Open Archive TOULOUSE Archive Ouverte (OATAO)

OATAO is an open access repository that collects the work of Toulouse researchers and makes it freely available over the web where possible.

This is an author-deposited version published in: http://oatao.univ-toulouse.fr/
Eprints ID: 14275

To link to this article: DOI:10.1016/j.polymer.2015.02.024
URL: http://dx.doi.org/10.1016/j.polymer.2015.02.024

To cite this version: Chen, Si and Cardozo, Andrés Fernando and Julcour, Carine and Blanco, Jean-François and Barthe, Laurie and Gayet, Florence and Lansalot, Muriel and D'Agosto, Franck and Delmas, Henri and Manoury, Eric and Poli, Rinaldo Amphiphilic core-cross-linked micelles functionalized with bis(4-methoxyphenyl)phenylphosphine as catalytic nanoreactors for biphasic hydroformylation. (2015) Polymer, vol. 72. pp. 327-335. ISSN 0032-3861

Any correspondance concerning this service should be sent to the repository administrator: staff-oatao@listes-diff.inp-toulouse.fr
Amphiphilic core-cross-linked micelles functionalized with bis(4-methoxyphenyl)phenylphosphine as catalytic nanoreactors for biphasic hydroformylation

Si Chen a,c, Andrés F. Cardozo b,c, Carine Julcour b,c, Jean-François Blanco b,c, Laurie Barthe b,c, Florence Gayet a,c, Muriel Lansalot d, Franck D’Agosto d, Henri Delmas b,c, Eric Manoury a,c,* Rinaldo Poli a,c,e,*

a CNRS, LCC (Laboratoire de Chimie de Coordination), 205 route de Narbonne, BP 44099, F-31077, Toulouse Cedex 4, France
b CNRS, LGC (Laboratoire de Génie Chimique), 4 Allée Emile Monso, BP 64234, 31030 Toulouse Cedex 4, France
c Université de Toulouse, UPS, INPT, F-31077, Toulouse Cedex 4, France
d Université de Lyon, Univ. Lyon 1, OPE Lyon, CNRS, UMR 5265, C2P2 (Chemistry, Catalysis, Polymers & Processes), Team LCPP Bat 308 F, 43 Bd du 11 Novembre 1918, 69616, Villeurbanne, France
e Institut Universitaire de France, 101, bd Saint-Michel, 75005, Paris, France

Abstract

Core-cross-linked micelles (CCM) functionalized at the core with covalently linked bis(p-methoxyphenyl)phenylphosphine (BMOPPP) ligands have been synthesized by a three-step one-pot radical polymerization in emulsion, using the polymerization-induced self-assembly (PISA) strategy and reversible addition-fragmentation chain transfer (RAFT) as the controlling method. The CCM are obtained by chain extending in water poly(methacrylic acid-co-polyethylene oxide methyl ether methacrylate) (P(MAA-co-PEOMA), degree of polymerization of 30, MAA/PEOMA units molar ratio of 50:50) synthesized in a first step by RAFT with a 95:5 M mixture of styrene and 4-[bis(p-methoxyphenyl)phosphino]styrene (BMOPPS) units. The resulting micelles exhibiting a core composed of P(S-co-BMOPPS) segments with a degree of polymerization of 300 are then crosslinked in a third step with a mixture of di(ethylene glycol) dimethacrylate (DEGDMA) and styrene. The resulting BMOPPP@CCM exhibit a narrow size distribution (PDI = 0.16) with an average diameter of 81 nm in water and swell in THF or by addition of toluene to the latex. The addition of [Rh(acac)(CO)2] to the toluene-swollen latex results in metal coordination to the phosphine ligands. 31P{1H} NMR spectroscopy shows that the Rh centers undergo rapid intraparticle phosphine ligand exchange. Application of these nanoreactors to the aqueous biphasic hydroformylation of 1-octene shows excellent activity and moderate catalyst leaching.

Keywords:
Catalytic nanoreactor
Biphasic hydroformylation catalysis
Core-cross-linked micelle (CCM)

1. Introduction

The power of controlled radical polymerization and the wide choice of monomers and polymerization mechanisms have made it possible to design and synthesize quite complex functionalized macromolecular architectures that were unimaginable only 20 years ago [1]. Among many possible uses of functional polymers, the area of nanoreactors for catalytic applications has developed rather recently but is now rapidly expanding [2–4]. By careful design, it is now possible to generate macromolecular architectures where the catalyst is confined in a specific part of the macromolecule conferring specific properties to the catalyst environment (affinity for the reaction substrate, size selectivity, site confinement, etc.) while other parts of the macromolecule are responsible for the nanoreactor compatibility with its environment (homogeneous dispersion, response to stimuli such as heat, pH, magnetic fields etc.). As notable examples, Fréchet et al. have shown the principle of catalyst site confinement for a cascade organic transformation involving one acid-catalyzed and one base-catalyzed step [5]. O’Reilly et al. have turned self-assembled catalytic micelles into unimolecular polymer particles by shell-cross-linking [6].
Sawamoto et al. have built thermoresponsive phosphine-containing microgels and applied them to a few transformations under aqueous biphasic conditions [7,8], and Resmini et al. have used the principle of molecular imprinting to develop shape-selective catalytic nanogels [9,10]. In most cases, the developed nanoreactors were applied to catalytic transformations under homogeneous conditions. Occasionally, the catalyst was recovered and recycled by ultrafiltration, by precipitation, or by thermoregulation.

