PERSPECTIVES OF FULLERENE DERIVATIVES IN PDT AND RADIOThERAPY OF CANCERS

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ABSTRACT
Nanoparticles of fullerenes and their water-soluble derivatives have been firmly introduced into solution of medical problems. Although there are still debates about their toxicity and long-term consequences of their application in the clinic, the success of fullerenes application in some sections is undeniable, in particular, in photodynamic therapy (PDT) of cancer tumors. Besides there are interesting data on radiotherapy where fullerenes appear to be more transporters than drugs, but due to own cytoprotective properties, the fullerene adducts can also participate in the combined treatment. This review evaluates the status of these sections of fullerene chemistry in terms of development and recent trends.

KEY WORDS: fullerenes, PDT, ionizing radiation, proliferation, cancer chemotherapies
Introduction

Due to its unique spherical structure, C\textsubscript{60} has a possibility to accept up to 6 electrons \([1]\) and possesses extended \(\pi\)-system. When C\textsubscript{60} is subjected to action of light, then an electron moves to higher energetic level, thereby producing an excited singlet C\textsubscript{60} that reacts with O\textsubscript{2} to form a singlet oxygen (\(^1\text{O}_2\)) \([2,3]\). Fullerenes are exclusively effective generators of the singlet oxygen with quantum yield of \(^1\text{O}_2\) close to one \([4]\). They absorb in UV- and in visible spectrum \([5]\) that allows their usage in photodynamic therapy (PDT). Originally formed the singlet excited state (\(^8\text{C}_{60}\)) undergoes intercombination transition into a triplet state (\(^3\text{C}_{60}\)). The triplet excited state is an excellent acceptor of electrons and potential producer of the superoxide anion radical \(\text{O}_2^-\) (Fig. 1). A latter way is more pronounced in organic solvents, predominantly, in polar ones, especially, in the presence of reducers such as NADH \([6]\).

Increasing number of functional groups added to fullerene, leads to decreasing of quantum yield of the singlet oxygen. Toxicity is significantly decreased or even is fully absent in hydrophilic fullerenes \([7]\). Under UV- irradiation the fullerenes demonstrate the ability to generate reactive oxygen species (ROS) with rate of 10 nmol mL\(^{-1}\) min\(^{-1}\) as confirmed by ESR method \([8]\). The efficiency of fullerene photocytotoxicity depends on the degree of modification as well as on the size of the fullerene cage \([9]\). More extended \(\pi\)-system generates a higher phototoxicity. That is why the photodynamic activity of C\textsubscript{70} is greater than C\textsubscript{60}-fullerene.

Recently the development of nanotechnology caused a careful investigation of medical capabilities of carbon nanoparticles, in particular, fullerene derivatives (adducts). Nanoparticles of 1-100 nm in size have principally others physical and chemical parameters, mechanisms and biomedical application. This is due to location of majority of their atoms at interphase of outer surface of the particles that, in turn, leads to new quantum and mechanical mechanisms of their action. To imagine the size of nanoparticles let’s take a height of a child of 4 years old to be equal to 1, then a diameter of the red blood cell will be in the range of \(10^{-5}-10^{-6}\), diameter of hepatitis C virus is \(\sim10^{-7}\), of carbon nanotube - \(\sim10^{-8}-10^{-9}\), and of water molecule – \(10^{-10}\) \([10]\). In the scale of nanoparticles the fullerenes are included to the levels from individual molecules up supramolecules and assemblies.

Now there is understanding of general parameters that are of fundamental importance to medical applications of fullerenes. Among these parameters there are a size of nanoparticles and an ability of self-aggregation due to the structure of the side chains and the formation of
hydrogen bonds. The size and shape of the nanoparticles define the morphology of the aggregates which depends on the method of fullerene preparation. As a rule smaller nanoparticles demonstrate higher cytotoxicity (although there are exceptions) which have caused by various factors, in particular, by the ability of small nanoparticles to integrate into proteins, membranes and DNA helix [11-14] affecting signaling pathways of apoptosis [15] that represented in Fig. 2.

Fullerenes are able to activate two contrary actions in the body, namely, a generation of apoptotic or necrotic cell death due to the appearance of ROS or a protective of cells acting as antioxidant agents (called sponges of radicals), in some cases, at the level of SOD-mimetics. In the first case, the fullerenes can be considered as potential anti-tumor agents [16-18] or preparations for PDT. As PDT agents fullerene derivatives may be a part of the conjugates, aggregates and supramolecules. Development of strategy of fullerene based nanoplatforms with drugs of different actions represents promising possibilities for carbon nanomedicine [19], especially, in cancer therapy.

A number of books that comprehensively describe an organic chemistry of fullerenes [2, 20] and their application in biochemistry and medicine [2,21,22] have been published. Many reviews appeared [23-28] to describe the investigation of synthesis of new compounds based fullerene, and their possible medical usage [29-35], in particularly, in PDT [27, 36]. However, the trends in PDT as well as in the emergence of new fullerene derivatives are changing very rapidly and this requires analysis of their medical options as effectors of the apoptosis of normal and cancer cells. Two aspects of development of fullerene chemistry are described in this work: new trends in PDT with features of biological usage and possibilities of applying of fullerenes in radiotherapy and radiodiagnostics.

I. PHOTOTOXICITY OF FULLERENES. PHOTODYNAMIC THERAPY

Photodynamic therapy as a treatment method involves the application of a photosensitive substance or its metabolic precursor, followed by irradiation with a focused light of a certain wavelength [28]. This results in the photochemical generation of singlet oxygen and/or ROS in targeted tissues. The active species initiate oxidation of sensitive biomolecules, resulting in the damage of cells, start of signaling pathways of apoptosis, autophagy, anoikis or necrosis. An ability to significant accumulation in the tumor, to stability and rapid elimination (clearance) as well as to the absence of general toxicity, synergistic effects with other drugs and to the
availability for the action of a specific wavelength light irradiation are necessary properties of photosensitizers [29].

