

PAPER

[View Article Online](#)
[View Journal](#) | [View Issue](#)

Cite this: *Polym. Chem.*, 2020, **11**, 7235

Defect engineering of polyethylene-like polyphosphoesters: solid-state NMR characterization and surface chemistry of anisotropic polymer nanoplatelets†

Jens C. Markwart,^a Oksana Suraeva,^a Tobias Haider,^a Ingo Lieberwirth,^a Robert Graf[✉] and Frederik R. Wurm[✉]

Anisotropic materials with very high aspect ratios, such as nanoplatelets, are an exciting class of materials due to their unique properties based on their unilamellar geometry. Controlling their size and surface-functionality is challenging and has only been achieved in some cases for synthetic polymers. We present a general approach to prepare polymer-nanoplatelets with control over the lateral size and basal functionality, by simple polycondensation of precisely spaced and functional phosphate groups in polyethylene-like polymers. Because of the relatively large size of the phosphate groups, they are expelled from the bulk crystal to the basal surface. This allows to control the chain-folding during crystallization, which we analyzed via solid-state NMR and TEM. Furthermore, we present chemistry “on the surface” of the platelets: the pendant ester group at the phosphate offers the possibility to introduce functional groups accessible for further chemical modification on the crystal surface. This is shown by introducing a 2-acetylthioethyl ester group and subsequently cleaving this 2-acetylthioethyl ester group to the free P–OH. Together, these results render polyethylene-like polyphosphoesters a versatile platform for soft-matter nanoplatelets as functional colloids.

Received 18th September 2020,
Accepted 28th October 2020

DOI: 10.1039/d0py01352h

rsc.li/polymers

Introduction

Due to the pioneering work of Karl Ziegler and Giulio Natta, the synthesis of polyethylene and propylene at mild temperatures and pressure became possible, which has led to more than hundred million tons of polyethylene (PE) and polypropylene (PP) produced per year.^{1–3} The application of these polymers is determined by their bulk properties like melting or glass transition temperature. Today, PE and PP are known as commodity material with various applications.^{4–7}

In contrast to these bulk materials, planar materials with very high aspect ratios are an exciting material class due to their unique properties based on their unilamellar geometry, allowing anisotropic material properties. Examples for such materials are graphene or inorganic platelets, which are often

used to reinforce bulk polymers.⁸ To date, only very few reports on chemically functional platelet-shaped nanomaterials based on commodity plastics like polyethylene were reported.^{9–11}

We utilized phosphorus chemistry and simple polycondensation reactions to introduce precisely spaced phosphoester defects in a polymer chain to achieve polyethylene-like polyphosphoesters (PPEs).¹² PPEs have been a focus of our research in recent years ranging from biomedical to material science applications.^{13,14} For the PE-like PPEs, the phosphates can introduce additional functionality but act as defects for the crystallization, as they are too large to be incorporated into the lamellar crystal. This led to a layered morphology of crystallites in the bulk.^{12,15} Besides, the phosphate units also confine the thickness of the polymer lamellae, which resulted in the formation of anisotropic polymer platelets, when crystallized from dilute solution¹⁵ or at the air–water interface.¹⁶ In addition, the pendant ester group at the phosphate gives the possibility for the introduction of functional groups which are accessible for further chemical modification on the crystal surface after crystallization from solution. Ramakrishnan's group followed a related concept for the preparation of liquid crystals from bulk by preparing polyethylene-like polyesters,

^aMax Planck Institute for Polymer Research, Ackermannweg 10, 55128 Mainz, Germany. E-mail: graf@mpip-mainz.mpg.de

^b“Sustainable Polymer Chemistry”, MESA+ Institute for Nanotechnology, Faculty of Science and Technology, Universiteit Twente, PO Box 217, 7500 AE Enschede, The Netherlands. E-mail: frederik.wurm@utwente.nl

†Electronic supplementary information (ESI) available. See DOI: 10.1039/d0py01352h



carrying long-chain alkyl segments in the backbone and either poly(ethylene glycol) (PEG) or fluorinated pendant groups at precisely placed positions.¹⁷

The crystallization of PE from dilute solution was used already more than 50 years ago to prepare PE-single crystals.^{18,19} The conditions for crystallization from solution of polymers influence the crystal properties and can be controlled for example by the rate of cooling,^{20–22} concentration,²³ or techniques like self-seeding.^{19,24} However, with all techniques it is challenging to adjust the lateral size and thickness of the materials at the same time. Precisely control over the chain folding by defect engineering allows eliminating the variable of varying thickness and focusing on lateral size control during crystallization from dilute solution.

