Aqueous Dispersions of Nonspherical Polyethylene Nanoparticles from Free-Radical Polymerization under Mild Conditions

Etienne Grau, Pierre-Yves Dugas, Jean-Pierre Broyer, Christophe Boisson,
Roger Spitz, Vincent Monteil

► To cite this version:

Etienne Grau, Pierre-Yves Dugas, Jean-Pierre Broyer, Christophe Boisson, Roger Spitz, et al.. Aqueous Dispersions of Nonspherical Polyethylene Nanoparticles from Free-Radical Polymerization under Mild Conditions. *Angewandte Chemie International Edition*, 2010, 49 (38), pp.6810-6812. 10.1002/anie.201001800 . hal-00948692

HAL Id: hal-00948692

<https://hal.science/hal-00948692>

Submitted on 20 Nov 2019

HAL is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers.

L'archive ouverte pluridisciplinaire **HAL**, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.

Aqueous Dispersions of Non Spherical Polyethylene Nanoparticles from Free Radical Polymerization under Mild Conditions**

*Etienne Grau, Pierre-Yves Dugas, Jean-Pierre Broyer, Christophe Boisson, Roger Spitz, and Vincent Monteil**

Polyethylene, the first manufactured polymer by volume, is usually synthesized from low pressure and temperature catalytic processes^[1,2] or a high temperature ($> 200^{\circ}\text{C}$) and pressure (> 1000 bar), highly energy consuming free-radical polymerization process.^[3-5] In the latter case a branched, low density, polyethylene is produced (LDPE) in contrast to Ziegler Natta catalysis which enables the synthesis of high density polyethylene (HDPE) exhibiting higher crystallinities and melting temperatures. These well established polymerization processes require improvement: reduction of energy consumption and/or Volatile Organic Compounds (VOC) are important targets. The VOC issue has been greatly solved for low pressure catalytic Ziegler-Natta polymerizations by using solvent-free gas-phase processes. For slurry polymerization new catalysts compatible with “green” diluents such as supercritical CO_2 ^[6,7] or water^[8-11] have been developed recently.

Recently we successfully produced polyethylene (PE) by a radical pathway under less energy consuming conditions: medium pressure < 250 bar and low temperature 70°C using organic solvents (toluene or THF).^[12] PE was synthesized in high yields and exhibited intermediate melting points and crystallinities in comparison to HDPE and LDPE ($115^{\circ}\text{C} < T_m < 119^{\circ}\text{C}$; crystallinity of 55-70%). However, polymer molecular weights remained low ($M_n < 5000$ g/mol, $\text{PDI} \approx 2$) because of frequent transfer reactions to the solvent.

Transposition to an emulsion polymerization in aqueous dispersed medium (benefiting from the compartmentalization of radicals and from the low transfer ability of water) should be useful to increase both molecular weight and yield and at the same time to solve the VOC issue.

Only few works on free radical polymerizations (FRP) of ethylene in aqueous dispersed media have been reported^[13-17] at relatively high pressures ($P > 300$ bar) and a wide range of temperatures. The interpretation of the results in these early works (from 1945 to 1975) is not easy because of the lack of analytical tools to study colloidal properties of the polymer dispersions

obtained.

Note that the emulsion process for ethylene polymerization can not be a classical one. Ethylene is a supercritical gas and consequently no ethylene droplets exist during the polymerization and no liquid unreacted monomer can remain in the latex. In addition PE is a crystalline material contrary to most conventional polymers produced by FRP.

In the present paper, FRP of ethylene in emulsion under mild conditions has been investigated, representing an innovative low energy consuming “green” efficient way to produce PE by a free radical mechanism. The transposition of ethylene polymerization process to aqueous medium has been achieved by using a cationic water soluble initiator, 2,2-azobis(2-amidinopropane)-dihydrochloride (AIBA). FRP of ethylene was performed in water at 70°C with and without a standard cationic surfactant (CTAB, CetylTrimethylAmmonium Bromide) to assist nucleation and particle stabilization. In all cases ethylene was polymerized with significant yields and stable dispersions of PE particles were obtained for ethylene pressure up to 250 bar (Figure 1). Interestingly, PE can be synthesized by this FRP process down to 50 bar of pressure.

