

1 Supplementary materials

2 Thermal properties

3 The presence of GnP in polymer matrix is expected to influence the crystallization degree of
 4 nanocomposites [1,2]. Therefore, the impact of GnP nanofiller on composite crystallinity was
 5 examined using Mettler Toledo DSC instrument. DSC analyzes were performed on 2 ÷ 5 mg samples
 6 from -50 °C to 150 °C with a scan rate of 10 °C/min in a heating-cooling-heating cycle under N₂
 7 atmosphere. The degree of crystallization was calculated as follows:

$$\chi_c(\%_{crystallinity}) = \frac{\Delta H_m}{\Delta H_0} * 100\%, \quad (S1)$$

8 Where: ΔH_m - is the melting enthalpy of LDPE (for the composites multiplied by the content of
 9 pure LDPE), and ΔH_0 - is a theoretical value of the melting enthalpy of 100% crystalline LDPE. The
 10 value $\Delta H_0 = 293$ J/g was used in the crystallinity calculations [3].

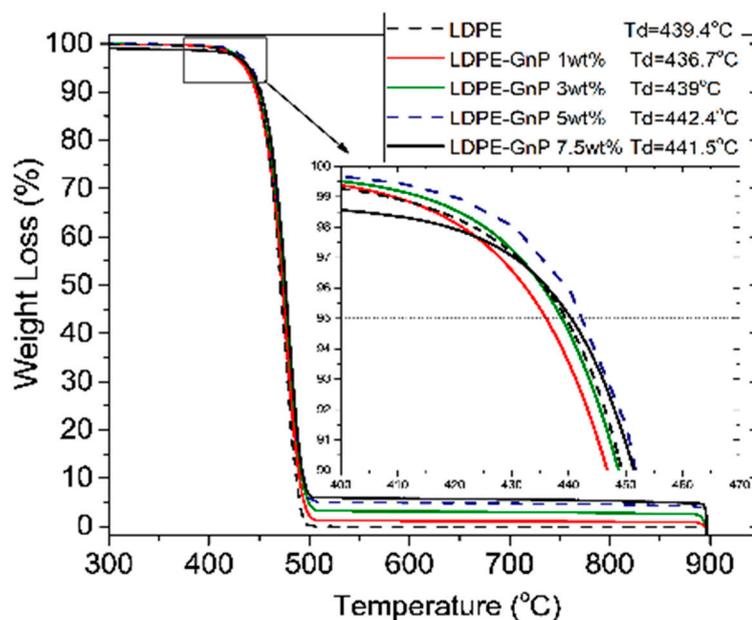
11 Table 1 shows crystallization and melting temperatures determined from DSC thermograms as
 12 well as crystallization and melting enthalpies. One can notice that the crystallinity degree reduces
 13 slightly, only for sample containing 7.5 wt.% of GnP, whereas for other composites its change is
 14 negligible. The same behavior is visible when considering the filler influence on crystallization and
 15 melting temperatures.

16 **Table S1.** The crystallization and melting parameters determined from DSC thermograms of
 17 LDPE-GnP composites.

Sample	T _c (°C) Crystallization temperature	ΔH _c (J/g) Crystallization enthalpy	T _m (°C) Melting temperature	ΔH _m (J/g) Melting enthalpy	χ _c (%) Crystallinity degree
LDPE	98.2	132.5	110.6	134.5	45.9
GnP 1wt%	103.6	124.3	107.9	131.8	45.0
GnP 3wt%	103.8	120.6	109.2	133.4	45.5
GnP 5wt%	103.8	126.5	108.0	134.6	45.9
GnP 7.5wt%	104.0	120.4	108.8	126.9	43.3

18 The filler content and the degradation temperature of all specimens were studied by means of
 19 TGA/DSC 3+ (Mettler Toledo, Inc., Greifensee, Switzerland). TGA measurements were carried out in
 20 N₂ atmosphere with a heating rate of 20 °C/min. 2 ÷ 5 mg samples were heated starting from 30 °C
 21 up to 900 °C and kept at 900 °C for 10 min in O₂ atmosphere.

22 TGA analyses have confirmed the content of nanofiller as well as allowed to evaluate the
 23 degradation temperature T_d of the nanocomposites. The extracted T_d values, defined as the
 24 temperature of 5 % weight loss, are presented in the insert of Figure 1. This parameter is a very
 25 important feature of polymeric materials concerning their applications. It is expected that
 26 incorporation of fillers with high degradation temperature influences the degradation temperature
 27 of nanocomposites. This effect is not visible for 1% and 3% of filler content, however it starts to be
 28 visible for samples filled with 5 wt. % and 7.5 wt. % of GnP where the curves are shifted to the higher
 29 temperatures.
 30
 31



32 **Figure S1.** TGA curves measured for investigated nanocomposites. The insert provides deduced
 33 degradation temperatures T_d .

34 References

- 35 1. Kalaitzidou, K.; Fukushima, H.; Askeland, P.; Drzal, L.T. The nucleating effect of exfoliated
 36 graphite nanoplatelets and their influence on the crystal structure and electrical
 37 conductivity of polypropylene nanocomposites. *Journal of Materials Science* **2008**, *43*, 2895-
 38 2907.
- 39 2. Jiang, X.; Drzal, L.T. Multifunctional high-density polyethylene nanocomposites produced
 40 by incorporation of exfoliated graphene nanoplatelets 2: Crystallization, thermal and
 41 electrical properties. *Polymer Composites* **2012**, *33*, 636-642.
- 42 3. Morawiec, J.; Pawlak, A.; Slouf, M.; Galeski, A.; Piorkowska, E.; Krasnikowa, N.
 43 Preparation and properties of compatibilized ldpe/organo-modified montmorillonite
 44 nanocomposites. *European Polymer Journal* **2005**, *41*, 1115-1122.

45