The method of PDT is widely used in cancer treatment. Two types of effects are distinguished [31], where reacting agents are superoxide radicals or singlet oxygen (Fig.1). For fullerenes the method is based on the selective accumulation of photosensitizer in tumor cells and its ability to generate both agents (singlet oxygen as well as superoxide radical) under irradiation with light beam of definite wavelength. Thus, cellular uptake of a preparation and selectivity of its action, namely, on cancer cells play a certain role in damaging of a tumor.

Such selective necrosis of cancer cells was observed in the presence of the fullerene adduct with some polyester chains and irradiation by light ($\lambda=400-505$ nm) [32].

Till now photosensitizers on the basis of carbon nanostructures did not pass clinical trials and are not allowed for their practical use [30]. Nevertheless, investigations are widely carried out with hope for their further usage. When PDT is fullerene mediated it can be used to damage multiple types of cancer cells including head and neck, breast and esophageal cancers which poorly respond to other kinds of cancer therapy.

Fullerene adducts that exhibited the phototoxic effect against cancer cells are shown in Table. 1.

Colloid solutions of C$_{60}$ and C$_{70}$, water-soluble fullerenes functionalized by amino acids, folic or malonic acids, some copolymers, inclusion compounds, conjugates, dyads and triads, some fullerols etc. are used for this purpose. The mechanism of action may be various. Incubation of tumor cells with fullerenes with subsequent irradiation by light causes apoptosis in different types of cancer through 4-6 hours after irradiation [34]. Most of fullerene derivatives induce oxidative stress, which, depending on the cellular uptake of a particular compound, as well as on the type of cells, leads to a selective or non-selective damage of cells or tumor tissue by apoptosis or necrosis mechanisms. Typically, these events are accompanied by the development of mitochondrial dysfunction as well as by the decline of activity of antioxidant enzymes and the growth of MDA level. Antioxidant functions of fullerenes in the conditions of a strong oxidative stress may acquire greater significance and to contradict the anti-tumor activity of the compound. Mono-adducts of C$_{60}$ (in comparison with bis- or tris-adducts) demonstrated the greatest activity in relation to cancer cells due to their high cell uptake and enhanced localization in mitochondria.
Phototoxicity of fullerene adducts to HeLa cells.

The aminoacid-C₆₀ derivatives (Fig.3): phenylalanine-C₆₀, L-arginine-C₆₀ (Fig.3.3) and folic acid-C₆₀ (Fig.3.4) - in concentration of 5 μg/mL did not cause cytotoxicity in the dark during long time and possessed selectivity for cancer HeLa cells [33]. The decrease of mitochondrial membrane potential, of cell viability and of SOD, CAT, Gpx activity and the increase of MDA level were observed after irradiation by visible light. Finally, the apoptosis proceeded through enhancement of caspase-3 activity.

Cationic adduct C₆₀ (Fig.4) incorporated into the liposome, was demonstrating a high efficiency of cell destruction in the PDT [37].

By using BMPF (Table) it was shown [38] that ROS together with calcium ions are involved in the process of sensibilization of apoptosis and necrosis. The presence of extracellular calcium promotes activity of fullerene derivatives, but its removal does not interrupt their ability of membrane damaging. That is indicating to the existence of both calcium-dependent and calcium-independent manners in the process. Simultaneously, BMPF enhances lipid peroxidation in dependence on dose and duration of irradiation [31].

Photodynamic activity of C₇₀-γ-cyclodextrine (C₇₀/γ-CD) inclusion compound was sufficiently greater than those for C₆₀ in relation to HeLa cells [39]. Similar picture was observed at comparison of photodynamic activity (λ>400 nm) of C₆₀ and C₇₀ incorporated into lipid membrane [40]. Authors consider that this is linked with a simple difference in their ability to generate singlet oxygen, due to asymmetry of C₇₀ [9]. Similar highly photoreactive properties were showed by C₇₀ included in surface-cross-linked liposomes (cerasomes). The photosensitizer was obtained by inclusion C₇₀/CD into a lipid membrane using the reaction of exchange [41] and C₇₀ is capable of acting without the release from the cerasomes of membranes. Dynamics of spontaneous penetration of pristine C₆₀ and N-substituted fullero[C₆₀]pyrrolidines into phospholipid membranes and its distribution there have been described [42]. The nature of fullerenes as well as a method of their introduction into liposomes play a significant role [43]. Fullerene-containing liposomes with the highest potential for photo cleavage of DNA were obtained in photo triggered the exchange reaction, but not in case of heat or microwave irradiation.

Fullerene-C₇₀, modified by malonic acid, caused a much more active cell death by necrosis than the C₆₀ derivative. This process accompanied by blebbing that is rare in PDT [9].
Co-polymers of C$_{60}$-N-vinylpyrrolidon showed the best phototoxicity in relation to HeLa and murine osteogenic sarcoma cells (Tabl.) in PDT [44]. Cationic micelles of block-copolymers of poly(N-vinylcaprolactam)-C$_{60}$ demonstrated themselves as a potential antivirus drug [45] and, simultaneously, as strong photosensitizers under UV-irradiation [46].

*Phototoxicity of fullerene adducts to leukemic cells.*

Action of C$_{60}$ colloid solution and irradiation (in UV- and visible ranges) produced cytotoxic effects on leukemic Jurkat cell line (but not on thymocytes) [47]. The process was accompanied by elevated activation of caspase-3. C$_{60}$-pyrofoeoforbids with different amount of substitutes showed varying phototoxicity in relation to Jurkat cells [35]. Effects were determined by difference in their physical and chemical properties and, as consequence, by different intracellular uptake.

*Photoinduced DNA-cleavage.*

Photoinduced cleavage of super-helix DNA in pBR322 (a plasmid used E. coli cloning vectors) predominantly at guanine bases [31] was observed after incubation with carboxy-fullerenes-C$_{60}$ and irradiation by visible light (but no in the darkness). CD/C$_{60}$ is also effective photosensitizers of DNA cleavage in PDT [48]. Effectiveness are increased in order of α-, β-, γ-CD/C$_{60}$. The difference is due to varying aggregation behavior of the inclusion complexes in aqueous solution, and correlates with the size of the cavity of host molecules - cyclodextrin (174 <262 <474 Å$^3$, respectively). Moreover, photodynamic activity of γ-CD/C$_{70}$ complex was much higher than in the case of C$_{60}$ [39].