Taken together, long-chain aliphatic polyphosphoesters are a versatile platform to prepare nanoplatelets with very high aspect ratio and distinct chemical functionality on their surface. The pendant ester group was used for the first time to tailor a multi-step reaction “on surface” of the nanoplatelets. The versatility of the chemistry in the pendant group renders the nanoplatelets as highly functional colloidal platform which can be utilized in various future applications.

Results and discussion

Acyclic diene metathesis polymerization (ADMET) allows the preparation of PE-like materials; in our group, it was extensively used for the preparation of phosphate-containing polymers.^{15,25–30} Here, we synthesized two different monomers equipped with two undecenyl chains for the olefin metathesis starting from POCl₃. **M1**, a phosphodiester with an ethoxy side group were prepared according to literature.^{31,32} In addition, a new monomer, **M2**, was prepared by the reaction of 10 equivalent phosphorus oxychloride with 1 equivalent of 2-bromoethanol and in a second step with 2 equivalents of

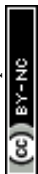
10-undecen-1-ol, in order to install the alkyl-bromide functionality as a side chain. The excessive 10-undecen-1-ol was removed by flash chromatography over alumina. The resulting monomer is a liquid at room temperature and is soluble in aromatic and halogenated solvents (*e.g.* toluene, dichloromethane, chloroform) and insoluble in water. The alkyl-bromide in the pendant chain acts as versatile electrophilic functionality, as it can be functionalized by various nucleophilic substitutions after the synthesis or after polymerization (Scheme 1 shows all chemical structures and reactions).

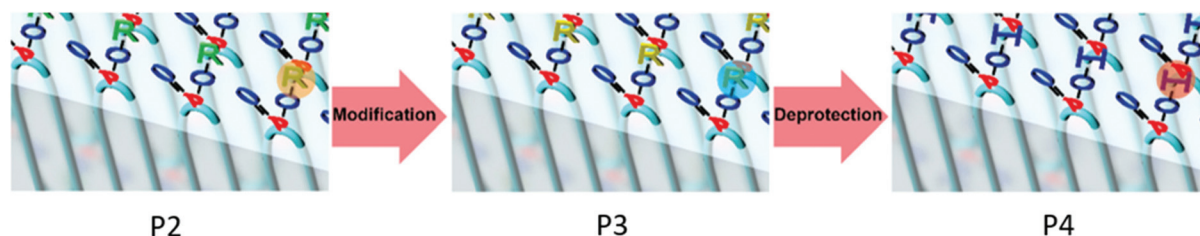
The phosphate groups in the polymer chain are precisely spaced after the ADMET polymerization. The aliphatic methylene spacer between them was adjusted to 20 units (–(CH₂)₂₀–) to guarantee their ability to crystallize. If crystallized from dilute solution, this resulted in lamellar crystals with the phosphate groups on two opposing sides. With the possibility of installing chemical functionality in the phosphate unit, this strategy gives access to polymer platelets bearing functional groups on their surface which are accessible for further surface chemistry (Scheme 2).

To prove that the phosphate groups are expelled from the crystal and arranged in a single plain, **P1** was synthesized from **M1** according to literature³² and analyzed by transmission electron microscopy (TEM). Previous studies on the crystallization of PPEs with varying lengths of methylene spacers between the phosphate units revealed a transition from pseudo hexagonal an to a orthorhombic phase with increasing chain-length.³³ From the electron diffraction pattern in Fig. 1a the only observed lattice distance d_{hkl} amounts to 4.1 Å. When assuming an orthorhombic unit cell, as it is common for polyethylenes, the (110) and the (200) (110) $\approx$ (200). The lattice constants extracted from this assumption amount to $a = 8.2$ Å and $b = 4.7$ Å, where a becomes nearly equal to $b\sqrt{3}$ and hence the unit cell becomes pseudo-hexagonal.³⁴ Using the unit cell parameters obtained by transmission electron microscopy studies, different models for the local planar arrangement of



Scheme 1 Overview of the monomers used to synthesize the functional PE-like polymers by ADMET polycondensation with subsequent hydrogenation and functionalization.





