INTRODUCTION
Oil and petroleum products (PeP) spills on the surface of water bodies and soils are one of the most dangerous types of man-made environmental disasters. Spilled toxic substances destroy animals and plants and make the contaminated areas uninhabitable for many years. Ecological consequences of oil spills are very difficult to estimate as oil contamination disrupts many natural processes and interrelations, significantly changes the habitability conditions of all types of living organisms and is accumulated in the biomass.

Oil is a long-term decaying product and, when a man-made disaster happens in water areas, it spreads very quickly over the water surface forming a dense layer of oil film which obstructs the access of air and light. The International Petroleum Industry Environmental Conservation Association states that such disasters do not result in immediate mass mortality in fishes, reptiles, animals and plants. However, in the medium- and long-term perspective the influence of oil spills is extremely negative. The effect of spills is the most severe for the organisms living in the coastal waters, especially the ones living on the seabed or on the surface.

Modern technologies for fighting large-scale oil spills are not always effective. Floating booms, special oil collecting vessels, sorbents, chemicals capable to "decompose" the oil or turn it into a gel (to make the collection process easier) are used. However, certain chemical agents applied for fighting such disasters are extremely toxic themselves. Nowadays, active developments are carried out in the area of creating new synthetic sorbents which meet (to a maximum degree) the integrated "price vs efficiency" criterion. Besides, the equipment allowing to carry out effective gathering of petroleum products from water surface and to directly separate (at site) water from oil is being developed.
1. PETROLEUM PRODUCTS GATHERING METHODS

Various methods are used to gather and extract oil and PeP from the water surface. They can be conventionally divided into mechanical, thermal, physical-chemical and microbiological methods (Fig. 1).

**Thermal method** which is based on oil layer burning out is used when the layer thickness is sufficient and immediately after contamination before the emulsions with water are formed. As a rule, it is combined with other methods for spill elimination. Burning of spilled oil is the oldest way of cleaning the water surface.

**Mechanical method** is the most rational way to remove petroleum products from the surface of not only reservoirs, but water streams as well. Mechanical gathering and pumping of oil and petroleum products is made by oil-collecting devices or directly by pumps from natural or artificial cavitations of the ground surface, wells or trenches (oil traps). This is the most effective way to remove the dissolved light hydrocarbons.

**Physical-chemical methods** using dispersants and sorbents are viewed to be effective when mechanical gathering of petroleum products is impossible, for example, when film thickness is small or when the spilled petroleum products pose real threats to the most ecologically vulnerable areas [8].

**Biochemical treatment** of the water surface to remove spilled oil is still not widespread due to poor biodegradability of oil components and long duration of decomposition process (up to 6 months) [9-11]. However, this trend has gained momentum in recent years [12]. Nowadays, the microbiological method for elimination of large-scale contamination of soils and water bodies with oil is used in combination with mechanical gathering and physico-chemical methods of hydrocarbons localization when the film thickness is not less than 0.1 mm. At the same time microorganisms are capable to utilize the emulsified and dissolved petroleum products. Long duration of biological degradation process is the limitation to the application of microbial neutralization of the oil-contaminated environments. The main advantages of microbiological destruction over other methods are better treatment, ecological safety, relatively low costs, possibility to continuously process the "chronic" pollutions and possibility to use it in the areas that are difficult to access.
Assessment of the existing methods allows to identify the most perspective directions. This is, first of all, PeP extraction using sorbents. The use of sorbing materials is the most effective way to eliminate the effects of oil contamination. For the oil film with thickness of less than 1-2 mm or a shallow water body, sorbents allow to clean the water surface from oil in a very short time at low cost. Quality of sorbents is determined, mainly, by their sorbent capacity with respect to oil, degree of hydrophobicity, floatability (both in the initial state and after sorption of petroleum products), possibility of oil desorption, regeneration and disposal of sorbent, technological effectiveness of their production and application.

2. SORBENTS FOR GATHERING OF OIL AND PETROLEUM PRODUCTS FROM THE SURFACE OF WATER BODIES

Materials used for gathering of oil and petroleum products from the surface of water bodies are commonly referred to as oil sorbents, as well as oil collectors and oil absorbers. Three main indicators are used to determine the quality of oil sorbents: oil absorption, water absorption and floatability. Efficiency of sorbents for oil gathering is estimated, first of all, by the value of oil absorption (oil capacity). High water absorption can be eliminated practically for all materials by additional hydrophobicity. Materials with low floatability can be effectively used in products with the reinforcing shell – booms, mats, napkins, etc. Various raw materials are used for oil sorbents production.