Dynamics of spontaneous penetration of pristine C$_{60}$ and N-substituted fullero[C$_{60}$]pyrrolidines into phospholipid membranes and its distribution there have been described [42]. The nature of fullerenes as well as a method of introducing them into liposomes is playing a significant role in DNA damaging [43]. Fullerene-containing liposomes with the highest potential for photo cleavage of DNA were obtained in photo-triggered exchange reaction, but not by heat or microwave irradiation. The liposomes were much higher photoreactive due to the lack of self-quenching of photoexcited fullerenes. That is resulted in the introduction of only isolated fullerenes because of controlling their self-aggregation.
Effect of fullerene adducts against different tumors in vivo.

The dominant necrotic effects over apoptosis were observed under treatment by N-methyl-pyrrolidine-C\textsubscript{60} placed in Cremophor-EL-micelles with simultaneous irradiation with white light through the abdominal wall in case of intraperitoneal carcinomatosis [49].

Multimeric (cross-linked) PEG-C\textsubscript{60} - nanoparticles [50] are used in photothermal/photodynamic therapy of mice with inoculated KB-tumor. At the laser irradiation ($\lambda = 670 \text{ nm}$, $10 \text{ min}$) the temperature on the surface of tumor was increased to $44^0\text{C}$ and caused high level of singlet oxygen leading to tumor regression.

Conjugates of fullerene with glycol chitosan (30 nm) were photoreactive in relation to cells of KB-tumor human cervical carcinoma. The selective accumulation of photosensitizer in the tumor and high generation of singlet oxygen were observed [51].

A system of 5,10,15,20-tetrakis (4-phenyl) porphyrin-polyvinylpyrrolidin-C\textsubscript{60} (TPP/PVP/C\textsubscript{60}) was tested on rats with subcutaneously inoculated Walker 256 carcinoma. Under the irradiation with red light ($\lambda = 685 \text{ nm}$) after intraperitoneal injection of 10 mg of preparation per kg of body weight the oxidative stress, lipid peroxidation and changes in antioxidant systems were observed in the tumor tissue that resulted in the destruction of cancer cells [19].

A role of substitutes.

An investigation of phototoxicity of three fullerene hexa-cis-adducts in combination with different amount of introduced photosensitizers: bis (3\textsuperscript{1},3\textsuperscript{2}-didehydrophytochlorine) fullerene [5:1]-hexa-adduct (FHP1), fullerene [5:1]-hexa-adduct with six 3\textsuperscript{1},3\textsuperscript{2}- didehydrophytochlorine groups (FHP6) and fullerene[6:0]-hexa-adduct with 12-3\textsuperscript{1},3\textsuperscript{2}- didehydrophytochlorine units (FHP12) was performed and FHP6 was recognized as the most prospective one [52]. The degree of intracellular uptake, which depended on the size and asymmetry of the fullerene complex through a changing of quantum yield of the singlet oxygen, had the greatest meaning.

A role of aggregation.

Dendro[C\textsubscript{60}]fullerene [53] inhibited growth of cells in the darkness and was slightly phototoxic under UV-irradiation due to independent formation of aggregates. A method of preparation of fullerene nanoparticles played a great role in its phototoxic activity.
Photoactivity of inclusion compounds

It is considered that PEG-fullerenes \( C_{60} \) possess good potential to be used in PDT with very small side effects for normal cells [54]. PEG-fullerene-C\( _{60} \) demonstrated phototoxicity (\( \lambda = 400-600 \) nm, 140 J/cm\(^2\)) in relation to human fibrosarcoma cells HT1080, significantly decreasing their viability. Normal fibroblasts in the same conditions kept viability of 85-93%. PEG-conjugated fullerenes, containing Gd\(^{3+} \) ion, were used for photodynamic therapy in combination with magnetic resonance tomography (theranostics) [55]. Samples of PEG with different terminal structures and molecular weights were conjugated with \( C_{60} \) by covalent binding. All studied conjugates demonstrated in vitro equal ability to generate the superoxide anion and anti-tumor activity in PDT. However, \( C_{60}\)-PEG conjugates in vivo possess a tumor suppression activity, the longest half-life in blood circulation and the highest accumulation in the tumor if they have the methyl-terminal PEG only [56].

Water-soluble derivative (\( \gamma\)-CD)\(_2\)/\( C_{60} \) is used in two ways: as drug transporter through eye barriers and photosensitizer in PDT for treatment of tumors [57]. Starting with 2 \( \mu \)mol L\(^{-1}\) (\( \gamma\)-CD)\(_2\)/\( C_{60} \) was highly toxic to the HLE B-3 cells under UVA irradiation. In the meantime, aggregated nanoparticles did not provide such an effect even at 30 \( \mu \)mol/L, and the effect was not observed under irradiation by a visible light. Obviously the singlet oxygen is an important intermediate product of phototoxicity of monomer (\( \gamma\)-CD)\(_2\)/\( C_{60} \) and \( ^1\)O production decreases with an aggregation of the particles.

Fullerols as photosensitizers.

Fullerol (\( C_{60}(OH)_{19}(ONA)_{17}18H_2O \)) also exhibited a high phototoxicity in water solutions in relation to different cancer cells and cellular compounds [58,59] that was explained by authors [279] as a result of possible accumulation of ROS products. Fullerol \( C_{60}(OH)_{24} \) is a powerful photosensitizer [60] at the expense of superoxide radicals and singlet oxygen generation through effective resonance transfer of energy [61]. \( C_{60}(OH)_{24} \) is used in PDT in ophthalmology, because it is able to penetrate eye barriers, and some correlation between intracellular distribution of \( C_{60}(OH)_{24} \) and progression in damage of human lens and retina in vivo is observed [62]. Fullerols demonstrated phototoxicity against human retina pigment epithelial cells causing the appearance of early apoptosis [63].
Conjugates and supramolecules with fullerene adducts in PDT.

Fullerene-carbohydrate hybrids (Fig. 5) can produce the selective degradation of HIV-1 protease [64]. However, these compounds did not demonstrate cytotoxicity against normal fibroblasts [65], besides; these hybrids are able to inhibit a lipid peroxidation in blood plasma [66].