Scheme 2 Schematic representation of the multi-step reaction, which was conducted “on surface” of the polymer platelets: 1. step: modification of the Br-group to a thioacetate-group. 2. step: cleavage of the 2-acetylthioethyl protective group.

the ^{31}P sites relative to the orthorhombic/pseudo-hexagonal lattice of the aliphatic chains can be derived (Fig. 2 top) and compared to structural information gathered by solid-state NMR.

Commonly used solid-state NMR methods like ^{31}P magic angle spinning (MAS) NMR or $^{31}\text{P}\{^1\text{H}\}$ CP-MAS NMR may probe the local chemical environment of the ^{31}P sites in the solid-state, however, they will not provide any information on spatial proximities of the chemically isolated ^{31}P sites in **P1**. Utilizing double-quantum (DQ) MAS-NMR methods, it is possible to probe molecular proximities in the range of 0.5 nm and thus gaining information regarding conformations and packing.³⁵ In addition to ^{31}P - ^{31}P double-quantum single-quantum (DQ-SQ) correlation experiments, probing directly spatial proximities between chemically distanced ^{31}P sites, this method can probe as well ^{31}P - ^{31}P DQ dipolar coupling between chemically equivalent ^{31}P sites based on measurements of build-up curves. These build-up curves (Fig. 1c) are sensitive to P-P inter-nuclear distances, as the nuclear dipolar

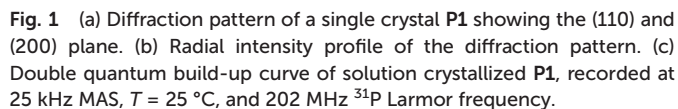
coupling constant $D_{ij} = \frac{\mu_0 \hbar^2 \gamma_i \gamma_j}{4\pi^2 r_{ij}^3}$ between two spins i and j is proportional to the inverse cube of the inter-nuclear distance r_{ij} , and the strongest ^{31}P dipolar coupling, thus the closest spatial proximity, determines predominantly the initial t^2 -proportional rising of the double quantum build-up curve (with μ_0 the vacuum permittivity; $\hbar$ the Planck constant; γ_i and γ_j are the nuclear gyro-magnetic ratios of the two spins involved in the dipolar coupling). Moreover, the long term behavior of DQ build up curves provides qualitative information on the distribution of the ^{31}P dipolar couplings and spatial proximities. In crystalline arrangements with well-defined distinct distances between neighboring ^{31}P sites, oscillatory behavior of the DQ build-up curves is observed (see Fig. 6b, in the publication by Saalwächter *et al.*³⁶). For very long excitation times, these oscillations vanish in powder samples due to the orientation dependence of the DQ excitation efficiency. In less controlled assemblies with broad distributions of distances and thus dipolar couplings, the oscillatory behavior of the DQ build-up curve vanishes due to destructive interferences of the different oscillation frequencies of discrete dipolar coupling values completely. In these cases, a second moment description of the dipolar coupling assuming a dipolar interaction of a single site with continuously distributed surrounding ^{31}P sites is

more appropriate and provides a better description of the DQ build-up behavior. The smooth DQ build-up curve with its quadratic initial behavior obtained for the local packing of the ^{31}P sites in crystallized **P1** samples shown in Fig. 1 indicates that the second moment analysis of the local packing should be appropriate (*i.e.* indicating a crystalline, planar arrangement of P on the crystal surface). Please note, that the fluctuations occurring in the DQ build-up curve for longer DQ excitation times result from increasing relative experimental uncertainties with increasing DQ excitation times and thus do not indicate the presence of discrete dipolar coupling values resulting from a crystalline lattice (which should vanish for long DQ excitation times) like the oscillations mentioned above.

In Fig. 2, we have drawn three possibilities, how the polymer chains may propagate along with the unit cell of the crystallites, with the phosphate groups positioned between the lattice positions of the alkyl chain in an alternating pattern as illustrated in Fig. 2. For clarity, only the phosphorus atoms of the top surface are shown. In Model 1 (Fig. 2a) the polymer chain propagates diagonal to the unit cell along the (110) direction of the crystal. In Model 2 (Fig. 2b), the polymer chain propagates along the b axis of the unit cell, but the neighboring polymer chain is shifted by $b/2$. In Model 3 (Fig. 2c), the polymer chains propagate along the b axis, but the shift alternates between $+$ and $-b/2$.