Efficient catalyst recovery and recycling is of great interest in industrial homogeneous catalysis, particularly when using expensive metals and/or ligand systems, the aqueous biphasic approach being the most attractive one because of the simplicity of the necessary equipment and operating procedures [11]. The prime example of successful industrial application is the Rhone-Poulenc/Ruhrlchemie hydroformylation of propene [12], where the Rh catalyst is completely confined in the aqueous phase by triphenylphosphine trisulfonate (TPPTS), but this process is unfortunately inefficient for the higher olefins because of their insufficient water solubility. Among the investigated strategies for circumventing this problem, micellar catalysis is the most attractive one.

Core-functionalized micelles, resulting from the self-assembly of surfactants or amphiphilic diblock copolymers where the catalyst is anchored to the hydrophobic part, are the simplest possible type of nanoreactor and many applications of biphatic catalysis have been described [13–18]. The dynamic nature of micelles, however, results in two major obstacles to large scale industrial implementation: uncontrolled swelling leading to the formation of stable emulsions and loss of the free surfactants, even when the critical micelle concentration (CMC) is very low [19–24]. For instance, anchoring of a rhodium complex to a poly(norbornene)-based amphiphilic diblock copolymer with CMC = 2.2 × 10⁻⁶ M and application to the aqueous biphasic 1-octene hydroformylation led to excellent turn over frequency (TOF) but also to leaching with 9 ppm of Rh detected in the organic product phase [25], which is an intolerably high loss for large scale production.

In order to remove both problems, we have recently introduced a new approach, which consists of cross-linking amphiphilic block copolymer micelles at the core to generate unimolecular nanooobjets [26,27]. These core-cross-linked micelles (CCM) have been assembled by an efficient one-pot procedure by a polymerization-induced self-assembly (PISA) in water [28,29], using reversibladdition-fragmentation chain transfer (RAFT) polymerization in a convergent approach. The ligand needed to bind the catalytic metal was incorporated statistically as a phosphine functionalized conomonmer, 4-diphenylphosphinostyrene (DPPS), at low loading (5–25% molar) in the hydrophobic core. This monomer carries a triphenylphosphine (TPP) ligand for metal coordination. The resulting TPP@CCM latex could readily be charged with the hydroformylation precatalyst, [Rh(acac) (CO)]²⁺, and the resulting catalytic nanoreactors, [Rh(acac) (CO)]²⁺ (TPP)@CCM, were successfufully used in the aqueous biphasic hydroformylation of 1-octene with excellent activity, catalyst recyclability, and low metal leaching (down to 1.8 ppm) [26].

On the basis of this initial success, we have embarked in a more systematic study of the CCM approach to aqueous biphasic catalysis. Initial questions concerned the modification of the CCM structure in terms of catalyst density and size of the hydrophobic core and the hydrophilic shell (degree of polymerization) and how such changes affect the catalytic performance [27]. Another point of interest is access to CCMs with other ligand functionalities in the hydrophobic core for wider applications in catalysis. In this contribution, we report the synthesis of a CCM functionalized with the bis(4-methoxyphenyl)phosphine (BMOPPP) ligand, BMOPPP@CCM, its physical characterization, its coordination chemistry with [Rh(acac) (CO)]²⁺, and the application of the [Rh(acac) (CO)]²⁺-loadened nanoreactors, [Rh(acac) (CO)]²⁺ (BMOPPP)@CCM, to aqueous biphasic hydroformylation. This specific ligand was chosen for the first CCM chemical modification because of its structural similarity with TPP and consequently of the expected similar polymerization behavior, relative to DPPS, of the suitable ligand-functionalized monomer, 4-[bis(4'-methoxyphenyl)phosphino]stylene (BMOPPS). While the Rh activity in the presence of p-OMe-substituted arylphosphines is very similar to that in the presence of the unsubstituted analogs for hydroformylations conducted with soluble complexes under homogeneous or biphasic conditions [30,31], the p-OMe substitution increases the phosphine binding ability toward Rh [32]. Therefore, lower leaching may be expected if the leaching mechanism involves loss of metal from the nanoreactor core.

2. Experimental

2.1. Materials

All manipulations were performed under an inert atmosphere of dry argon by using Schlenk line techniques. 4,4’-azobis(4-cyanopentanoic acid) (ACPA, >98%, Fluka), methacrylic acid (MAA, 99.5%, Acros), poly(ethylene oxide) methyl ether methacrylate (PEOEMA, Mn = 950 g mol⁻¹, Aldrich), di(ethylene glycol) dimethacrylate (DEGDMA, 95%, Aldrich), 1,3,5-trioxane (Aldrich, >99%), acrylamidododecyldecylrhodium(I), ([Rh(acac) (CO)]²⁺), 99% Strem), chloro(1,5-cyclooctadiene) rhodium(I) dimer ([Rh(COD) Cl]², >98%, Strem), triphenylphosphine (PPh₃ or TPP, >98.5%, Fluka), chlorobis(4-methoxyphenyl) phosphine, (>98%, Alfa), and 4-bromostyrene (98%, stab. with 0.1% 4-tert-butylicatetole, Alfa) were used as received. Styrene (5%, 99%, Acros) was purified by passing through a column of active basic aluminum oxide to remove the stabilizer. The RAFT agent 4-cyano-4-thiothiopropylsulfanyl pentanoic acid (CTPPA) was synthesized as described previously [33]. Mg turnings was washed with HCl (1 M) until the metallic color appearing then washed with diethylether. Solvents were dried by standard procedures and distilled under argon prior to use.