Recently, the conjugates of C₆₀ and lipophilic meso-aryl-porphyrins with long chain substituents were obtained for using in PDT [67].

Among fullerene nanomaterials a porphyrin-fullerenes play an important role due to high bioavailability and relative non-toxicity [68]. Therefore, photosensitizers in the form of dyads and triads, which include porphyrin and fullerene derivatives, are produced. Dyad C₆₀-porphyrin (C₆₀-P) was used in PDT against human cells Hep-2 cancer of the larynx, causing apoptosis in the caspase-dependence manner [69]. These dyads, being included (C = 1 μmol L⁻¹) into liposomal vesicles, have caused under light irradiation the death of 80% of the cells [70].

Apoptosis without participation of caspase-3 was observed when the human lymphoblast cell line (K562) was treated by TPP/PVP/C₆₀ [2]. The usage of TPP is linked with its ability to generate singlet oxygen with high quantum yield (0.63) [71]. Three types of interactions were registered (Fig. 3) in this dyad: electrostatic, hydrogen bonds in TPP/PVP and the donor-acceptor bonds between fullerene and other components [27]. Here the high ability of these compounds to the formation of photo-induced state with divided (isolated) charges was first noted. Cell survival was dependent on the level of illumination and high phototoxic effect persisted even in an atmosphere of argon. Depending on the microenvironment of the sensitizer site localization, the tissue is damaged either through the mechanism of ¹O₂ - mediated photoreaction process or through ROS attack at a low concentration of oxygen. Apoptosis by caspase-3-dependent pathway (58% of apoptotic cells) has been replaced by predominant necrotic phenomena in anaerobic conditions.

Complex drugs containing fullerene derivatives.

By using deca cationic C₈₄O₂-malonate quaternary ammonium iodide/triiodide salts, it has been shown [72] that the generation of hydroxyl radicals can be enhanced by application of short excitation wavelength in the presence of a combination of electron donors of amine and low concentrations of ascorbic acid (AA). AA acts as an effective reducer of tertiary amine cation radicals formed during photoinduced intramolecular electron transfer from the hexa-bis
(aminoethyl) amidated malonic donor fragment to the fullerene cage. AA electron reduction can regenerate neutral deca-tertiary aminoethyl fragments and, therefore, continue photo-induced oxidation-reduction cycles that ultimately leads to an increase of hydroxyl radicals generation. C₈₄O₂-fullerene derivatives exhibit the properties of effective photosensitizers.

The photoelectric effect can be further amplified by using the light-harvesting liposomal system with introduced light-harvesting antenna molecule [32]. To improve the photodynamic activity of liposomal photosensitizer against cancer cells, the construct as "paddle" with the light-harvesting molecules 1,10-diocadecyl-3,3,30,30-tetra-methyl-indodicarbocyanine (DiD) and C₆₀ in bilayer lipid membranes was developed [73]. The light energy (λ = 610-740 nm) is absorbed by the antenna and transmitted to the C₆₀ molecules for the ROS generation as shown in Fig. 6.

Two naphthalene-diimide-C₆₀ (NDI-C₆₀) dyads in which alkyl amino substituents acted as light-harvesting antennas, and C₆₀ served as a spin converter for intercombination transition from the singlet excited to the triplet excited state were proposed for PDT [74].

The idea of association of fullerene and porphyrin structures is considered to create new photosensitizers with enhanced generation of singlet oxygen and more simple entry into tumors. Some polyads have been synthesized [75], but they have not been tested in medicine yet [76]. Calculations carried out by DFT-method, have shown [77] that some of the metal-containing fullerenes (especially, bis-fullerenes) have even more powerful potential as photosensitizers.

Two sugar-functionalized (glucose- and lactose) water-soluble cyclotrimeratrylene derivatives that are capable to form photoreactive supramolecular compounds with C₆₀ are described [78]. To enhance the photoactivity of drugs, fullerene derivatives are covalently linked with well-known photosensitizers, for example, with zinc phthalocyanines. Compounds of C₆₀-pyrrolidine tris-acid ethyl ester (PyC₆₀) are obtained with a series of photosensitizers-zinc phthalocyanines (ZnPc) among them unsubstituted ZnPc, zinc - 1,4,8,11,15,18,22,25 – octa butoxy-29H, 31H-phthalocyanine and zinc - 2,3,9,10,16,17,23,24 - octakis - (octyloxy) - 29H, 31H-phthalocyanine [79]. Also there are similar compounds for C₇₀ [80]. Theoretical calculations of the photoexcitation mechanism showed that interaction between ZnPc and PyC₆₀ is regulated by electrostatic mechanism, but not dispersive forces associated with π-π interaction. Photodynamic activity of fullerenes can be used not only in the treatment of cancer, but also for the sensitization of antimicrobial action [81]. The inactivation rate of MS2 bacteriophage and the
rate of generation of singlet oxygen increased in the following order of four suspensions of fullerene photosensitizers [82]: aqu-\(\text{nC}_{60}<\text{C}_{60}(\text{OH})_{6}\cong\text{C}_{60}(\text{OH})_{24}<\text{C}_{60}(\text{NH}_{2})_{6}\). In all cases the inactivation mechanism has involved a destruction of capsid by \(\text{C}_{60}(\text{NH}_{2})_{6}\), an infringement of the secondary structure and oxidation of the capsid proteins.

Some modern aspects of fullerene application in PDT, in particular, therapeutic perspectives of their using, are considered in the review [83]. It is necessary to account that fullerenes may be more effective in PDT of hypoxia tumors (where the oxygen level is low), because they are able to change a mechanism of cell damage.

Thus, photo-sensitizing effect of fullerene derivatives may be performed at the expense of \(^1\text{O}_2\) (\(\text{C}_{60}\)) and ROS enhancing (\(\text{C}_{60}(\text{OH})_{18}\)). It is necessary to take into account the result of differential interaction with various structures, for example, binding to molecules of the lipid membrane, increasing of OH-generation etc. The effect may be additionally enhanced by using of light gathering antenna molecule and conjugates involving components reinforcing each other effect. The greatest phototoxicity is characteristic for mono-substituted \(\text{C}_{60}\) adducts that have no ability to self-aggregation, especially in the presence of calcium ions. The process is a dose- and concentration-dependent and the method of preparation of fullerene nanoparticles is of a great importance.