Materials capable to absorb and retain oil and PeP on its surface are increasingly applied in the worldwide practice to remove petroleum products from water. These materials can be classified by different categories, in particular: natural and synthetic, mineral and organic, natural and modified [13].

Different types of clays, diatomite rocks (mainly, friable diatomaceous earth), sand, zeolites, tuffs, pumice stone, etc. refer to inorganic sorbents for absorption of oil and its components [14-16]. Clay and diatomites constitute the majority of products in the sorbents’ market due to their low cost and possibility of large-scale production. Sand used as a filling material for combating small spills of oil and petroleum products can also be related to this category. However, quality of inorganic sorbents is not at the high level from the point of view of ecological safety. First of all, their oil capacity is low (70-150%) and they are not good at retaining light fractions, such as gasoline, kerosene, diesel fuel. When eliminating oil spills on water, the inorganic sorbents go down together with oil without addressing the problems of water treatment to remove pollutants. Finally, washing of these sorbents with extractive agents or water with SAS, as well as their burning-out are virtually the only methods of their disposal [17].

Besides, the materials produced by processing rocks and sedimentary rocks (expanded perlite, expanded-clay aggregate, basalt cotton wool) are related to mineral sorbents [16]. In order to achieve the required hydrophobicity properties at the surface of such sorbents, they are often processed with chemical agents: silicone fluids, vapors of mineral oils and paraffins, organic amines with aliphatic or aromatic hydrocarbon chain. However, when using the non-modified mineral sorbents there is a danger that their particles will sink together with the absorbed oil, and decomposition of the sunken oil is extremely slow and has negative impacts on the flora and fauna of water bodies [1, 18].

Natural organic and organo-mineral sorbents are a promising type of sorbents for oil pollution elimination. Wood chips, sawdust, modified peat, dried grain products, wool, waste paper are the most frequently used materials. Wool is one of the best natural sorbents which is comparable in its oil capacity to the modified peat. It can absorb up to 8-10 kg of oil per kilogram of its mass; at the same time, natural elasticity of wool allows to squeeze out much of the light fractions of oil. However, after several such squeezings the wool turns into the bituminous felt and becomes unsuitable for use. Because of its high price, insufficient quantity and strict requirements to storage (wool is very attractive to rodents, insects and undergoes biochemical transformations) the wool cannot be considered as a promising large-scale oil sorbent at all. Besides, natural organic sorbents are significantly inferior to inorganic and synthetic sorbents on sorbing capacity. However, their application is linked with the need to deliver them in large quantities to the place of localization which affects both the final costs and the time for starting the activities. Their limited floatability (which is considerably inferior to floatability of inorganic and synthetic sorbents) is also an important factor.

The synthetic sorbing materials are mostly produced based on carbamide formaldehyde resins, polyvinyl chloride, nylon, lavsan, Nitron, polypropylene, as well as the natural and synthetic latexes, rubber resins and rubbers made in the form of powder, fibrous material, various foam plastics [19-25]. Hydrophobization is used to increase floatability and oil capacity of this type of sorbents. Some existing methods of hydrophobization for fibrous materials are rather difficult to perform, technological process requires significant amounts of flammable solvents, thus increasing fire and explosion hazards of the processes. Runoff of the part of absorbed petroleum products (sometimes - up to 60-70%) when the sorbent is extracted from the
water surface is a significant drawback of the majority of fibrous sorbents [26-29].

Synthetic oil sorbents are used in the countries with the advanced petrochemical industry (USA, EU countries, Japan). Most commonly, they are made of the polypropylene fibers which are formed into non-woven rolled fabrics of different thickness. Besides, polyurethane in the spongy or granulated form, formed polyethylene with polymer fillers and other types of plastics are used. At the same time, their application in the form of high-dispersed powders to increase efficiency of the use on thin films is inadmissible due to the risk of cancerogenic diseases. On the other hand, the use of different synthetic materials which have (along with the large pore volume) a rigid three-dimensional lattice pattern is of great interest for wastewater treatment. Polyurethane foam (PPU) polyfoams possess rather high sorption properties [21, 30, 31]. Their oil capacity is 6-8 kg/kg. One of their drawbacks is that polyurethane polyfoams absorb water in the conditions of increased salinity.