2.2. Characterization techniques

2.2.1. Nuclear magnetic resonance

¹H NMR and ³¹P NMR spectra were recorded in 5 mm diameter tubes at 297 K in D₂O, DMSO-d₄ or THF-d₈ solution (the polymerization medium aliquots were directly dispersed in the solvent) using a Bruker Avance 400 spectrometer. ¹H and ¹³C chemical shifts were determined using the residual peak of deuterated solvent as internal standard and are reported in ppm (δ) relative to tetramethylsilane. ³¹P chemical shifts are reported relative to external 85%H₃PO₄. The solid-state ¹³C NMR experiment was recorded on a Bruker Avance 400 spectrometer equipped with a 3.2 mm probe. The sample was spun at 16 kHz at the magic angle using ZrO₂ rotors, using a small flip angle (~30°) with a recycle delay of 5 s and a contact time of 2 ms. Peaks are labeled as singlet (s), doublet (d), triplet (t), multiplet (m) and broad (br). The aromatic C positions are labeled as C₁ (ipso, P bonded), C₀ (ortho), C₉ (meta) and C₈ (para, bonded to OMe or to CH=CH₂). For the CCM characterization, the chemical shift scale was calibrated on the basis of the solvent peak (δ 2.50 for DMSO, 3.58 and 1.73 for THF), and 1,3,5-trioxane was used as an integration reference (δ 5.20).

2.2.2. Size exclusion chromatography

Size exclusion chromatography (SEC) measurements were performed in THF (with butylhydroxytoluene (BHT) as a flow rate
2.2.3. Dynamic light scattering
The intensity-average diameters of the latex particles (Dₐ) and the dispersity factor (PDI, polydispersity index) were measured at 25 °C on a Malvern Zetasizer NanoZS. After filtration through a 0.45 μm pore-size membrane, deionized water or THF was used to dilute the latex sample. Solutions were analyzed without further filtration to ensure that undesired populations were not removed. Data were analyzed by the general-purpose non-negative least squares (NNLS) method. The typical accuracy for these measurements was 10–15%.

2.2.4. Transmission electron microscopy
The morphological analysis of the copolymer nano-objects was performed with a JEOL JEM 1011 transmission electron microscope equipped with 100 kV voltage acceleration and tungsten filament (Service Commun de Microscopie Electronique TEMSCAN, plateforme de l’Université Paul Sabatier, Toulouse, France). Diluted latex samples were dropped on a formvar/carbon-coated copper grid and dried under vacuum.

2.2.5. Mass spectrometry
The mass spectral analyses were performed with a high resolution electrospray XevoG2QT Waters instrument by Service Commun de Spectrométrie de Masse of the Université Paul Sabatier, Toulouse, France.

2.3. Synthesis of 4-[bis(4'-methoxyphenyl)phosphino]styrene (BMOPPS)
Chlorobis[4-methoxyphenyl]phosphine (1 g, 3.56 mmol) was added slowly at 0 °C to a solution of the Grignard reagent prepared from 4-bromostyrene (0.65 g, 3.56 mmol) and Mg (0.104 g, 4.275 mmol) in dry THF (4 mL). After the addition was complete, the reaction mixture was stirred at room temperature for 3 h. The reaction mixture was then diluted with ethyl acetate (30 mL) and washed sequentially with water (2 × 50 mL), 10% aqueous HCl (2 × 50 mL), saturated aqueous NaHCO₃ (2 × 50 mL), and brine (2 × 50 mL). The organic layer was dried over MgSO₄, filtered, and concentrated in vacuum. The crude product was purified by silica gel chromatography (5% EtO/hexane, then EtO) to afford 1 as a white solid (0.66 g, 53%). ¹H NMR (400 MHz, CDCl₃): δ 7.39–7.22 (m, 8H, CH₃), 6.93–6.90 (m, 4H, CH₂), 6.72 (dd, J = 10.9 Hz, 1H, J = 17.6 Hz, CH₂), 5.78 (d, J = 17.6 Hz, CH₂), 5.29 (d, J = 10.9 Hz, CH₂), 3.83 (s, 6H, CH₃ (OMe)), 3.15 (s, 1H, CH₂), 2.31 (s, 1H, CH₂). ¹³C¹H NMR (101.5 MHz, CDCl₃): δ 160.3 (s, C₅(OMe)), 138.2 (d, J = 101.0 Hz, C₁, 1C), 137.5 (s, 1C, C₆(CH₂)), 136.5 (s, 1H, CH = CH₂), 135.3 (d, J = 19.3 Hz, C₉(OMe)), 133.3 (d, J = 19.3 Hz, C₈(CH₂)), 128.2 (d, J = 81.0 Hz, C₂, 2C), 126.2 (d, J = 61.0 Hz, C₅(CH₂)), 114.4 (s, CH = CH₂), 114.2 (d, J = 81.0 Hz, C₉(OMe)), 55.2 (s, OCH₃). HR El-MAL spectral values were 348.1356 and 348.1348. M.p.: 123 °C. Elemental analysis for C₂₉H₂₁O₂P: % C 74.38, H 5.99, found % C 74.57, H 5.51.

