Do you or someone you know have newly diagnosed Higher-Risk Myelodysplastic Syndrome (MDS)?

Consider enrolling in the VERONA study—a clinical research study evaluating venetoclax in combination with azacitidine, versus azacitidine alone. The study is evaluating the effect of these treatments on Higher-Risk Myelodysplastic Syndrome (MDS).

Participants must meet the following criteria:

- 18 years of age or older
- Newly diagnosed with Intermediate, High-Risk or Very High-Risk (Higher-Risk) MDS
- Have not received prior treatment for MDS
- Are ineligible for a stem cell transplant OR are eligible for a stem cell transplant, but have not yet identified a donor or arranged for the transplant

If you meet these criteria and are interested in participating, please contact your doctor to discuss the VERONA study and your eligibility.

For more information, visit ClinicalTrials.gov and search NCT04401748.

Venetoclax is an investigational drug that is not approved by the FDA or other global health authorities in MDS. Safety and efficacy have not been established in MDS.