Actually, sorbents, both natural and synthetic (e.g. perlite, vermiculite, zeolite) can absorb in their porous structure only up to 0.2-0.3 g of oil per 1 g of sorbent, however, owing to adhesion the amount of oil retained at the sorbents increases manifoldly. The sorbents can be modified to improve their adhesive properties. So, the perlite treated with organosilicon compounds collects up to 6-9 g of oil. One gram of hydrophobizated basalt fiber is capable to retain up to 50-60 g of light petroleum products. Heavy sorbents, such as zeolite and vermiculite, can be used for oil sinkage, while the light ones – for formation (on the surface of water and soil) of an oil-saturated sorbent layer with subsequent removal of this layer and its concentration in the places of oil disposal and destruction.

Sorbents based on different wastes of agricultural and wood-processing industries are represented by various products. They are produced on the basis of sawdust, buckwheat peel and many other crops, peat and other vegetable-based materials [32-37]. All these sorbents possess rather high oil capacity (3-5 kg/kg and higher), they are inexpensive and have a large source of raw materials. However, their main components are cellulose and lignine which impart the surfaces with strong hydrophilic properties. Therefore, sorbents, which possess the extended surface, easily and quickly absorb oil and petroleum products; at the same time, they absorb moisture even better which has an adverse effect on the oil capacity. Thus, it is necessary to impregnate the adsorbent after its deep drying with waterproofing compounds. Hydropobization of the surface by reactant or physico-chemical treatment considerably improves technological parameters but makes the sorbent more expensive. The time of protective action of coatings, and, therefore, the sorbent shelf life is usually limited to less than two years. Sufficiently complete regeneration of the exhausted sorbents of this type is difficult, as squeezing (which allows to perform regeneration at 70-85%) breaks the sorbent structure and application of thermal methods is limited to the material thermal stability (usually 250-300 °C).

Properties of some materials which are used at gathering of oil or form the basis for producing oil sorbents are given in Table 1 [38].

| Material | Oil absorption, g/g | Water-absorption, g/g | Oil press ratio, % |
|----------|---------------------|-----------------------|-------------------|
| **Natural organic materials** | | | |
| Wheat straw (broken grain) | 4.1 | 4.3 | 36 |
| Broken cane: | | | |
| - leaves | 6.1 | 4.6 | 31 |
| - stalks | 2.7 | 3.9 | 17 |
| Buckwheat peel | 3.0-3.5 | 2.2 | 44 |
| Aspen / pine bark | 0.5/0.3 | 0.8/0.8 | 25/0 |
| Sawdust | 1.7 | 4.3 | 10-20 |
| Lignin hydrolised | 1.5-3.0 | 4.1 | 25 |
| Wastes of cottonwool production | 8.3 | 0.26 | 60 |
| Peat | 17.7 | 24.3 | 74 |
| Dry moss | 3.5-5.8 | 3.1-3.5 | - |
| Wool | 8.0-10.0 | 4.5 | 87 |
| Crushed brown coal | 1-2 | 0.2 | - |
| Macroporous technical carbon | 4.0-4.5 | 0-1.0 | 10-81 |
| **Synthetic organic materials** | | | |
| Expanded polystyrene: | | | |
| - granules | 9.3 | 4.5 | 0 |
| - fiber | 7.0-12.0 | 6.0-11.5 | 80-90 |
3. SORBENTS BASED ON FIBROUS-POROUS POLYMER MATERIALS

Fibrous materials represent a system of chaotically arranged thin threads which are freely distributed in space. As a rule, their structure is not spatially oriented, thus allowing the pollutants to contact a large surface per time unit. Cotton wool, felt, fabrics, polyester batting, basalt fiber, etc. are typical fibrous materials for oil collection. In the course of oil absorption, the sorbent fibers are capable to move apart, thus creating a specific sorbent-PEP structure which (after gathering is completed) starts to gradually compress by gravity and to strain off up to 20-25% of the collected petroleum product. According to the data in Table 1, all fibrous sorbents are characterized by high ratio of the absorbed oil squeezing. Some absorbers with fibrous structure demonstrate a rather high water absorption (polyester batting, sheet foam plastic with thickness of 18 mm) which is caused by low hydrophobicity of the surface. This disadvantage can be eliminated by introduction of special hydrophobization additives.

Emergency cleaning of water bodies is an important practical application of fibrous melt-blown materials (i.e. the ones produced by pneumoextrusion method) [39, 40]. Oil-retaining capacity of such materials (which is defined by the nature and structural parameters of fibrous matrix) reaches 10 g/g and more. The melt-blown sorbents made in the form of rolls, mats, floating pillows serve as a means to remove oil from the water surface, protect the water body coasts and gather the petroleum products which have spilled onto the soil up to several hundred square meters. High degree of extraction of the emulsified petroleum oil (60-90%) is observed even in the tough dynamic modes of filtering the oil-contaminated water through the melt-blown material.

