MPN
RESEARCH
FOUNDATION

MPN
progression™
registry

Progress over progression

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

Dear [Nurse Name],

As a nurse caring for patients with myeloproliferative neoplasms (MPNs: ET, PV, or MF), your role is critical — not only in delivering care, but in guiding patients through decisions that affect their long-term health and well-being.

We're inviting you to help share an opportunity with your patients that lets them contribute directly to research: the **MPN PROGRESSion Registry**, a new initiative from the MPN Research Foundation.

This patient-driven registry collects longitudinal data — securely linking electronic medical records with brief, optional surveys — to help researchers better understand how MPNs progress, how treatments work in real life, and what symptoms patients face over time.

Why it's important:

- **Gives patients a voice** in shaping the future of MPN care
- **Gathers vital real-world evidence**, especially from diverse practice settings

How you can help:

- **Share information about the MPN PROGRESSion Registry** with interested patients during visits
- **Let them know their privacy is fully respected** — HIPAA-compliant, IRB oversight, secure encrypted platform, and all data is de-identified for research use
- **Let them know their participation is voluntary and easy** — and can be done from home
- **Direct them to our Registry webpage** to learn more and enroll at: **mpnresearchfoundation.org/mpnprogressionregistry**

Access more resources: We've developed nurse-friendly materials to make sharing this information easy. Download resources here: **mpnresearchfoundation.org/registryprovidertoolkit.**

Thank you for all that you do — and for helping us move MPN research forward, one patient at a time. 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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