also be provided. Articles should consist of around 10 double-spaced pages of text (including references, tables, and legends). The liberal use of schemes and figures is encouraged.

2. Full Papers

Full papers must either be of current general interest or of great significance to a more specialized readership. All contributions will be judged on the criteria of originality, quality, and novelty. Papers that are suitable for consideration will be sent to a minimum of two referees. Based on the referees' recommendations, the Editor will make a decision on whether to accept a contribution. Manuscripts may not have been published previously, except in the form of a preliminary communication (reprint requested).

Chemistry does not publish full papers that consist mainly of results reported in a previous communication with an added experimental section.

3. Correspondence

Comments on publications in *Chemistry—A European Journal* are welcome if they contribute to the scientific discussion.

Manuscript and Disk Preparation

A *disk* (MS DOS or Macintosh format) should be submitted with the final accepted version of the manuscript; the text should be saved as a single file, graphics should be sent on a separate disk (further details will be provided on acceptance of the manuscript). Not all graphics programs are suitable for use by our printer; please consult the *Chemistry* Checklist (printed immediately after the Notice to Authors or available as a PDF file on <http://www.chemeurj.org>). If possible, the standard Symbol font should be used to create Greek letters, rather than special characters or graphics embedded in the text. Use a tabulator to indicate new paragraphs. If you use L^AT_EX, please send standard L^AT_EX files only; please do not include your own style sheets or macros.

Organization of Full Papers

We can process your manuscript faster if you lay it out as described below.

Title page: Title; authors' names with academic titles, an asterisk denotes the correspondence author; affiliations of all the authors, including the full postal address, fax number, and e-mail address of the correspondence author; series title, number, and reference to previous paper in the series, if applicable; dedication, if applicable.

Abstract should be brief (600–2000 characters) and not too technical. An additional version of the abstract in the authors'

native language may also be supplied. When written in a non-Roman alphabet, this must be in camera-ready form: the text must fit into a single column 8.5 cm wide; type size, 2 mm (7 point); about 4 mm between lines (single-spaced).

Keywords: A maximum of five keywords to appear in the printed and online indexes should be given in alphabetical order. At least two keywords from the core keyword list, which is also available on the WWW (see section on Keywords at the end of the Notice to Authors), should be included to aid online searching.

Introduction should include relevant references.

Results and Discussion may be combined or kept separate and may be further divided by subheadings. This section should not be cluttered with technical details. Abbreviations and acronyms should be used sparingly and consistently. Where they first appear in the text, they should—apart from the most common ones such as NMR, IR, and UV—be defined; you may prefer to explain large numbers of abbreviations and acronyms in a footnote on the first page.

Experimental Section should be given in sufficient detail to enable others to repeat your work. In theoretical papers, technical details such as computational methods should likewise be confined to an appropriately named section.

In so far as practical, authors should use a systematic name (IUPAC or Chemical Abstracts) for each title compound in the Experimental Section. Please do not use computer programs to generate elaborate systematic names or use long multi-line compound names; for the sake of clarity general descriptors such as compound **2**, dendrimer **3**, or alcohol **4** should be used.

Equipment and conditions used for the measurement of physical data should be described at the beginning of the Experimental Section. Quantities of reactants, solvents etc. should be included in parentheses rather than in the running text (e.g., triphenylstannyl chloride (0.964 g, 2.5 mmol) in toluene (20 mL)). Physical data should be quoted with decimal points and negative exponents (e.g., 25.8 JK⁻¹ mol⁻¹). The purity of all new compounds should be verified by elemental analysis, to an accuracy of within ±0.4%. In special cases, for instance, when the compound is unstable or not available in sufficient quantities for complete analysis, the exact relative molecular mass obtained from a high-resolution mass spectrum and a clean ¹³C NMR spectrum (as additional material for inspection by the referees) should be supplied.

