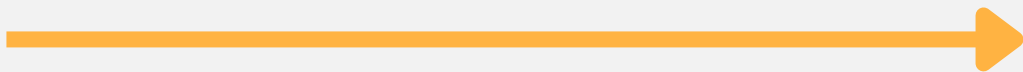


Social Media Toolkit 2026



FSGS
AWARENESS DAY



Make Your Voice Heard on June 10th!

Thank you for raising awareness about FSGS.

Why Post About FSGS Awareness Day?

Focal segmental glomerulosclerosis (FSGS) Awareness Day is an opportunity to come together and shine a light on this rare kidney disease and the real impact it has on patients, families, and caregivers around the world.

This year, awareness matters more than ever. [With the first FDA-approved treatment now available for FSGS](#), there is new hope for patients — but many people still remain undiagnosed, misdiagnosed, or unaware of the treatment options and support available to them.

By posting on social media and sharing your story, you can help others recognize the importance of early diagnosis, encourage meaningful conversations about rare kidney disease, and connect patients to information, resources, and care that could make a difference in their lives.

Use your platform to raise awareness and help ensure the FSGS community is seen, heard, and supported.

Together, we can drive awareness, improve access to care and treatment, and build a stronger future for everyone affected by FSGS.

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Story Posts

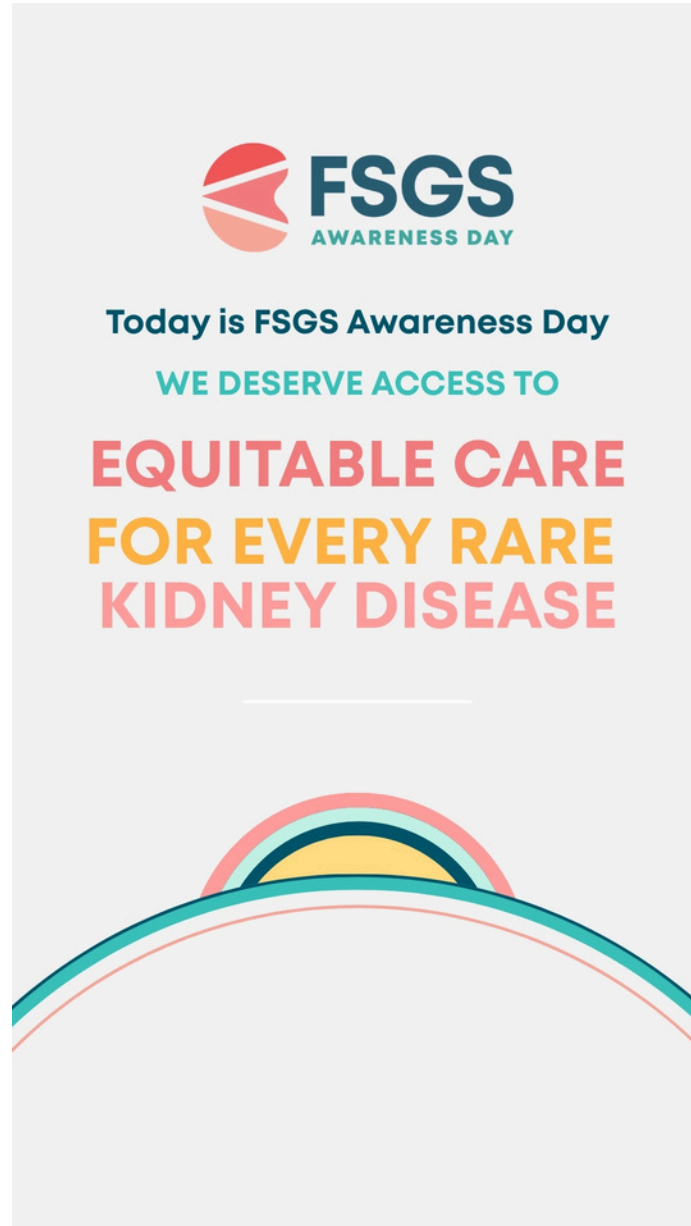


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Story Posts



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Sample Captions

Quick Caption:

Today I'm recognizing FSGS Awareness Day for everyone living with this rare kidney disease and the people who love and support them. Raising awareness may seem small, but it helps bring understanding, research, hope, and reminds others they're not fighting alone.

Sample Captions

Expanded Caption:

This FSGS Awareness Day, take a moment to learn about a rare kidney disease that impacts children, adults, and families around the world. Focal segmental glomerulosclerosis (FSGS) can lead to serious kidney damage and, for many patients, uncertainty about the future.

For years, people living with FSGS have faced limited treatment options, difficult side effects, and the fear of dialysis or transplant. Today, there is hope. The first FDA-approved treatment for FSGS marks an important step forward for the community and a reminder that research, awareness, and advocacy truly matter.

If you or someone you love is living with FSGS, know that you are not alone. Visit [NephCure.org](https://www.NephCure.org) and follow @NephCure to learn more about resources, support, research, and treatment options available for the FSGS community.

Resources to Share

Click the Links Below

- [FAQ's about the new FDA approved treatment, sparsentan \(FILSPARI\)](#)
- [FSGS informational webpage](#)
- [Press release about the FDA's approval of sparsentan for FSGS](#)

Please Tag Us @NephCure



**Thank you for using your voice this FSGS
Awareness Day!**

**Whether you share a graphic, a personal
story, or a simple message of support—
every post makes a difference.**

**Tag @NephCure so we can repost and
amplify your voice.**