

# Follistatin is a key player in embryo implantation

June 2 2017

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Human Embryo. Credit: Ed Uthman, MD/Wikipedia

Looking to improve the success rate of assisted reproductive technologies, researchers at Baylor College of Medicine investigated in more detail the mechanism involved in successful embryo implantation, an essential component of female fertility. They discovered that the protein follistatin plays a key role in establishing receptivity of the uterus to embryo implantation in an animal model. The results, published in the *Proceedings of the National Academy of Sciences*, contribute to the understanding of embryo implantation and provide an animal model in which to study human embryo implantation failure.

"Embryo implantation into the [uterus](#) wall is a highly coordinated process that involves many proteins and communication between the embryo and the mother. If this communication fails, the embryo won't attach to the uterus and a new life won't develop," said co-author Dr. Diana Monsivais, postdoctoral fellow in pathology & immunology at Baylor College of Medicine.

The complex process of [embryo implantation](#) in the uterus has raised interest among researchers in the field in part because it fails in about half of in vitro fertilization procedures. A better understanding of the process would help raise the odds of successful attempts at in vitro fertilization.

## **A mouse model helps understand embryo implantation**

In this study, the researchers genetically engineered mice to lack follistatin in the uterus and determined the effect on mouse [female fertility](#).

"Follistatin was already known to be important after implantation. It promotes the decidualization of the uterus, that is, the changes in the

uterus that are necessary to support and nurture the embryo as it develops," Monsivais said. "When decidualization occurs, follistatin levels rise, so we expected the consequences of deleting the follistatin gene would be in decidualization. They were, but we were surprised to also see effects even earlier than decidualization; they occurred in implantation, which had not been described before."

Female mice that lacked follistatin in the uterus produced fewer pups per female and fewer litters than normal mice. Careful assessment of implantation in these mice revealed that the [embryos](#) do not attach to the uterine wall. They float inside the uterus. Because they fail to attach to the uterine wall, the embryos do not trigger decidualization.

"The discovery that follistatin is an important protein that is required for successful mouse embryo implantation can help us develop strategies that might improve the success rate of human embryo implantation procedures in the assisted [reproductive technologies](#) clinic," said senior author Dr. Martin Matzuk, director of the Center for Drug Discovery, and Stuart A. Wallace Chair, Robert L. Moody, Sr. Chair and professor of pathology & immunology at Baylor College of Medicine.

"I think that embryo implantation is like a black box, and a mouse model is a great tool we can use to get insights on how that black box might work in humans," Monsivais said. "This study opens the possibility of identifying biomarkers, that is, compounds, such as follistatin, that could work as clues that tell us the best time to transfer an embryo to a woman attempting in vitro fertilization, and thus improve the chances of a successful implantation."

**More information:** Paul T. Fullerton et al, Follistatin is critical for mouse uterine receptivity and decidualization, *Proceedings of the National Academy of Sciences* (2017). [DOI: 10.1073/pnas.1620903114](https://doi.org/10.1073/pnas.1620903114)

Provided by Baylor College of Medicine

Citation: Follistatin is a key player in embryo implantation (2017, June 2) retrieved 20 March 2024 from <https://medicalxpress.com/news/2017-06-follistatin-key-player-embryo-implantation.html>

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