2.4. Preparation of the BMOPP@CCM latex by one-pot RAFT polymerization in water

2.4.1. Step 1: preparation of the P(MAA-co-PEOMA)-TTC macromolecular RAFT agent (macroRAFT) in water
A stock solution containing ACPA (30 mg mL⁻¹) in deionized water (1 mL) containing also NaHCO₃ (30 mg) was prepared. 100 μL of this stock solution (3 mg ACPA, 0.0108 mmol), 15 mg of CTPPA (0.054 mmol), 75 mg of MAA (0.87 mmol), 0.79 g of PEOMA (0.83 mmol) and 4.2 g of deionized water (including the water amount of the ACPA solution) were added into a 25 mL flask equipped with a magnetic stirrer bar, which was then sealed with a rubber septum. 1,3,5-trioxane was also added into the flask as an internal reference for the determination of the monomer conversion by ¹H NMR. The solution was purged for 45 min with argon and then heated to 80 °C in a thermostated oil bath with stirring. After 120 min, 0.15 mL of solution was taken to determine the monomer conversion and the molar mass of the macroRAFT product. The overall monomer molar conversion was about 98% as determined by ¹H NMR spectroscopy in DMSO-D₆. The molar mass was analyzed by size exclusion chromatography (SEC) in THF (experimental M₀ = 11,200 g mol⁻¹; D ≈ 1.19).

2.4.2. Step 2: chain extension of the macroRAFT with S and BMOPPS in water
During Step 1, a suspension containing 1.57 g of S (15 mmol), 0.269 g of BMOPPS (0.772 mmol, 5% mol relative to S) in 4.62 g of deionized water, to which was also added the ACPA stock solution (100 μL containing 3 mg of ACPA, 0.0108 mmol) were purged separately for 45 min with an argon stream at 0 °C. This mixture was quickly injected into the first flask under argon at 80 °C after the end of Step 1. The separated organic/aqueous phases became one opaque suspension phase at around 1 h. After 2 h of further stirring for the resulting suspension, a 0.5 mL sample was withdrawn for analysis and the polymerization was quenched by immersion of the flask in iced water. The overall conversion of S (94%) was determined by gravimetric analysis and that of BMOPPS (100%) was measured by ³¹P NMR in THF-d₈. SEC: M₀ = 43,900 g mol⁻¹ (D = 1.3), D₀ (DLS) = 78 nm (D = 0.2).

2.4.3. Step 3: cross-linking
In the flask containing the P(MAA-co-PEOMA)-b-(P’S-co-BMOPPS) latex from Step 2, 0.48 g of S (4.6 mmol), 0.137 g of DEG DMA (0.565 mmol, 10% mol relative to S), 90 μL of the ACPA stock solution (2.7 mg of ACPA, 0.009 mmol) and 2.43 g of deionized water were further added. The mixture was purged for 1 h with argon at 0 °C, and the flask was then placed in an oil bath thermostated at 80 °C. After 90 min, the polymerization was quenched by immersion of the flask in iced water. The overall conversion of the comonomers (97%) was determined by ¹H NMR and ³¹P NMR in THF-d₈. DLS (H₂O): D₂ = 81 nm (PDI = 0.16). DLS (THF): D₀ = 207 nm (PDI = 0.20).

2.5. Metal complexation to the phosphine ligand within the nanoparticle core
[Rh(acac)₂(CO)₃] (31.7 mg, 101 eq) in toluene (1 mL) was added to a previously swollen nanoparticles latex (5 mL, 0.1 mg mL⁻¹) prepared from the BMOPP@CCM latex (2 mL) diluted in D₂O (3 mL) by addition of toluene (0.5 mL). The swelling was very rapid (<1 min upon stirring at room temperature) as confirmed by visual disappearance of the toluene phase and by the ³¹P NMR observation of the core phosphine resonance (see Results and Discussion). The reaction mixture was stirred at room temperature for 10 min until the latex color changed to yellow. The resulting latex was...
washed by toluene (2 × 1 mL) under argon to remove any excess of the Rh precursor; both toluene washings were colorless. The [Rh(acac)(CO)2] (BMOPPP)@CCM latex was collected after decantation for further NMR studies. 31P{1H} NMR (162 MHz, CDCl3, 298 K): δ 44.5 (d, J = 176 Hz) for the polymer linked [Rh(acac)(CO)2] (BMOPPP) complexes.