Detailed presentation of physical data: $R_f = 0.38$ (CHCl₃/MeOH 9:1); m.p. 20 °C; $[\alpha]_D^{20} = -13.5$ ($c = 0.2$ in acetone); ¹H NMR (200 MHz, [D₈]THF, 25 °C, TMS): $\delta = 1.3$ (q, ³J(H,H) = 8 Hz, 2H; CH₂), 0.9 (t, ³J(H,H) = 8 Hz, 3H; CH₃); IR (Nujol): $\tilde{\nu} = 1790$ cm⁻¹ (C=O); UV/Vis (*n*-hexane): $\lambda_{\max}(\epsilon) = 320$ (5000), 270 nm (12000 mol⁻¹ dm³ cm⁻¹); MS (70 eV, EI): m/z (%): 108 (20) [$M^+ - H$], 91 (100) [$C_7H_7^+$]; elemental analysis calcd (%) for C₁₂H₁₁AsOS (278.2): C 51.85, H 3.99, As 26.93, S 11.53; found: C 51.59, H 3.96, As 27.07, S 11.39.

Acknowledgments

References: In the text the numbers should be typed in square brackets as superscripts (e.g., Wittig^[3]) and, if applicable, after punctuation. If you use the automatic reference collation system of your word-processing program, please convert the references into text before submitting the disk; otherwise they may disappear when typeset. Journal titles should be abbreviated according to the Chemical Abstracts Service Source Index (CASSI). Please follow the examples below.

Journals: [1] a) B. M. Trost, *Chem. Eur. J.* **1998**, *4*, 2405–2412; b) H. J. Ache, *Angew. Chem.* **1989**, *101*, 1–21; *Angew. Chem. Int. Ed.*

Engl. **1989**, *28*, 1–20; c) H. Frey, *Angew. Chem.* **1998**, *110*, 2313–2318; *Angew. Chem. Int. Ed.* **1998**, *37*, 2193–2197.

[2] a) A. Kraft, *Chem. Commun.* **1996**, 77–79, and references therein; b) *Sci. Am.* **1984**, *250*(4), 7–8; c) B. Krebs, H. U. Hürter, *Acta Crystallogr. Sect. A* **1981**, *37*, 163.

Books: Without editor: [3] E. Wingender, *Gene Regulation in Eukaryotes*, VCH, Weinheim, **1993**, p. 215. With editor: [4] T. D. Tullius in *Comprehensive Supramolecular Chemistry*, Vol. 5 (Eds.: J. L. Atwood, J. E. D. Davies, D. D. MacNicol, F. Vögtle, K. S. Suslick), Pergamon, Oxford, **1996**, pp. 317–343.

Miscellaneous: [5] a) C. R. A. Botta (Bayer AG), DE-B 2235093, **1973** [*Chem. Abstr.* **1974**, *80*, 55356c]; b) A. Student, PhD thesis, University of Newcastle (UK), **1991**; c) G. Maas, *Methoden Org. Chem. (Houben-Weyl)* 4th ed. 1952, Vol. E21/1, pp. 379–397; d) “Synthesis in Biochemistry”: R. Robinson, *J. Chem. Soc.* **1936**, 1079; e) S. Novick, “Biography of Rotational Spectra for Weakly Bound Complexes”, to be found under <http://www.wesleyan.edu/chem.bios/vdw.html>, **1999**; f) G. M. Sheldrick, SHELXS-96, Program for the Solution of Crystal Structures, University of Göttingen, Göttingen (Germany), **1996**.

Legends: Each figure and scheme should have a legend; these should be listed together at the end of the reference section of the text file rather than being included with the drawings in the graphics files.

Tables must have a brief title and should only be subdivided by three horizontal lines (head rule, neck rule, foot rule). Footnotes in tables are denoted [a], [b], [c], etc.

Illustrations (structural formulas, figures, schemes) must be submitted camera-ready on separate sheets and on disk and should, if possible, be designed for reduction to a one-column format (8.5 cm wide). The maximum width is the two-column format (17.5 cm wide). For optimum reproduction, illustrations should be larger than the desired final size. We recommend: font for script, Helvetica; size of lettering, 3–3.5 mm; total maximum width, 14 cm (or 28 cm for two-column width) for 60% reduction.

