

Meet an EverVibe Tribe member

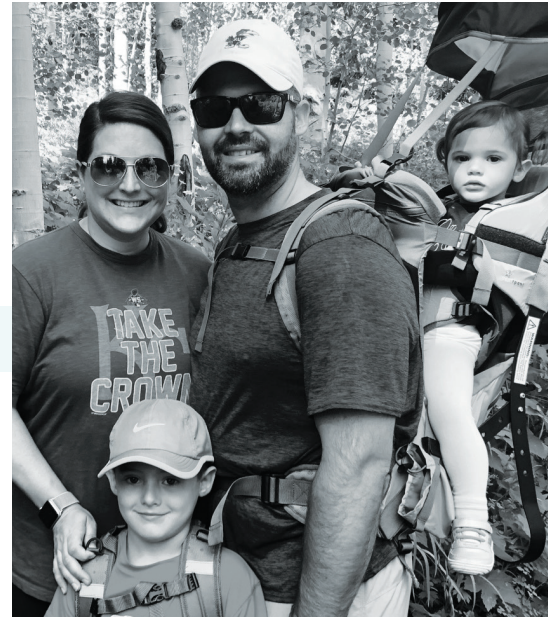
# Megan Casey

Lifelong advocate

**Occupation:** National Manager, JDRF One Walk Program

**Lives in:** Orlando, Florida

**Fun facts:** Was a youth ambassador for JDRF; married her high school sweetheart; junk TV is her guilty pleasure at the end of a crazy day; has two kids and two dogs



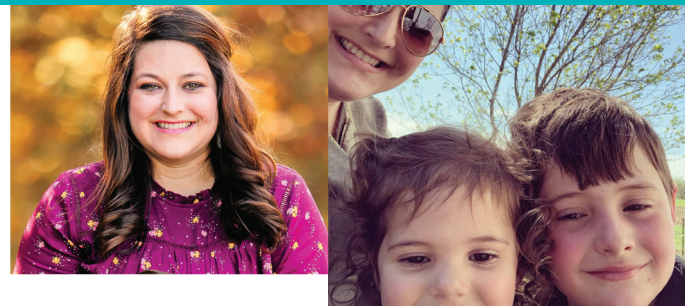
## About Megan

*"Diabetes? I'm going to die?" At six years old, that's all Megan could think about after hearing of her type 1 diagnosis. 'My doctor was like, "No. No. No. This is very manageable.'*

*"Our family had to dive in from the beginning and learn to manage the disease. I was very responsible at a young age, always wanted to do the right thing, and so we all learned it together," she said.*

*Megan has always been one to take charge of situations. "From the get-go I gave my own shots. I poked my own finger. I just was one of those who was going to take the bull by the horns."*

*She went on to become a teen advocate for juvenile diabetes—and hasn't stopped her advocacy work since. "The reason I'm so passionate about advancing the medical devices, the treatments, and the research is because I don't want to have to worry about either of my children developing type 1."*



**On Eversense:** Megan had tried other CGMs. "Before—when I wore my CGM on my stomach—my daughter would constantly bump it when I carried her around. So I'd bleed and bruise a lot, and then the CGM would become inaccurate."

What made Eversense so different for her is its ease of use. "It's so easy to be able to charge the receiver for ten minutes at night, and you put your adhesive on and that's it. There's no insertion and it doesn't hurt."

It's also "incredibly accurate," Megan says. "And that was one of the issues I had with my previous CGMs."

**"I don't feel like a pincushion. I feel like myself."**