From the different models, the distance r_{ij} between two phosphate groups in close spatial proximity is known and thus can be used to calculate the ^{31}P dipolar second moment (M_2) $\left(M_2 = \frac{3}{5} \gamma^4 \hbar^2 I(I+1) \sum_j r_{ij}^{-6}\right)^{37}$ and compare it to the M_2 obtained by the experimental DQ build-up curves of 0.197 kHz^2 (for details on data acquisition, processing and analysis see Solid-state NMR Experimental section). For the calculation of M_2 , five unit cells in each direction were taken into account. Note that the integral contribution of more remote ^{31}P sites to the value of M_2 is less than 0.1%, as their contribution to the second moment decreases with the 6th power of the distance. For all 3 models, the calculation yields a M_2 of $\sim 0.14 \text{ kHz}^2$ with only minor variations, which is significantly lower than the M_2 value obtained from the experimental DQ build-up curves. However, the crystals were measured in a dry state, so that the stacking of multiple crystalline layers is possible.





The chemical accessibility of the pendant chains in the phosphates at the surface of the polymer platelets was studied using a multi-step reaction, which was conducted “on surface” (Scheme 2). The polymer platelets were prepared by solution crystallization of **P2**, which was prepared as followed: **M2** was polymerized *via* ADMET (**P2a**, $M_n = 20\,500\text{ g mol}^{-1}$, $M_w/M_n = 2.07$), the purified polymer was then hydrogenated using a Pd/C catalyst to give **P2** ($M_n = 15\,900$, $M_w/M_n = 1.67$). To prepare the dispersion of the polymer platelets of **P2**, the polymer was dissolved in ethyl acetate at 60 °C at a concentration of 1 mg mL⁻¹. Then the solution was cooled in a temperature bath, which was set to 20 °C. A part of the cooled solution was transferred to a separate vial as a reference for later TEM images. The first reaction step “on surface” was the nucleophilic substitution of the bromides in the polymer platelets of **P2** by potassium thioacetate to **P3**. Afterwards, a small amount of sample was taken and the solvent was removed at reduced pressure. The dried polymer platelets were then dissolved in deuterated chloroform and ¹H NMR spectroscopy proved the characteristic resonances at 3.17 ppm