Italicize symbols of physical quantities, but not their units (e.g., *T* for temperature, in contrast to *T* for the unit Tesla, but *K* as unit; *J*, but Hz; *a*, but nm), stereochemical information (*cis*, *Z*, *R*, etc.), locants (*N*-methyl, α -amino) and symmetry (*C*_{2v}). Chemical formulas should be numbered with boldface Arabic numerals (e.g., **1**). Labels of axes are to be separated from their units by a slash (e.g., *T/K*). Abbreviations such as Me, Et, *n*Bu, *i*Pr, *s*Bu, *t*Bu and Ph (not ϕ) may be used in formulas. General substituents should be indicated by R¹, R² (not R₂, which means 2R) or R, R' (not R'). The spatial arrangement of the substituents should be indicated by hatched lines (||||) and a wedge. A minus sign must be as long as the crossbar of a plus sign.

Part of the additional cost for the reproduction of color figures must be paid by the author (details will be provided after acceptance of the manuscript). Good prints should be submitted for the referees and editors; for reproduction, originals should be provided as slides or on disk (please contact the editorial office for details on the latter option; data transmission by e-mail is very costly and should be avoided if possible).

Cover picture/frontispiece: We encourage authors to submit pictures for the cover page and for frontispieces.

Graphical Abstract: A short text for the table of contents should be included as the last page and saved as part of the main text. It

should tempt browsers to read the article, and so need not summarize the entire paper. Graphics (formulas, part of a figure) should be kept small.

Crystal Structure Analysis

If a crystal structure analysis is not an essential part of the paper, only a footnote is required indicating where the detailed results can be found. Otherwise, the following data should be given in the manuscript: crystal dimensions, crystal system, space group, unit cell dimensions and volume, ρ_{calcd} , $2\theta_{\text{max}}$, radiation, wavelength, scan mode, temperature of measurement, no. of measured and independent reflections, no. of reflections included in refinement, σ limits, whether and how Lorentzian polarization and absorption corrections were performed (μ , min/max transmission), method of structure solution and program, method of refinement and program, no. of parameters, treatment of H atoms. R , wR , whether refined against $|F|$ or $|F^2|$, residual electron density. An ORTEP-type plot will not be reproduced when it merely serves to confirm the structure of a synthetic intermediate.

Authors should deposit their data before submitting their manuscripts or update data already available, so that referees can retrieve the information directly from the database.

For organic and organometallic compounds: send your data (**without** structure factors) to the Cambridge Crystallographic Data Centre as an ASCII file by e-mail or as a hard copy (CCDC, 12 Union Road, Cambridge CB2 1EZ (UK); tel: (+44) 1223-336-408; fax: (+44) 1223-336-033; e-mail: deposit@ccdc.cam.ac.uk; WWW: <http://www.ccdc.cam.ac.uk>). The data will be assigned a registry number, which should be included with the following standard text in the manuscript: "CCDC-. . . contain(s) the supplementary crystallographic data for this paper. These data

can be obtained free of charge via www.ccdc.cam.ac.uk/conts/retrieving.html (or from the Cambridge Crystallographic Data Centre, 12 Union Road, Cambridge CB2 1EZ, UK; fax: (+44) 1223-336-033; or e-mail: deposit@ccdc.cam.ac.uk)."

For inorganic compounds: The Fachinformationszentrum Karlsruhe only accepts data deposited in electronic form. Send the data by e-mail, on disk, or by FTP (in the last case, contact FIZ beforehand) to FIZ, 76344 Eggenstein-Leopoldshafen (Germany); tel: (+49) 7247-808-205; fax: (+49) 7247-808-666; e-mail: crysdata@fiz-karlsruhe.de; FTP: ftp.fiz-karlsruhe.de (path: /pub/csd); WWW: <http://crystal.fiz-karlsruhe.de> under "Products and Services". You will be given a CSD number, which should be included with the following standard text in the manuscript: "Further details of the crystal structure investigation(s) can be obtained from the Fachinformationszentrum Karlsruhe, 76344 Eggenstein-Leopoldshafen, Germany, (fax: (+49) 7247-808-666; e-mail: crysdata@fiz.karlsruhe.de) on quoting the depository number CSD-. . ."

