

SWOG 9007: Cytogenetic Studies in Leukemia Patients, Ancillary

FAST FACTS

PATIENT ELIGIBILITY

1. Patients must be registered on one of the following SWOG treatment protocols: 8326, 8600, 8612, 9034, 9108 and all new leukemia protocols approved as of 1990.
2. All patients must be informed of the investigational nature of this study and must sign and give written informed consent in accordance with institutional and FDA guidelines.

PROCEDURE

Patients receive treatment as directed by the treatment protocols on which they are registered. The treatment protocols will specify when specimens are to be submitted for cytogenetic analysis (see appendix 10.4)