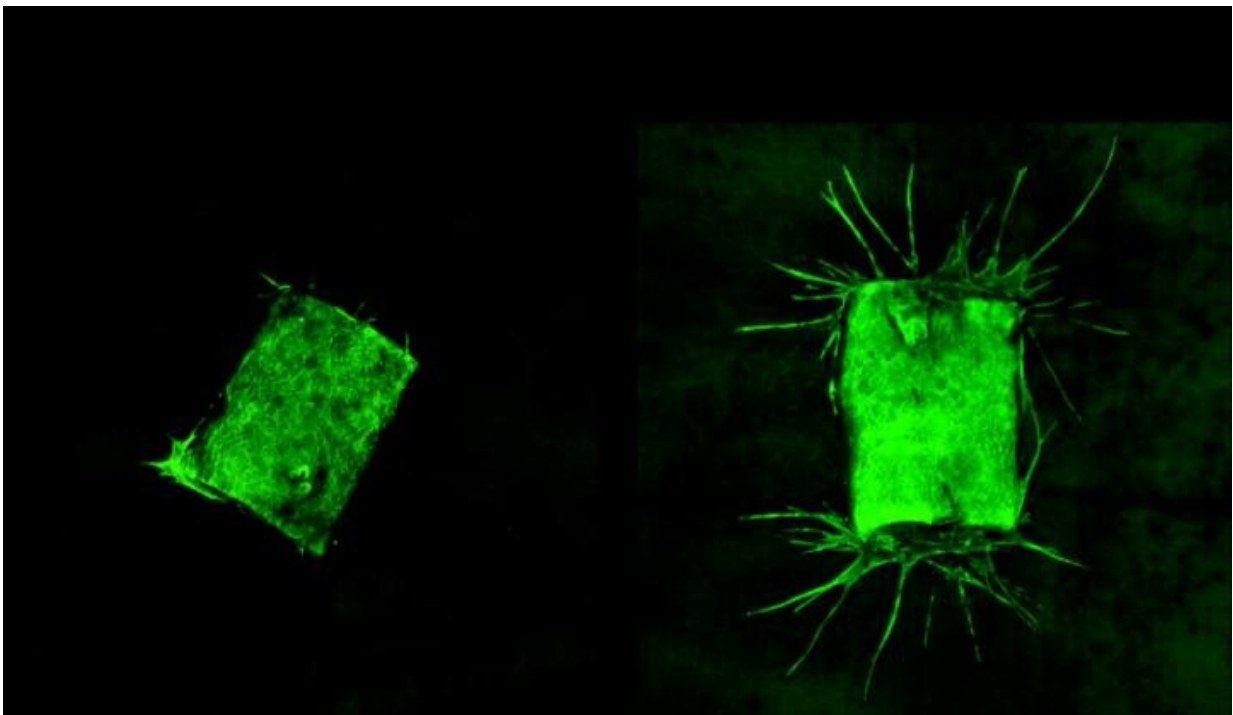


# Lesion of doom—how a parasitic bacterium induces blood vessel formation to cause lesions

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Aorta tissues that were not exposed to BafA (left) did not sprout new vessels, while BafA-exposed aorta tissues (right) did. Credit: Kentaro Tsukamoto

Bacteria of the genus *Bartonella* are parasites that can be transmitted to humans via insect bites and animal scratches, resulting in an infection known as bartonellosis. Cat-scratch disease and trench fever are forms of

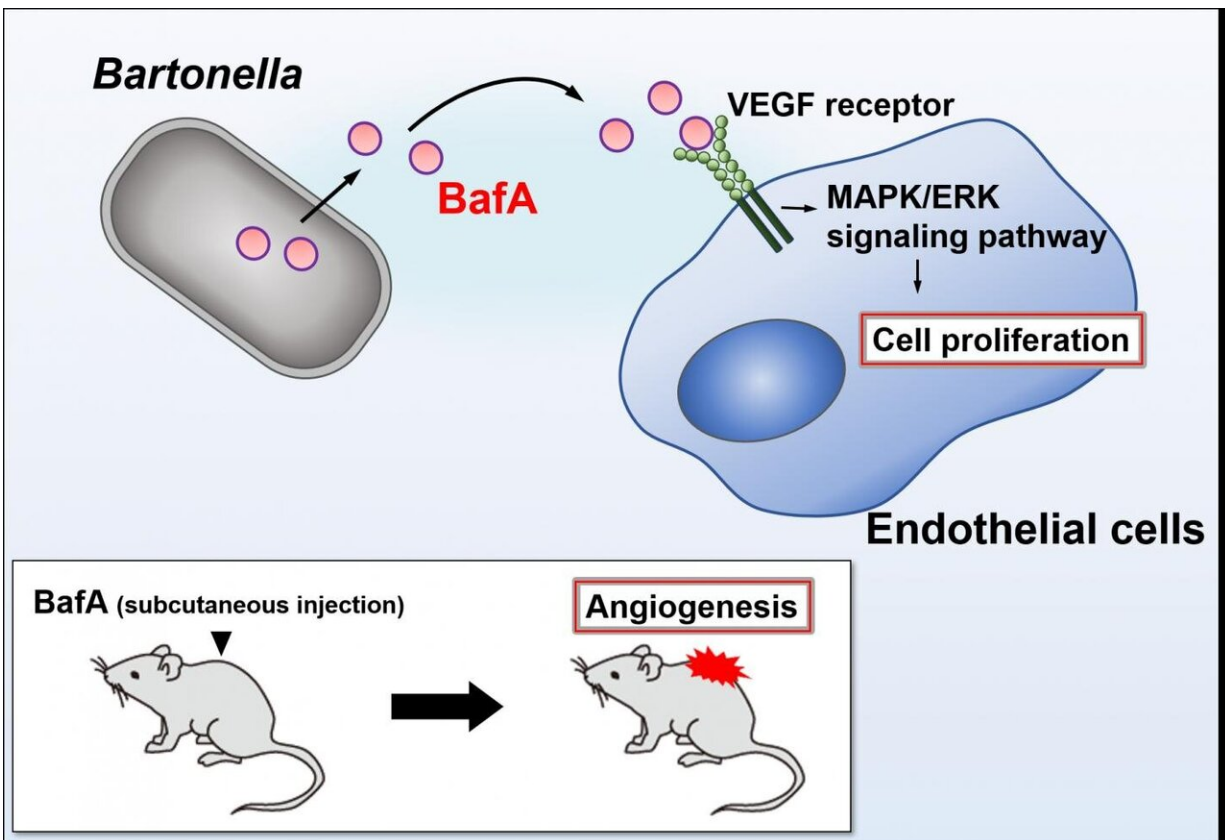
bartonellosis caused by different *Bartonella* species infecting humans. *Bartonella* bacteria can cause lesions to pop up in the skin and internal organs. To provide themselves with a safe habitat, the bacteria bring about the increase of the number of vascular endothelial cells (cells that line the interior of blood vessels), which hide themselves from the host immune system and stimulate the creation of new blood vessels, through a process called angiogenesis.

