Immobilization of Polymeric Luminophor on Nanoparticles Surface

Yuliia Bolbukh1*, Beata Podkoscielna2, Agnieszka Lipke2, Andrzej Bartnicki2, Barbara Gawdzik2 and Valentin Tertykh1

Abstract

Polymeric luminophors with reduced toxicity are of the priorities in the production of lighting devices, sensors, detectors, bioassays or diagnostic systems. The aim of this study was to develop a method of immobilization of the new luminophor on a surface of nanoparticles and investigation of the structure of the grafted layer. Monomer 2,7-(2-hydroxy-3-methacryloyloxypropoxy)naphthalene (2,7-NAF.DM) with luminophoric properties was immobilized on silica and carbon nanotubes in two ways: mechanical mixing with previously obtained polymer and by in situ oligomerization with chemisorption after carrier’s modification with vinyl groups. The attached polymeric (or oligomeric) surface layer was studied using thermal and spectral techniques. Obtained results confirm the chemisorption of luminophor on the nanotubes and silica nanoparticles at the elaborated synthesis techniques. The microstructure of 2,7-NAF.DM molecules after chemisorption was found to be not changed. The elaborated modification approach allows one to obtain nanoparticles uniformly covered with polymeric luminophor.

Keywords: Multiwalled carbon nanotubes, Silica nanoparticles, Polymeric luminophor, Chemisorption, Luminescence properties

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Background

Materials with luminophor properties are widely used in the production of lighting devices, sensors, detectors, bioassays or diagnostic systems. The most ambitious and fascinating application of luminescent nanomaterials is probably related to medicine and molecular biology. Luminescent nanomaterials are promising tags for optical imaging and fluorescent labelling to allow for novel techniques of non-invasive diagnosis and in vivo observation of complex vital functions [1]. The recent advances in nanotechnology have demanded to raise a new class of fluorescent labels. Researchers have developed several different types of luminescent nanomaterials, including semiconductor quantum dots, dye-doped nanoparticles, up-converting lanthanide-doped nanoparticles and carbon nanodots on silica surface [2]. From this point of view, the optical properties of carbon nanotubes have attracted much recent attention. For example, there have been extensive investigations of carbon nanotubes as optical limiters for attenuating pulsed laser irradiation [3]. Apparently, the functionalization of the nanotubes resulted in more intense luminescence emissions [4]. Along with existing nanoscale materials [5, 6], modified nanotubes with luminescence properties have potential as materials for bioanalytical assays and luminescent imaging [7]. That is why the development of composites based on polymeric luminophor and nanoparticles attracted a considerable interest. One of the priorities is the development of polymer luminophors with reduced toxicity. Such polymers can extend the luminophor applications in biotechnology and medicine, including the creation of biomarkers. Recent advances in surface grafting of polymers onto carbon materials, such as carbon black, graphite powder and carbon nanotubes, and the application of polymer-grafted carbon materials as novel functional materials have been mainly reviewed [8–10]. The grafting of polymers onto the surface can be achieved by (1) grafting onto process [11], (2) grafting from process [12], (3) polymer reaction process [13] and (4) stepwise growth by dendrimer synthesis methodology.

* Correspondence: yu_bolbukh@yahoo.com
1 Chuiiko Institute of Surface Chemistry of National Academy of Sciences of Ukraine, 17 General Naumov Str., 03164 Kyiv, Ukraine
Full list of author information is available at the end of the article

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For polymer anchorage, the surface functional groups, such as carboxyl and phenolic hydroxy groups, were used as grafting sites. For example, in the living cationic polymerization initiated by carbon black/ZnCl₂ system, carboxyl groups on the surface acted as initiating and grafting sites. The in situ atom transfer radical polymerization (ATRP) grafting from approach was successfully applied to graft poly(methyl methacrylate) (PMMA) onto the convex surface of multiwalled carbon nanotubes (MWCNTs). The thickness of the coated polymer layers can be conveniently controlled by the feed ratio of MMA to preliminarily functionalized MWCNT (MWCNT-Br). Moreover, the approach has been extended to the copolymerization system, affording novel hybrid core–shell nano-objects with MWCNTs as the core and amphiphilic poly(methyl methacrylate)-block-poly(hydroxystyryl methacrylate) (PMMA-b-PHEMA) as a shell.