**II. PHOTOPROTective ROLE OF FULLERENES.**

Some works are devoted to dual role of fullerene as pro-oxidant as well as antioxidant in UV-induced damages of different tissues, i.e. synergic or cumulative effects in PDT [84]. This problem is very important in production of cosmetic anti-sunburn means and also in some skin diseases. It is known that under UV-irradiation a set of events, leading to pro-apoptotic alterations (generation of ROS, cell rounding, bubbling of its surface etc.) and anoikis, is observed at the cellular level [85]. In mice, sebaceous glands fulfill a role of a main site for ROS generation in UVB-irradiation [86]. Use of \(\text{C}_{60}\) did not lead to emergence of toxicity, but index of ROS and index of apoptosis were decreased. More significant decrease was detected at simultaneous use of ascorbic acid and fullerene that, possibly, is provided by binding of fullerene with AA and decreasing of the Fenton reaction yield due to intercalation of AA to the heme pocket. Thus, use of a sum of fullerene+AA in combination with UV-irradiation is an effective remedy against oxidative damage of a skin.
C\textsubscript{60}, incorporated into phospholipid membrane (74.5 nm), in the case of its injection before or after UVA irradiation, restores viability of cells by decreasing of 30% the level of ROS \cite{87}. Liposome-fullerene under full absence of own toxicity significantly inhibited damage caused by chronic UVA irradiation of skin (Table.) \cite{88}. Water-soluble nanoparticles containing fullerene-C\textsubscript{60} incorporated into liposomes (75.6 nm) showed dose-dependent protective effect HaCaT cells against OH-radicals, emerging under UVA- and UVB- irradiation. Prevention of cell degeneration was observed and any protective effect exists in the presence of C\textsubscript{60} only \cite{89}.

Carboxyl-fullerene-C\textsubscript{60} sufficiently reduced a blocking of proliferation of human keratinocytes induced by UVB-irradiation \cite{90} and simultaneously decreasing an amount of cells with depolarized mitochondria. The mechanism of action had included interference of the preparation to the process of ROS generation by depolarized mitochondria, however, without participation of Bcl-2.

Tris-malonyl-C\textsubscript{60} (C3) is able to protect, selectively, cells against intracellular and/or membrane changes in UV-irradiation \cite{91}. It suggested that protection of epithelial cells A431 is linked with ability of namely this compound to maintain a network of cytoskeleton components and the integrity of coordination linkage, but only in the case, when fullerene derivative presents during irradiation. It is not excluded that the ability of a preparation to capture the superoxide radical before its conversion into OH- radical is a substantial component of the antioxidant action. C3 actually have been localized in the cell membrane.

Skin keratinocytes (HaCaT) were used for approbating of different fullerols: C\textsubscript{60}(OH)\textsubscript{44} 8H\textsubscript{2}O (SHH-F), C\textsubscript{60}(OH)\textsubscript{6-12} (LH-F) and C\textsubscript{60}(OH)\textsubscript{32-34} 7H\textsubscript{2}O (HH-F) \cite{92} in conditions of UVA and UVB-irradiation. Acceptor activity in relation to ROS was higher for HH-F and SHH-F, than for LH-F that determined a degree of their cytoprotective effect, which was greater in the case of SHH-F. Protective effect of SHH-F in relation to UVB- induced damages was higher than those in relation to UVA. Thus, SHH-F is a highly effective cytoprotector under UV-irradiation.

PVP/C\textsubscript{60} \cite{93} repressed the changes caused by UVA in HaCaT cells in the form of translocations of the transcription factor NF-κB into cytoplasm to the nucleus of keratinocytes. Protective effect and a blockage of abnormal signaling pathways were observed.
III. IONIZING RADIATION AND FULLERENES

It is known that impact of radiation in great degree is determined by generation of large amounts of ROS. Cytotoxicity or, on the contrary, antioxidant effects of fullerenes may play a substantial role in these conditions, increasing or compensating impact of ionizing radiation on the organism. Moreover, properties of the fullerenes may also be changed under action of ionizing radiation. For example, nanoC$_{60}$ in certain conditions inhibits tumor cells growth, expressing properties of a sensitizer under radiotherapy, enhancing apoptosis [94]. However, $\gamma$-irradiation is able to influence on cytotoxicity of n-C$_{60}$ (THF) [95]. Irradiated fullerenes not only did not cause oxidative stress and induce ERK-dependent death of different mammalian cells, but, on the contrary, they are protecting cells against oxidative stress, induced by THF-nC$_{60}$ or hydrogen peroxide. Thus, $\gamma$-irradiation is able to alter physical and chemical properties of n-C$_{60}$, leading to full loss of its cytotoxicity and transforming it into a cytoprotective agent. In this case, it is not possible to exclude a possibility of functionalization of fullerene surface by products of water radiolysis that, in addition to a possible role of the residual THF in cytotoxicity, may cause the loss of cytotoxicity and acquisition of new properties by the fullerene.

Radioprotective (anti-radical) activity of hydrated fullerene C$_{60}$HyFn and its labile nanosize clusters at concentrations of $10^{-11}$-$10^{-6}$ mole/L was observed under X-ray irradiation of DNA (1-7 Gy in vitro) [96]. C$_{60}$HyFn at concentrations of $10^{-7}$-$10^{-6}$ mole/L protects nucleic acids against radical-induced damage. An optimal radioprotective concentration of C$_{60}$HyFn was equal to 1 mg/kg in vivo (mice, intraperitoneally, 1 hour before or 15 min. after irradiation) at the lethal dose of 7 Gy. Lysine-C$_{60}$ was used for preliminary treatment of human lymphoblastic cells AHH-1 before $\gamma$-irradiation (>400 mg/L) and did not show visible toxicity. This sufficiently increased the cell survival after irradiation by decreasing the apoptosis level in a dose-dependent manner [97].

