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Review article

Recent advances in click chemistry reactions mediated by transition metal based systems

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ABSTRACT

The recent advances of click chemistry reactions have shown that catalysts based on Cu(I) ions and nanoparticles have the potential to mediate and activate the alkyne-azide cycloaddition reactions in mild conditions and with remarkable yields. This review is focused on the recent progress recorded, over the last five years, by the transition metal catalysts in the development of novel methods and applications in the field of click chemistry reactions. Their catalytic efficiency, along with their different applications in materials, drug discovery and biological environments are also presented.

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Aurel Tăbăcaru (born in 1984 in Galați, Romania) received his B.Sc. degree in 2008 and Master degree in 2010 from "Dunărea de Jos" University of Galați. In 2013, he obtained his PhD in Chemistry from the University of Camerino (Italy), working in the field of coordination polymers and metal-organic frameworks based on poly(pyrazole)-type ligands, under the supervision of Prof. Claudio Pettinari. His research career was continued with a postdoctoral fellowship at "Dunărea de Jos" University of Galați, working in the field of Nanotechnology, under the supervision of Prof. Viorica Muşat. In 2014, he joined the Department of Chemistry, Physics and Environment, from "Dunărea de Jos" University of Galați, as Teaching Assistant. His research is focused on the synthesis and study of coordination compounds (with porous, luminescent, catalytic, and antibacterial properties) and surface-modified metal oxide nanoparticles for optoelectronic applications.

Abbreviations: AOC, azide-olefin cycloaddition; APSiO₂, 3-aminopropyl-functionalized silica; AscNa, sodium ascorbate; aterpy, 4'-azido-2,2':6',2"-terpyridine; bipy, 4,4'bipyridine; BMIM BF4, 1-butyl-3-methylimidazolium tetrafluoroborate; bpoh, (N1E,N2E)-N1,N2-bis(phenyl(pyridin-2-yl)methylene)oxalohydrazide; BTRB, BisTriazole-Rhodamine B; CA153, cancer antigen 153; CL, ε-caprolactone; CH₂Cl₂, dichloromethane; CPMV, cowpea mosaic virus; CuAAC, copper-catalyzed azide-alkyne cycloaddition; CV, crystal violet; DIBO, dibenzocyclooctyne; DO3A, 1,4,7-tris(carboxymethyl)-1,4,7,10-tetraazacyclodecane; DMF, dimethylformamide; DOTA, 1,4,7,10-tetrakis(carboxymethyl)-1,4,7,10-tetrabacyclodecane; DMF, dimethylformamide; DOTA, 1,4,7,10-tetrakis(carboxymethyl)-1,4,7,10-tetrabacyclodecane; DMF, dimethylformamide; DOTA, 1,4,7,10-tetrabacyclodecane; DMF, dimethylformamide; DMF, boxymethy1)-1,4,7,10-tetraazacyclododecane; DMSO, dimethylsulfoxide; DTT, dithiothreitol; 6eBMPA, 6-ethynyl-bis(methylpyridyl)amine; 18F-ArBF₃-, aryltrifluoroborate; Gly, glycine; Hacac, acetylacetone; FcMeTrz-6eBMPA, triazolyl-6-ethynyl-bis(methylpyridyl)amine bearing the ferocenylmethyl moiety; GC-MS, gas chromatographic mass spectrometry; GO, graphene oxide; GTCF, graphene transparent conductive film; H₃Eshi, 4-ethynylsalicylhydroxamic acid; Himdc, 4,5-imidazoledicarboxylate; ¹H NMR, proton nuclear magnetic resonance; H₃TTshi, 4-(2,2':6',2"-terpyridyl-1H-[1,2,3]triazol-4-yl)salicylhydroxamic acid; Ln, lanthanides; 12-MC-4, 12-metallacrown-4; MCH, 6-mercapto-1-hexanol; MNPs, magnetic nanoparticles; MOFs, metal-organic frameworks; MSN, magnetic silica nanoparticles; MVS, metal vapor synthesis; NaN3, sodium azide; NHC, N-heterocyclic carbene; NMe4, tetramethylammonium ion; oda, oxodiacetate; NPs, nanoparticles; PEG, polyethyleneglycol; PEI, polyethyleneimine; PHEMA, poly (2-hydroxyethyl methacrylate); PhPrTrz-6eBMPA, triazolyl-6-ethynyl-bis(methylpyridyl)amine bearing the phenylpropyl moiety; PS-b-PCL, polystyrene-blockpoly(ε-caprolactone); PS-OH, hydroxyl-modified polystyrene; PS-N₃, azide-modified polystyrene; PS-C22-Cul, polystyrene resin supported copper(I) iodide-cryptand-22 complex; PPh₃, triphenylphosphine; PPi, pyrophosphate ion; PVC-N₃, azide-modified polyvinyl chloride; PVBC, polyvinylbenzyl chloride; PVBAz, polyvinylbenzylazide; RB, glycosyl rhodamine B; ROS, reactive oxygen species; RuAAC, ruthenium-catalyzed azide-alkyne cycloaddition; SAMs, self-assembled monolayers; siRNA, small interfering ribonucleic acid; SMMs, single magnetic molecules; TBTA, tris(benzyltriazolylmethyl)amine; Thi-Fc, ferrocene-appended thiophene; tiRNA, transcription initiation ribonucleic acid; TRGO, thermally reduced graphene oxide; TMV, tobacco mosaic virus; THPTA, tris(hydroxypropyltriazolyl)methylamine; VLP, virus-like particles.

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Rodica Mihaela Dinică (born in 1968 in Tecuci, Romania) received her B.Sc. degree in 1991 from "Alexandru Ioan Cuza" University of Iași (Romania). In 1999, she obtained her PhD in Chemistry from the same university, working in the field of N-heterocyclic chemistry. In 1999, she had a THEMPUS – PHARE fellowship at the Department of Chemical Sciences, University of Camerino (Italy), under the supervision of Prof. Claudio Pettinari. Her research career was continued in 2001–2002 with a postdoctoral fellowship at "Joseph Fourier" University of Grenoble, France (Laboratoire d'Etudes Dynamiques et Structurales de la Selectivite), working in the field of acridinic antitumoral compounds, under the supervision of Dr. Martine Demeunynck. From 1995, she joined the Department of Chemistry, Physics and Environment from "Dunărea de Jos" University of Galați. Since 2014, she is full professor and in 2016 she obtained the Habilitation qualification. Her research is focused on the synthesis and study of N-heterocyclic compounds with biologic activities, using click chemistry reactions, biocatalysts, ultrasound and microwave techniques.

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