



The mutational landscape of chronic lymphocytic leukemia and its impact on prognosis and treatment

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The typical genome of chronic lymphocytic leukemia (CLL) carries ~2000 molecular lesions. Few mutations recur across patients at a frequency >5%, whereas a large number of biologically and clinically uncharacterized genes are mutated at lower frequency. Approximately 80% of CLL patients carry at least 1 of 4 common chromosomal alterations, namely deletion 13q14, deletion 11q22-23, deletion 17p12, and trisomy 12. Knowledge of the CLL genome has translated into the availability of molecular biomarkers for prognosis and treatment prediction. Prognostic biomarkers do not affect treatment choice, and can be integrated into prognostic scores that are based on both clinical and biological variables. Molecular predictive biomarkers affect treatment choice, and currently include *TP53* disruption by mutation and/or deletion and *IGHV* mutation status. *TP53* disruption by gene mutation and/or deletion associates with chemoimmunotherapy failure and mandates treatment with innovative drugs, including ibrutinib, idelalisib, or venetoclax. The mutation status of *IGHV* genes represents a predictive biomarker for identifying patients that may benefit the most from chemoimmunotherapy with fludarabine, cyclophosphamide, and rituximab. Assessment of these biomarkers at the time of treatment requirement is recommended by most current guidelines for CLL management. Other molecular predictors are under investigation, but their application in clinical practice is premature.

Learning Objectives

- Understand that the major advances in the understanding of CLL genomics has led to the clarification of the mutational landscape of the disease and of its clonal evolution over time and under the selection pressure of treatment
- Learn that such molecular knowledge has expanded the availability of prognostic and predictive biomarkers that can be used for patient counseling, treatment tailoring, and clinical trial design

Molecular genetics of chronic lymphocytic leukemia

The typical genome of unselected chronic lymphocytic leukemia (CLL) carries ~2000 molecular lesions, of which, however, only ~20 are nonsynonymous mutations and only ~5 are gross structural abnormalities.^{1,2} Few mutations recur across patients at a frequency >5%, whereas a large number of biologically and clinically uncharacterized genes are mutated at lower frequency. In total, >40 recurrently mutated driver genes have been identified in CLL (Figure 1).^{1,2} Recurrent mutations are not homogeneously spread across the CLL genome, but, rather, affect genes that can be integrated into a small set of pathways (Figure 2). These include microenvironment-dependent signaling through NOTCH (*NOTCH1*, *FBXW7*), inflammatory receptors (*MYD88*), MAPK–extracellular signal-regulated kinase (*BRAF*, *KRAS*, *NRAS*, *MAP2K1*) and NF-κB pathways (*BIRC3*, *TRAF3*, *NFKBIE*), as well as intracellular programs

such as DNA damage and cell cycle control (*ATM*, *TP53*, *SAMHD1*, *POT1*), chromatin modification (*HIST1H1E*, *CHD2*, *ZMYM3*), transcription (*EGR2*, *IRF4*, *BCOR*, *MED12*), and ribosomal processing (*XPO1*, *SF3B1*, *RPS15*).^{1,2}

Approximately 80% of CLL patients carry at least 1 of 4 common chromosomal alterations, namely deletion 13q14, deletion 11q22-23, deletion 17p12, and trisomy 12. Deletion 13q14 is the most frequent genetic lesion of CLL occurring in 50% to 60% of cases.³ The minimal deleted region on 13q14 contains the *miR15A* and *miR16A* microRNAs.⁴ In normal cells, *miR15A* and *miR16A* inhibit the expression of key regulators of apoptosis and cell cycle. Deletion of *miR15A* and *miR16A* abrogates this inhibitory effect, favors the constitutive survival and cycling of tumor B cells, and leads to CLL development in mouse models.⁵ B-cell lymphoma 2 (*BCL2*) codes for an antiapoptotic protein and is one of the genes upregulated in CLL as a consequence of *miR15A*/*miR16A* deletion. Consistent with the central contribution of *BCL2* activation in the pathogenesis of CLL, selective inhibition of *BCL2* through the BH3 mimetic venetoclax results into high response rates and deep remissions.⁶

Although genes encoding for components of the B-cell receptor (BCR) signaling machinery are usually not targeted by somatic mutations in CLL, the critical role of this pathway in disease pathogenesis is supported by the therapeutic success of targeted agents that interfere with the BCR. These agents are commonly termed as BCR inhibitors,

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Off-label drug use: Pembrolizumab and Venetoclax.