In the surfactant-free system, yield is lower than that obtained using the same amount of initiator in THF but higher than in toluene.^[12] The stabilization of PE particles is assumed to result from the cationic fragments of the initiator attached at the chain end which induce electrostatic repulsion. Average particles diameters (D_p) measured by DLS (dynamic light scattering) increase with the ethylene pressure (and consequently with the yield) from 30 nm to 110 nm. Polydispersity indexes remain very low ($\text{PI} \sim 0.05$) indicating the monodisperse character of particle size distribution. In addition, the yield/ D_p^3 ratio, standing for the number of particles, remains constant whatever the ethylene pressure.

When polymerizations were performed in the presence of a standard cationic surfactant (CTAB) at 1 g/L (above the critical micelle concentration: 0.2 g/L at 25°C), much higher activities were observed (Figure 1). This emulsion system is even more efficient than the polymerization in THF.^[12] In these non-optimized conditions, up to 40% of solid content are obtained (after degassing the 250 bar of ethylene). Average particle diameters seem to reach a plateau at 50 nm when increasing ethylene pressure. This indicates that the number of particles increases with the yield. Polydispersity indexes measured by DLS remain surprisingly higher ($\text{PI} \sim 0.5$) than for the surfactant-free process.

The PE produced exhibits a low melting point ($T_m \sim 100^{\circ}\text{C}$) and low crystallinity (30-40%). Highest values were obtained in the case of the surfactant-free polymerization process (see Supporting Information Table S1). As expected high molecular weights PE (M_n from 10^4 up to 10^5 g/mol) were produced. The number of PE chains synthesized is greater with CTAB indicating a possible transfer to surfactant (Supporting Information Figure S1). The PE obtained are moderately branched under both conditions (Figure S1: 30 branches/1000C without surfactant and 37 branches/1000C with CTAB) as determined by ^{13}C NMR^[18] which is in agreement with the crystallinities and melting temperatures measured. This higher branching level in water than in an organic solvent (THF:

[*] E. Grau, P.-Y. Dugas, J.-P. Broyer, Dr. C. Boisson, Dr. R. Spitz, Dr. V. Monteil
UMR 5265 Laboratoire de Chimie Catalyse Polymères et Procédés (C2P2), LCPP team
Université de Lyon, Univ. Lyon 1, CPE Lyon, CNRS
Bat 308F, 43 Bd du 11 novembre 1918, 69616
Villeurbanne, France
Fax: (+) 33 4 72 43 17 68
E-mail: monteil@lcpp.cpe.fr

[**] E.G. thanks the “Ministère de la Recherche et de l'Enseignement Supérieur” for his fellowship. The authors thank Mettler-Toledo for the thermal analysis. We are indebted to Mr C. Graillat and Prof. B. Charleux for their valuable contributions to this research

9 branches/1000C or Toluene: 7 branches/1000C)^[12] can be explained by the compartmentalization of the PE growing chains which increases transfer reactions to the polymer. The proportion of short chain branches is lower in emulsion (25% vs 35% in organic solvent) due to favored intermolecular over intramolecular transfer reactions in a confined environment.

Figure 1. Free radical polymerization of ethylene in aqueous dispersed medium: —■— yield and □— average particle diameter* vs ethylene pressure (80 mg AIBA, 50 mL water, 4 h at 70°C under ethylene pressure); —▲— yield and △— average particle diameter* vs ethylene pressure (80 mg AIBA, 50 mL water with 1 g/L of CTAB, 4 h at 70°C under ethylene pressure), * determined by DLS

Figure 2. Reaction profile for of free radical polymerization of ethylene in aqueous dispersed medium: —■— yield and □— average particle diameter* vs time (80 mg AIBA, 50 mL water at 70°C under 100 bar); —▲— yield and △— average particle diameter* vs time (80 mg AIBA, 50 mL water with 1 g/L of CTAB at 70°C under 100 bar), * determined by DLS