Earlier, we elaborated the approach that allows to obtain carbon nanotube-based nanomaterials with tailor-made structure and properties. The MWCNTs were modified with monomer 2-hydroxyethylmethacrylate using radical process or polycondensation reaction to obtain different functional groups, namely hydrophobic (methacrylic) or hydrophilic (hydroxyl), in the boundary layer. Along with silica surface functionalization, the modification of MWCNTs with functional silanes was evaluated and optimized based on accumulated experience.

Background of the Study
For polymeric luminophors, the fluorescent light mostly appears in a thin surface layer without participation of the polymer in volume. Obtaining of the polymeric surface layer on a surface of nanoscale particles while maintaining its luminescent properties is of considerable practical interest. The high specific surface of nanoparticles supposed should promote better contact of luminophor with light radiation, and thus, luminescence should increase at lesser amounts of polymer. Moreover, utilization of the nanoparticles with immobilized luminophor has to provide target localization light-sensitive markers. The aim of this study was to develop a method of immobilization of the monomeric luminophor on a surface of nanoparticles based on elaborated approaches and investigation of the structure of the grafted layer in the modified products.

Methods
The monomer 2,7-(2-hydroxy-3-methacryloyloxypropoxy)naphthalene (2,7-NAF.DM) was used for immobilization. The oxidized H₂O₂ mult-walled nanotubes produced in the Chuiko Institute of Surface Chemistry (Ukraine) using the catalytic vapour deposition method and pyrogenic silica with a specific surface area of 260 m²/g (A-300, Kalush, Ukraine) were utilized as carriers. In order to improve monomer–nanoparticle interaction, nanotubes and silica were activated via attaching of the vinyltriethoxysilane (Fluka) that was used without addition treatment. Analytical grade solvents were purchased from Sigma-Aldrich.

Synthesis of Monomer 2,7-NAF.DM
The 2,7-NAF.DM was obtained in the two-step reaction. In the first step, naphthalene-2,7-diol (2,7-NAF) was reacted with epichlorohydrin (EP) in the two-phase liquid/liquid system including organic and aqueous phases. Next, esterification of the obtained diglycidyl ether was carried out using methacrylic acid (MA) in the presence of hydroquinone (polymerization inhibitor) and triethylbenzylammonium chloride as a catalyst. Finally, the aromatic monomer (2,7-NAF.DM) with photoluminescent properties was prepared.

Luminophor 2,7-NAF.DM Immobilization
Liquid-phase adsorption and chemisorption were applied for 2,7-NAF.DM immobilization on the surface of nanoparticles. Before modification with 2,7-NAF.DM, the silica (SiO₂) and MWCNTs were activated by treatment with vinyl-containing trialkoxysilane (VTES). The modification of silica or carbon nanotubes was carried out in toluene solution of VTES with ethylenediamine as initiator of condensation of surface silanols (or phenolic groups on nanotube surface) with alkoxy groups. The process was stimulated by ultrasonic treatment and heating at 90 °C for 6 h in a closed reactor. Then, vinylated nanoparticles were dried at 110 °C until the complete solvent removal. As expected, the vinyl groups on the
nanoparticles surface provide better affinity between the carriers and luminophor (2,7-NAF.DM) [24].

The modification process with 2,7-NAF.DM was carried out from toluene solution of monomer using azobisisobutyronitrile (AIBN) as attachment and polymerization initiator with heat and ultrasonic stimulation. In brief, demanded amount of the vinylated silica was dispersed in toluene solution of monomer by ultrasonic treatment (ultrasonic bath with frequency 44 kHz). After this, the dissolved initiator was introduced into dispersion and sonication was repeated. The composites with different carrier/monomer ratios were prepared (Table 1). Modification process was carried out at 80 °C for 24 h after sonication for 30 min. Then, samples were filtered, washed with toluene and acetone and dried at 110 °C until the complete solvent removal. The mechanical mixtures were prepared via milling of nanoparticles with cured solid polymer using Micro Ball Mill.

The carrier/monomer ratio was established experimentally based on the reaction of the obtained materials on lighting of UV lamp. Assessment of the prospects of the samples was performed with a commercial 12-W UV lamp with radiation range 315–280 and 300–400 nm. For silica (white powder) that was used as a test sample, a luminescence under UV radiation is observed at smaller monomer (polymer) content in comparison with nanotubes (black powder), where luminescence was detected at higher concentrations of 2,7-NAF.DM.