Figure 1. Genes and chromosomal regions affected by molecular lesions in CLL. The word cloud shows the genes that are reported as mutated in CLL by the Catalogue of Somatic Mutations in Cancer (COSMIC; v77) and the gross chromosomal abnormalities that are recurrent in CLL. The size of the font is proportional to the frequency of the molecular lesion.

although they should be more properly defined as kinase inhibitors because they inhibit many different pathways. Functional evidence indicates that BCR pathway activation in CLL results from contacts between tumor cells and antigens, which is influenced, among others, by the somatic hypermutation (SHM) load of the rearranged immunoglobulin heavy-chain variable (*IGHV*) genes.⁷ CLLs carrying unmutated *IGHV* genes display enhanced BCR signaling compared with CLLs carrying mutated *IGHV* genes, reflecting functional differences in the type of antigenic interaction through the BCR. The BCRs of CLLs carrying unmutated *IGHV* genes maintain poly-reactivity because of the lack of SHM, and thus are capable of binding multiple epitopes and are more prone to sustained signaling.⁷ Conversely, CLLs carrying mutated *IGHV* genes fail to respond to BCR stimulation because of anergy, or because the antigen specificity has

been extremely narrowed by the accumulation of mutations through the SHM process.⁷

Definition and use of prognostic and predictive biomarkers in CLL

Biomarkers are variables that can be measured and associated with treatment or disease outcome. Biomarkers can be single variables or classifiers composed of multiple variables (ie, scores).

Prognostic biomarkers usually reflect the underlying biology and natural history of CLL, and thus define prognosis in the absence of treatment, or, for patients requiring treatment, independent of the treatment received.^{8,9} Clinical endpoints captured by prognostic biomarkers include: (1) death; (2) treatment requirement, if “watch and wait” is the proposed management; and (3) other clinically relevant events such as Richter syndrome. In clinical practice, the impact of CLL prognostic biomarkers is limited. In the field of clinical research, prognostic biomarkers allow for identification of those CLL patients appropriate for early intervention studies. Predictive biomarkers, on the other hand, provide information on the likely benefit from a specific treatment. Consistently, the efficacy of a given treatment is different in biomarker-positive vs biomarker-negative patients. In clinical practice, predictive CLL biomarkers can significantly impact treatment decisions for a particular patient.

Variables can be defined as predictive biomarkers on statistical grounds or on biological grounds.^{8,9} On statistical grounds, at least 2 comparison groups are needed (ie, 2 different treatment arms in a randomized trial) to determine whether a biomarker is potentially predictive, and a formal test for an interaction between the biomarker and the treatment group needs to be performed. Conversely, to establish whether a biomarker is purely prognostic requires the demonstration that there is a significant association between

