Nanocomposites and nanomaterials

Comparative analysis of the C₆₀ Fullerene and Cytostatic 5-Fluorouracil antineoplastic activity *in vivo*

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The antitumor activity of water-soluble C_{60} fullerenes (C_{60} FAS) compared to traditional cytostatic drug 5-fluorouracil (5-FU) was investigated and analyzed in detail using the model of colorectal cancer induced by 1,2-dimethylhydrazine (DMH) in rats. Male Wistar rats (180-200 g) were divided into following groups: 1 – control; 2-4 - animals got weekly s.c. injections of DMH (20 mg/kg) for 20 weeks; 3 and 4 - starting from 21 week animals got weekly s.c. injections of C_{60} FAS (2 mg/kg) and 5-FU (45 mg/kg), accordingly, for 7 weeks. Animals were sacrificed and examined for intestinal tumors. The number, size and location of the tumors were recorded and the pathology was examined.

The tumors were detected in all parts of the colon of experimental rats, but in a majority they were observed in the distal part of colon, that is typical for this model of carcinogenesis. The tumors had different size and shape, exophytic and endophytic type of growth. Histological studies revealed adenoma and adenocarcinoma. Animals treated by $C_{60}FAS$ in a majority had adenomas. It was found that administration of $C_{60}FAS$ contributes significantly reducing the number of tumors and total lesion area in the cecum on 57% and 65%, respectively, compared with control. The 5-FU drug caused only a slight tendency to reduce these indexes. Same time, administration of $C_{60}FAS$ and 5-FU helped reduce of

the number of tumors on 31% and 28%, and total lesion area on 42% and 43%, respectively, in the colon compared with control. The results of the research clear indicate that the biocompatible and bioavailable C_{60} fullerenes at low doses are perspective agents in the therapy for colorectal cancer.