**Characterization of the Samples**

The structure of the grafted surface layer was studied by means of thermal (differential scanning calorimetry (DSC), TGA) and spectral analysis. DSC thermograms were obtained using a DSC NETZSCH 204 calorimeter (NETZSCH, Günzbung, Germany). All thermal scans were carried out in aluminium pans with a sample weight of ~5–10 mg under nitrogen atmosphere (20 mL min⁻¹). Dynamic scans were performed at the heating rate of 10 K min⁻¹ in the temperature range 30–1000 °C. Based on DSC and TGA results, the decomposition temperatures, the final decomposition mass, final decomposition temperature and enthalpy of process (ΔHₜₐ₉) were evaluated.

Room temperature UV–Vis absorption spectra were recorded using a V-660 JASCO spectrophotometer, and emission spectra were obtained with a Photon Technology International spectrofluorimeter equipped with a continuous wave xenon arc (Xe-arc) lamp as a light source. Spectral resolution was maintained at 1 nm. Liquid samples were measured in a quartz cuvette with a 2-mm optical path and solid samples—in appropriate adapters. UV–Vis reflectance spectra were registered using a horizontal sampling integrating sphere (Model PIV-756) connected to the V-660 spectrophotometer. The decay curves of polymer samples were excited at 450 nm by an Nd:YAG pulse laser (Opolette System, Quantel, USA) connected to the Photon Technology International spectrofluorimeter.

**Results and Discussions**

**Thermal Analysis of Nanocomposites Based on Silica**

The polymer thermal decomposition goes through three mass loss stages with the total weight loss of 99 %
(Fig. 1a). The first stage at 90–150 °C corresponded to solvent removal; the second stage at 150–300 °C was attributed to decomposition of terminal groups and ungrafted monomer evaporation; and the stage at 300–450 °C is caused by main chain and benzene ring decomposition (Fig. 1b) [25–27]. The modified VTES silica has the total mass loss of 16 %. Decomposition process consists of a residual solvent removal (98 °C), alkoxy group and terminal vinyl group thermal oxidative destruction (130–250 °C) and degradation of the attached organosilicon layer (250–350 °C) (Fig. 1a, b).

The composite obtained by the mechanical mixing of vinylated silica (sample 3, Table 1) and polymer prepared by radical polymerization in a block (sample 4, silica/polymer ratio 2:1) loses 36.4 % of weight during thermooxidation. The initial decomposition temperature ($T_{\text{in}}$) was observed at 230 °C that is higher than for pure polymer. The thermooxidation goes via two main stages with extrema of the mass removal rate at 308, 360 °C and maximum at 520 °C that can be attributed to the elimination of the polymer (or its degradation products) from the silica surface. The stages of silica surface ethoxy and terminal vinyl group thermooxidation as well as degradation of the terminal methacrylic, ester or hydroxyl groups of polymer in the interval 100–250 °C were not detected. It can be caused by interaction of the functional groups on silica surface with polymer during heating that promotes more thermally stable organic–inorganic composition capable of carbonizing. For the silica with immobilized polymer obtained via in situ polymerization in toluene solution, the thermooxidation process took place in two stages. At a low-temperature stage (300–400 °C) on the DTG curve, there are two peaks of the mass loss with maxima at 305 and 350 °C corresponding to the weight loss due to thermooxidation of the terminal groups [25, 26] and condensed benzene rings during grafted polymer depolymerisation [28]. Unlike degradation of pure polymer, the mass loss peaks of the main stage of thermooxidation were shifted to the lower temperatures.

At the same time, a narrow peak at 471 °C and a broad peak ranging from 450 to 660 °C with extremum at 540 °C (attributed to oxidation of the polymer layer directly on the silica surface) were detected on the DTG thermogram of sample 4 (Fig. 1b). After immobilization on the silica surface (via mechanical milling or by chemisorption), the polymer thermooxidation was observed at the lowest temperatures (Fig. 1c) probably due to silica surface’s ability to promote a thermooxidative
degradation process [25]. The heat of process ($\Delta H$) above 460 °C is decreased, and polymer destruction is completed at the lower temperature.

It should be noted that the degradation rates at the main decomposition stages (closely to 300 and 350 °C) for mechanically mixed and chemisorbed polymer (samples 3 and 4, Fig. 1b) differed, which indicates changes in composite polymeric structure. According to the DSC data (Fig. 2), the mechanically mixed with vinylated silica polymer after elimination of the volatile products at 83 °C has phase transition in the interval 150–270 °C with maximum on the endotherm at 230 °C corresponding to the melting process, which is not accompanied by a weight loss on the TG curve. In the range 300–460 °C on the DSC curve, exotherms with maxima at 330 and 410 °C were observed. These exotherms apparently correspond to chemical interaction of silica surface with products of the polymer thermal decomposition or post-curing of polymer into the surface layer. For chemisorbed polymer, the weak change in heat capacity at 210–230 °C (Fig. 1c, sample 4) can be attributed to glass transition and it can testify an amorphous structure of the attached layer. Also for sample 4, the heat of exothermal process is larger for mechanically mixed polymer, especially at 330 °C.