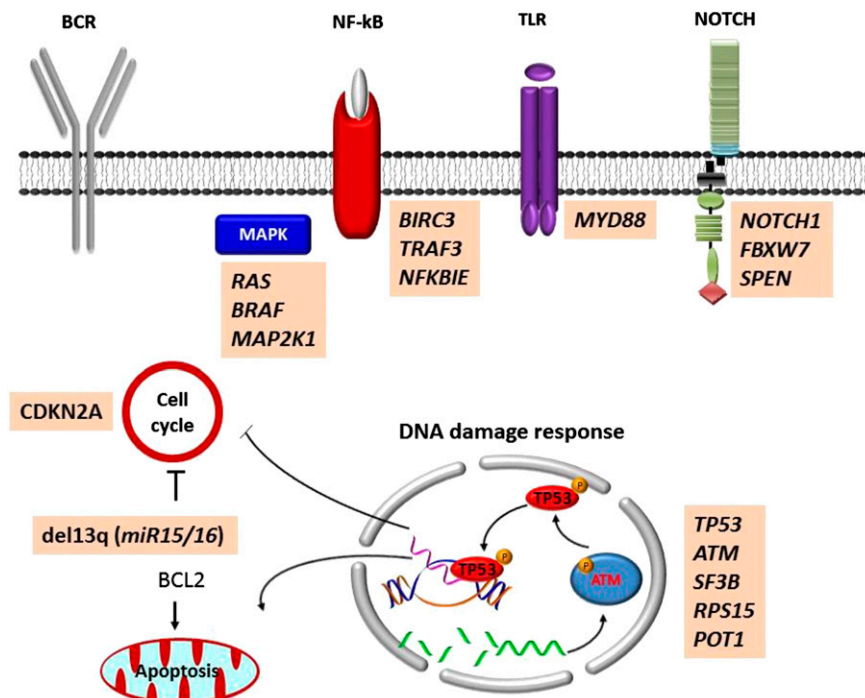


Figure 2. Mutated pathways in CLL. Cellular programs that are affected by the most recurrent molecular lesions are represented. Boxes show the genes that are recurrently mutated in each cellular program.

the biomarker and outcome, regardless of treatment, and that treatment effects do not depend on the biomarker. A biomarker may have both predictive and prognostic implications. On biological grounds, a predictive biomarker is one whose presence is required for the treatment to work. Thus, in this context, the term predictive should be reserved for situations in which the biomarker is the target of the therapy and needs to be present for the treatment response to occur (eg, *BCR/ABL1* transcript in chronic myeloid leukemia treated with tyrosine kinase inhibitors).

A number of molecular biomarkers have been proposed in CLL that fulfill the definition of prognostic factors, though only a few of them have been consistently validated. Conversely, a few molecular biomarkers meet the statistical definition of predictive factors, and none the biological definition (Table 1).

Molecular prognostic biomarkers in CLL

Immunoglobulin gene mutations

The *IGHV* genes of CLL can accumulate variations as a consequence of the SHM process. The prevalence of mutated *IGHV* genes (defined as a <98% identity compared with the germ line nucleotide sequence) is higher among newly diagnosed and asymptomatic CLL patients (~60%), whereas the prevalence of unmutated *IGHV* genes (defined as ≥98% identity compared with the germ line nucleotide sequence) is higher among progressive (~50%-60%) and relapsed/refractory (~70%-80%) CLL patients. Patients with mutated *IGHV* genes, when compared with patients with unmutated *IGHV* genes, experience longer time to first treatment when managed with “watch and wait,” longer progression-free survival (PFS) when treated with chemoimmunotherapy, lower risk of transformation, and thus, in the end, better overall survival (OS) (Table 2).¹⁰⁻²⁹ The robust and consistent prognostic value of *IGHV* mutation status has been formally validated by a systematic review and meta-analysis.³⁰ Apart from the mutation status, the usage of specific *IGHV* genes may also affect prognosis, including progression (ie, *IGHV3-21* in stereotyped subset 2) and transformation to Richter syndrome (*IGHV4-39* in stereotyped subset 8).^{17,19}

Genetic lesions

Over the last few years, several genetic prognostic biomarkers have been identified that are significantly associated with CLL OS, time to first treatment in cases managed with “watch and wait,” or PFS in treated cases.

In a relevant fraction of CLL patients (~25%), deletion of 13q14 occurs in the absence of any concomitant driver genetic lesion. Patients harboring solely 13q14 deletion have an excellent clinical outcome with a progression rate of <1% per year, longer PFS after therapy, and low risk of transformation, which overall translate into an expected survival only slightly lower than that of the general population.³¹

Deletion of 11q22-23 always includes *ATM* and occurs in <10% of newly diagnosed CLL, whereas its prevalence rises to ~20% at the time of first treatment and 30% at disease relapse. Mutations of the *SF3B1* and *NOTCH1* genes are observed in 10% to 15% of newly diagnosed CLL, whereas their prevalence rises to ~20% in progressive or relapsing CLL. Deletion of 11q22-23, *SF3B1* mutations and *NOTCH1* mutations identify patients with intermediate-risk disease (~40% of cases are alive at 10 years) whose survival is ~50% less than that expected for the general population.³¹ Patients with deletion of 11q22-23, *SF3B1* mutations, or *NOTCH1* mutations have short time to progression if managed with “watch and wait,”

Table 1. Prognostic and predictive biomarkers commonly used in clinical practice

	Prognostic	Predictive
Age	●	●
CIRS	●	●
Stage	●	●
β2-microglobulin	●	●
CD49d	●	●
CD38	●	●
ZAP70	●	●
<i>IGHV</i> mutation	●	●
17p13 deletion	●	●
11q22-23 deletion	●	●
Trisomy 12	●	●
13q14 deletion	●	●
<i>TP53</i> mutation	●	●
<i>SF3B1</i> mutation	●	●
<i>NOTCH1</i> mutation	●	●

