

Together, individual drops of data can make an **ocean of change**.



you



Help advance MPN research!

ARE YOU AN MPN PATIENT LIVING IN THE U.S.?

Join the new MPN PROGRESSION Registry™

Dear MPN Patient,

We're inviting you to take part in a groundbreaking effort to shape the future of MPN research: the **MPN PROGRESSION Registry™**, a new patient-powered initiative from the MPN Research Foundation. The MPN PROGRESSION Registry is designed to help researchers better understand how and why myeloproliferative neoplasms (MPNs) — including essential thrombocythemia (ET), polycythemia vera (PV), and myelofibrosis (MF) — change over time, to ultimately improve patient care and outcomes.

By joining the Registry and sharing your health journey over time, you can help answer some of the most urgent questions in MPN research — what causes progression, how symptoms affect daily life, and what treatments work best.

How does it work?

Sign up online and securely connect your medical records through our easy-to-use health portal. Then, complete brief, optional surveys about your symptoms, treatment experiences, and living with MPN.

Why join?

- **It's safe and secure.** All the data you share is securely stored and HIPAA-compliant. You control what you share, and all identifying information is removed before any research use.
- **It's easy and flexible.** Participate from home and on your schedule.
- **It's impactful.** Your data will help researchers learn how MPNs progress — and how to better treat them.

Every experience is different. Every patient counts.

To get started or learn more, visit: mpnresearchfoundation.org/mpnprogressionregistry

Need materials to share with your care team or family?

We've created patient-friendly flyers and FAQs to help explain how the MPN PROGRESSION Registry works. Download resources here: mpnresearchfoundation.org/registryresources

If you have questions — especially about how your data is protected — please reach out to us via email at research@mpnrf.org.

Thank you for considering being part of this important effort.

Warmly,

MPN Research Foundation



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