Supporting Information

Supporting information may be included for deposition on the WWW; the author must keep a copy to make available to readers who do not have access to the WWW. This material undergoes the peer review process and must therefore be included as hard copies with the original submission. Succinct text and the necessary graphics should then be sent as a single MS Word (preferably Word 7)/MS Word for Macintosh file with the revised version on a separate disk marked "supporting information for" + MS number + correspondence author name. In this case only, graphics should be present as imports in the file and not as separate files to which the document refers. Supporting Information should **not include** crystallographic data that are available from CCDC or FIZ.

Basic Keyword List

An interjournal browsing facility has been developed for the readers of the Wiley-VCH journals *Advanced Synthesis & Catalysis*, *Angewandte Chemie International Edition*, *Chemistry—A European Journal*, *European Journal of Inorganic Chemistry*, *European Journal of Organic Chemistry*, *Journal für praktische Chemie* and *Zeitschrift für anorganische und allgemeine Chemie*: in the Internet you will be able to jump between lists of thematically related contributions by mouse click.

To be able to offer this option on the WWW we have compiled a common keyword catalogue that is printed here and is also available online (see <http://www.chemeurj.org/>). To assist you in finding keywords, they are listed according to category. As with all such records, a few guidelines facilitate its use, and these are briefly explained below:

- At least two of the maximum of five keywords assigned to an article must come from this list.
- Named reactions will be incorporated only in exceptional cases. Generally the reaction type is selected instead. For example, Diels–Alder reactions will be found under “cycloadditions” and Claisen rearrangements under “rearrangements”.
- Heteroanalogues of compounds are mainly classified under the C variants, for example, (hetero)cumulenes, (hetero)dienes. A few aza and phospho derivatives are exceptions.
- Compounds with inorganic components that are central to the article are listed under the element, for instance, iron complexes under “iron” and the type of ligand. Some group names like “alkali metals” exist alongside the names of important members of the group like “lithium”. In such cases the group name is used for these members only when comparative studies are described. The members not appearing separately are also categorized under the group name.
- A keyword in the form “N ligand” is only chosen if a considerable portion of the paper deals with the coordination of any ligand ligating through the atom concerned (in the example, nitrogen).
- Spectroscopic methods are assigned as keywords only if the article is about the method itself or if the spectroscopic technique has made an important contribution to the problem under investigation.
- “Structure elucidation” is intended only if the crux of the paper is a structural elucidation or if a combination of several spectroscopic techniques were needed for conclusive solution of the structure.
- An attempt has been made to avoid synonyms and to select more general concepts rather than specialized terms. Thus the term “double-decker complexes” is excluded in favor of “sandwich complexes”. See also points 2. and 3. in these guidelines.
- Enzymes should be assigned to one of the six main enzyme classes.

This list will be a “living” catalogue to be flexible enough to absorb the new developments in chemistry. We therefore welcome all suggestions from our readers and authors that might improve its user-friendliness. The current version may be found in the World Wide Web under the following address: <http://www.wiley-vch.de/vch/journals/keyword.php>

Analytical Chemistry and Spectroscopic Methods

Analytical methods	Gas chromatography	Moessbauer spectroscopy	Surface plasmon resonance
Circular dichroism	High-throughput screening	Neutron diffraction	Trace analysis
Cyclic voltammetry	Ion chromatography	NMR spectroscopy	UV/Vis spectroscopy
Electron diffraction	Ion exchange	Photoelectron spectroscopy	Vibrational spectroscopy
Electron microscopy	IR spectroscopy	Plasma chemistry	X-ray absorption spectroscopy
Electrophoresis	Isotopic labeling	Raman spectroscopy	X-ray diffraction
ENDOR spectroscopy	Laser spectroscopy	Rotational spectroscopy	ZEKE spectroscopy
EPR spectroscopy	Liquid chromatography	Scanning probe microscopy	
EXAFS spectroscopy	Luminescence	Sensors	
Fluorescence spectroscopy	Mass spectrometry	Surface analysis	