A green circle indicates yes, a red circle indicates no, and an orange circle indicates not yet fully validated. CIRS, Cumulative Illness Rating Scale; *IGHV*, immunoglobulin heavy variable gene.

and faster relapse if treated with chemoimmunotherapy, but not if treated with ibrutinib (Table 2).^{3,19,21,23,26-29,31-44} The risk of transformation is specifically affected by *NOTCH1* mutations, but not by 11q22-23 deletion or by *SF3B1* mutations.^{19,34,42}

At diagnosis, the *TP53* gene is disrupted in 4% to 8% of unselected CLL by deletions, mutations, or a combination of both. The incidence rises to 10% at the time of first-line treatment, 30% to 40% in relapsed/refractory CLL, and 50% to 60% in Richter syndrome because *TP53* abnormalities can be acquired/selected during the course of the disease.^{13,19,23,26-28,31,37,39,43} *TP53* abnormalities mark the worst outcome in CLL, with an estimated median OS of 3 to 5 years, which is ~70% less than that expected for the general population.³¹ Patients having *TP53* abnormalities have short time to progression if managed with “watch and wait,” and generally fail chemoimmunotherapy (Table 2).^{13,19,23,26-28,31,37,39,45} The risk of transformation is also increased among patients with *TP53* abnormalities.⁴² The robust and consistent prognostic value of *TP53* abnormalities in CLL has been formally validated by a systematic review and meta-analysis.³⁰ The negative impact of *TP53* abnormalities is only partly mitigated, but not overcome, by ibrutinib, idelalisib, or venetoclax, at least in the relapsed/refractory setting.^{6,44,46,47}

A number of other gene mutations recurring at lower frequency have been signaled for their association with CLL outcome, including *BIRC3*, *SAMHD1*, *RPS15*, *NFKBIE*, *EGR2*, *KRAS*, and *POT1*. However, their annotation as CLL prognostic biomarkers is controversial or awaits validation studies.⁴⁷

Table 2. Clinical outcome of unselected, previously untreated CLL from study cohorts

