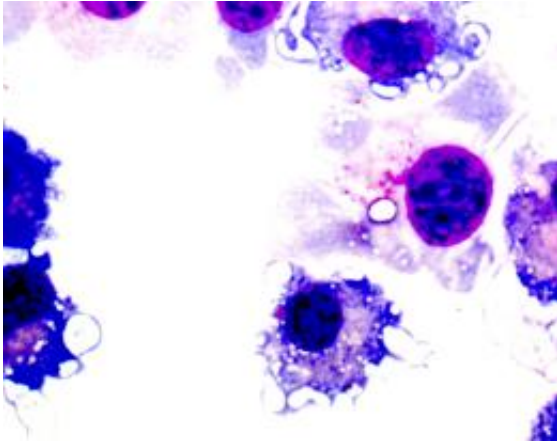


Using Viagra to combat malignant melanoma

8 November 2011



MDSC (myeloid-derived suppressor cells) suppress the tumor-specific immune response Credit: Dr. Alexandra Sevko, German Cancer Research Center

Many tumors cause chronic inflammations, which, in their turn, suppress specific attacks against the tumor by the immune system. Scientists at the German Cancer Research Center and Medical Faculty Mannheim at Heidelberg University have now shown in mice with melanoma that sildenafil - the active ingredient in Viagra - cancels the suppression of the specific immune response. Cancerous mice treated with the drug survived more than twice as long as untreated fellow animals.

At first it sounds like good news: The body's own [immune system](#) gets active in almost every cancer - however, not necessarily for the benefit of the patient. "We distinguish between two different types of immune response," says Professor Dr. Viktor Umansky, immunologist at DKFZ and University Medical Center Mannheim. "On the one hand, cells of the immune system specifically attack tumor cells. On the other, however, almost every tumor causes in its microenvironment a chronic inflammatory immune response which suppresses the specific antitumor immunity."

In an inflammatory immune response, immune cells of a certain type migrate into the tumor environment and release characteristic immune mediators. "Our aim is to reduce the chronic inflammations and, thus, to support the immune system in actively fighting the cancer," Umansky said.

He and his team have now studied chronic inflammation as it is caused by malignant melanoma. For their research, they used genetically-modified (transgenic) [mice](#) that spontaneously develop a type of skin cancer which is very similar to human melanoma. In the tumor environment and in metastatic lymph nodes of the animals, the investigators detected inflammatory mediators such interleukin-1- γ and interferon- γ . These immune mediators attract what are called myeloid-derived suppressor cells (MDSC). These immune cells are known to inhibit the [immune](#) system's most important tumor-specific fighters, the T cells.

What exactly happens when T cells get under the influence of MDSC? When the researchers exposed T cells obtained from the spleen of a healthy mouse to MDSC in the culture dish, the T cells stopped proliferating. Moreover, the T cells reduced the level of an important activating molecule. "This shows that tumor-specific T cells are actively suppressed by myeloid-derived suppressor cells," said Umansky, explaining their findings.

The melanoma mice were then given sildenafil. This substance, which is better known under its trade name, [Viagra](#)®, had already been reported to improve [tumor](#) immunity in experimental animal models several times before. Of the mice that had been given the substance with their drinking water, more than twice as many were still alive after about seven weeks as of their untreated fellows. In the animals that had been treated, both the number of tumor-specific T cells and the level of activating molecules had returned to normal. This means that

sildenafil successfully neutralizes the chronic inflammation in the melanoma environment and combats the immunosuppressive activity of MDSC.

"Our research approach is special because the disease takes a very similar course in mice as melanoma does in humans," Viktor Umansky said explaining the medical relevance of his findings. "Therefore, it is very well possible that sildenafil can also inhibit the immunosuppressive effects of inflammation and thus improve antitumor immunity in people with melanoma. In this way, the drug may contribute to achieving better treatment results in malignant [melanoma](#)."

More information: Christiane Meyer, Alexandra Sevko, Marcel Ramacher, Alexandr V. Bazhin, Christine S. Falk, Wolfram Osen, Ivan Borrello, Masashi Kato, Dirk Schadendorf, Michal Baniyash and Viktor Umansky: Chronic inflammation promotes myeloid-derived suppressor cell activation blocking antitumor immunity in transgenic mouse melanoma model. *Proceedings of the National Academy of Science (PNAS) USA*, 2011; DOI: [10.1073/pnas.1108121108](https://doi.org/10.1073/pnas.1108121108)

Provided by Helmholtz Association of German Research Centres

APA citation: Using Viagra to combat malignant melanoma (2011, November 8) retrieved 31 May 2022 from <https://medicalxpress.com/news/2011-11-viagra-combat-malignant-melanoma.html>

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