Oil trapping occurs not only due to their adsorption on the surface of fibers, but also by retaining the coagulating microdrops in the material pore space. Melt-blown sorbents are produced by using granulated polyethylene, polypropylene, polyethylene terephthalate, as well as secondary thermoplastics (product of recycling the plastic vessels, utensils, packages, etc.).

### 3.1. Melt blowing technology

The melt blowing technological process (Fig. 2) traditionally includes operations on processing the granulated thermoplastic polymer in an extruder, dispersion of the melt being formed with a flow of compressed gas and application of the fiber-porous layer onto the rotating substrate (drum). The fibers generator, manipulator and the mechanism for material removal are the main technological nodes used at melt blowing. The fibers generator which includes an extruder and a pneumatic spraying head is intended for processing the granulate into the melt, forming the fibers out of it and transferring them to the forming substrate. The manipulator ensures rotary and reciprocating movement of the substrate which is required to produce fiber-porous products of the given configuration. Usually, melt blowing is exercised under temperature ranges which ensure the melt viscosity at the extrusion head outlet corresponding to the melt flow index of not less than 20 g/10 min. This value is used as a criterion to establish distribution of temperatures along the extruder zones [41]. The most important operation is the dispersion (spraying) of the melt and fiber drawing with the gas flow. When the temperature conditions are stabilized, the spraying is controlled by adjusting the temperature and compressed air pressure. External electrization may be applied to the fibers being created.
Figure 2 – Diagram of the melt blowing technological process: 1 – engine; 2 – channel for bunker zone cooling; 3 – channel for the melt discharge to the profiling head; 4 – cylinder heaters; 5 – cylinder; 6 – screw; 7 – filling hopper; 8 – bunker; 9 – thrust bearing; 10 – reduction gear; 11 – spraying head; 12 – compressed air branch pipe; 13 – gas-polymer flow; 14 – shape-generating mandrel. Zones of screw: I – feeding (loading); II – compression (plasticization); III – squeezing (dosing). Electrization zone: 15 – high-voltage electrode; 16 – grounding electrode.

Polymer fibrous materials (PFM) represent a set of the polymer fibers which are cohesively bonded in the places of contact and which form a fiber mass. Availability of the cohesive bond between the fibers release from the need to use additional processes (needlepunching, cross-linking, etc.) in PFM production. The melt blowing technology allows to impart the sorbing elements with shape stability and structural definiteness.

Density and diameter of fibers are the main PFM parameters which define their sorption characteristics. The material density can be adjusted within the range of 0.05 – 0.5 g/cm³, while the diameter of fibers – 5 to 500 microns (Fig. 3).

Figure 3 – Electron microscopic images of different areas of cross-sections of polymer fibrous melt-blown material

In certain cases, parameters of filter-adsorption treatment are determined by the nature of PFM fibrous matrix. For example, melt-blown materials consisting of thin lipophilic fibers (polyethylene, polypropylene) are excellent oil adsorbents [42]. High adsorptive capacity, adjustable distribution of fibers diametrically and by stacking density, large volume of cavities between fibers, permeability for liquids and gases are their characteristic features. Oil-retaining capacity of sorbents made of PFM in static conditions reaches 10 kg of oil per one kilogram of adsorbent, which considerably exceeds the same parameter of composite materials intended to gather petroleum products [43]. PFM made in the form of rolls, hoses and floating pillows serve as a means to remove oil from the water surface, protect the water body coasts and gather the petroleum products which have spilled in emergency situations at the enterprises and during transportation [44].

Adsorption capacity of prototypes of melt-blown PFM based on polyolefins is as good as (and sometimes exceeds) the foreign composite non-woven fabrics specifically intended to collect petroleum products. So, the melt blowing technology was used to produce the material based on thermoplastic polymer which consists of chaotically arranged fibers with a diameter of 5-20 microns and density of 0.01-0.2 g/cm³ [45]. Absorption and
Impact Factor:

|            | ISRA (India) | SIS (USA) | ICV (Poland) | PIF (India) | JIF | IBI (India) |
|------------|--------------|-----------|--------------|-------------|-----|-------------|
| ISI (Dubai, UAE) | 0.829       |           |              |             |     |             |
| GIF (Australia) | 0.564       |           |              |             |     |             |
| JIF        | 1.500        |           |              |             |     |             |
| SISF (Morocco) | 2.031       |           |              |             |     |             |
| РИНЦ (Russia) | 0.207       |           |              |             |     |             |
| ESJI (KZ)  | 4.102        |           |              |             |     |             |
| SJIF (Morocco) | 4.102       |           |              |             |     |             |