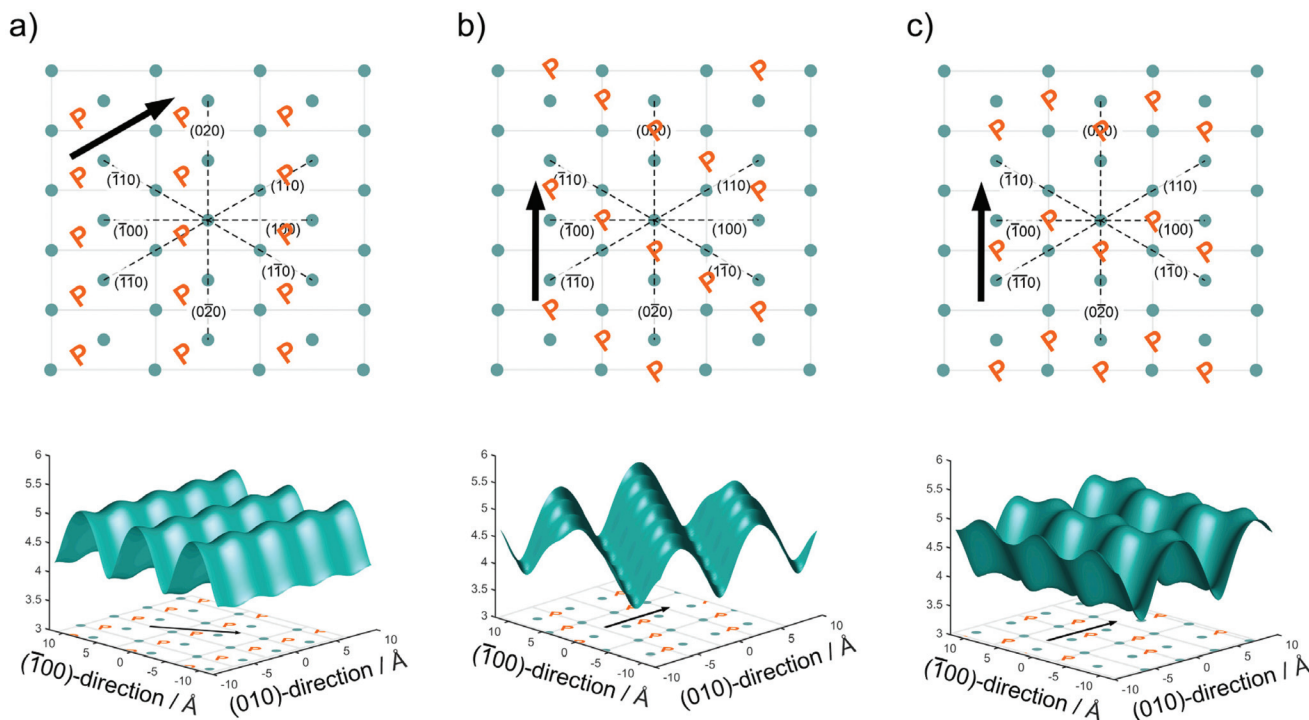


Fig. 2 Arrangement of the P atoms on the crystal surface depending on the direction of the chain folding along the crystal lattice, indicated by the black arrow (a) polymer chain propagating along the (110) direction of the crystal lattice (Model 1). (b) Polymer chain propagating along the *b* axis of the crystal shifted by *b*/2 (Model 2). (c) Polymer chain propagating along the *b* axis of the crystal shifted by *b*/2 alternating (Model 3). (bottom) Plane on which M_2 equals the experimentally obtained value of 0.197 kHz².

(methylene group next to the thioacetate group) and 2.35 ppm (methyl group of the thioacetate group) for **P3** (Fig. S11†) for a successful modification. When the same reaction was conducted in solution with **P2**, the NMR spectra of the product proved the same resonances for **P3** (Fig. S9†). The ³¹P NMR spectrum shows a single signal at −1.13 ppm (Fig. S10†) confirming only one phosphorus species present. The TEM images in Fig. S12 and S13† prove the unchanged lateral dimensions of the polymer platelets of **P2** and **P3** before and after the “on surface” reaction. In DSC measurements, **P3** shows a T_g of −49 °C and a T_m of 58 °C (Fig. 3). The second reaction step was the hydrolysis of the 2-acetylthioethyl protective group, which was conducted “on surface” as well. The 2-acetylthioethyl ester group is a protective group for phosphoric acid (P–OH-groups) and can be cleaved under acidic or basic conditions to release P–OH functionality, which should alter the surface properties of the polymer platelets.³⁸

The dispersion was cooled to 0 °C for 15 min before starting stirring and adding the hydrazine in ethanol. After *ca.* 45 min, aggregation of the crystals was visible. The solvent was allowed to evaporate while keeping the temperature at 0 °C. The product (**P4**) was not soluble in any solvent, as expected for a polyphosphodiester.³¹

Fig. 4 shows the TEM images of the crystal dispersion before (**P3**) (Fig. 4a and b) and after (**poly6**) (Fig. 4c and d) the addition of hydrazine. Before the treatment, the crystals exhibited a size of around 128 ± 40 nm and are crystalline as it is

apparent from the diffraction pattern. After the addition of the hydrazine, the crystal size remained at around 132 ± 46 nm and the diffraction pattern still proves their crystallinity. The P–OH groups on the crystal surface induced hydrogen bonds, which lead to the stacking of the crystals, with P–OH as an H-bond donor and P=O as an H-bond acceptor.³¹ Fig. 4c shows a TEM image of the crystals stacking on top of each other.

IR spectroscopy proved the successful cleavage of the thioacetyl-protective group and the formation of the P–OH groups as shown in Fig. 5. The C=O stretching frequency at 1694 cm^{−1} and the C(O)–S stretching frequency at 625 cm^{−1}, which are characteristic for thioacetates,³⁹ vanished and a new resonance at 1200 cm^{−1} appeared, which is characteristic for P=O stretching in acidic phosphates.⁴⁰

Experimental section

Materials and methods

All chemicals were purchased from commercial suppliers as reagent grade and used without further purification.