Ref.	Time to first treatment		OS	
	<i>TP53</i> ^{mut}	<i>TP53</i> ^{wt}	<i>TP53</i> ^{mut}	<i>TP53</i> ^{wt}
37	4.8 y (median)	7.5 y (median)	60.9% (at 5 y)	86.8% (at 5 y)
19	—	—	4.6 y (median)	13.9 y (median)
31	—	—	50.7% (at 5 y)	76.5% (at 5 y)
28	—	—	54.6% (at 5 y)	81.8% (at 5 y)
23	0.3 y (median)	5.3 y (median)	4.6 y (median)	12.5 y (median)
39	1.5 y (median)	3.5 y (median)	—	—
26	—	—	31.9% (at 5 y)	74.6% (at 5 y)
45	—	—	66.2% (at 5 y)	85.2% (at 5 y)
27	—	—	43.5% (at 5 y)	68.0% (at 5 y)
13	1.8 y (median)	Not reached	2.5 y (median)	Not reached
	<i>NOTCH1</i> ^{mut}	<i>NOTCH1</i> ^{wt}	<i>NOTCH1</i> ^{mut}	<i>NOTCH1</i> ^{wt}
37	3.5 y (median)	7.6 y (median)	75.7% (at 5 y)	85.1% (at 5 y)
19	—	—	3.5 y (median)	13.9 y (median)
31	—	—	56.3% (at 5 y)	75.2% (at 5 y)
28	0.2 y (median)	3.4 y (median)	63.6% (at 5 y)	85.5% (at 5 y)
23	0.4 y (median)	5.3 y (median)	5.5 y (median)	12.8 y (median)
39	3.1 y (median)	3.1 y (median)	—	—
26	—	—	36.3% (at 5 y)	72.5% (at 5 y)
27	—	—	69.9% (at 5 y)	76.1% (at 5 y)
35	3.6 y (median)	7.5 y (median)	—	—
34	1.8 y (median)	6.0 y (median)	64% (at 10 y)	35% (at 10 y)
36	0.2 y (median)	1.2 y (median)	55% (at 5 y)	75% (at 5 y)
38	45% (at 5 y)	55% (at 5 y)	75% (at 5 y)	95% (at 5 y)
33	—	—	10.4 y (median)	Not reached
21	0.4 y (median)	4.9 y (median)	5.5 y (median)	Not reached
40	—	—	38% (at 5 y)	70% (at 5 y)
	<i>SF3B1</i> ^{mut}	<i>SF3B1</i> ^{wt}	<i>SF3B1</i> ^{mut}	<i>SF3B1</i> ^{wt}
37	3.8 y (median)	8.0 y (median)	64.7% (at 5 y)	86.7% (at 5 y)
31	—	—	60.3% (at 5 y)	73.85 (at 5 y)
28	0.2 y (median)	2.0 y (median)	—	—
23	0.1 y (median)	4.8 y (median)	5.3 y (median)	12.2 y (median)
39	1.9 y (median)	3.0 y (median)	—	—
26	—	—	53% (at 5 y)	70.8% (at 5 y)
27	—	—	64.9% (at 5 y)	77.7% (at 5 y)
21	0.2 y (median)	4.7 y (median)	5.3 y (median)	Not reached
40	—	—	46% (at 5 y)	68% (at 5 y)
29	0.8 y (median)	1.1 y (median)	—	—
32	—	—	30.0% (at 10 y)	77% (at 10 y)
41	1.2 y (median)	5.0 y (median)	—	—
	<i>IGHV</i> ^{unmut}	<i>IGHV</i> ^{mut}	<i>IGHV</i> ^{unmut}	<i>IGHV</i> ^{mut}
19	—	—	11.7 y (median)	Not reached
28	0.3 y	7 y (median)	74.6% (at 5 y)	87.3% (at 5 y)
23	0.9 y (median)	9.9 y (median)	6.7 y (median)	Not reached
26	—	—	55.6% (at 5 y)	76.4% (at 5 y)
27	—	—	6.4 y (median)	Not reached
13	1.5 y (median)	9.2 y (median)	81% (at 5 y)	88% (at 5 y)
21	0.8 y (median)	9.9 y (median)	6.4 y (median)	Not reached
29	0.3 y (median)	5.4 y (median)	—	—
10	—	—	9.7 y (median)	24.5 y (median)
11	—	—	9.0 y (median)	—
22	—	—	7.0 y (median)	Not reached
12	—	—	8.0 y (median)	Not reached
25	—	—	43% (at 10 y)	87% (at 10 y)
16	5.0 y (median)	Not reached	3.2 y (median)	—
17	—	—	7.0 y (median)	Not reached
20	—	—	41% (at 10 y)	75% (at 10 y)
24	2.4 y (median)	19 y (median)	9.8 y (median)	17.9 y (median)
18	4.6 y (median)	Not reached	13.0 y (median)	23.0 y (median)
15	1.1 y (median)	Not reached	7.0 y (median)	Not reached
14	2.8 y (median)	11.0 y (median)	9.7 y (median)	Not reached

Clinical outcome of unselected, previously untreated CLL from study cohorts including >200 patients stratified according to the *TP53*, *NOTCH1*, *SF3B1*, and *IGHV* mutation status. —, not available; Ref., reference.

Score systems

Hierarchical prognostic models integrating together multiple genetic features of CLL and partitioning the risk according to the prognostic weight of the genetic biomarkers effectively stratify CLL survival. In 2000, this notion was unequivocally documented by the seminal study of Döhner et al who established a hierarchical prognostic model for CLL based on 5 risk categories.³ CLL cases harboring 17p13 deletion independent of concomitant abnormalities had the worst prognosis (median survival, 32 months), followed by cases carrying 11q22-23 deletion (median survival, 79 months), trisomy 12 (median survival, 114 months), normal fluorescence in situ hybridization (FISH) (median survival, 111 months), and 13q14 deletion (median survival, 133 months).³ Though the FISH hierarchical model was developed in the chemotherapy era, it stands also when applied to stratify the survival of patients treated with ibrutinib.⁴⁸

