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Our <u>class action lawyers (http://www.classaction.com</u>) are <u>investigating claims</u> (<u>/class-action-lawyers/</u>) that the drug Actos may be linked to bladder cancer. According to a FDA safety announcement concerning **Actos and bladder cancer**, patients taking pioglitazone, an ingredient found in these diabetes drugs, for more than a year may have an increased risk of developing cancerous tumors of the bladder.

It has also been alleged that **drug maker Takeda Pharmaceutical withheld information about the Actos bladder cancer risk** and failed to provide adequate warnings with its diabetes drug.

Actos Bladder Cancer Link

On June 15, 2011, the FDA released a safety announcement concerning <u>Actos and</u> <u>bladder cancer (/blog/actos-bladder-cancer-lawsuits-consolidated-in-louisianacourt/). The agency released the Actos bladder cancer warning following a review of data from a five-year analysis of an ongoing study of the drug by the manufacturer, Takeda Pharmaceuticals.</u>

While there was no overall increased risk of Actos bladder cancer among users, patients with the longest exposure to pioglitazone had a greater chance of developing the disease, according to the results. Specifically, the study revealed that patients taking the drug for more than a year had a 40% higher risk of developing **Actos bladder cancer**.

In light of the link between Actos and bladder cancer

(https://www.fda.gov/Drugs/DrugSafety/ucm259150.htm), the FDA has advised that pioglitazone should not be prescribed to individuals who have bladder cancer, and should be used with caution in patients with a history of the disease. The agency also advised users to be on alert for signs which may possibly indicate **Actos bladder cancer**, such as:

- Blood in urine
- Back or lower abdominal pain
- Urgent need to urinate
- Pain during urination

Everything you need to know about Class Actions (/faq/class-actions/)

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