Dendrite adduct of [C$_{60}$]fullerene, containing 18 carboxyl groups (C$_{60}$DF) [98], is able to protect human lymphocytes and intestine cells from consequences of impact of high doses of $\gamma$-irradiation. The process includes decrease of ROS level, inhibition of radiation-induced apoptosis and cells necrosis, DNA damages, oxidative stress. However, the process is not selective, because no difference between modest protective effect of the dendro-fullerene on normal fibroblasts (dose modifying factor=1.1 [99]) and on tumor cells was observed in vivo. LD$_{50/30}$ for mice, received 300 mg/kg of the dendro-fullerene before irradiation, was equal to 10.09 Gy in comparison with 8.29 Gy for control group. No protective effect was described for a
dose of the drug of 200 mg/kg. Linked with antioxidant effects of C\textsubscript{60}DF protective effect also was observed under irradiation of \textit{zebrafish} embryos in a dose of 20 Gy and 40 Gy \cite{100}.

Observed protective effect of polyhydroxylated derivatives of fullerene attracts great attention, however, the mechanism of action is not clarified up to now, excluding its antioxidant action. It is known \cite{101,102} that effects of fullerene derivatives are predominantly linked with mitochondria. It is supposed that nanosize particles may be captured by reticule-endothelial cells \cite{103}. Namely, the specific localization of fullerene derivatives, may serve as decisive factor of their effector activity. The using of C\textsubscript{60}(OH)\textsubscript{24} \cite{104} in a dose of 40 mg/kg for 2 weeks before irradiation of the mice with a lethal dose of \gamma-irradiation (\textsuperscript{60}Co) has led to a decrease in mortality due to increase of immunity, reduction of oxidative damages and improvement of mitochondria functioning (by restoring of mitochondrial membrane potential) in comparison with control group of animals.

Fullerols demonstrated radioprotective effect during X-ray irradiation of animals (8 Gy) \cite{105} exhibiting protective effect in relation to heart and liver against chronic toxicity induced by doxorubicin \cite{106}.

C\textsubscript{60}/PVP and \gamma-CD/C\textsubscript{60}, as antioxidants, are able to stabilize radioprotective properties of \beta-carotene by preventing its oxidation \cite{107}. This fact increases the interest to inclusion complexes as compensating drugs during cancer treatment by using radiotherapy.

The combined effect of sulfa-containing radioprotectors and C\textsubscript{60}-derivatives resulted in reduction of side effects of the original drug (for example, amifostine) \cite{108}.

Fullerenes, serving as drugs, additionally can be transporters of radiomarkers and radioagents and as such intended to be used in radiotherapy and radiodiagnostics. For example, complex C\textsubscript{60}(OH)\textsubscript{20} with \textsuperscript{99m}Tc(CO)\textsubscript{3} \cite{109} rapidly reaches all tissues of the body except the brain and is retained there for 3 h without losing activity. Clearance is 24 h.

Complex \textsuperscript{212}Pb@C\textsubscript{60}-malic acid \cite{110} was stable under the beta decay of \textsuperscript{212}Pb to \textsuperscript{212}Bi and caused a weakening of \textsuperscript{212}Pb myelotoxicity. \textsuperscript{177}Lu\textsubscript{3}Lu\textsubscript{3-x}N@C\textsubscript{80}, conjugated with IL-13 \cite{111}, was interesting for radioimmunotherapy (RIT).
To prove the efficiency of $C_{60}(\text{OH})_x$ for RTD, $^{125}\text{I}-C_{60}(\text{OH})_x$ was used, which demonstrated more rapid and steady accumulation in cancer tumors in mice [112] compared to normal tissues. The combined use of fullerene derivatives and radiotherapy is still insufficiently explored, but in future it could be a far more efficient procedure than even the PDT.

The synergistic effects in radiotherapy described earlier [113] becomes all more urgent now due to the development of nanomedicine.

**Conclusion**

Recent trends in the use of fullerene derivatives in medicine are related to development of nanoplatforms that contain drugs of different composition and are capable to carry out selective delivery of them to specific organs. The main medicinal targets are cancer cells of different types. It is believed that in this aspect the fullerenes are of great interest because of their opportunity to participate in the composition of such nanoplatforms in several roles: cytotoxic agent as well as, conversely, an antioxidant (these roles may change depending on accumulation in different organs and tissues) ones; as transporter of drugs; as photo- or radiosensitizer (or protector).

However, the complexity of the problem is that until now there is no predictive model of action of fullerene derivatives under concrete conditions for a specific cell type. Moreover, the set of possible mechanisms of the effect of fullerenes on the signaling pathways of apoptosis varied (Fig.2) and depends on many factors that are difficult to administrate. All this hinders the active using of fullerene nanoparticles in medical practice. However, the individual success of some fullerene derivatives in a particular application, such as action against HIV, the selectivity to certain lines of cancer cells without damaging normal tissue, the possibility of using in theranostics [114] suggest good perspectives of fullerenes in the field of nanomedicine.

Apparently, porphyrin fullerenes of different composition that act well in transportation (drug delivery), PDT and targeting are highly successful.
FIGURE LEGENDS

Fig.1. Mechanisms of generation of singlet oxygen and superoxide radical by fullerenes.

Fig.2. Effects of fullerenes (Fs) on some signaling pathways of apoptosis

Fig.3. Aminoacid-C_{60}: Ala-C_{60}(1), Cystine-C_{60}, Arg-C_{60}(3). Folic-C_{60} (4).

Fig.4. Complex C_{60}-liposome having different surface charges due to compounds 1-4 [76].

Fig.5. Carbohydrate substituted [C_{60}] fullerenes [62,64].

