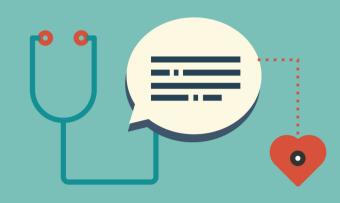
MPN PATIENT BILL OF RIGHTS

This Bill of Rights helps establish and promote awareness of core elements that are crucial to quality MPN care. It is our hope that patients will use this document to learn about their disease and participate actively in their care.

AS AN MPN PATIENT, YOU HAVE THE RIGHT TO:

RECEIVE A CLEAR,
UNDERSTANDABLE DIAGNOSIS,
AND TO SEEK A SECOND OPINION
FROM AN MPN SPECIALIST.





ASK YOUR **DOCTOR** TO COMMUNICATE YOUR **TREATMENT PLAN** (AND IT'S POTENTIAL IMPLICATIONS) IN A WAY YOU UNDERSTAND.

BE INFORMED AS TO HOW YOUR DIAGNOSIS AND TREATMENT MAY IMPACT YOUR REPRODUCTIVE HEALTH.





ACCESS MEDICATIONS AND TREATMENTS DEEMED APPROPRIATE BY YOUR DOCTOR, NOT YOUR INSURANCE.

FIND SOURCES OF FINANCIAL INFORMATION AND ASSISTANCE, UNDERSTAND HOW YOUR INSURANCE COMPANY PRIORITIZES CARE, AND HOW YOU CAN SUBMIT MULTIPLE APPEALS IF YOUR INSURANCE DENIES A TEST, TREATMENT OR APPOINTMENT.





ACCESS ACCURATE INFORMATION
FROM REPUTABLE SOURCES
INCLUDING ACADEMIC INSTITUTIONS,
HEALTHCARE PROFESSIONALS, PATIENT
ADVOCACY ORGANIZATIONS, ONLINE
PATIENT COMMUNITIES AND ELSEWHERE.



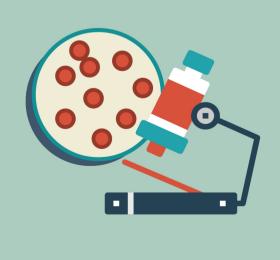




8 INCLUDE YOUR CAREGIVER(S), FAMILY, AND FRIENDS IN CONSULTATIONS WITH HEALTHCARE TEAMS.

DISCLOSE OR NOT DISCLOSE YOUR DIAGNOSIS TO YOUR EMPLOYER(S), AND PREVENT OR STOP UNLAWFUL DISCRIMINATION OR JUDGEMENT DUE TO YOUR CONDITION.





ACCESS CLINICAL TRIAL INFORMATION AND PARTICIPATE IN THOSE TRIALS IF YOU MEET THE ELIGIBILITY CRITERIA.



LEARN MORE AT MPNRESEARCHFOUNDATION.ORG/PBOR