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Data Article

Data of PCL-*b*-P(MMA-DMAEMA)₂ characterization and related assays

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ABSTRACT

The data presented here are related to the research paper entitled “PCL-*b*-P(MMA-co-DMAEMA)₂ new triblock copolymer for novel pH-sensitive nanocapsules intended for drug delivery to tumors” by Franco et al. [1]. Characterization data of PCL-diol, macro-initiator Br-PCL-Br, homopolymers (PMMA and PDMAEMA) and copolymers (batch 1 and batch 2) analyzed by FTIR, SEC and NMR, as well as, characterization of PCL-NS formulation by laser diffraction and DLS analysis, initial nanocapsule formulations and 1C-NC and 2C-NC formulations, including hydrodynamic diameter at different pH media, and DMSO cytotoxicity.

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Specifications Table

Subject Area	Chemistry, Biology, Pharmacy
More specific sub- ject area	Copolymer synthesis and pH-sensitive nanocapsules
Type of data	Tables and figures
How data was acquired	FTIR spectrometer (Varian 640 FT-IR, USA), SEC by GPCMax tripletdetector (Viscotec, Marvel Instruments Ltd, England, UK, columns of Styragel 10 ⁴ , 10 ⁵ , and 10 ⁶ Å), ¹ H NMR (300 MHz) and ¹³ C NMR (75 MH) by INOVA-300 (Varian, USA), laser diffraction (Malvern Mastersizer [®] 2000, Malvern Instruments, UK), dynamic light scattering (DLS, Malvern Zetasizer instrument - NanoZS, Malvern Instruments, UK) and cytotoxicity in MCF-7 cells (ATCC [®] -HTB-22 TM Rockville, MD, λ of 570 and 630 nm - SpectraMax M2, Molecular Devices)
Data format	Raw, analyzed
Experimental factors	Synthesis and products isolation by filtration and purification by impurities dissolution. Nanocapsules were analyzed as produced, without pre-treatment
Experimental features	Chemical characterization and identification of modifications induced by synthesis procedures or by formulation of materials
Data source location	Commercial reagent: PCL, MMA and DMAEMA
Data accessibility	Data is provided with this article

Value of data

- Characterization spectra of the materials were compared with data from other works when developing a similar delivery system or copolymer synthesis.
- SEC and NMR data provided information on the efficiency of the copolymer synthesis and were useful for their identification.
- Nanocapsules parameters and its response to different pH media is innovative for scientific community since the copolymer maintains its integrity and expands upon acid pH.
- The bromide end-group of the copolymer permit application as active targeting system after covalent binding with ligands.

1. Data

The data presented in [Section 1.1](#) is the ¹H NMR analysis of the homopolymers PMMA and PDMAEMA ([Fig. 1](#)). [Section 1.2](#) involves the profiles by laser diffraction of PCL-NS and its parameters ([Fig. 2](#), [Table 1](#)). The data presented in [Section 1.3](#) includes the synthesis of the macroinitiator and the characterization by FTIR and SEC analysis of the PCL-diol and Br-PCL-Br ([Fig. 3](#)), ¹H NMR ([Fig. 4](#)) and ¹³C NMR ([Fig. 5](#)). [Section 1.4](#) brings data referent to the copolymers (batch 1 and batch 2) with FTIR, SEC ([Fig. 6](#)), ¹H NMR ([Fig. 7](#)) and ¹³C NMR ([Fig. 8](#)). The data contained in [Section 1.5](#) is related to the characterization of nanocapsules formulations, as size distribution profiles of initial nanocapsule formulations ([Fig. 9](#)) and 1C-NC and 2C-NC formulations ([Fig. 10](#)), including its parameters ([Table 2](#)) and the DLS profile ([Fig. 11](#)) and its behavior in different pH ([Fig. 12](#)). [Section 1.6](#) presented the DMSO cytotoxicity data ([Fig. 13](#)).

1.1. ¹H NMR spectra of homopolymers PMMA and PDMAEMA

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