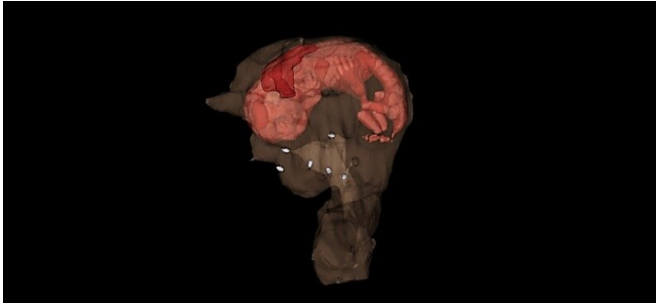


Scientists prove bioengineered uteri support pregnancy

29 June 2020



A CT image of a bioengineered rabbit uterus with a fetus, proving the organ supported fertilization, fetal development. Credit: Wake Forest Institute for Regenerative Medicine (WFIRM)

In new research from the Wake Forest Institute for Regenerative Medicine (WFIRM), scientists have shown that bioengineered uteri supported fertilization, fetal development, and live birth with normal offspring. With further development, this approach may someday provide a regenerative medicine solution for women with the inability to get pregnant due to uterine dysfunctional infertility

Published today in the journal *Nature Biotechnology*, the scientists were able to show that bioengineered uteri in an [animal model](#) developed the native tissue-like structures needed to support normal reproductive function.

"The study shows that engineered uterine tissue is able to support normal pregnancies, and [fetal development](#) was normal, with offspring size and weight being comparable to those from a normal [uterus](#)," said Anthony Atala, MD, study principal investigator and director of WFIRM. "With further development, this approach may provide a pathway to pregnancy for women with an abnormal uterus."

WFIRM scientists are world leaders in the field of

[regenerative medicine](#) and a number of the basic principles of tissue engineering and regenerative medicine were first developed at the institute. Their strategy to bioengineer functional tissues using a patient's own cells seeded onto biodegradable scaffolds has been effectively explored in preclinical studies and applied successfully in human patients to restore function in tubular and in hollow non-tubular organs. Regenerative medicine and tissue engineering technologies have emerged as an attractive option for overcoming donor organ shortages and other limitations of transplantation from donors. The scientists used the same bioengineering strategy to engineer the uterus, a more complex organ with higher functional requirements involving support of embryo implantation and fetal development.

For this study, rabbits were randomly assigned to four groups: (1) a tissue-engineered uteri group that received a cell-seeded [scaffold](#) using the animals' own cells; (2) a non-seeded scaffold group, that received a polymer scaffold only; (3) a subtotal uterine excision-only control group, where the subtotal excision was repaired by suturing and (4) a normal [control group](#), where animals underwent a sham laparotomy.

Rabbits have long been used in reproductive biology research and are ideal for uterine tissue regeneration studies as they have a relatively large uterus compared with other laboratory animals. The female rabbit's uterus is formed by two separated, functional uterine horns and cervixes, each with a capacity to carry a pregnancy.

The biodegradable polymer scaffold constructs were custom-made for each animal. The cells needed to regenerate the uterine tissue and seed the scaffolds were cultured and grown from the uterine structures of each rabbit. Six months after undergoing the scaffold implantation procedure, the rabbits were naturally mated with fertile male rabbits.

"The rabbits with cell-seeded constructs had normal pregnancies in the reconstructed segments of the uteri," said co-author Renata S. Magalhaes, MD, Ph.D., BCMAS. "This research introduces new avenues for potentially creating tissue substitutes derived from a patient's own cells to treat uterine defects."

Uterine transplantation from deceased and living donors to patients has shown promise as a treatment for permanent uterine infertility although it requires the use of anti-rejection therapies.

To date, there have been about 70 uterus transplants performed worldwide. In the United States, fewer than 10 babies have been born via a transplanted uterus. This proposed strategy, creating [uterine](#) tissue with a patient's own cells, avoids the need for a transplanted organ from a deceased or living donor, and avoids the risk of rejection and the need for antirejection drugs.

"Our results indicate that the [tissue](#)-engineered uteri responded to the expansion and mechanical strains that occur during pregnancy," said co-author Kouidy Williams, DVM. "Further preclinical studies are being planned before clinical trials are contemplated."

More information: A tissue-engineered uterus supports live births in rabbits, *Nature Biotechnology* (2020). [DOI: 10.1038/s41587-020-0547-7](https://doi.org/10.1038/s41587-020-0547-7) , www.nature.com/articles/s41587-020-0547-7

Provided by Wake Forest University Baptist
Medical Center

APA citation: Scientists prove bioengineered uteri support pregnancy (2020, June 29) retrieved 4 May 2021 from <https://medicalxpress.com/news/2020-06-scientists-bioengineered-uteri-pregnancy.html>

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