



runx1

The *RUNX1* Research Program



WE ARE THE *RUNX1* RESEARCH PROGRAM

Founded by a *RUNX1*-FPD patient family, we are the only non-profit advocacy organization focused on ***RUNX1* Familial Platelet Disorder** (*RUNX1*-FPD, FPDMM, FPD/AML). Our mission is to serve patients, increase awareness of the disorder, create a global team of stakeholders and fund research toward finding a cure.

We are rare. The more patients we know about, the better we can advocate. If you are a patient, please connect with our Director of Patient Engagement and Clinical Programs, **Amanda Eggen**.

We are here to help. We are in this together.

CONNECT WITH THE COMMUNITY

www.runx1-fpd.org/connect

LEARN ABOUT *RUNX1*-FPD & RESEARCH

www.runx1-fpd.org/learn
www.runx1-fpd.org/research

GET SUPPORT

Questions about living with *RUNX1*-FPD? Looking for resources? Contact us!

GET INVOLVED

Join our efforts. Volunteer.
Fundraise. Spread the word.

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www.runx1-fpd.org