NMR. ¹H and ³¹P nuclear magnetic resonance (NMR) spectra were recorded on Bruker AV 300 at 300 MHz (¹H) and 121.50 MHz (³¹P) or Bruker AV 700 spectrometers at 700 MHz (¹H). The temperature of measurement is indicated in the corresponding figure captions. Chemical shifts are reported in

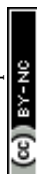




Fig. 3 (a) Transmission electron microscope micrograph of the single **P3** crystals (0.1 mg mL^{-1}). (b) Differential scanning calorimetry (baseline corrected) of **P3** showing the second heating and cooling curve with a T_g at -49°C and T_m at 58°C determined in the second heating curve.



Fig. 4 (a) EM micrograph of 1 mg mL^{-1} **P3**. (b) EM diffraction pattern of 1 mg mL^{-1} **P3**. (c) Transmission electron microscopy image of 1 mg mL^{-1} **P3**. (d) Transmission electron microscopy diffraction pattern of 1 mg mL^{-1} **P4**.

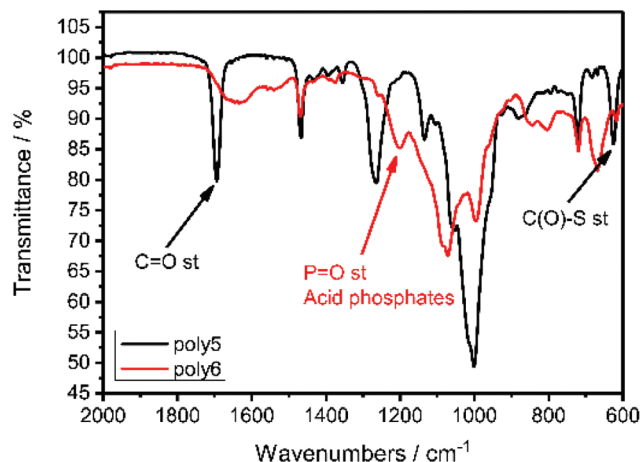


Fig. 5 FTIR spectrum of **P3** and **P4** (after two step reaction) highlighting the important frequencies.

ppm using the residual non-deuterated solvent signals as an internal reference.

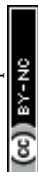
DSC. The thermal properties of the synthesized polymers have been measured by differential scanning calorimetry (DSC) on a Mettler Toledo DSC 823 calorimeter. Three scanning cycles of heating/cooling were performed in a nitrogen atmosphere (30 mL min^{-1}) with a heating and cooling rate of $10^\circ\text{C min}^{-1}$.

SEC. Size exclusion chromatography (SEC) measurements were performed in THF on an Agilent Technologies 1260 instrument consisting of an autosampler, pump and column oven. The column set consists of 3 columns: SDV $10^6 \text{ \AA}$, SDV $10^4 \text{ \AA}$ and SDV $500 \text{ \AA}$ (PSS Standards Service GmbH, Mainz, Germany), all of $300 \times 8 \text{ mm}$ and $10 \text{ \mu m}$ average particle size were used at a flow rate of 1.0 mL min^{-1} and a column temperature of 30°C . The injection volume was $100 \text{ \mu L}$. Detection was accomplished with an RI detector (Agilent Technologies). The data acquisition and evaluation were performed using PSS WINGPC UniChrom (PSS Polymer Standards Service GmbH, Mainz, Germany). Calibration was carried out by using polystyrene provided by PSS Polymer Standards Service GmbH (Mainz, Germany).

Transmission electron microscopy (TEM). The crystal morphology, thickness, and crystal structure were determined using an FEI Tecnai F20 transmission electron microscope operated at an acceleration voltage of 200 kV .

The crystallization behavior of these polymers was studied by drop-cast TEM measurement. A TEM specimen of solution-grown polymer crystals was prepared from a 1 mg mL^{-1} or 0.1 mg mL^{-1} dispersion in ethyl acetate. The solution was heated and cooled according to the different temperature profiles. One drop of each dispersion was applied to a carbon-coated TEM grid, excess liquid was blotted off with a filter paper and the specimen was allowed to dry under ambient conditions.