Fig.6. Light-harvesting liposomal photosensitizer, which is a lipid membrane containing C_{60} and antenna molecule (1,10-dioctadecyl-3,3,30,30-tetramethylindodicarbocyanine - DiD). Copied from [81].
Table. Effect of light irradiation in the presence of fullerenes

### I. Fullerene adducts as a protectors against the photoirradiation

<table>
<thead>
<tr>
<th>Fullerene adduct</th>
<th>Cells</th>
<th>Conditions</th>
<th>Effect</th>
<th>Ref.</th>
</tr>
</thead>
<tbody>
<tr>
<td>C&lt;sub&gt;60&lt;/sub&gt;(OH)₄₄ 8H₂O (I)</td>
<td>Skin keratinocytes</td>
<td>UVA, UVB</td>
<td>ROS acceptors: (I)&gt;(III)&gt;(II). (I) shows higher protective effect in UVB than in UVA</td>
<td>70</td>
</tr>
<tr>
<td>C&lt;sub&gt;60&lt;/sub&gt;(OH)₆₋₁₂ (II)</td>
<td>HaCaT</td>
<td>UVB</td>
<td>Reduction of blocking cell proliferation. Without the participation of Bcl-2 family</td>
<td>71</td>
</tr>
<tr>
<td>C&lt;sub&gt;60&lt;/sub&gt;(OH)₃₂₋₃₄ 7H₂O (III)</td>
<td>HaCaT</td>
<td>UVA</td>
<td>Blocks the signaling pathways of apoptosis</td>
<td>73</td>
</tr>
<tr>
<td>Carboxyl-fullerene[C&lt;sub&gt;60&lt;/sub&gt;]</td>
<td>HaCaT</td>
<td>UVB</td>
<td>Restoration of blocking cell proliferation.</td>
<td>74</td>
</tr>
<tr>
<td>PVP/C&lt;sub&gt;60&lt;/sub&gt;</td>
<td>HaCaT</td>
<td>UVA</td>
<td>Prevention of cell morphological degeneration from OH radicals</td>
<td>75</td>
</tr>
<tr>
<td>C&lt;sub&gt;60&lt;/sub&gt; incorporated into the phospholipid membrane</td>
<td>HaCaT</td>
<td>UVA, 10 J sm⁻², 74.5 nm, C=150ppm</td>
<td>Inhibition of cell damages</td>
<td>77</td>
</tr>
<tr>
<td>C&lt;sub&gt;60&lt;/sub&gt; integrated in liposomes</td>
<td>HaCaT</td>
<td>UVA (12 J sm⁻²), UVB (500 mJ sm⁻²), 75.6 nm</td>
<td>Prevention of cell morphological degeneration from OH radicals</td>
<td>77</td>
</tr>
<tr>
<td>Tris-malonyl-C&lt;sub&gt;60&lt;/sub&gt;</td>
<td>A431</td>
<td>UV</td>
<td>The maintaining a network of cytoskeleton components is carried out only in the presence of the substance during irradiation</td>
<td>72</td>
</tr>
</tbody>
</table>

