

09 June 2023

FIGHTING FOR EVERY BREATH

Last week we introduced you to 19 yr old, Zahra.

Zahra courageously shared [her story](#) with us and allowed us to share this with you.



Whilst Zahra is incredibly happy for the Australian CF Community and the increased access to 'life-changing' treatments, it has also been bittersweet for her. Zahra lives with two rare gene mutations, meaning she is ineligible for these treatments.

Every day, Zahra is fighting for every breath.

Let's support her in ensuring her voice is heard.

There are only three weeks until the end of the financial year.

Now is a key time to invest in the future activities of CFA as we plan and budget for all the critical work that must take place. This is work that will ultimately drive outcomes so that people living with CF can have longer and healthier lives. These projects include advancing our advocacy so that more people can have access to more therapies, also the CF standards of care and peer review process that exist to increase and improve clinical care outcomes, as well as supporting CF research.

We want all people with CF to have long healthy lives, and we need your partnership to do this work.

Please do not delay in supporting this vital work. Your generosity will help benefit the lives of thousands of people living with CF.

[Donate Today](#)

REMINDER: UPCOMING CONSUMER CONNECT SESSION



Join our connect session

RARE GENE MUTATIONS



6:30 PM - 7:30 PM (AEDT)

22 JUNE, 2023



Speaker

CEO Jo Armstrong



Speaker

Prof Adam Jaffe



Speaker

Dr. Shafagh Waters



Speaker

Prof Peter Wark

ASK QUESTIONS

Online

Free to attend

Save the Date

Be part of the discussion and learn more about what is happening in the space of rare gene mutations.

Learn more [HERE](#)