Solid-state NMR. The solid-state NMR measurements have been performed on a Bruker Avance III console operating at



P4: A vial was charged with **P3** (30 mg, 59.0 μ mol, 1 eq.) and ethyl acetate (30 mL). The dispersion was heated to 60 $^{\circ}$ C and allowed to cool to R.T. for 4–5 h. Then the crystal dispersion was cooled to 0 $^{\circ}$ C and hold at this temperature for 15 min. The stirring was turned on and 1 M hydrazine solution in ethanol (2 mL, 2 mmol, 40 eq.) was added while keeping the temperature at 0 $^{\circ}$ C. After 24 h the solvent was evaporated with a flow of nitrogen while still keeping the temperature at 0 $^{\circ}$ C.

Conclusion

We utilized phosphorus chemistry and simple polycondensation to introduce precisely spaced defects in polyethylene like polymers. We could demonstrate, that the relatively large size of the phosphate groups allow the control of the chain-folding during crystallization. The molecular arrangement in the crystal was studied by TEM and solid-state NMR, which proved a pseudohexagonal crystal structure with the phosphate groups emanating from the two opposing surfaces of the crystal. With chemical functionality in the pendant phosphoester groups, we pave the way for conducting chemistry “on surface” of anisotropic nanoplatelets, which was proven by nucleophilic substitution and hydrolysis reaction. The long-chain PE-like polyphosphates can be used as a general platform to design chemically functional anisotropic nanomaterials for various applications from the biofield to materials science, with the possibility of degradation of the phosphoester bonds combined with the crystallinity of polyethylene.

Author contributions

The manuscript was written through the contributions of all authors. All authors have approved the final version of the manuscript.

Conflicts of interest

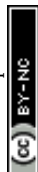
There are no conflicts to declare.

Acknowledgements

The authors thank the Deutsche Forschungsgemeinschaft (DFG WU 750/8-1) for funding. The authors thank Christine Rosenauer for the DLS measurements. The authors thank Prof. Dr Katharina Landfester (MPI-P, Germany) for her continuous support. Open Access funding provided by the Max Planck Society.