Previous studies on *Bartonella henselae* (*B. henselae* for short), the bacterium responsible for cat-scratch disease, have shown that it can directly 'inject' proteins that inhibit programmed [cell death](#) (apoptosis) into the endothelial [cells](#). However, *B. henselae* can also promote angiogenesis without directly contacting endothelial cells, which implies that the bacterium can secrete a bioactive substance that takes on the duty of kick-starting angiogenesis.

In a new study published in *Nature Communications*, a team of scientists led by Senior Assistant Professor Kentaro Tsukamoto and Professor Yohei Doi of Fujita Health University, Japan, have identified that this bioactive substance is actually a [protein](#). They have also named this protein as *Bartonella* angiogenic factor A, or BafA for short. This is the very first report of a vascular endothelial growth factor (VEGF for short)-like protein produced by [bacteria](#).

The scientists started their project by introducing *B. henselae* into human endothelial cells in [petri dishes](#), and observed that the bacteria caused the endothelial cells to multiply. To identify the genes that give *B. henselae* this ability, the researchers began inducing random mutations in the DNA of the bacteria and seeing whether the mutated bacteria could still make the endothelial cells multiply. Through these experiments, the scientists determined that *B. henselae* can stimulate angiogenesis in human endothelial cells only if it possesses a functional copy of the gene that codes for, or guides the synthesis of, the BafA protein. They also

observed that exposing human endothelial cells to the isolated BafA protein caused the cells to multiply.



Study identifies the molecular mechanism by which Bartonella bacteria cause lesions to appear over a patient's body. Credit: Kentaro Tsukamoto

Then, to confirm that BafA stimulates angiogenesis, the scientists extracted samples of a major blood vessel called the aorta from mice and placed the samples in gels that did or did not contain BafA. As can be seen in the image below, the aorta samples that were not exposed to BafA did not sprout new blood vessels, but the aorta samples that were exposed to BafA grew vessels that extended into the gel. The scientists

also found that surgically placing a BafA-containing gel plug into living mice led to blood vessels growing from the surrounding tissue into the gel.

Further experiments with human endothelial cells in petri dishes indicated that BafA activated cell surface receptors that recognize VEGF. By binding to these receptors, BafA triggered the activation of a process inside the cells, involving proteins called mitogen-activated protein kinase (MAPK) and extracellular signal-regulated kinases (ERKs). The MAPK/ERK pathway plays an important role in the multiplication of endothelial cells and angiogenesis. "In the last set of experiments, we performed similar studies in a related bacterium called *Bartonella quintana*, the bacterium that causes trench fever, and we found that it produces its own version of BafA that also causes human [endothelial cells](#) to multiply," explains Dr. Tsukamoto.

These findings provide valuable insights into the mechanisms by which infectious bacteria can produce lesions in their hosts. "We believe that BafA proteins can be leveraged as tools for studying angiogenesis, and we also consider potential medical benefits," reports Prof Doi. "Most importantly," he elaborates, "BafA is a potential target for the development of diagnostic and therapeutic strategies for bartonellosis."

The scientists also speculate that BafA proteins could be used in regenerative medicine, which is a highly specialized branch of medicine that deals with replacing or regenerating lost or damaged parts of the body. Further research is needed to confirm the scientists' findings, but needless to say, BafA proteins will certainly be of immense interest to the scientific community.

**More information:** Kentaro Tsukamoto et al, The *Bartonella* autotransporter BafA activates the host VEGF pathway to drive angiogenesis, *Nature Communications* (2020). [DOI:](#)

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