2.6 Biphasic hydroformylation catalysis

The catalytic aqueous phase was prepared under a nitrogen atmosphere by diluting the CCM latex in Milli-Q water (for a phosphine equivalent of 0.65 mmol in 25 mL), then swelling the hydrophobic nanoparticle core with 3 mL of decanal, prior to the addition of the Rh precursor, [Rh(acac)(CO)2] (43 mg, 0.163 mmol), dissolved in 3 mL of decanal. At each step, the resulting mixture was vigorously stirred for a few minutes. This colloidal suspension was poured into the autoclave, then immediately covered by additional decanal (56 mL) and by 1-octene (81.6 mmol). The reactor was flushed three times with 15 bar of nitrogen, then four times with 15 bar of syngas. It was subsequently heated under low syngas pressure (2 bar) and slow stirring speed (300 rpm, well below gas self-induction) to generate the catalytic species in situ, albeit hindering the start of the reaction. When the desired reaction temperature (363 K) was achieved (after about half an hour), stirring was stopped and the autoclave was pressurized and constantly fed with syngas at the desired pressure (20 bar). A sample was withdrawn to evaluate the amount of products formed during the heating period.

Then, the data acquisition was started and the stirring speed was set to 1200 rpm. Both temperature and pressure of the reactor and the gas ballast were recorded on-line, in order to measure the instantaneous syngas consumption. After a few hours of reaction, a final sample of the organic phase was withdrawn for the chromatographic analysis (using anisole as internal standard). Acquisition and heating were stopped and the autoclave was cooled slowly at low stirring speed (200 rpm). Once the autoclave was cold, stirring was stopped and the reactor was depressurized and purged four times with nitrogen. The whole reaction mixture was left to settle overnight under a nitrogen atmosphere. Finally, the contents of the autoclave were then taken out and separated. An aliquot of the recovered organic phase was diluted into water (with a volumetric dilution factor of 103) for the Rh ICP/MS analysis.

3. Results and discussion

3.1 Monomer and copolymer synthesis

The BMOPPP@CCM latex was prepared by an analogous procedure to that recently described for the similar TPP@CCM latex [26], by replacing the DPPS comonomer with the related 4-[(bis(4'-methoxyxyphenyl)phosphine)styrene] (BMOPPS). This new ligand-functionalized monomer was prepared from chlorobis(4-methoxyxyphenyl)phosphine and 4-bromostyrene following the Scheme 1. Procedure used for the synthesis of BMOPPS.

methyly ether methacrylate (PEOMA) with an average of 19 ethylene oxide units was statistically copolymerized using 4-cyano-4-thiothiopropylsulfanyl pentanoic acid (CTPPA) as the controlling agent and 4,4'-azohis(4-cyanopentanoic acid) (ACPA) as the radical source. Full conversion was attained within 2 h (see Fig. 1, left). The overall monomer/controlling agent ratio being set at 30, the polymer chains obtained at complete conversion have an experimentally determined (SEC analysis) number average molar mass, Mn = 11200 g mol⁻¹, not far from the expected 30 monomer units per chain (Mn,th = 15,625 g mol⁻¹) and a low dispersity (Đ = 1.19).

The resulting water-soluble HOOCCH₂CH₂C(CN)(CH₃)₁₈-P(MAA₁₅-co-PEOMA₁₅)ₘSC(S)SPr was used in step 2 as macro-controlling agent, upon further addition of ACPA, for the chain extension with the mixture of styrene (S) and BMOPPS (95:5 M ratio corresponding to 300 monomer units per chain). During this step, the solid BMOPPS monomer is dissolved in the styrene yielding a single liquid phase that is added to water. The emulsion polymerization starts rather slowly with an induction period (Fig. 1, center) that corresponds to the time required for the P(MAA₁₅-co-PEOMA₁₅)ₘ hydrophilic chains to add sufficient hydrophobic monomer units to start to self-assemble. Once micelles are formed, the polymerization is fast and a complete conversion is obtained within 2 h. The polymer SEC analysis confirms the efficiency and control of the chain extension. The experimentally determined molar mass (Mₚₐ = 43,900 g mol⁻¹) of the polymer analyzed after this step increased relative to the starting P(MAA₁₅-co-PEOMA₁₅)ₘ chains while the molar mass distribution remained narrow (Đ = 1.3) although slightly broader than for P(MAA₁₅-co-PEOMA₁₅)ₘ (Đ = 1.19). This appears related to the presence of a certain amount of dead P(MAA₁₅-co-PEOMA₁₅)ₘ chains, see Fig. S2 in the Supporting Information. Nevertheless, P(MAA₁₅-co-POEMA)ₘ-b-(P(S-POEMA)ₘ-P(PEOMA)ₘ)-b-(P(S-POEMA)ₘ-P(PEOMA)ₘ)-b-(P(S-POEMA)ₘ-P(PEOMA)ₘ) amphiphilic block copolymer micelles exhibiting narrowly distributed sizes were obtained after step 2 (see characterization below). Using a proportion of the phosphine-functionalized monomer greater than 5% results in a three-phase system with undissolved solid BMOPPS and the polymerization does not yield well-dispersed micelles of narrow size distribution. Thus, the chemistry involving BMOPPP is less flexible in terms of degree of ligand functionality relative to the previously reported TPP@CCM synthesis, where the hydrophobic core could be charged with up to 25% of the phosphine-functionalized monomer DPPS [26,27].