**Thermal Analysis of Nanocomposites Based on Multiwalled Carbon Nanotubes**

The thermooxidation process of polymer mechanically milled with modified VTES carbon nanotubes (Table 1, sample 5) occurs in three stages with maxima of weight loss rates at 223, 340 and 404 °C with total mass loss of 96 % (Fig. 3a). It should be noted that for oxidized nanotubes, the starting decomposition temperature is 470 °C and the total mass loss is 99 % [29]. After nanotubes modification with silanes, the total mass loss was close to 96–98 % (Fig. 3c). A low content of a residual pitch after thermooxidation of modified with silane nanotubes was due to relatively low temperature of siloxane chains and attached silane oligomer degradation that occurs at temperatures below the nanotubes’ structural benzene ring destruction (closely to 250–320 °C) [30].

The splitting of main degradation peak testifies two kinds of polymer states, namely without and with interaction with the nanotube boundary layer. For composites with nanotube/polymer ratios 1:10 (sample 6) and 1:40 (sample 7), the strong changes in thermooxidation process were not observed (Fig. 3a). But, the unlike silica-based composite, destruction of samples 6 and 7 (Table 1) in the range 250–410 °C occurs in one step with a maximum weight loss at 340 °C. This may indicate uniformity of the grafted layer. The different character of thermooxidation curves of mechanically mixed and chemisorbed polymer above 400 °C (Fig. 3a, curves 5 and 6) indicates the different nature of the polymer/carbon interaction, which leads to the formation of two types of thermooxidation products that decompose at 511 and 673 °C. For the pure polymer, there are no significant changes in the rate of mass loss in this temperature range (Fig. 3a, curve 2). But exotherms at 570 and 630 °C were noted (Fig. 3b, curve 2), whereas for the chemisorbed polymer, the exotherms were observed at about 520 and 660 °C. These data suggest that unlike polymer chemisorbed on silica, not only the interaction with grafted vinyl groups is realized in the surface layer of nanotubes but the orientation of condensed benzene rings of the polymer in the plane of the structural benzene rings of nanotubes is occurred [31, 32].

As well as for the mechanical blend of the polymer with silica (Fig. 2), at heating of carbon nanotubes mixed with the polymer in the region near 223 °C on the DSC curve, the endotherm is detected (Fig. 4, curve 5), but in this case, the process is followed by the weight loss on the TGA curve (Fig. 3, curve 5). However, the endothermic peak at 258 °C (Fig. 4, curve 5) can be attributed to the polymer melting. In the temperature region of the polymer depolymerization, the exotherm at 330 °C, earlier marked for the composite with silica (sample 3, Table 1), was not observed and the exothermal process at 403 °C corresponding to post-curing of polymer goes into endothermic decomposition reaction (Fig. 4, curve 5). The endotherm at 430 °C corresponds to benzene ring degradation. At the polymer chemisorption in the low-temperature region (Fig. 4, curves 6 and 7), appearance of exotherms on the DSC curves testifies the chemical
interaction in the grafted layer. At temperatures above 300 °C, the character of the DSC curves of samples 6 and 7 is similar to that of composite 4 (polymer chemisorption on the silica surface), but the heat of the process is increased. With the grafted polymer content increasing (samples 6 and 7), the heat of the exothermic process at 100–200 °C decreased with extremum shift towards high temperatures (Fig. 4, curve 7), which may be due to compaction of the polymeric layer grafted to the surface.

Implementation of chemisorption with simultaneous polymerization of the monomer on the surface of the vinylated nanotubes results in formation of homogeneous polymeric layer. It was confirmed by TEM images (Fig. 5c). For mechanical mixture, nanotube/poly2,7-NAF.DM material contains a separated phase of polymer and carrier (Fig. 5b).

