In recent years, nanotechnology has become a key player in multiple biomedical fields, drug delivery being one of the domains where nanotechnological innovations are highly applied. Nanocarrier systems proved to be very useful in improving the physicochemical and pharmacological properties of different compounds, thus augmenting their effectiveness as therapeutic agents. Betulinic acid (BA) is a pentacyclic lupane-type triterpene of natural origin that exerts a plethora of biological effects, including antitumor, antiviral, anti-inflammatory, immunomodulatory, anti-angiogenic, hepatoprotective, etc. The main handicap of BA consists in very low water solubility which limits its use in vivo. To adjust this flaw and to improve its bioavailability, we prepared a nanoformulation of BA using silver and silver PEGylated (PEG) nanoparticles and verified its cytotoxic effects in vitro against a panel of tumor cell lines, as: human (A375) and murine melanoma (B16F10), lung (A549), breast (MCF-7 and MDA-MB-231) and hepatic carcinoma (HepG2) and on a healthy cell line – HaCaT – human immortalized keratinocytes. The obtained nanoformulation were characterized in terms of physicochemical properties by applying standard methods as transmission electron microscopy (TEM) and UV-VIS, that confirmed the development of stable solutions. The cytotoxicity was evaluated by the means of MTT (3-(4,5-Dimethylthiazol-2-yl)-2,5-Diphenyltetrazolium Bromide) and Alamar blue techniques, and the impact on cell migration and proliferation was measured using scratch assay. The nanoformulations of BA managed to inhibit the proliferation of all tumor cells at a higher extent as compared with the solution of BA in DMSO (dimethyl sulfoxide) used as the standard, whereas in the case of a healthy cell line, the toxic effect was minimum. The migration of tumor cells was also impaired by the nanoformulations. These preliminary results indicate that the antiproliferative effect of BA was improved.

Further studies are required to establish the mechanism of action of this nanoformulation and to prove its effectiveness in vivo.

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