Biomarkers Provide Insight into a Patient's Risk Stratification and Potential Response to CLL Therapies

Testing for CLL biomarkers



Established biomarkers with prognostic value³



LOXO@Lill

Prognosis	Cytogenetic marker	Molecular marker			
Favorable	 del(13q) alone—55% of patients⁴ 	• Mutated <i>IGHV</i> -60% of patients ³	31%	11%	11%
Intermediate	 Trisomy 12–10%-20% of patients⁴ Normal karyotype³ 		Chromosomal abnormalities (FISH)	TP53 mutations	<i>IGHV</i> mutational status
Unfavorable ^a Chemotherapy-naïve.	 Del(17p)-5%-8% of patients^{4,a} Del(11q)-10% of early disease and 25% of advanced disease^{4,a} Complex karyotype-14%-34% of patients⁵ 	 TP53 mutation—8% of untreated patients³ Unmutated <i>IGHV</i>—40% of patients³ 	Includes testing in both previously untreated and relapsed/refractory pati Data from Mato AR, et al. ⁶		

Awareness and integration of biomarkers into clinical practice is important to improve the diagnosis, prognosis, and treatment of patients with CLL⁶

CLL, chronic lymphocytic leukemia; del(11q), deletions of the long arm of chromosome 11; del(13q), deletions of the long arm of chromosome 13; FISH, fluorescence in situ hybridization; del(17p), deletions of the short arm of chromosome 17; *IGHV*, immunoglobulin heavy chain variable region gene; *TP53*, tumor protein 53.

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