The resulting solution was then treated in a third step with a 90:10 M mixture of styrene and di(ethylene glycol) dimethacrylate (DEGDMA) used as cross-linking agent. This step takes place very rapidly without induction time (Fig. 1, right) to yield the desired BMOPPP@CCM. Higher content of DEGDMA would not induce the formation of individual core-shell nano-objects but rather a macrogel formation, as described previously [26].

3.2 BMOPPP@CCM characterization

The full 1H and 31P{1H} NMR characterization was carried out in THF-d₈, which is a good solvent for all the polymer constituents.
The $^1$H NMR spectrum, see Fig. 2a, clearly shows the core aromatic protons in the $\delta$ 7.5–6.2 region. The shell PEO side chains give rise to a sharp resonance at $\delta$ 3.63 (overlapped with one of the two THF-$d_8$ residual proton resonances) for the CH$_2$ protons and a smaller resonance at $\delta$ 3.34 for the OMe chain end. The backbone aliphatic protons of both core and shell are visible as two broad features at $\delta$ 2.2–1.2. The aromatic protons of the BMOPPP moiety overlap with those of PS. The BMOPPP methoxy protons probably give rise to the small resonance at $\delta$ 3.82, to the left of the large PEO methylene resonance (cf. $\delta$ 3.83 for the monomer in CDCl$_3$), providing direct evidence for the incorporation of BMOPPS in the CCM. Indeed, this peak is absent in the $^1$H NMR spectrum of the analogous TPP@CCM, the two spectra being otherwise identical [26]. The extensive resonance overlap hampers the quantitative measurement of the core and shell monomers, however more clear information was obtained from the NMR of the swollen latex (next section).

Furthermore, a solid state $^{13}$C($^1$H) MAS-NMR analysis with quantitative integration reveals the correct area ratio expected for the resonances of the aromatic C atoms at $\delta$ 125–130 and of the PEG C atoms at $\delta$ 70 (see SI, Fig. S3). Further analyses were performed by $^{31}$P($^1$H) NMR (Fig. 2b), which confirmed the incorporation of the phosphorus functionalized styrene monomer in the CCM. The phosphorus resonance of the BMOPPP monomer at $\delta$ –8.9 is slightly broadened and shifted to $\delta$ –11.4 once incorporated in the polymer structure. This displacement was a convenient probe to monitor the BMOPPS consumption during the polymerization.

Additional NMR investigations will be shown below, in reference to the complexation studies.

The particle size was measured by DLS in water both before and after cross-linking, as well as in THF for the final cross-linked BMOPPP@CCM particles. The $D_z$ values are ca. 80 nm in water for both the micelles and the CCMs, with a narrow size distribution (PDI ca. 0.2). The CCMs swelled by a factor of ca. 2.5 in diameter (17 in volume) when placed in a THF solution, see Fig. S4 in the SI (cf. 10 in volume for the related TPP@CCM, both with 5% and with 10% molar fraction of phosphine functionalized monomer in the core [26,27]). The spherical morphology, dimensions, and narrow polydispersity were in all cases confirmed by TEM observations (Fig. 3).

### 3.3. Swelling and metal coordination

As shown in the previous section, all nano-object flexible components (shell and core linear arms) are well solvated in THF-$d_8$. The corresponding $^1$H NMR spectrum in D-enriched water (obtained upon directly diluting the latex with D$_2$O, Fig. 4(a)) only shows the water-solvated hydrophilic shell: PEO methylene resonance at $\delta$ 3.63 and small resonance at $\delta$ 3.32 for the terminal OCH$_3$ group (better visible in the expansion of Fig. 4C). The small and irregular shoulder upfield of the more intense CH$_2$ resonance (at ca. $\delta$ 3.55) is probably caused by the PEO CH$_2$ groups located close to the shell/core interface. Water is a non-solvent for the polystyrene-
Fig. 2. a) $^1$H NMR spectrum of BMOPPP@CCM in THF-$d_8$. The resonance marked with ○ belongs to $\text{H}_2\text{O}$, that marked with □ to silicone grease and those marked with △ to the THF-$d_8$ residual proton resonances. b) $^{31}$P($^1$H) NMR spectra of the BMOPPS monomer (blue) and of BMOPPP@CCM (brown) in THF-$d_8$. (For interpretation of the references to color in this figure legend, the reader is referred to the web version of this article.)