Biological Chemistry and Chemical Biology (including Biochemistry, Bioorganic Chemistry, Bioinorganic Chemistry, Medicinal Chemistry, and Molecular and Cell Biology)

Allosterism	Antifungal agents	Azapeptides	Bioorganic chemistry
Amino acids	Antigens	Azasugars	Biosensors
Angiogenesis	Antisense agents	Bioinformatics	Biosynthesis
Antibiotics	Antitumor agents	Bioinorganic chemistry	Biotransformations
Antibodies	Antiviral agents	Biomimetic synthesis	C-glycosides

Carbohydrates	Genomics	Metalloproteins	Protein models
Carbon dioxide fixation	Glycoconjugates	Micelles	Protein modifications
Carotenoids	Glycolipids	Molecular evolution	Protein structures
Cell adhesion	Glycopeptides	mRNA	Proteins
Cell recognition	Glycoproteins	Mutagenesis	Proteomics
Cerebrosides	Glycosides	Natural products	Proton transport
Chaperone proteins	Glycosylation	Neurochemistry	Radiopharmaceuticals
Cobalamines	Growth factors	Neurotransmitters	Receptors
Cofactors	Helical structures	Nitrogen fixation	Redox chemistry
Combinatorial chemistry	Heme proteins	Nitrogenases	Ribonucleosides
Cyclitols	Hormones	Nucleic acids	Ribozymes
Cyclodextrins	Hydrolases	Nucleobases	RNA
Cytokines	Immobilization	Nucleosides	RNA recognition
DNA	Immunoassays	Nucleotides	RNA structures
DNA cleavage	Immunochemistry	Oligonucleotides	Sensitizers
DNA damage	Immunology	Oligosaccharides	Sequence determination
DNA methylation	Inhibitors	Oxidoreductases	Sialic acids
DNA recognition	Ion channels	Peptide nucleic acids	Siderophores
DNA replication	Ionophores	Peptides	Signal transduction
DNA structures	Isomerases	Peptidomimetics	Sphingolipids
Dopamines	Ligases	Pheromones	Steroids
Drug delivery	Lipids	Phospholipids	Structure – activity relationships
Drug design	Lipophilicity	Photoaffinity labeling	Terpenoids
Electron transport	Lipoproteins	Photosynthesis	Toxicology
Enzyme models	Liposomes	Phytochemistry	Transferases
Enzymes	Lyases	Polyketides	tRNA
Fibrous proteins	Medicinal chemistry	Polymerase chain reaction	Vesicles
Fluorescent probes	Membrane proteins	Prostaglandins	Vitamins
Gene expression	Membranes	Protein design	
Gene sequencing	Metabolism	Protein engineering	
Gene technology	Metalloenzymes	Protein folding	

Catalysis

Asymmetric catalysis	Catalytic antibodies	Heterogeneous catalysis	Phase-transfer catalysis
Autocatalysis	Enzyme catalysis	Homogeneous catalysis	Supported catalysts
Biphasic catalysis			

Coordination Chemistry: Compound Classes

Cage compounds	Cuprates	Metallacycles	Polyoxometalates
Chelates	Dendrimers	Metallocenes	Sandwich complexes
Clathrates	Heterometallic complexes	Nitrogen oxides	Ylides
Cluster compounds			

Coordination Chemistry: Ligand Classes

Alkene ligands	Carboxylate ligands	Macrocyclic ligands	P ligands
Alkyne ligands	Carbyne ligands	N ligands	Si ligands
Allyl ligands	Cyclopentadienyl ligands	N,O ligands	S ligands
Arene ligands	Diene ligands	N,P ligands	Tridentate ligands
As ligands	Dioxygen ligands	O ligands	Tripodal ligands
Bridging ligands	Fluorinated ligands	Oxo ligands	Vinylidene ligands
Carbene ligands	Hydride ligands	Peroxo ligands	
Carbonyl ligands	Isocyanide ligands	Phosphane ligands	

Coordination Chemistry: Methodology and Reactions

Carbon dioxide fixation	Matrix isolation	Oxidation	Solvolysis
Chemical vapor deposition	Metathesis	Radical reactions	Substituent effects
Chiral resolution	Neighboring-group effects	Reduction	Template synthesis
Crystal engineering	Nitrogen fixation	Ring-opening polymerization	
Ligand design	O-O activation	Solvent effects	

Coordination Chemistry: Structure

Agostic interactions	Electron-deficient compounds	Hydrogen bonds	Noncovalent interactions
Aurophilicity	Electronic structure	Inclusion compounds	Pi interactions
Charge transfer	Electrostatic interactions	Isolobal relationship	Stacking interactions
Cooperative effects	Fluxionality	Jahn-Teller distortion	Structure elucidation
Coordination modes	Helical structures	Ligand effects	Through-bond interactions
Donor-acceptor systems	Host-guest systems	Metal-metal interactions	Through-space interactions
		Multiple bonds	

Elements and Element Groups

Actinides	Copper	Magnesium	Scandium
Alkali metals	Deuterium	Manganese	Selenium
Alkaline earth metals	Fluorine	Mercury	Silicon
Aluminum	Gallium	Molybdenum	Silver
Antimony	Germanium	Neon	Sodium
Argon	Gold	Nickel	Strontium
Arsenic	Group 13 elements	Niobium	Sulfur
Barium	Group 14 elements	Nitrogen	Tantalum
Beryllium	Hafnium	Noble gases	Technetium
Bismuth	Halogens	Osmium	Tellurium
Boron	Helium	Oxygen	Thallium
Bromine	Hydrogen	Palladium	Tin
Cadmium	Indium	Phosphorus	Titanium
Calcium	Iodine	Platinum	Tungsten
Carbon	Iridium	Pnicogens	Uranium
Cerium	Iron	Potassium	Vanadium
Cesium	Krypton	Rhenium	Xenon
Chalcogens	Lanthanides	Rhodium	Ytterbium
Chlorine	Lanthanum	Rubidium	Yttrium
Chromium	Lead	Ruthenium	Zinc
Cobalt	Lithium	Samarium	Zirconium

Environmental and Atmospheric Chemistry

Anions	Fluorine	Oxidation	Radicals
Atmospheric chemistry	Gas-phase reactions	Ozone	Reaction mechanisms
Cations	Green chemistry	Peroxides	Reactive intermediates
Chlorine	Halogenation	Photochemistry	Sensors
Computer chemistry	Kinetics	Photolysis	Toxicology
Cycloaddition	Molecular dynamics	Photooxidation	Trace analysis
Denitrification	Molecular modeling	Radical ions	Waste prevention
Desulfurization	Nitrogen oxides	Radical reactions	Water chemistry
Environmental chemistry			

Inorganic Chemistry

Alanes	Cations	Inclusion compounds	Perovskite phases
Allotropy	Chain structures	Intercalations	Peroxides
Alloys	Chromates	Intermetallic phases	Phosphaalkenes
Aluminosilicates	Clathrates	Isoelectronic analogues	Phosphaalkynes
Amalgams	Cluster compounds	Isomers	Phosphanes
Amorphous materials	Cyanides	Layered compounds	Phosphazenes
Anions	Electron-deficient compounds	Lewis acids	Platinates
Automerization	Fluorides	Lewis bases	Pnictides
Autoxidation	Halides	Main group elements	Polyanions
Azides	High-pressure chemistry	Metal-metal interactions	Polyocations
Bond theory	Host-guest systems	Mixed-valent compounds	Polychalcogenides
Boranes	Hydrates	Nitrides	Polyhalides
Borates	Hydrides	Nonstoichiometric compounds	Polymorphism
Carbene homologues	Hydrothermal synthesis	Organic-inorganic hybrid composites	Polyoxometalates
Carbides	Hypervalent compounds		Radical ions
Carboranes			Radicals

Silanes
Silicates
Sol-gel processes
Solid-phase synthesis
Solid-state reactions
Solid-state structures

Spinel phases
Stannanes
Subvalent compounds
Synthesis design
Titanates

Topochemistry
Transition metals
Transuranium elements
Valence isomerization
Vanadates

Zeolite analogues
Zeolites
Zincates
Zintl anions
Zintl phases

Materials Science: General

Alloys
Amorphous materials
Automerization
Block copolymers
Ceramics
Charge carrier injection
Chemical vapor deposition
Chemical vapor transport
Clays
Cluster compounds
Colloids
Conducting materials
Copolymerization
Crystal engineering
Crystal growth

Cyclooligomerization
Cyclotrimerization
Dendrimers
Energy conversion
Fullerenes
Gels
Glasses
Holography
Imprinting
Intercalations
Interfaces
Intermetallic phases
Ladder polymers
Layered compounds

Liquid crystals
Materials science
Mechanical properties
Membranes
Mesophases
Mesoporous materials
Metal-metal interactions
Metallomesogens
Micelles
Microporous materials
Monolayers
Nanostructures
Nanotechnology
Nanotubes

Nonlinear optics
Polymerization
Polymers
Ring-opening polymerization
Scanning probe microscopy
Semiconductors
Sensitizers
Sensors
Superconductors
Surface chemistry
Thin films
Vesicles
Zeolite analogues
Zeolites

Miscellaneous

History of Science

Organic Chemistry: Compound Classes

Alcohols
Aldehydes
Alkaloids
Alkanes
Alkenes
Alkynes
Allenes
Allylic compounds
Amides
Amines
Amino acids
Amino alcohols
Amino aldehydes
Amphiphiles
Anhydrides
Anions
Annulenes
Arenes
Arynes
Azides
Azo compounds
Azomethine ylides
Betaines

Biaryls
Calixarenes
Carbanions
Carbenes
Carbenoids
Carbocations
Carbocycles
Carbohydrates
Carboxylic acids
Carotenoids
Catenanes
Cations
Cavitands
Crown compounds
Cryptands
Cumulenes
Cyanides
Cyanines
Cyclodextrins
Cyclophanes
Dendrimers
Diazo compounds
Dyes/Pigments

Enols
Enones
Enynes
Fatty acids
Fragrances
Fullerenes
Fused-ring systems
Heterocycles
Hydrazones
Hydrides
Hydrocarbons
Ketones
Lactams
Lactones
Ladder polymers
Macrocycles
Mannich bases
Medium-ring compounds
Metallacycles
Natural products
Nitrogen heterocycles
Oxygen heterocycles
Peroxides

Pheromones
Phosphorus heterocycles
Phthalocyanines
Polycycles
Polymethines
Porphyrinoids
Quinodimethanes
Quinones
Radical ions
Radicals
Rotaxanes
Schiff bases
Small ring systems
Spiro compounds
Steroids
Sulfonamides
Sulfur heterocycles
Surfactants
Terpenoids
Ylides
Zwitterions

Organic Chemistry: Methodology and Reactions

Acylation
Aldol reaction
Alkylation
Allylation
Amination

Annulation
Aromatic substitution
Aromaticity
Asymmetric amplification
Asymmetric catalysis

Asymmetric synthesis
Automerization
Autoxidation
Biomimetic synthesis
C-C activation