In order to link solvent and emulsion processes, the influence of addition of organic solvents to water (water-miscible (THF) or non-miscible (toluene)) was investigated. PE molecular weights dropped in the presence of solvents (see supporting information Table S2). Mn dropped from 50500 g/mol in water to 8300 g/mol and 2350 g/mol for toluene/H₂O (1/4) and THF/H₂O mixtures respectively in the presence of CTAB. This decrease can be related to an increased frequency of transfer reactions to solvent (contrary to water, THF and toluene exhibit high transfer abilities)^[12] which has been confirmed by NMR analysis (see supporting information Figure S2). With THF, the transfer reaction should take place in the

continuous aqueous phase or at the particle surface and not in the particles, because THF is not an efficient swelling agent for amorphous PE (same Dp is observed before and after removal of THF by partial reduced pressure evaporation). For toluene, the Dp drops by about 10 nm by removing the organic solvent (toluene is a swelling solvent for PE) so transfer could additionally take place inside the particles.

Surfactant-free and classical emulsion polymerization processes were compared by investigating the reaction profile at 70°C under 100 bar of ethylene (Figure 2). For the surfactant-free system, particle diameters increase with yield and the yield/D_p³ ratio remains constant, thus no renucleation or aggregation took place during the polymerization.

In the presence of CTAB the behavior is quite different. Initially particles with large diameters are formed and seem to disappear with time to generate only small particles (D_p≈30 nm) after 2h. The mechanism for the extinction of large particles to generate very small particles still remains unknown, but preliminary experimental results suggest a primordial role of the surfactant itself. For example if 1g/L of CTAB is added to surfactant-free PE latex, after stirring at 70°C small particles are ejected (see supporting information Figure S3).

Figure 3. TEM pictures of PE latex: a) standard particles without CTAB, 100 bar, 4h at 70°C b) standard particles with 1 g/L CTAB (included with a tilting angle of 60°), 100 bar, 4h at 70°C.

The PE latexes were also characterized using TEM analysis (Figure 3). In the surfactant-free process quasi-spherical particles were observed. The rigid lamellas of semi-crystalline PE prevent the formation of spheres (as already observed for latex prepared by catalytic emulsion polymerization).^[19] Nevertheless, particles show a relative diameter homogeneity which is in agreement with DLS measurements (low PI).

In the presence of CTAB, TEM pictures show a low contrast for the surface of particles which could originate from flat particles. This has been confirmed by tilting the sample since disks were observed at 0° and ellipses at 60°. Note that no significant changes were observed during tilt with PE latex obtained from surfactant free

polymerization. From the hypothesis of cylinder-like particles, the dimensions of these objects were estimated (average disks diameters about 35 nm and thickness about 3-4 nm). From these findings we can better explain the high values of PI (~ 0.5) obtained by DLS (autocorrelation function of DLS being calculated for a size distribution of spherical particles) which were not in agreement with the apparent homogeneity in the diameters of particle sizes observed by TEM.^[20]

In summary, compartmentalization in water from emulsion processes (with or without the presence of surfactant) is a very promising way to produce high molecular weight polyethylenes in the low to very low density range by FRP under mild conditions from a water soluble cationic initiator. PE yields are higher than for the solvent processes previously developed. From a colloidal point of view, FRP in emulsion exhibits unexpected original behaviors. In the presence of surfactant, very small PE cylinder particles are generated while larger quasi-spherical particles were formed in the surfactant-free process. Investigations of the coating properties of these attractive PE nanoparticles are in progress.

The experimental details are provided in the Supporting Information part.

Keywords: nanoparticles • polymerization • free radical polymerization • polyethylene • emulsion

-
- [1] R. Mulhaupt, *Macromol. Chem. Phys.* **2003**, *204*, 289-327.
 [2] D.L. Beach, Y.V. Kissin, in *Encyclopedia of Polymer Science and Engineering*, 2nd ed., Vol. 6 (Eds: H.F. Mark, N. M. Bikales, C. G.