Swelling of the CCM latex by toluene leads to several changes in the $^1$H NMR spectrum, see Fig. 4. A first one is the reappearance of the core constituents as broad features at δ 6–7 for the aromatic protons (Fig. 4B) and at δ 1–2 for the backbone protons (Fig. 4D). These broader resonances overlap with the sharper ones of the free toluene located inside the CCM core (only the latex phase was carefully selected for the NMR measurement after swelling and decanting), at δ 6.5–6.8 (aromatic) and 1.75 (methyl). The small resonance at δ 2.9 (see Fig. 4C) is assigned to the BMOPPP methoxy protons. Although this resonance is upfield shifted by almost 1 ppm from the value in the monomer and in the THF-$d_8$-solvated polymer (vide supra), which may be related to the different solvation, there are no other resonances expected for the polymer in this region and the integrated intensity relative to the PEO resonances in is relatively good agreement with the expected value (observed ratio of 0.061 vs. a theoretical value of 0.076). For comparison, the OMe resonance in the BMOPPS monomer also shifts upfield from δ 3.83 to 3.27 when the solvent is changed from CDCl$_3$ to toluene-$d_8$. Note that no backbone resonance is visible for the unswollen sample, suggesting that only the PEO chains are solvated by water, whereas the polymer chain backbone and the MAA methyl groups of the shell compartment remain solidary with the hydrophobic core in the pristine (unswollen) latex. The core phosphine functions become equally visible, after swelling, by $^{31}$P($^1$H) NMR with a resonance at δ ~ 9.7, Fig. 5b. A second change caused by swelling in the $^1$H NMR spectrum is the splitting of the PEO CH$_2$ and CH$_3$ resonances into a pair of smaller and sharper resonances on one hand, located at the same positions as in the unswollen latex (cf. (a) and (b) in Fig. 4C) and therefore assigned to water-solvated PEO chains, and a pair of more intense and broader resonances on the other hand, shifted upfield to δ 3.52 (CH$_2$) and 3.20 (CH$_3$). The latter are assigned to PEO chains that are folded back into the swollen core. Hence, the polystyrene core solvation by toluene makes this environment compatible with the PEO chains. The deconvolution of the PEO methylene resonance as the sum of two Lorentzian functions (see details in the SI, Fig. S5) yields a chain distribution of (23.2 ± 0.1)% in water and (76.8 ± 0.1)% in the core. This illustrates a slightly greater tendency of core confinement for toluene-swollen BMOPPP@CCM than for the analogous TPP@CCM (10% of triphenylphosphine molar content in the core), for which the water/core ratio was determined by the same technique as 30.3:69.7 (±0.1)% [26].

Integration of the $^1$H spectrum for the toluene-swollen sample affords the expected intensities of the aromatic and backbone protons relative to the PEO protons when considering only the protons in the flexible chains (i.e. excluding the cross-linked nucleus, presumably characterized by slow tumbling even after swelling) and also allows a rough estimate of the amount of toluene, although these measurements are quite imprecise because of overlap. The solvent amount is in the range of 730–810 molecules per chain, depending on whether the calculation is based on the methyl or the aromatic resonance intensity. This amount is similar to that measured for the swelling of the related TPP@CCM [26].

After swelling, the hydroformylation precatalyst, [Rh(acac) (CO)$_2$], could be readily introduced into the CCM core by stirring the latex with a toluene solution of the metal complex, as demonstrated by $^{31}$P($^1$H) NMR. The reaction leads to replacement of one CO ligand by the polymer-anchored phosphine with formation of [Rh(acac) (CO) (BMOPPP@CCM)], as indicated by the disappearance of the metal-free BMOPPP@CCM resonance at δ ~ 11.4 and its replacement with a doublet at δ 44.5 (J$_{PRh}$ = 176 Hz) when one equivalent of Rh per P atom is introduced in the CCM, see Fig. 6. The analogous molecular complex [Rh(acac) (CO) (BMOPPP)] has not been reported to the best of our knowledge, but the resonances of related complexes [Rh(acac) (CO) (TPP)] and [Rh(acac) (CO) (TMOPP)] [TMOPP = tris(4-methoxyphenyl)phosphine], which are reported respectively at δ 48.6 (J$_{PRh}$ = 179.7 Hz) [35] and 43.5 (J$_{PRh}$ = 175.6 Hz) [36], bracket the resonance attributed to [Rh(acac)...

Fig. 3. TEM images of micelles (after Step 2 of the synthesis), left, and final core-cross-linked micelles (BMOPPP@CCM), right.
When only half of the phosphine ligands are metal bonded (Rh/P = 0.5), however, no signal is visible in the $^{31}$P{H} NMR spectrum. This behavior is identical to that observed for the analogous [Rh(acac) (CO) (TPP@CCM)] and is assigned to a rapid exchange between the coordinated and the free phosphine ligands [26]. This observation indicates that the exchange rate for the BMOPP ligand, like that of the TPP ligand, is in the appropriate range to yield coalescence at room temperature.

### 3.4. Hydroformylation catalysis

The BMOPPP@CCM latex was investigated as nanoreactor for the biphasic Rh-catalyzed hydroformylation of 1-octene, using the same operating conditions as for the related TPP@CCM [26,27]: 363 K, 20 bar of syngas pressure and 20% molar as initial concentration of 1-octene in decanal. The volume ratio of organic mixture to aqueous latex suspension was set to 3:1 to keep the catalytic phase dispersed into the organic phase after swelling of the CCM. Table 1 compares the performance of this new ligand in terms of activity, selectivity and Rh leaching with respect to the reference TPP@CCM. As stated in section 3.1, the BMOPPP@CCM latex could only be prepared with a molar fraction of functionalized monomer in the hydrophobic core of 5% (BMOPPS:S = 1:19), whereas the corresponding TPP@CCM latex could also be prepared with higher molar fractions [26]. The most useful comparison in terms of performances related the two CCM with the same functional monomer content [27]. The BMOPPP@CCM based catalyst yielded a quite similar performance as the TPP homolog, with a marginally greater initial TOF and a marginally smaller l/b ratio. Similar activities and selectivities were also reported in homogeneous catalysis in the presence of $p$-OMe-substituted and non-substituted arylphosphines [30,31]. However, the BMOPPP@CCM based catalyst unexpectedly resulted in twice more rhodium leaching in the organic phase. This leaching does not appear related to loss of rhodium.

