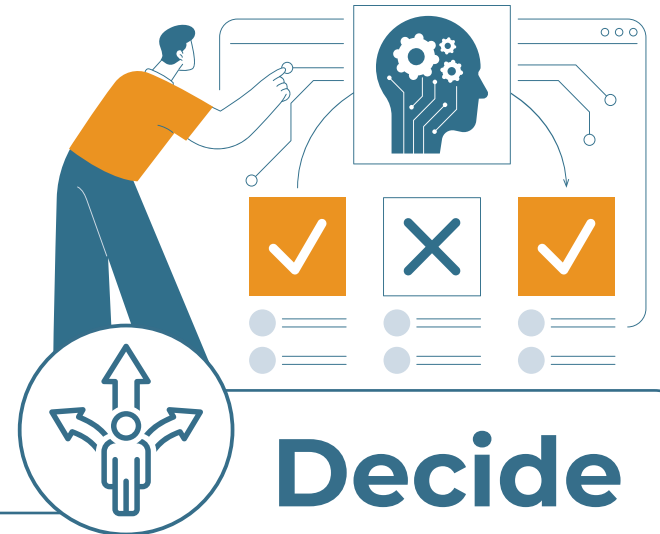
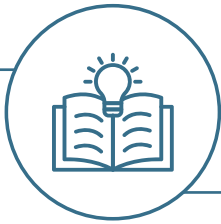


How to get in the know about myelofibrosis (MF)

What patient advocates say you should learn, decide, ask, and consider when it comes to your journey with MF

Learn

- What MF is
- How MF may affect you
- Your risk of MF progressing over time
- What symptoms you may have



Decide

- What is most important to you when discussing MF treatments with your doctor, such as:
 - Working quickly
 - Few side effects
- How comfortable you are making decisions and having conversations about MF
- Which symptoms most affect your daily life

Ask

- What blood counts mean and why they matter
- Which treatment is best if you have low platelet counts (thrombocytopenia)
- What treatment options you have and how they differ
- About a holistic approach to treatment to learn how you can manage symptoms in addition to medication
- How long it takes a treatment to start working
- What side effects you may experience
- How to know the difference between a symptom of MF and a medication side effect
- How to manage any side effects you may have
- What resources may be available to you



Consider

- Tracking symptoms to see how they change over time
- A second opinion if you feel it may be useful
- Bringing a support person with you to health care visits
- Joining a support group, using patient counseling, or finding other community resources



Stay IN THE KNOW with MF.

Learn more and find resources through the patient advocacy groups below: