

The Mitsunobu Reaction

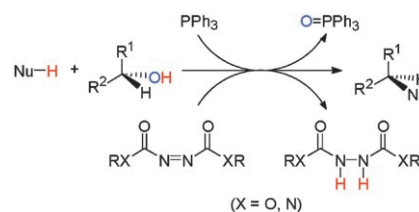
T. Y. S. But, P. H. Toy*

The Mitsunobu Reaction: Origin, Mechanism, Improvements, and Applications

Chem. Asian J.

DOI: 10.1002/asia.200700182

Classics never fade away: The Mitsunobu reaction is a widely used and versatile method for the dehydrative oxidation–reduction condensation of an acid/pronucleophile with an alcohol mediated by phosphine and azo reagents. The history, mechanism, and recent developments of this stereoselective workhorse condensation reaction are reviewed.



Photoactivation

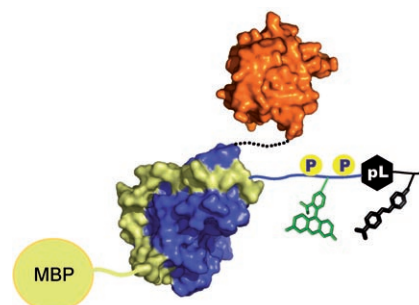
M. E. Hahn, J.-P. Pellois,
M. Vila-Perelló, T. W. Muir*

Tunable Photoactivation of a Post-translationally Modified Signaling Protein and its Unmodified Counterpart in Live Cells

ChemBioChem

DOI: 10.1002/cbic.200700404

Strike a pose... Expressed protein ligation was used to prepare caged analogues of the signaling protein Smad2 (see illustration). The function and fluorescence of the analogues could be photocontrolled in a correlated fashion. This strategy permitted the titration of the cellular levels of active phosphorylated Smad2 in its biologically relevant, full-length form.



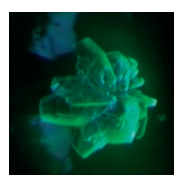
Lanthanide Luminescence

A. D'Aléo, G. Pompidor, B. Elena,
J. Vicat, P. L. Baldeck, L. Toupet,
R. Kahn, C. Andraud,* O. Maury*

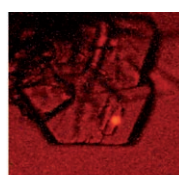
Two-Photon Microscopy and Spectroscopy of Lanthanide Bioprobes

ChemPhysChem

DOI: 10.1002/cphc.200700375



Single-photon



Two-photon
Microscopy

Lanthanide bioprobes: The first two-photon microscopy imaging experiments using tris-dipicolinate terbium complexes as a probe were carried out on derivative protein crystals. Whereas one-photon irradiation results in green luminescence of the entire crystal aggregate, two-photon excitation gives a three-dimensionally resolved spot corresponding to the confocal volume (see picture).

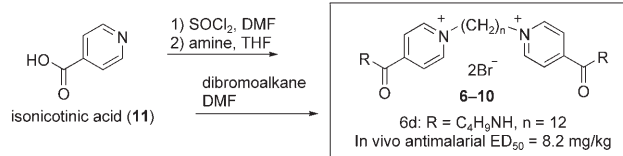
Synthetic Methods

K. Motoshima, Y. Hiwasa,
M. Yoshikawa, K. Fujimoto, A. Tai,
H. Kakuta,* K. Sasaki

Antimalarial Cation-dimers Synthesized in Two Steps from an Inexpensive Starting Material, Isonicotinic Acid

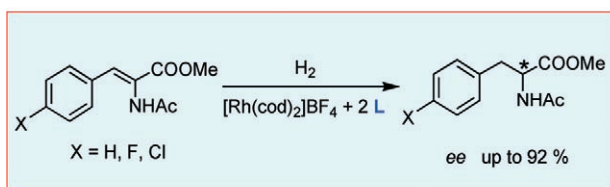
ChemMedChem

DOI: 10.1002/cmdc.200700107



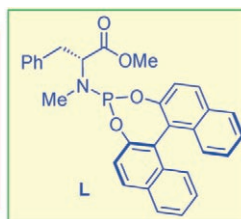
Managing malaria. As the area affected by malaria includes a large proportion of developing countries, there is a need for new antimalarials that can be synthesized and supplied inexpensively. In this

study, bis-cation dimers, MAP series 6–10 synthesized from an inexpensive isonicotinic acid (11) in just two steps, were designed. MAP-412 (6d) exhibited a potent in vivo antimalarial activity.



A series of chiral monodentate ligands combining a 3,5-dioxa-4-phosphacycloheptadinaphthyl unit either with a phenylalanine- or with an alanine-derived

fragment were synthesised and tested in the hydrogenation of 2-(acetylamino)-3-(aryl)propenoic methyl esters.



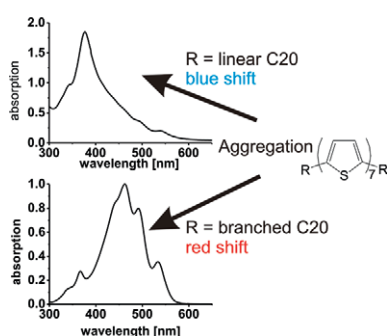
Asymmetric Catalysis

L. Eberhardt, D. Armspach,* D. Matt,*
L. Toupet, B. Oswald

Synthesis of Chiral, Monodentate Aminophosphane and Phosphoramidite Ligands Derived from Amino Acid Esters: Application in Rh-Catalysed Asymmetric Olefin Hydrogenation Reactions

Eur. J. Inorg. Chem.

DOI: [10.1002/ejic.200700474](https://doi.org/10.1002/ejic.200700474)



α - and α,ω -substituted oligothiophenes show a strong aggregation behaviour in solution depending on the geometry of the alkyl substituents which is reflected by a distinct blue or red shift, respectively, in the absorption spectra. A strong influence of the substituent is also found in the morphology of the aggregates and of thermal properties in the solid state.

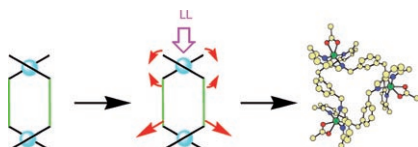
Aggregation of Oligothiophenes

S. Ellinger, A. Kreyes, U. Ziener,*
C. Hoffmann-Richter, K. Landfester,
M. Möller

Aggregation Phenomena of Long α - and α,ω -Substituted Oligothiophenes – the Effect of Branched vs. Linear End-Groups

Eur. J. Org. Chem.

DOI: [10.1002/ejoc.200700566](https://doi.org/10.1002/ejoc.200700566)



Taken in with open arms! Coordination chemistry is used to destabilise a linear array and turn it into a circular helicate (see figure). Changes in the geometry of the metal centre allow the product to be tailored to form dimers or circular helicates.

Supramolecular Chemistry

J. Hamblin, F. Tuna, S. Bunce,
L. J. Childs, A. Jackson, W. Errington,
N. W. Alcock, H. Nierengarten,
A. Van Dorsselaer, E. Leize-Wagner,
M. J. Hannon*

Supramolecular Circular Helicates
Formed by Destabilisation of
Supramolecular Dimers

Chem. Eur. J.

DOI: [10.1002/chem.200700848](https://doi.org/10.1002/chem.200700848)



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