References

- 1 R. Geyer, J. R. Jambeck and K. L. Law, *Sci. Adv.*, 2017, **3**, e1700782.
- 2 T. E. Nowlin, *Business and Technology of the Global Polyethylene Industry: An In-depth Look at the History, Technology, Catalysts, and Modern Commercial Manufacture of Polyethylene and Its Products*, Wiley, 2014.
- 3 T. M. Karlsson, L. Arneborg, G. Broström, B. C. Almroth, L. Gipperth and M. Hassellöv, *Mar. Pollut. Bull.*, 2018, **129**, 52–60.
- 4 J. M. Kelly, *J. Macromol. Sci., Part B: Phys.*, 2002, **42**, 355–371.
- 5 S. Hambir and J. P. Jog, *Bull. Mater. Sci.*, 2000, **23**, 221–226.
- 6 B. M. Sole and A. Ball, *Tribol. Int.*, 1996, **29**, 457–465.
- 7 G. Liu, Y. Chen and H. Li, *Wear*, 2004, **256**, 1088–1094.
- 8 L. A. Goettler, K. Y. Lee and H. Thakkar, *Polym. Rev.*, 2007, **47**, 291–317.
- 9 A. Osichow, C. Rabe, K. Vogtt, T. Narayanan, L. Harnau, M. Drechsler, M. Ballauff and S. Mecking, *J. Am. Chem. Soc.*, 2013, **135**, 11645–11650.
- 10 B. Scheinhardt, J. Trzaskowski, M. C. Baier, B. Stempfle, A. Oppermann, D. Wöll and S. Mecking, *Macromolecules*, 2013, **46**, 7902–7910.
- 11 K. Busse, C. Fuchs, N. Hasan, M. Pulst and J. Kressler, *Langmuir*, 2018, **34**, 12759–12763.
- 12 Y.-R. Zheng, H. T. Tee, Y. Wei, X.-L. Wu, M. Mezger, S. Yan, K. Landfester, K. Wagener, F. R. Wurm and I. Lieberwirth, *Macromolecules*, 2016, **49**, 1321–1330.
- 13 K. N. Bauer, H. T. Tee, M. M. Velencoso and F. R. Wurm, *Prog. Polym. Sci.*, 2017, **73**, 61–122.
- 14 C. Pelosi, M. R. Tinè and F. R. Wurm, *Eur. Polym. J.*, 2020, **141**, 110079.
- 15 T. Haider, O. Shyshov, O. Suraeva, I. Lieberwirth, M. von Delius and F. R. Wurm, *Macromolecules*, 2019, **52**, 2411–2420.
- 16 N. Hasan, C. Schwieger, H. T. Tee, F. R. Wurm, K. Busse and J. Kressler, *Eur. Polym. J.*, 2018, **101**, 350–357.
- 17 J. Mandal, S. Krishna Prasad, D. S. S. Rao and S. Ramakrishnan, *J. Am. Chem. Soc.*, 2014, **136**, 2538–2545.
- 18 P. H. Till Jr., *J. Polym. Sci.*, 1957, **24**, 301–306.
- 19 D. J. Blundell, A. Keller and A. J. Kovacs, *J. Polym. Sci., Part B: Polym. Lett.*, 1966, **4**, 481–486.
- 20 S. A. E. Boyer and J. M. Haudin, *Polym. Test.*, 2010, **29**, 445–452.
- 21 T. Miyata and T. Masuko, *Polymer*, 1998, **39**, 5515–5521.
- 22 N. Bosq, N. Guigo, E. Zhuravlev and N. Sbirrazzuoli, *J. Phys. Chem. B*, 2013, **117**, 3407–3415.
- 23 L. Mandelkern, *J. Appl. Phys.*, 1955, **26**, 443–451.
- 24 G. Guerin, G. Molev, D. Pichugin, P. A. Rupa, F. Qi, M. Cruz, I. Manners and M. A. Winnik, *Macromolecules*, 2019, **52**, 208–216.
- 25 F. Marsico, M. Wagner, K. Landfester and F. R. Wurm, *Macromolecules*, 2012, **45**, 8511–8518.
- 26 T. Steinbach, E. M. Alexandrino, C. Wahlen, K. Landfester and F. R. Wurm, *Macromolecules*, 2014, **47**, 4884–4893.
- 27 T. Steinbach, E. M. Alexandrino and F. R. Wurm, *Polym. Chem.*, 2013, **4**, 3800–3806.
- 28 M. Steinmann, J. Markwart and F. R. Wurm, *Macromolecules*, 2014, **47**, 8506–8513.
- 29 H. Mutlu, L. M. de Espinosa and M. A. R. Meier, *Chem. Soc. Rev.*, 2011, **40**, 1404–1445.
- 30 H. Li, L. Caire da Silva, M. D. Schulz, G. Rojas and K. B. Wagener, *Polym. Int.*, 2017, **66**, 7–12.
- 31 H. T. Tee, K. Koynov, T. Reichel and F. R. Wurm, *ACS Omega*, 2019, **4**, 9324–9332.
- 32 K. N. Bauer, H. T. Tee, I. Lieberwirth and F. R. Wurm, *Macromolecules*, 2016, **49**, 3761–3768.



- 33 T. Haider, O. Suraeva, M. L. O'Duill, J. Mars, M. Mezger, I. Lieberwirth and F. R. Wurm, *Polym. Chem.*, 2020, **11**, 3404–3415.
- 34 O. Ruiz de Ballesteros, F. Auriemma, G. Guerra and P. Corradini, *Macromolecules*, 1996, **29**, 7141–7148.
- 35 K. Saalwächter, in *Modern Magnetic Resonance*, ed. G. A. Webb, Springer International Publishing, Cham, 2018, pp. 755–781, DOI: 10.1007/978-3-319-28388-3_59.
- 36 K. Saalwächter, F. Lange, K. Matyjaszewski, C.-F. Huang and R. Graf, *J. Magn. Reson.*, 2011, **212**, 204–215.
- 37 P. A. Abragam and A. Abragam, *The Principles of Nuclear Magnetism*, Clarendon Press, 1961.
- 38 J. C. Markwart and F. R. Wurm, *Tetrahedron*, 2018, **74**, 7426–7430.
- 39 A. M. M. El-Assar, C. P. Nash and L. L. Ingraham, *Biochemistry*, 1982, **21**, 1972–1976.
- 40 G. Socrates, *Infrared and Raman Characteristic Group Frequencies: Tables and Charts*, Wiley, 2004.
- 41 K. Saalwächter, *Prog. Nucl. Magn. Reson. Spectrosc.*, 2007, **51**, 1–35.

