



5 November 2020

Be Part of an International Study

Dear Cystic Fibrosis Community

As many of you know Cystic Fibrosis Australia (CFA) works very closely with the Cystic Fibrosis Foundation (CFF) in the United States and we are thrilled to support a recent CFF initiative to benefit the worldwide CF community.

An international group of cystic fibrosis (CF) and transplant clinicians have begun work on determining best practice models of post-lung transplant CF care.

They want to learn from a broad audience about current practices, barriers, and ideal models of post-lung transplant CF care and we ask that you complete one of the surveys below - **Clinician Survey** or **Patient and Family Survey** - depending on your experience.

Please feel free to share the **Clinician Survey** or **Patient and Family Survey** with colleagues or family networks who have a connection to lung transplantation.

Both surveys will close on **November 16th**.

- **Clinician Survey:** <https://www.surveymonkey.com/r/WYT3HKY>
- **Patient and Family Survey:** <https://www.surveymonkey.com/r/HZ5VTGR>

Kind regards

A handwritten signature in black ink that reads "Nettie Burke". The signature is written in a cursive, flowing style.

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