**Photoluminescent Properties**

UV–Vis absorption spectra of the materials under this study in their powder form and dispersed in toluene are given in Fig. 6. The strong absorption in the UV region, up to 350 nm, obviously originates from naphthalene...
The 450-nm band observed for pure polymer [23] and probably originates from the $\pi$ conjugation in the naphthalene-diepoxymethacrylate moiety was not detected for 2,7-NAF.DM chemisorbed on the silica surface (Fig. 6). The 250-nm peak can be attributed to saturated compounds containing atoms with lone pairs (non-bonding electrons) capable of $n \rightarrow \sigma^*$ transitions. Also, the aromatic residues (mostly phenyl) show bands near 250 nm [33]. Isomerization of polymer can result in 250-nm peak appearances too [34].

For composites based on carbon nanotubes, the presence of the bands at 240, 280 and 322 nm in the spectrum is characteristic, which are more evident in the spectra of dispersions (Fig. 6). Investigation of powders showed that only for chemisorbed polymer the strong bands, namely at 312 and 327 nm, were observed in the spectrum (Fig. 6, spectra 6 and 7).

It has been found that the polymer chemisorption on the modified silica does not worsen its luminescent properties (Fig. 7, curve 4), whereas chemisorption on the modified nanotubes significantly reduces the intensity of excitation and emission bands at wavelengths characteristic for the pure polymer. Mechanical mixing of nanotubes with cured polymer has little effect on its luminescence but does not provide a sufficient composite’s uniformity.

Low intensity of the excitation and emission spectra of the polymer chemisorbed on carbon nanotubes is apparently attributable to a significant influence of the nanotube surface on the mechanism of electron transitions. The nature of surface interactions and conformation of grafted macromolecules are also important. Consequently, to achieve the luminescence intensity, it is necessary to hold the obtained degree of uniformity of the grafted polymer layer and ensure a certain orientation of polymeric macromolecules in the surface layer. It is known that increased interaction between the polymer molecules reduces the intensity of luminescence [23]; at the same time, it is not clear how the nanotube surface interact with the lumino- phor molecules. This aspect, as well as study of conditions...
that allow achieving positive effect of nanotubes surface on light intensity, demands more detailed investigation that will be carried out soon.

**Conclusions**

The methods have been elaborated for luminophor chemisorption on fumed silica and MWCNTs by in situ attaching of the monomer via surface functional groups with simultaneous polymerization of 2,7-NAF.DM. Obtained results confirm the chemisorption of luminophor on the nanotubes and silica nanoparticles. After chemisorption the microstructure of 2,7-NAF.DM molecules is not changed, as opposed to the mechanical mixture. It was shown that the presented method allows one to obtain a homogeneous amorphous polymeric layer on the surface of nanoparticles. The polymer chemisorbed on the silica surface was found to retain its fluorescent properties (emission intensity neat and chemisorbed 2,7-NAF.DM was $3 \times 10^6$ and $2.7 \times 10^6$ a.u., respectively). It was noted that the polymer chemisorbed on MWCNTs has low intensity luminescence and no significant excitation in the wavelength range characteristic of pure polymer due to the peculiarities of orientation of the macromolecules in the surface layer of the carrier. The findings are the basis for optimizing the approaches to polymeric luminophor immobilization to achieve the desired structure of the graft polymer layer in order to improve the luminescence properties of composites based on carbon nanotubes or other carbon nanomaterials.

**Abbreviations**

2,7NAF: naphthalene-2,7-diol; 2,7-NAF.DM: 2,7-(2-hydroxy-3-methacryloyloxypropoxy)naphthalene; AIBN: azobisisobutyronitrile; ATRP: atom transfer radical polymerization; DSC: differential scanning calorimetry; DTA: differential thermal analysis; DTG curve: differential thermogravimetric curve; EP: epichlorohydrin; MA: methacrylic acid; MWCNTs: multiwalled carbon nanotubes; PMMA: poly(methyl methacrylate); PMMA-b-PHEMA: poly(methyl methacrylate)-block-poly(hydroxyethylmethacrylate); TGA: thermogravimetric analysis; UV: ultraviolet radiation; UV–Vis absorption: ultraviolet-visible absorption; VTES: vinyl trialkoxysilane; $\Delta H^\circ$ enthalpy of process.

**Competing Interests**

The authors declare that they have no competing interests.

**Authors’ Contributions**

YuB generated the idea, elaborated the synthetic approach, carried out the immobilization of luminophor and drafted the manuscript. BP synthesized the monomeric and polymeric luminophor. AL studied the photoluminescent properties of the samples. AB designed the reactor construction and synthesized the monomer and polymer. BG coordinated the analytical part of the study. VT coordinated the scientific part of the study and edited the manuscript. All authors read and approved the final manuscript.