- Overberger, G. Menges), WILEY INTERSCIENCE, New York, **1985**, pp 454-489.
 [3] K.W. Doak, in *Encyclopedia of Polymer Science and Engineering*, 2nd ed., Vol. 6 (Eds: H.F. Mark, N. M. Bikales, C. G. Overberger, G. Menges), WILEY INTERSCIENCE, New York, **1985**, pp 386-428.
 [4] S.L. Aggarwal, O.J. Sweeting, *Chem. Rev.* **1957**, *57*, 665-742.
 [5] S.P. Ehrlich, G. A. Mortimer, *Adv. Polymer Sci.*, **1970**, *7*, 386-448.
 [6] M. Kemmere, T. de Vries, M. Vorstman, J. Keurentjes, *Chem. Eng. Sc.* **2001**, *56*, 4197-4204.
 [7] A. Bastero, G. Francio, W. Leitner, S. Mecking, *Chem. Eur. J.* **2006**, *12*, 6110-6116.
 [8] L. Kolb, V. Monteil, R. Thomann, S. Mecking, *Angew. Chem.* **2005**, *117*, 433-436.
 [9] A. Held, F.M. Bauers, S. Mecking, *Chem. Comm.* **2000**, 301-302
 [10] R. Soula, C. Novat, A. Tomov, R. Spitz, J. Claverie, X. Drujon, J. Malinge, T. Saudemont, *Macromolecules* **2001**, *34*, 2022-2026.
 [11] S. Mecking, *Coll. Polym. Sci.* **2007**, *285*, 605-619.
 [12] E. Grau, J.P. Broyer, C. Boisson, R. Spitz, V. Monteil, *Macromolecules* **2009**, *42*, 7279-7281.
 [13] H. Hopff, R. Kern, *Modern Plastics* **1946**, 153-220.
 [14] A.F. Helin, H.K. Stryker, G.J. Mantell, *J. App. Polym. Sc.* **1965**, *9*, 1797-1805.
 [15] A.F. Helin, H.K. Stryker, G.J. Mantell, *J. App. Polym. Sc.* **1965**, *9*, 1807-1822.
 [16] T. Suwa, H. Nakajima, M. Takehisa, S. Machi, *Polym. Let. Ed.* **1975**, *13*, 369-375.
 [17] H.K. Stryker, G.J. Mantell, *J. Polym. Sci. C Polym. Symp.* **1969**, *27*, 35-48.
 [18] G.B. Galland, R.F. de Souza, R.S. Mauler, F.F. Nunes, *Macromolecules* **1999**, *32*, 1620-1625.
 [19] F.M. Bauers, R. Thomann, S. Mecking, *J. Am. Chem. Soc.* **2003**, *125*, 8838-8840.
 [20] High PI can also be partially due to a wide distribution in cylinder thickness.

Entry for the Table of Contents (Please choose one layout)

Layout 1:

Polyethylene Nanoparticles

E. Grau, P.-Y. Dugas, J.-P. Broyer, C. Boisson, R. Spitz, V. Monteil

Page – Page

Aqueous Dispersions of Non Spherical Polyethylene Nanoparticles from Free Radical Polymerization under Mild Conditions

Stable PE Nanoparticles Dispersions

Free radical polymerizations of ethylene usually require severe conditions. We investigate the efficiency of this reaction under mild conditions (<250 bar) in water in order to produce PE stable aqueous dispersions. Latexes of PE nanoparticles with various shapes (cylinder or sphere) and solid contents up to 40% were prepared.

Supporting Information

Table of Content :

Experimental Section	S2
Figure S1 : Typical ^{13}C NMR of polyethylene prepared in water	S4
Figure S2 : Typical ^{13}C NMR of polyethylene prepared in water/THF	S5
Figure S3 : TEM picture of PE latex before (a) and and after 4h at 70°C and additional CTAB (b)	S6
Table S1 : Influence of ethylene pressure on free radical polymerization of ethylene in aqueous dispersed medium	S7
Table S2 : Influence of additional organic solvent on free radical polymerization of ethylene in aqueous dispersed medium	S8
Table S3 : Influence of polymerization time on free radical polymerization of ethylene in aqueous dispersed medium	S9

Experimental section

All chemicals were purified using standard Schlenk procedures and handled under argon atmosphere. Solvents (THF and toluene) were distilled or degassed under argon. Water was purified using Milli-Q academic system (Millipore Corporation) and degassed under argon. Ethylene (purity 99.95%) was purchased from Air Liquide, AIBA and CTAB from Acros and used without any further purification.

