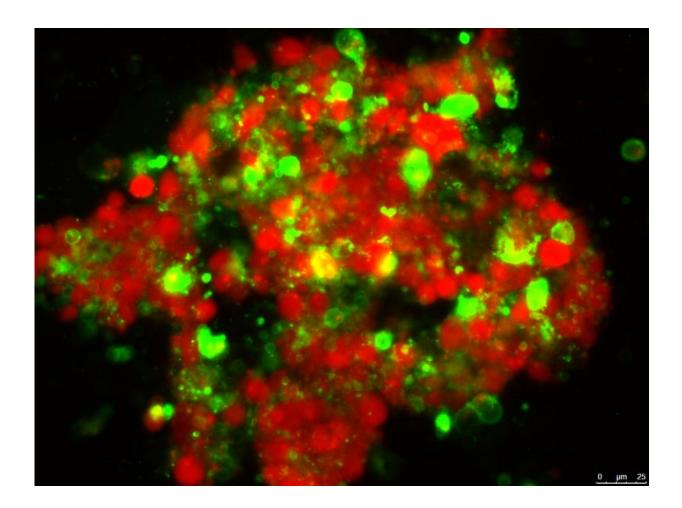


## Probing the mystery of how cancer cells die

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A photograph of colon cancer cells. The ones in red are unharmed. The ones in green are undergoing a highly orchestrated form of cellular death called apoptosis. A new study investigates the role that lipids play in this dying process. Credit: Virginia del Solar



You've probably never heard of 'sphingolipids' before. But these curiously named organic compounds play a vital role in one of humanity's most well-known diseases: cancer.

Sphingolipids are a type of <u>organic molecule</u> found inside the body. Their odd appellation comes from the chemical 'sphingosine,' which forms one of the lipids' building blocks and has been said to be named after the riddle-telling sphinx. (The famed scientist Johann Ludwig Wilhelm Thudichum wrote in 1884 that he chose to call the compound sphingosine "in commemoration of the many enigmas which it presented to the inquirer.")

One mystery that remains today is the role that sphingolipids play in cancer—in particular, the way in which <u>cancer cells</u> die.

A new study by University at Buffalo scientists sheds light on this topic, tracing how levels of various sphingolipids spike inside cancer cells when the cells are undergoing a highly organized form of cellular death called <u>apoptosis</u>.

In <u>normal cells</u>, apoptosis - akin to cellular suicide—is carried out when a cell is damaged. The self-destruction is healthy: It's thought to help the body rid itself of dysfunctional cells.

Cancer cells are adept at evading apoptosis, which is one reason they multiply out of control. But in the new study, the scientists forced the cells to commit suicide, then carefully tracked what happened inside the cells as they self-destructed.

"A better understanding of the machinery of apoptosis, at the fundamental level, can perhaps help us find new therapeutics for cancer," said G. Ekin Atilla-Gokcumen, PhD, the study's senior author and an assistant professor of chemistry in the UB College of Arts and



Sciences.

Previous studies have established that sphingolipid levels rise during apoptosis in cancer cells, but the new research—which will be published on Dec. 17 in the journal *Chemistry & Biology*—makes two important contributions.

First, the scientists found that the sphingolipid spikes may be unique to cancer. The team observed the phenomenon in three types of cancer cells (colon, ovarian and breast) during apoptosis, but not in two types of noncancerous cells (colon fibroblast cells, and retinal epithelial cells) undergoing the same process, Atilla-Gokcumen said.

Second, the scientists were able to create a detailed portrait of how the levels of many different sphingolipid compounds changed over time as death unfolded.

"We show that dying cancer cells appear to carefully tune the levels of individual sphingolipids as apoptosis progresses," Atilla-Gokcumen said. "Different compounds accumulate at different rates. Our results build on evidence that numerous sphingolipids are involved in managing this process of cellular death."

The study used a powerful technique called untargeted lipodomics to analyze the entire lipodome—the collection of all lipids within cells. Lipids, which include compounds like fats and waxes, are one of the major classes of molecules found in the body, and there are thousands of them.

Despite this huge diversity, the study found that sphingolipids accounted for nearly 90 percent of lipids that accumulated in colon cancer cells during apoptosis.



The findings emphasize what previous studies have found: that sphingolipids, which have been linked to some of the body's most mysterious processes, such as cell proliferation and differentiation, appear to be highly active during cellular suicide.

The research also raises new questions, including what accounts for the differences in sphingolipid changes inside cancerous and noncancerous cells. The UB team found that noncancerous colon cells already had high levels of sphingolipids prior to apoptosis, whereas colon cancer cells had lower levels. This could help explain why sphingolipids spiked in the cancer cells during apoptosis but not in noncancerous cells, but more research must be done before scientists can understand the implications of this finding.

Provided by University at Buffalo

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