



Progress over progression

Invite your patients to join the **MPN PROGRESSION Registry™** —
a new research opportunity

Dear [Provider],

We are reaching out to invite you to join us in advancing MPN research by encouraging your patients to participate in the **MPN PROGRESSION Registry**, a patient-powered research initiative from the MPN Research Foundation.

This national registry is designed to improve our collective understanding of disease progression in myeloproliferative neoplasms (MPNs: ET, PV, and MF) by combining real-world clinical data and patient-reported outcomes. Your support in connecting eligible patients to the MPN PROGRESSION Registry can help drive forward the science behind MPNs and inform future care.

How it works:

Patients enroll online and securely link their electronic medical records through a HIPAA-compliant portal. They are then invited to complete optional surveys about symptoms, quality of life, and treatment experiences.

Why it matters:

- **Enables longitudinal tracking** of MPN progression using real-world data
- **Supports hypothesis generation and clinical insights** into therapeutic effectiveness and unmet needs
- **Helps ensure research is informed by the full patient experience** — including those outside academic centers

What we ask of you:

- **Share information about the MPN PROGRESSION Registry** with eligible patients during visits
- **Display or distribute our clinician-facing materials** in your practice
- **Reassure patients that their participation is voluntary**, and only anonymous health data — never identifying information — will be shared for research

Access the provider toolkit:

Download resources here: mpnresearchfoundation.org/registryprovider toolkit

Thank you for supporting patient-centered research. If you would like to learn more or request printed materials for your practice, please reach out to us via email at research@mpnrf.org.

Warmly,

MPN Research Foundation



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