Latex characterizations

Particle size was determined by dynamic light scattering (DLS) using a Malvern Zetasizer 1000 HAS autosizer apparatus with a detection angle of 90° at 25°C. The measurements were performed on highly diluted samples in order to rule out interaction and multiple scattering effects. The intensity average diameter was computed from the intensity autocorrelation data using the cumulant analysis method. The final data was the average of 5 measurements for each sample. TEM analysis was performed after placing a drop of the particle suspension on a copper grid (3.05mm copper grid with Formvar/Carbon support Film, 200 mesh (Agar Scientific)) and dried before analysis. TEM was performed in a Philips CM120 transmission electron microscope, at an acceleration voltage of 80 kV (Centre Technologique des Microstructures (CTμ) - Plateforme d'Imagerie Integrative (PI²), Claude Bernard University, Lyon, France).

Polymer characterizations

Molecular weights of polyethylenes were determined by size exclusion chromatography (SEC) using a Waters Alliance GPCV 2000 instrument (columns: PLgel Olexis); two detectors (viscosimeter and refractometer) in trichlorobenzene (flow rate: 1 mL/min) at 150°C. The system was calibrated with polystyrene standards using universal calibration. Differential scanning calorimetry (DSC) was performed on a Mettler Toledo DSC1 at a

heating rate of 5 K/min. Two successive heating and cooling of samples were performed. We have considered data (T_m values, crystallinity) obtained during the second heats. High-resolution liquid NMR spectroscopy was carried out with a Bruker DRX 250 spectrometers operating at 250 MHz for ^1H . Spectra were obtained with a 5-mm QNP probe. PE samples were examined as 10–15 % (w/v) solutions using a mixture of tetrachloroethylene (TCE) and perdeuterobenzene (C_6D_6) (2/1 v/v) as solvent at 363 K. Chemical shift values (δ) are given in ppm in reference to internal tetramethylsilane (TMS).

Standard polymerization procedure

Caution, all polymerizations involve high pressure and explosive gas.

Ethylene polymerizations were done in a 160 mL stainless steel autoclave (equipped with safety valves, stirrer, oven) from Parr Instrument Co.. The 2,2-azobis(2-amidinopropane) dihydrochloride (AIBA) was dissolved in the desired volume of water (or a solution of cetyltrimethylammonium bromide (CTAB) at 1 g/L) in a Schlenk tube under argon. The mixture was introduced through cannula into the reactor. Ethylene was introduced and the mixture was heated at the desired temperature under stirring (250 rpm). To manage safely polymerization over 50 bar of ethylene we use a 1.5 L intermediate tank. The tank was cooled down to -20°C to liquefy ethylene at 35 bar. When thermodynamic equilibrium was reached, the intermediate tank was isolated and heated to reach up to a pressure of 300 bar. This tank was used to charge the reactor. After 4 hours of polymerization the reactor was slowly cooled down and degassed. A stable dispersion of polyethylene nanoparticles was obtained. A fraction of the latex was then dried under vacuum at 70°C to determine solid content. Polymer analyses (by NMR, DSC, SEC) were performed on dried samples washed with water.

Figure S1: Typical ^{13}C NMR of polyethylene prepared in water (notation from Galland et al ref 18 of the article) * a, b signals are representative of CTAB. The presence of CTAB can be explained by two different hypotheses: CTAB is linked to PE after transfer to surfactant or has cocrystallized with PE.

S5

Figure S3: TEM picture of PE latex before (a) and after 4h at 70°C and additional CTAB (b)

Table S1 : Influence of ethylene pressure on free radical polymerization of ethylene in aqueous dispersed medium^a (experimental data of Figure 1)

Run	Ethylene Pressure (bar)	CTAB (g/L)	Yield (g)	Melting point (°C) ^b	Crystallinity (%) ^b	Mn (g/mol) ^c	PDI ^c	Dp (nm) ^d	PI ^d
1	50	0	0.3	70.3	12	10800	3.4	32 (±2)	0.07 (±0.01)
2	100	0	1.3	96.5	35	21600	6.0	89 (±1)	0.04 (±0.01)
3	150	0	1.8	100.9	30	31100	7.3	104 (±1)	0.06 (±0.03)
4	200	0	2.2	106.0	40	20800	6.1	109 (±1)	0.03 (±0.01)
5	250	0	2.5	104.6	37	21000	7.9	113 (±1)	0.03 (±0.02)
6	50	1	0.7	69.5	10	38800	2.8	97 (±3)	0.71 (±0.03)
7	100	1	4.6	92.8	31	50500	8.7	24 (±1)	0.52 (±0.08)
8	150	1	7.9	97.3	31	60500	7.4	51 (±1)	0.17 (±0.02)
9	200	1	12.3	99.4	29	73800	8.1	52 (±1)	0.22 (±0.02)
10	250	1	19.9	97.0	31	119000	6.3	46 (±2)	0.33 (±0.09)