Integration of the most recurrent mutations onto the backbone of the FISH hierarchical model has allowed an improvement in outcome discrimination.³¹ Four groups of patients are hierarchically classified by the integrated mutational-cytogenetic model: (1) high-risk patients, harboring *TP53* and/or *BIRC3* abnormalities independent of co-occurring genetic lesions, who account for ~15% to 20% of newly diagnosed CLL and show a 10-year survival of 29%; (2) intermediate-risk patients, harboring *NOTCH1* and/or *SF3B1* mutations and/or del11q22-23 in the absence of *BIRC3* and *TP53* abnormalities, who account for ~15% to 20% of newly diagnosed CLL and show a 10-year survival of 37%; (3) low-risk patients, harboring +12 or normal genetics, who account for ~40% of newly diagnosed CLL and show a 10-year survival of 57%; and (4) very-low-risk patients, harboring del13q14 only in the absence of any additional abnormality, who account for ~25% of newly diagnosed CLL and have a nearly normal life expectancy with a 10-year survival of 69%.³¹

With the broadening of the number of prognostic biomarkers, it has become difficult for the practicing clinician to determine which factors are most important to counsel individual patients, and how to combine the results of multiple prognostic tests when some suggest an aggressive clinical course and others a more indolent one in the same patient. Systematic reviews and meta-analyses illustrated the robust and consistent prognostic value of *IGHV* mutation status and FISH results for the 11q22-23 and 17p13 chromosomal abnormalities in patients with previously untreated CLL.³⁰ These studies also indicated that *IGHV* mutation status and FISH karyotyping provide complementary information with respect to outcome anticipation. Accordingly, recent efforts have focused on combining *IGHV* mutation status and genetic biomarkers into scoring systems that may anticipate CLL survival.

The German CLL Study Group (GCLLSG) developed a comprehensive prognostic index by taking advantage of a large population of prospectively collected patients within its clinical trials.⁴⁹ The prognostic index score is calculated by giving points to 17p13 deletion, 11q22-q23 deletion, and *IGHV* mutation status among molecular biomarkers, and sex, age, ECOG performance status, serum thymidine kinase, and β 2-microglobulin among clinical factors (Table 3). Patients are then grouped into low (5-year survival, 95%), intermediate (5-year survival, 86%), high (5-year survival, 67%), and very-high (5-year survival, 18%) risk groups.⁴⁹ Among biomarkers, the GCLLSG score includes serum thymidine kinase. Thymidine kinase is a key cellular enzyme of the salvage pathway of DNA

Table 3. GCLLSG Prognostic Index

	Score	5-y survival, %
Biomarker		
Age, y		
≤60	0	
>60	1	
Sex		
Female	0	
Male	1	
ECOG PS		
0	0	
>0	1	
β 2-microglobulin, mg/L		
≤1.7	0	
1.7-3.5	1	
>3.5	2	
Serum thymidine kinase, U/L		
≤10	0	
>10	2	
<i>IGHV</i>		
Mutated	0	
Unmutated	1	
11q22-q23		
Nondeleted	0	
Deleted	1	
17p13		
Nondeleted	0	
Deleted	6	
Risk group		
Low risk	1-2	95
Intermediate risk	3-5	86
High risk	6-10	67
Very-high risk	11-14	18

ECOG PS, Eastern Cooperative Oncology Group performance status; GCLLSG, German CLL Study Group.

synthesis whose activity is cell cycle dependent, and therefore it represents a general proliferation marker. Serum thymidine kinase is available as a routine clinical assay in some European countries, whereas it is less used in the United States. Therefore, a slightly modified version of the model based on 7 prognostic variables, instead of 8, and not including serum thymidine kinase has been proposed.⁵⁰

More recently, an international collaboration used information from ~3500 patients enrolled in clinical trials or institutional/population-based cohorts to develop a comprehensive CLL-International Prognostic Index (IPI), which was subsequently validated in independent CLL populations (Table 4).^{27,51,52} The CLL-IPI score incorporates both molecular and clinical CLL aspects, and is based on 5 robust and widely used prognostic biomarkers (age, clinical stage, 17p13 deletion and/or *TP53* mutation, *IGHV* mutation status, and β 2-microglobulin), and documents that *TP53* disruption is the sole recurrent genetic abnormality that shows independent prognostic information after adjusting for *IGHV* mutation status and for the most important clinical variables. Conversely, 11q22-q23 deletion and mutations of *NOTCH1* and *SF3B1* do not carry independent prognostic value in this model. The CLL-IPI identifies low-, intermediate-, high-, and very-high-risk patients showing significantly different survival at 5 years (93%, 79%, 63%, 23%, respectively). Although the CLL-IPI has been primarily developed to prognosticate CLL survival, it also anticipates time to first treatment in “watch and wait” patients.²⁷ A prognostic system based only on