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**Author details**

1. Chuiko Institute of Surface Chemistry of National Academy of Sciences of Ukraine, 17 General Naumov Str., 03164 Kyiv, Ukraine.
2. Faculty of Chemistry, Maria Curie-Skłodowska University, pl. Maria Curie-Skłodowskiej 3, 2-031 Lublin, Poland.

**References**

1. Feldmann C (2011) Luminescent nanomaterials. Nanoscale 3:1947–48
2. Shang L, Dong S, Nienhaus GU (2011) Ultra-small fluorescent metal nanoclusters: synthesis and biological applications. Nano Today 6:401–18
3. Vivien L, Lanc P, Riehl D, Hache F, Anglaret E (2002) Carbon nanotubes for optical limiting. Carbon 40:1789–97
4. Lin Y, Zhou B, Martin RB, Henbest KB, Harruff BA, Riggs JE, Guo Z-X, Allard LF, Sun Y-P (2005) Visible luminescence of carbon nanotubes and dependence on functionalization. J Phys Chem B 109(31):14779–82
5. Wang F, Tan WB, Zhang Y, Fan X, Wang M (2006) Luminescent nanomaterials for biological labelling. Nanotechnology 17(1)–13
properties of new copolymers of naphthalene-2,7-diol dimethacrylate and N-vinyl-2-pyrrolidone: synthesis, characterization and properties. J Therm Anal Calorim 116:785–793
21. Sementsov V, Melezhik AV, Prirkhodko GP et al (2007) Synthesis, structure, physicochemical properties of carbon nanomaterials. In: Shak AP, Gorbik PP (eds) Physics and chemistry of nanomaterials and supramolecular structures, vol 2. Naukova Dumka, Kyiv, pp 116–158
22. Podkoščina B, Gawdzik B (2010) Influence of diluent compositions on the porous structure of methacrylate derivatives of aromatic diols and divinylbenzene. Appl Surf Sci 256:2462–67
23. Podkoščina B, Lipke A, Gawdzik B, Majdan M (2015) Synthesis, characterization and luminescent properties of new copolymers of dimethacrylate derivatives of naphthalene-2,7-diol. Polym Adv Technol 26:176–181
24. Homenick CM, Lawson G, Adronov A (2007) Polymer grafting of carbon nanotubes using living free-radical polymerization. Polym Rev 47(2):265–70
25. Yu B, Kuzema P, Tertykh V, Laguta I (2008) Thermal degradation of polyethylene containing antioxidant and hydrophilic/hydrophobic silica. J Therm Anal Calorim 94:727–36
26. Shlenksy OF (1983) Thermal decomposition of polymeric materials: characteristics of kinetics at non-isothermal heating. J Therm Anal Calorim 28(2):421–30
27. Beyer CL, Hirschler MM (2001) Thermal decomposition of polymers, in SFPE handbook of fire protection engineering 2, section 1, chapter 7., pp 110–31
28. Bolbukh Y, Kuzema P, Tertykh V, Gawdzik B (2007) Effect of surface hydroxide, vinyl, and methyl groups on thermal stability of modified silica-divinylbenzene-dimethacryloyloxyethyl-nanophthalene composites. Int J Polym Mater 56(8):803–23
29. Bolbukh YM, Gun’ko GS, Prirkhodko GP, Tertykh VA (2009) Multilayered carbon nanotubes/poly(styrene) composites. J Nanostruct Polym Nanocomp 5:14–22
30. Protsak IS, Kuzema PO, Tertykh VA, Bolbukh YM, Kozakevich RB (2015) Thermogavimetric analysis of silicas chemically modified with products of deoligomerization of polydimethylsiloxane. J Therm Anal Calorim 121:547–557
31. Star A, Han T-R, Gabriel J-C, P, Bradley K, Grüner G (2003) Interaction of aromatic compounds with carbon nanotubes: correlation to the Hammet parameter of the substituent and measured carbon nanotube FET response. Nano Letters 3(10):1421–3
32. Yellampalli S (2011) Polymer/carbon nanotube nanocomposites. In: Tech., p 396. ISBN 978-953-307-498-6
33. Nakamura K (2014) Photopolymers: photoresist materials, processes, and applications. CRC Press, United States, ISBN 97814665172882014
34. Kennedy JF, Phillips GJ, Williams PA, Piculell JL, editors (1993). Cellulose and cellulose derivatives. Cellucon ‘93 proceedings: physicochemical aspects and industrial applications. Woodhead Publishing Ltd, United Kingdom, Cambridge. ISBN 9781855731224