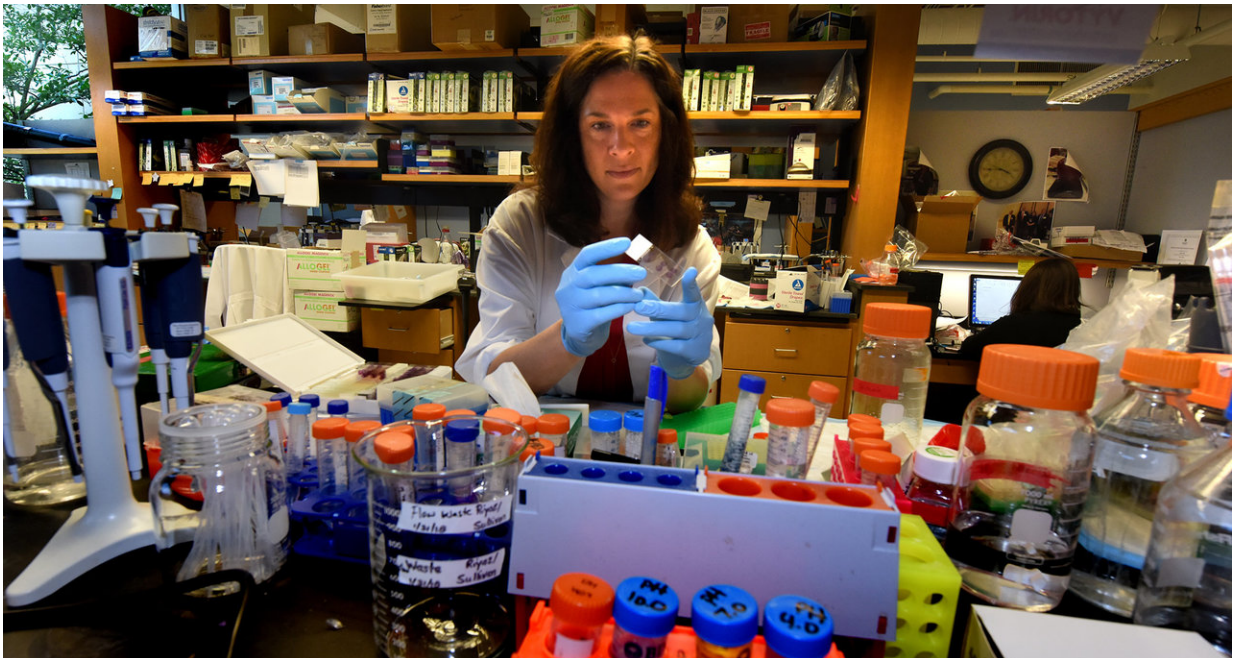


High oxidative stress hampers males' production of powerful blood vessel dilator

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Dr. Jennifer C. Sullivan, pharmacologist and physiologist in the Department of Physiology at the Medical College of Georgia at Augusta University. Credit: Phil Jones, Senior Photographer, Augusta University

Higher levels of oxidative stress in males results in lower levels of a cofactor needed to make the powerful blood vessel dilator nitric oxide, researchers report.

An antioxidant appears to help level the playing field between males and

[females](#) of the cofactor BH4 deep inside the kidneys—where the fine-tuning of our blood pressure happens—and restore similar production levels of protective nitric oxide.

Higher nitric oxide levels help reduce blood pressure both by enabling dilation of blood vessels and increasing the kidneys' excretion of sodium, which decreases the volume in those blood vessels.

"BH4 has to be there," says Dr. Jennifer C. Sullivan, pharmacologist and physiologist in the Department of Physiology at the Medical College of Georgia at Augusta University, who is exploring gender differences in hypertension. "We found that [oxidative stress](#) makes a big difference in BH4 levels."

The study in the journal *Bioscience Reports* is the first to look at sex differences of BH4 in a rodent model of hypertension.

Male humans generally have higher blood pressures and oxidative stress levels than females, at least until menopause. The findings provide more evidence that the cofactor might be a novel treatment target for both sexes, says Sullivan, the study's corresponding author.

BH4, or tetrahydrobiopterin, is required for the precursor nitric oxide synthase to make nitric oxide. Oxidative stress, which results from high levels of natural byproducts of oxygen use, is known to reduce BH4 levels, is implicated in high blood pressure and, at least before menopause, females tend to be less sensitive to it, possibly because of the protective effects of estrogen.

In an attempt to figure out why females, even in the face of hypertension, have more nitric oxide, the scientists measured BH4 levels in the inner most part of the kidney in male and female spontaneously hypertensive rats.

"We found BH4 levels were higher in the hypertensive females than the hypertensive males," Sullivan says. Females also had more nitric oxide and lower—but still high—blood pressures, and the males had more oxidative stress.

They had previously shown that young spontaneously hypertensive female rats have significantly more nitric oxide and nitric oxide synthase activity in the inner portion of their kidney than their male hypertensive counterparts, and that difference holds as the rats mature. The new work helps explain why.

"If we don't understand why females have more nitric oxide, we can't do things to potentiate our ability to make it," Sullivan says.

The scientists theorized—and found—that the elevated levels of oxidative stress in the males meant less BH4, and ultimately less nitric oxide compared to females.

They found that reducing oxidative stress improved BH4, levels and nitric oxide production and "normalized the playing fields between the two sexes," Sullivan says.

Pouring more BH4 on the situation on the other hand, didn't work without reducing oxidative stress.

"If you have a ton of oxidative stress, you can give as much BH4 as you want, and all you are going to get is more BH4," Sullivan says of BH4's destructive counterpart and the unhealthy, vicious cycle it helps create.

Without BH4, nitric oxide synthase becomes "uncoupled" and instead produces superoxide, which decreases nitric oxide production but also interacts with the nitric oxide that is available to form the oxidant peroxynitrite. Destructive peroxynitrite, in turn, targets the BH4 that is

present so it becomes BH², which further interferes with BH₄'s normal job of helping nitric oxide synthase make nitric oxide.

"You don't make the product you want nitric oxide synthase to make, which is nitric oxide," Sullivan says.

While it's not clear that females are any better of making BH₄, it is clear that the cofactor is easily altered by oxidative stress to become its unhealthy counterpart BH², Sullivan says.

Giving both males and females the synthetic antioxidant treatment Tempol for two weeks is what leveled the gender field.

Bottom line: the antioxidant treatment essentially eliminated the sex differences in BH₄ and [nitric oxide](#) synthase activity in that key region of the kidneys.

Males had a higher blood pressure at baseline and the antioxidant treatment had no effect on the blood pressure of either sex.

More work is needed to explore BH₄'s treatment potential in both sexes, Sullivan says.

BH² is widely available without a prescription, and its impact has been evaluated in a number of clinical trials including a current study at the University of Nebraska, Omaha, looking at its effect on [blood](#) flow and exercise capacity in patients with peripheral artery disease.

More information: Ellen E. Gillis et al. Oxidative stress induces BH₄ deficiency in male, but not female, SHR, *Bioscience Reports* (2018).

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