Retention of liquids in it occurs due to the capillaries formed by the fibers twisted in bundles and clusters. Content of such capillaries in the material reaches 60%, while the remaining 40% of fibers create the material base. Optimum absorbing ability of the material is created when the capillaries content in it is from 30% to 60%. This material is capable to absorb the amount of liquid which is 40-50 times greater than its own weight.

3.1. PFM modification with disperse sorbents

Fibers modification with porous adsorbents allows to increase the PFM adsorptive capacity. The specific feature of such PFM structure is that the adsorbent particles are adhesively fixed to the fiber surface (Fig. 4).

![Figure 4 – Electron microscopic images of PFM based on high pressure polyethylene modified with activated charcoal particles (14 wt.%).](image)

It is inexpedient to process the dispersed adsorbents together with a polymer, since porous particles are encapsulated by the binders and lose their adsorptive capacity. If (at PFM forming) the particles are injected into the gas-polymer flow, then they come into adhesive contact with the fibers in viscous-flow state and (at hardening) are firmly fixed on the surface of fibers. This technology provides for the possibility to obtain materials with high adsorptive characteristics. Thus, PFM containing 11-14 wt. % of activated charcoal is three orders of magnitude higher than the one of the unmodified analog, and these PFM contain a considerable amount of sorption-active micro- and mesopores [39, page 163].

On the surface of PFM fibers (which are in viscous-flow state) it is also possible to adhesively fix the solid particles of mineral sorbent (for which purpose the modified bentonite clay is used) [46]. Polypropylene (PP) with filler (silicon dioxide SiO₂, dispersion – 5-10 microns) with 8-16% concentration in the fiber and bentonite clay particles (adhesively fixed to the particles of bentonite clay) with dispersion of 5-10 microns in the amount of 4-21 wt. % which are modified by soap stocks of fatty acids were used for producing specimens. The disperse filler was preliminary processed in the corona discharge field (intensity – 8-21 kV/cm). As a result of polarization, an electret charge with the effective area density of \( \sigma_{\text{eff}} = 0.2-0.3 \, \text{nC/cm}^2 \) is formed on the material fibers.

To assess the sorption capacity of the produced PFM, an emulsion of oil in water (0.6%) had been created which was then passed through the developed multilayer PFM. The mass of emulsion was determined by weighing before and after filtration and, on this basis, the sorption capacity of the material was calculated (Fig. 5).
Figure 5 – The dependence diagram of oil sorption capacity \( C \), g/g of the combined PFM versus corona discharge field intensity at various contents of filler (SiO\(_2\)) and clay (wt. %/wt. %):

\[ 1 \rightarrow 8/4; 2 \rightarrow 10/5; 3 \rightarrow 12/10; 4 \rightarrow 15/20; 5 \rightarrow 16/21 \]

As the result of the research it was established that combination of the disperse filler encapsulated in the fibers in the form of silicon dioxide SiO\(_2\) particles treated in the corona discharge and the particles of the modified bentonite clay applied on the surface of fibers significantly increases the efficiency of sorption of the oil dispersed in water. The largest sorption capacity was demonstrated by two PFM specimens made of PP with filler (silicon dioxide SiO\(_2\)) concentration in the fiber – 15-16% polarized in the corona discharge field with intensity of 18 kV/cm and the particles of the modified bentonite clay which are adhesively fixed on fibers in the amount of 20-21 wt. % (curves 4 and 5 in Fig. 5).

Apparently, this is because of the fact that bipolar electret charge acquired by the fibers contributes to better wetting of the fiber surface due to deformation of solvation shells and disorientation of water dipole molecules. This results in improvement of hydrophobic properties of polymer fibers. Further increase in the amount of the applied fillers leads to an increase in sorbent weight and decreases its sorption capacity due to porosity reduction.

Thus, by using the additivity of properties, an effective combined sorbent [47] can be produced which summates the sorption properties of the modified polymer fibrous material and solid particles of bentonite clays adhesively fixed on the fiber surfaces.