Table 4. CLL-IPI

	Score	5-y survival, %
Biomarker		
Age, y		
≤65	0	
>65	1	
Stage		
Rai 0/Binet A	0	
Rai I-IV/Binet B-C	1	
β2-microglobulin, mg/L		
≤3.5	0	
>3.5	2	
<i>IGHV</i>		
Mutated	0	
Unmutated	2	
<i>TP53</i>		
No abnormalities	0	
Deletion and/or mutation	4	
Risk group		
Low risk	0-1	93
Intermediate risk	2-3	79
High risk	4-6	63
Very-high risk	7-10	23

IGHV mutation status and FISH cytogenetics that simplifies the CLL-IPI has also been proposed.⁵³

Importantly, these indices were developed in patients in the chemoimmunotherapy era, and will need to be validated in the novel agent era. The use of molecular biomarkers in CLL clinical practice will be discussed below in “Recommendations for clinical practice.”

Molecular predictive biomarkers in CLL

Immunoglobulin gene mutations

Among patients receiving chemoimmunotherapy, the *IGHV* mutation status affects the kinetics of relapse and thus PFS, which is longer in *IGHV*-mutated vs *IGHV*-unmutated patients. Accordingly, ~50% to 60% of patients with mutated *IGHV* genes who receive potent chemoimmunotherapy regimens maintain disease remission long-term, including persistent negativity of minimal residual disease in some instances, which translates into a plateau on the PFS curve, no relapses beyond 10 years, and an OS similar to the one expected in normal healthy subjects. Conversely, long-term, almost all *IGHV* unmutated CLL patients are projected to progress after chemoimmunotherapy.⁵⁴⁻⁵⁶

In contrast to chemoimmunotherapy, CLL patients benefit from ibrutinib and idelalisib independent of *IGHV* mutation status. Indeed, upon treatment with ibrutinib or idelalisib, the PFS of *IGHV*-unmutated patients is similar to that of *IGHV*-mutated cases.^{44,48,57} The lymphocytosis tends to resolve more quickly in the *IGHV*-unmutated group. On these bases, though the interaction between different treatments (chemoimmunotherapy vs novel targeted agents), *IGHV* mutation status, and outcome has never been tested in a clinical trial, cross-trial comparisons suggest that unmutated *IGHV* genes might fulfill the statistical definition of predictive biomarker for lack of benefit from chemoimmunotherapy.

Genetic lesions

TP53 codes for a central regulator of the DNA damage response pathway and is the target of the genotoxic effect of chemotherapy.

Upon chemotherapy-induced DNA damage, *TP53* is activated and CLL cells undergo apoptosis. Conversely, when *TP53* is non-functional because of deletion or mutation, the apoptosis of CLL cells cannot be triggered in response to chemotherapy. Consistently, CLL patients with 17p13 deletion or *TP53* mutation have a very poor response to chemoimmunotherapy regimens, including fludarabine and cyclophosphamide (FC) and rituximab (FCR), bendamustine and rituximab (BR), obinutuzumab and chlorambucil (G-Clb), ofatumumab and chlorambucil (O-Clb), or rituximab and chlorambucil (R-Clb).⁵⁸⁻⁶¹ Ibrutinib, idelalisib, and venetoclax do not exert their antileukemic activity through genotoxic mechanisms, and are therefore active irrespective of *TP53* dysfunction.⁶² Consistently, the response rate in *TP53*-disrupted patients is superimposable to that observed in patients with a wild-type *TP53* gene, and better than that observed in every previous historical control treated with chemoimmunotherapy.⁶³⁻⁶⁵ Notably, the interaction between treatment (chemoimmunotherapy vs novel targeted agents), *TP53* status, and outcome has never been formally tested in a clinical trial. However, cross-trial comparison of patients' outcome according to *TP53* status allows for indirectly suggesting that *TP53* disruption might fulfill the statistical definition of predictive biomarker for lack of benefit from chemoimmunotherapy in CLL.