C-C coupling
C-H activation
C1 building blocks
Carbonylation
Carboxylation

Chiral auxiliaries	Glycosylation	Michael addition	Regioselectivity
Chiral pool	Grignard reaction	Molecular diversity	Retro reactions
Cleavage reactions	Halogenation	Multicomponent reactions	Ring contraction
Combinatorial chemistry	Heck reaction	Nucleophilic addition	Ring expansion
Cracking	High-pressure chemistry	Nucleophilic substitution	Sigmatropic rearrangement
Cross-coupling	Hydroamination	Olefination	Solid-phase synthesis
Cyclization	Hydroboration	Oligomerization	Solvent effects
Cycloaddition	Hydroformylation	Oxidation	Solvolysis
Cyclotrimerization	Hydrogen transfer	Oxygenation	Steric hindrance
Dehydrogenation	Hydrogenation	Ozonolysis	Substituent effects
Dihydroxylation	Hydrolysis	Perfluorinated solvents	Synthesis design
Dimerization	Hydrosilylation	Pericyclic reaction	Synthetic methods
Domino reactions	Hydrostannylation	Phosphorylation	Template synthesis
Electrocyclic reactions	Hydroxylation	Photooxidation	Topochemistry
Electrophilic addition	Immobilization	Polymerization	Total synthesis
Electrophilic substitution	Insertion	Protecting groups	Transesterification
Elimination	Ionic liquids	Protonation	Umpolung
Ene reaction	Isomerization	Radical reactions	Wittig reactions
Epoxidation	Lithiation	Rearrangement	
Flash pyrolysis	Metalation	Reduction	

Organic Chemistry: Stereochemistry and Structures

Atropisomerism	Configuration determination	Enantioselectivity	Structure elucidation
Chemoselectivity	Conformation analysis	Hyperconjugation	Tautomerism
Chiral resolution	Conjugation	Kinetic resolution	Valence isomerization
Chirality	Diastereoselectivity	Strained molecules	

Physical Chemistry and Chemical Physics (including Electrochemistry, Kinetics, Photochemistry, Radiochemistry, Thermodynamics and Theoretical Chemistry)

Ab initio calculations	Electron transfer	Laser chemistry	Plasma chemistry
Absorption	ELF (Electron Localization Function)	Lewis acids	Radiochemistry
Acidity	Energy conversion	Lewis bases	Radiopharmaceuticals
Adsorption	Femtochemistry	Linear free energy relationships	Reaction mechanisms
Basicity	Fluorescence	Liquid crystals	Reactive intermediates
Bond energy	Fluorescent probes	Liquids	Redox chemistry
Bond theory	Fractals	Magnetic properties (including magnetochemistry)	Salt effect
Calorimetry	FRET (fluorescence resonant energy transfer)	Matrix isolation	Semiempirical calculations
CARS (Coherent Anti-Stokes Raman Scattering)	Gas-phase reactions	Mesophases	Single-molecule studies
Charge carrier injection	Gels	Metallomesogens	Singlet oxygen
Chemisorption	Glasses	Metastable compounds	Sol-gel processes
Chromophores	Heats of formation	Microrreactors	Solvatochromism
Colloids	High-pressure chemistry	Molecular dynamics	Spin crossover
Computer chemistry	High-temperature chemistry	Molecular modeling	Statistical mechanics
Conducting materials	Hot-atom chemistry	Monolayers	Statistical thermodynamics
Conical intersections	Hydrophobic effect	Nanotechnology	Structure-activity relationships
Crystal engineering	Imaging agents	Neighboring-group effects	Superacidic systems
Crystal growth	Ion pairs	Phase diagrams	Supercritical fluids
Cyclic voltammetry	Ion-molecule reactions	Phase transitions	Thermochemistry
Density functional calculations	Ionization potentials	Photochemistry	Thermodynamics
Donor-acceptor systems	Isotope effects	Photochromism	Time-resolved spectroscopy
Electrochemistry	Isotopes	Photolysis	Transition states
Electron microscopy	Kinetics	Physisorption	Voltammetry

Supramolecular Chemistry

Aggregation	Molecular evolution	Pi interactions	Self-assembly
Host-guest systems	Molecular recognition	Receptors	Supramolecular chemistry
Molecular devices	Nanostructures		