^a Reactions were carried out at 70°C under ethylene pressure during 4h with 80 mg AIBA in 50 mL of water; ^b determined by DSC, ^c determined by HTSEC, ^d determined by DLS

Table S2 : Influence of additional organic solvent on free radical polymerization of ethylene
in aqueous dispersed medium^a

Run	Water (mL)/ Solvent (mL) [solvent]	CTAB (g/L)	Yield (g)	Melting point (°C) ^c	Crystallinity (%) ^c	Mn (g/mol) ^d	PDI ^d	Dp (nm) ^e	PI ^e
11 (=2)	50/0	0	1.3	96.5	35	21600	6.0	89 (±1)	0.04 (±0.01)
12 (=7)	50/0	1	4.6	92.8	31	50500	8.7	24 (±1)	0.52 (±0.08)
13 ^b	0/50 [THF]	0	3.9	115.2	58	1190	1.9		
14	40/10 [THF]	0	1.3	105.0	44	1760	3.8	129 (±2)	0.03 (±0.02)
15	40/10 [THF]	1	3.2	103.1	40	2350	3.2	16 (±1)	0.39 (±0.07)
16 ^b	0/50 [toluene]	0	0.7	115.9	63	2340	1.9		
17	40/10 [toluene]	0	0.4	104.5	27	17100	2.6	72 (±5)	0.14 (±0.06)
18	40/10 [toluene]	1	0.8	99.9	22	8300	3.2	121 (±32)	0.88 (±0.11)

^a Reactions were carried at 70°C under 100 bar ethylene pressure during 4h with 80 mg AIBA in 50 mL of solvent; ^b In organic medium initiator is 50 mg of AIBN because AIBA is not soluble in this solvent see E. Grau, J.P. Broyer, C. Boisson, R. Spitz, V. Monteil, *Macromolecules* **2009**, *42*, 7279-7281, ^c determined by DSC, ^d determined by HTSEC, ^e determined by DLS

Table S3 : Influence of polymerization time on free radical polymerization of ethylene in aqueous dispersed medium^a (experimental data of Figure 2)

Run	Time (h)	CTAB (g/L)	Yield (g)	Melting point (°C) ^b	Crystallinity (%) ^b	Mn (g/mol) ^c	PDI ^c	Dp (nm) ^d	PI ^d
19	0.25	0	0.1	97.6	13	nd	nd	29 (±1)	0.04 (±0.01)
20	0.5	0	0.3	99.3	16	nd	nd	46 (±1)	0.04 (±0.02)
21	1	0	0.5	94.8	19	319700	6.2	57 (±1)	0.05 (±0.01)
22	2	0	0.7	96.3	24	74100	4.8	76 (±1)	0.03 (±0.03)
23 (=2)	4	0	1.3	96.5	35	21600	6.0	89 (±1)	0.04 (±0.01)
24	14	0	2.4	97.7	36	30500	5.4	122 (±2)	0.02 (±0.02)
25	0.25	1	0.2	91.4	15	55400	4.0	106 (±6)	0.82 (±0.11)
26	0.5	1	0.6	94.0	21	22400	7.3	121 (±3)	0.91 (±0.02)
27	1	1	1.2	88.2	23	107400	5.6	37 (±3)	0.52 (±0.03)
28	2	1	2.5	89.0	25	54200	7.7	28 (±1)	0.52 (±0.06)
29 (=7)	4	1	4.6	92.8	31	50500	8.7	24 (±1)	0.52 (±0.08)
30	14	1	6	91.1	30	43900	5.7	19 (±1)	0.48 (±0.01)

^a Reactions were carried at 70°C under 100 bar of ethylene with 80mg AIBA in 50 mL of water; ^b determined by DSC, ^c determined by HTSEC, ^d determined by DLS, nd not determined