### II. Fullerenes in PDT

<table>
<thead>
<tr>
<th>Fullerene adduct</th>
<th>Cells</th>
<th>Conditions</th>
<th>Effect</th>
<th>Ref.</th>
</tr>
</thead>
<tbody>
<tr>
<td>C&lt;sub&gt;60&lt;/sub&gt;</td>
<td>T-leukemic lymphocytes, Jurcat</td>
<td>UV, visible 12-50 nm, 50 μmole/L</td>
<td>Cytotoxic effect</td>
<td>35</td>
</tr>
<tr>
<td>C&lt;sub&gt;60&lt;/sub&gt; and C&lt;sub&gt;70&lt;/sub&gt; incorporated into lipid membranes</td>
<td>HeLa</td>
<td>λ&gt;400 nm</td>
<td>Phototoxicity of C&lt;sub&gt;70&lt;/sub&gt; is higher than C&lt;sub&gt;60&lt;/sub&gt;.</td>
<td>60</td>
</tr>
<tr>
<td>C&lt;sub&gt;70&lt;/sub&gt;, incorporated into surface-cross-linked liposome (cerasomes)</td>
<td>HeLa</td>
<td>UV</td>
<td>High photoreactivity</td>
<td>80</td>
</tr>
<tr>
<td>C&lt;sub&gt;60&lt;/sub&gt;-aminoacid: -Phe -L-Arg C&lt;sub&gt;60&lt;/sub&gt;-folic acid</td>
<td>HeLa,</td>
<td>5 μg/ml, visible light</td>
<td>The decrease of mitochondrial membrane potential, viability, activity of antioxidant enzymes: SOD, CAT, Gpx. An</td>
<td>32</td>
</tr>
<tr>
<td>Copolymer C$_{60}$-N-vinylpyrrolidone</td>
<td>HeLa, osteogenic sarcoma cells</td>
<td>100 µg/ml, 30% of the cells are killed. Damage of membrane cells.</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Bis-methano-phosphonate-C$_{60}$ (BMPF)</td>
<td>HeLa, osteogenic sarcoma cells</td>
<td>4 µmol/L, green light, An increase in MDA. The enhancement of lipid peroxidation.</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Carboxy-C$_{60}$ in present of Ca$^{2+}$</td>
<td>Superhelical DNA pBR 322</td>
<td>Visible light, Cleavage at guanine bases. Mono-substituted adducts are higher phototoxicity</td>
<td></td>
<td></td>
</tr>
<tr>
<td>C$_{60}$-pyropeophorbide</td>
<td>Jurcat</td>
<td>Phototoxicity</td>
<td></td>
<td></td>
</tr>
<tr>
<td>C$_{60}$/PEG with different terminal chains</td>
<td></td>
<td>PEG with methyl terminal structure produces the maximum accumulation in the tumor</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Gd$^{3+}@C_{60}$/PEG</td>
<td>PDT+MRT</td>
<td>Theranostics</td>
<td></td>
<td></td>
</tr>
<tr>
<td>multimeric (crosslinked) PEG/C$_{60}$</td>
<td>KB-tumor grafted into mice</td>
<td>λ=670 nm (10 min), A raising the temperature up to 44$^0$ and tumor regression</td>
<td></td>
<td></td>
</tr>
<tr>
<td>N-methyl pyrrolidine – C$_{60}$ in Cremophor EL-micelles</td>
<td>Irradiation through the abdominal wall</td>
<td>White light, Predominance of necrosis</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Cationic micelles of poly(N-vinyl-caprolactam)-C$_{60}$</td>
<td></td>
<td>Photosensitizer, antiviral drug</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Cationic adduct C$_{60}$ (fig.4) integrated into liposome</td>
<td>HeLa</td>
<td>Highly effective phototoxic agent</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Carbohydrate substituted fullerene-C$_{60}$</td>
<td>Cancer cells</td>
<td>UV and visible light λ=355 nm, laser irradiation, Selective degradation of cells and HIV-protease</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Glycol chitosan-C$_{60}$</td>
<td>KB-tumor of human carcinoma of the cervix</td>
<td>λ=670 nm, Selective accumulation in the tumor and high generation of singlet oxygen</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Compound</td>
<td>Emitted Light</td>
<td>Toxicity</td>
<td></td>
<td></td>
</tr>
<tr>
<td>--------------------------------</td>
<td>------------------------</td>
<td>------------------------</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Malonic acid-C&lt;sub&gt;70&lt;/sub&gt;</td>
<td>400-700 nm</td>
<td>Necrosis, blebbing</td>
<td></td>
<td></td>
</tr>
<tr>
<td>FHP1, FHP6, FHP12</td>
<td></td>
<td>Compound II is more phototoxic due to higher intracellular uptake</td>
<td></td>
<td></td>
</tr>
<tr>
<td>C&lt;sub&gt;60&lt;/sub&gt;(OH)&lt;sub&gt;x&lt;/sub&gt;</td>
<td>human retinal pigment epithelial cells</td>
<td>Visible light, 83 J sm&lt;sup&gt;-2&lt;/sup&gt;, &gt;5 μmole/L</td>
<td>Early apoptosis</td>
<td></td>
</tr>
<tr>
<td>C&lt;sub&gt;60&lt;/sub&gt;(OH)&lt;sub&gt;24&lt;/sub&gt;</td>
<td>Human lens and retina cells</td>
<td>penetrate eye barriers</td>
<td>51,53</td>
<td></td>
</tr>
<tr>
<td>C&lt;sub&gt;60&lt;/sub&gt;(OH)&lt;sub&gt;19&lt;/sub&gt;(ONa)&lt;sub&gt;17&lt;/sub&gt;18H&lt;sub&gt;2&lt;/sub&gt;O</td>
<td>Different cells</td>
<td>High phototoxicity</td>
<td>55,56</td>
<td></td>
</tr>
<tr>
<td>Deca cationic C&lt;sub&gt;84&lt;/sub&gt;O&lt;sub&gt;2&lt;/sub&gt;-Malonate Quaternary Ammonium iodide/triiodide Salts</td>
<td>The relationship of the wavelength to the cell photo-toxicity in the present of a donor in the side chain (a tertiary amine)</td>
<td>cyclic process enhancing the generation of OH radicals</td>
<td>50</td>
<td></td>
</tr>
<tr>
<td>(γCD)&lt;sub&gt;2&lt;/sub&gt;/C&lt;sub&gt;60&lt;/sub&gt;</td>
<td>HLE B3</td>
<td>UVA, visible, 2 μmole/L</td>
<td>1) high toxicity in the absence of aggregation 2) transport through eye barriers</td>
<td></td>
</tr>
<tr>
<td>α,β,γ-CD/C&lt;sub&gt;60&lt;/sub&gt;</td>
<td>DNA</td>
<td>400 nm</td>
<td>DNA cleavage: α&lt;β&lt;γ-CD</td>
<td></td>
</tr>
<tr>
<td>Conjugates C&lt;sub&gt;60&lt;/sub&gt; with lipophilic meso-aryl-porphyrins with long chain substituents</td>
<td></td>
<td></td>
<td>49</td>
<td></td>
</tr>
<tr>
<td>C&lt;sub&gt;60&lt;/sub&gt;-porphyrin incorporated into liposomal vesicles</td>
<td>Hep-2</td>
<td>1 μmole/L, D=54 J/sm&lt;sup&gt;2&lt;/sup&gt;</td>
<td>death of cancer cells (80%)</td>
<td></td>
</tr>
<tr>
<td>C&lt;sub&gt;60&lt;/sub&gt;-porphyrin</td>
<td>Hep-2</td>
<td></td>
<td>Caspase-dependent pathway of apoptosis</td>
<td></td>
</tr>
<tr>
<td>5,10,15,20-tetra-kis(4-phenyl)porphyropolyvinylpyrrolidone-C&lt;sub&gt;60&lt;/sub&gt; (TPP/PVP/C&lt;sub&gt;60&lt;/sub&gt;)</td>
<td>1) K562</td>
<td>436 nm, 1000 mJ/sm&lt;sup&gt;2&lt;/sup&gt;, 20-200 mW/sm&lt;sup&gt;2&lt;/sup&gt;, 0.05 μmole/L</td>
<td>Apoptosis without the participation of caspase-3</td>
<td></td>
</tr>
<tr>
<td></td>
<td>2)rat with</td>
<td>685 nm, 50</td>
<td>Oxidative stress leads to</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Walker 256 carcinoma inoculated subcutaneously</td>
<td>J/s m², t=15 min, 10 mg/kg intraperitoneally</td>
<td>the destruction of cancer cells</td>
<td></td>
</tr>
<tr>
<td>----------------</td>
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<td></td>
</tr>
<tr>
<td>Naphthalene-diimide-C₆₀</td>
<td></td>
<td></td>
<td>Photosensitizers with light-harvesting antenna</td>
<td></td>
</tr>
<tr>
<td>C₆₀-cycloveratrilen with glucose or lactose residues</td>
<td></td>
<td></td>
<td>Photoreactive supramolecules</td>
<td></td>
</tr>
</tbody>
</table>

82
86
Figure 1.

Type 1: $O_2^-$

$^1_{\text{PS}} \rightarrow ^3_{\text{PS}} \rightarrow RH$

$\text{PS} \leftarrow \text{PS}^-$

$O_2^-$

$^3O_2$

Type 2: $^1O_2^*$

$^1_{\text{PS}} \rightarrow ^3_{\text{PS}}$

$\text{PS} \leftarrow ^3O_2$

$^1O_2^*$

$hv$
Figure 2.
Figure 3.

1. ALFD

2. CYFD

3. ARFD

4.
Figure 4.
Figure 5.

\[ C_{60}-(Glc)1 \]

\[ C_{60}-(Glc)2 \]

\[ C_{60}-(6Glc)1 \]

\[ C_{60}-(6Glc)2 \]
Figure 6.
REFERENCES


