HEPATOBLASTOMA WEBINAR

In honor of Rare Disease Day 2026, this event will feature a parent and patient advocate perspective in the fight against hepatoblastoma.

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In collaboration with:

The Hepatoblastoma Resource Network

February 23, 2026

10.00-10:45AM PST

RSVP FOR THIS FREE WEBINAR

Includes live Q&A session!

Guest Speaker: Sarah Lansing

Sarah is a passionate parent and patient advocate in the fight against hepatoblastoma. In August 2021, her son Glenn was diagnosed with Stage 3 Hepatoblastoma. After several rounds of chemotherapy and an unsuccessful liver resection surgery, Glenn received a life-saving liver transplant in February 2022. Join us, as we hear their story, and her continued advocacy work in the hepatoblastoma patient community.