![Fig. 4. A: $^1$H NMR spectrum of BMOPPP@CCM in D$_2$O before (a) and after (b) swelling with toluene. B–D: Expansions in selected regions. The resonance marked with ○ is due to water, while those marked with Δ belong to the swelling toluene molecules.](image)

![Fig. 5. $^{31}$P{H} NMR spectrum of BMOPPP@CCM in D$_2$O before (a) and after (b) swelling with toluene.](image)

![Fig. 6. $^{31}$P{H} NMR spectra of [Rh(acac) (CO) (BMOPPP@CCM)] with different amounts of rhodium complex. The starred resonance is due to a minor amount of oxidized phosphine impurity.](image)
Results of the biphasic hydroformylation of 1-octene.

Table 1

<table>
<thead>
<tr>
<th>Entry</th>
<th>Investigated ligand</th>
<th>Initial rate&lt;sup&gt;a&lt;/sup&gt; (kmol m&lt;sup&gt;-3&lt;/sup&gt; s&lt;sup&gt;-1&lt;/sup&gt;)</th>
<th>TOF&lt;sub&gt;max&lt;/sub&gt;&lt;sup&gt;a&lt;/sup&gt; (h&lt;sup&gt;-1&lt;/sup&gt;)</th>
<th>l/b</th>
<th>Leaching&lt;sup&gt;b&lt;/sup&gt; [Rh]&lt;sub&gt;avg&lt;/sub&gt; (ppm)</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>BMOPPPP@CCM (BMOPPS:S = 1:19)</td>
<td>1.3 × 10&lt;sup&gt;-3&lt;/sup&gt;</td>
<td>742</td>
<td>2.7</td>
<td>4.5</td>
</tr>
<tr>
<td>2</td>
<td>TPP@CCM with (DPPPS:S = 1:19)</td>
<td>1.3 × 10&lt;sup&gt;-3&lt;/sup&gt;</td>
<td>695</td>
<td>3.3</td>
<td>1.8</td>
</tr>
</tbody>
</table>

* Operating conditions: [Rh(acac) (CO)]<sub>2</sub> = 6.5 × 10<sup>-2</sup> kmol m<sup>-3</sup> [1-octene] = 1.1 kmol m<sup>-3</sup>, P(Rh) = 4, V<sub>org</sub> = 75 mL (octene + decanal), V<sub>a,q</sub> = 25 mL (latex + water), T = 363 K, P<sub>org</sub> = 20 bar (CO/H<sub>2</sub> = 1), w = 1200 rpm.

<sup>a</sup> Initial reaction rate (with respect to non-swollen aqueous phase) and corresponding Turn Over Frequency calculated from the syngas consumption during the first 5 min of reaction after the gas absorption phase.

<sup>b</sup> Linear to branched aldehyde ratio determined from the GC/FID analysis of the final sample.

Replacement of diphenylphosphonostyrene with 4-[bis[5-methoxypheny]phosphino]styrene in our optimized synthesis of core-cross-linked micelles (CCM) by emulsion RAFT polymerization [26,27] leads to the successful fabrication of functionalized nanoreactors containing covalently linked bis[5-methoxyphenyl]phosphine (BMOPPP) ligands in the hydrophobic core, although incorporation of this ligand functionalized monomer is limited by its low solubility in styrene. These BMOPPP@CCM nanoreactors show the same behavior as the previously described TPP@CCM in terms of core swelling, coordination of [Rh(acac) (CO)]<sub>2</sub>, intra- particle phosphine ligand exchange, and catalytic activity in aqueous biphasic 1-octene hydroformylation. Catalyst leaching, however, is slightly greater, proving the point that this metal leaching process is not related to partial metal loss from the nanoreactor but rather to partial loss of the entire nanoreactor into the organic product phase. This phenomenon is presumably favored by a greater tendency of this more strongly bonding phosphine to form bigger and more lipophilic particle aggregates through interparticle cross-linking. On the basis of these hypotheses, improvement of this catalytic nanoreactor appears possible by turning to polymer-anchored bidentate ligands and efforts to assemble polymer architectures of this type are currently underway.

Acknowledgments

We are grateful to the Agence Nationale de la Recherche (ANR) for support of this work through grant “BIPHNASNOCAT” (ANR-11 BS07-025-01). Additional support from the Centre National de la Recherche Scientifique (CNRS) and from the Institut Universitaire de France (IUF) is also gratefully acknowledged.

Appendix A. Supplementary data

Supplementary data related to this article can be found at http://dx.doi.org/10.1016/j.polymer.2015.02.024.

References