4 THE USE OF SECONDARY POLYMERS TO PRODUCE FIBROUS SORBENTS

The repeated use of sorbents is possible only when gathering pure fractions of oil. Most often, after two-three regeneration cycles the sorbent capacity considerably decreases as its pores become clogged with dirt and heavy fractions and the structure of fibrous sorbent can be deformed. Change in the physical-chemical properties of the spilled oil as a result of its evaporation, oxidation, emulsification and other processes may be another reason for sorption deterioration. The increased content of gas, light fractions and emulsified water in oil often results in an increase in consumption of sorbent for the oil removal from the water surface.

Secondary raw materials can be used for sorbents production; in this case, such important quality indicator as retaining capacity of the sorbents made of secondary polymers may be even higher than the one of the sorbents made of primary raw materials.

The melt flow index (MFI) of the secondary polymer raw materials is very important for producing non-woven fibrous materials. Raw materials with low MFI values are used to produce coarse-fibered materials with high MFI values. Usually, these materials have low strength characteristics. Insignificant MFI reduction due to an increase in the gel fraction content is observed in the secondary polyethylene. At repeated processing of polypropylene, MFI goes up at the increase in the number of cycles; besides, at low processing temperatures (up to 220 °C) slow MFI growth is observed at the increase in number of processing cycles. When the processing temperatures exceed 250 °C, a drastic rise in MFI is observed (14-fold increase after 5 processing cycles at \( T = 290^\circ C \)) [48].

Sorption capacity of non-woven fabrics produced from the secondary PP (wastes of polypropylene bags) and used as sorbents is not worse than that of the materials made of primary PP. At the same time, retaining capacity of a sorbent produced from secondary PP is even better (up to 15%) than that of the materials made of primary raw materials.

One of the most important advantages of the sorbents produced from fibrous materials is their high regenerative capability without considerable loss in sorption capacity at subsequent applications. Figure 6 shows the change in sorption capacity of PFM specimens produced from wastes of secondary
Impact Factor:

| Journal | Impact Factor |
|---------|---------------|
| ISRA (India) | 1.344 |
| ISI (Dubai, UAE) | 0.829 |
| GIF (Australia) | 0.564 |
| JIF | 1.500 |
| PIF (India) | 1.940 |
| РИНЦ (Russia) | 0.207 |
| ESJI (KZ) | 4.102 |
| IBI (India) | 4.260 |
| SJIF (Morocco) | 2.031 |

thermoplastics depending on the number of regeneration cycles [49]. It can be seen that at repeated use their sorption ability remains at the level of 50-70% of the initial value.

![Figure 6](image)

**Figure 6** – Change in oil capacity (ΔC, %) of PFM produced from wastes of secondary thermoplastics at repeated (n) use: 1 – fibers from disposable medical syringes (polypropylene); 2 – fibers from plastic bottles (polyethyleneterephthalate); 3 – fibers from a mix of the used disposable syringe bodies and plungers (polyethylene and polypropylene)

**Conclusion**

It has been experimentally proven that meltblown materials of different types possess the oil-retaining capacity in static conditions up to 50 g/g and degree of extraction (from the water) of the emulsified oily substances up to 100%, which exceeds the properties of the majority of the known sorption materials applied both for filtering and gathering of the spilled petroleum products. The studied patterns allow outlining a perspective to increase the efficiency of polymer fibrous materials as sorbents for oil and petroleum products. The regulation of structural parameters of the fiber-porous materials produced by the melt blowing method, along with modification of polymer fibers in physical fields and introduction of target additives that will allow obtaining new highly efficient sorbents for gathering of oil and petroleum products may become a very promising direction.

The idea of modifying PFM in electric and magnetic fields is based on the existing technological possibility to impart the melt-blown materials with a number of additional properties, in particular, electret and magnetic ones. The field of electret charge contributes to electrostatic capture of the particles of contaminants and drops of oil emulsions, facilitates separation of organic liquids into polar and non-polar components and settling of pollutants, increases the system lipophilicity. The magnetic field generates the hydrodynamic pressure which breaks the solvate layers of drops of organic liquids and facilitates their coagulation. Oils and petroleum products trapping occurs not only by their adsorption on the surface of fibers, but also due to electrocipillary and magnetocapillary phenomena stimulated by the physical fields. Phase separation takes place as the result of disturbance in kinetic stability of water emulsions of organic substances and is accompanied with irreversible coagulation of drops and their strong retention in the pores of the sorption material. Therefore, it is necessary not only to take electric and magnetic effects into consideration, but also to initiate and apply them (in a targeted way) at development and creation of new sorbents.

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