

Parenting Cystinosis Support Group

A web-based peer support group for parents of children living with the rare disease, cystinosis facilitated by Kerry Heckman, MSW, LICSW

3rd Tuesday of every other month
(March 16th, May 18th, July 20th, Sept 21st, Nov 16th 2021)
2:30-3:30pm (PST)

To sign up, visit www.thecenterforchronicillness.org/groups

Contact us at [info @thecenterforchronicillness.org](mailto:info@thecenterforchronicillness.org)
or (425) 296-2705 with questions
www.thecenterforchronicillness.org
This program is free of cost.



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