Among new CLL genetic lesions, only *NOTCH1* mutations behave as a candidate predictive biomarker. Indeed, among CLL harboring *NOTCH1* mutations, treatment with FCR, R-Clb, or O-Clb does not result in the expected increase in PFS compared with treatment with FC or with chlorambucil alone.⁶⁶⁻⁶⁸ These observations point to *NOTCH1* mutations as a biomarker of resistance to the anti-CD20 monoclonal antibodies rituximab and ofatumumab in CLL. The outcome of CLL patients treated with obinutuzumab combined to chlorambucil is improved independent of *NOTCH1* mutation status, suggesting that the augmented cytotoxicity of obinutuzumab or the increased dose of the anti-CD20 antibody used in the obinutuzumab-chlorambucil schema overcomes *NOTCH1* mutation-associated resistance to rituximab.⁶⁷ The mechanism underlying the anti-CD20 refractoriness associated with *NOTCH1* mutations remains obscure.

Resistance to ibrutinib, idelalisib, or venetoclax is not fully captured by predictive biomarkers that are otherwise used in the setting of chemoimmunotherapy. On this basis, novel biomarkers are required to redefine high-risk CLL patients in the era of novel agents. Development of resistance to ibrutinib is due to acquired mutations in Bruton tyrosine kinase (*BTK*) at the binding site of ibrutinib, and gain-of-function mutations in phospholipase C γ 2 (*PLCG2*), a direct downstream target of *BTK*.⁶⁹ The reason why some CLL cases are prone to develop mutation-driven resistance to ibrutinib, whereas other cases persist in long-lasting remission, is unknown. The notion that *BTK* and *PLCG2* mutations associate with complex karyotype and with defects of the DNA damage response pathway (ie, *TP53* and *ATM* abnormalities) suggests that genetic instability may be involved in the development of resistance.

Recommendations for clinical practice

The great activity of FCR in patients with mutated *IGHV*, and the improvement of outcome among *IGHV*-unmutated patients when treated with ibrutinib, may announce the emergence of a therapeutic approach that is guided by the *IGHV* mutational status (Figure 3). Accordingly, the most recent guidelines support *IGHV* mutation analysis as desirable at the time of treatment requirement, and a consensus has been reached on the minimal technical requirements for a reliable and reproducible analysis of the rearranged *IGHV*

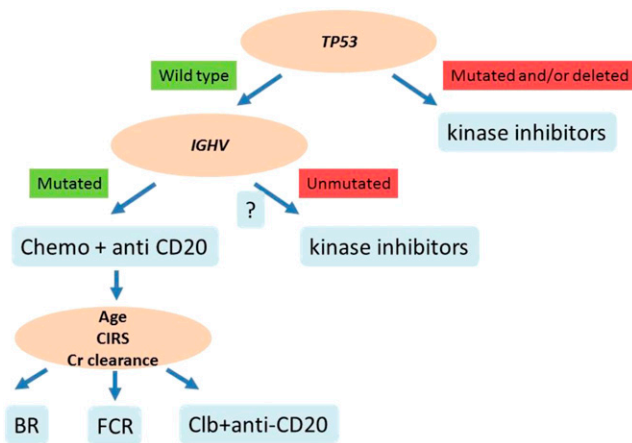


Figure 3. Biomarker-informed decision nodes in the management of newly presented CLL. At the time of treatment requirement, the presence of *TP53* mutation/deletion represents an indication for treatment with kinase inhibitors, for example, ibrutinib or, alternatively, idelalisib. Patients who require first-line treatment and carry a wild-type *TP53* gene may be candidate to chemoimmunotherapy with one of the available combinations, especially in the presence of mutated *IGHV* genes. In patients with wild-type, a *TP53* gene, and unmutated *IGHV* genes, the role of chemoimmunotherapy is under debate, and kinase inhibitors may represent a valuable option. CIRS, Cumulative Illness Rating Scale.

sequences.⁷⁰⁻⁷² Because the *IGHV* mutation status does not change over time, there is no need to repeat its analysis during disease course.

Given the opportunity to effectively treat *TP53*-disrupted patients with novel targeted agents, CLL guidelines recommend testing for 17p13 deletion and *TP53* mutations in patients requiring therapy (Figure 3). Mutations represent the most frequent form of *TP53* inactivation in CLL, and frequently (~70% of the cases harboring *TP53* disruption) pair with the loss of the second *TP53* allele through 17p13 deletion. A proportion of patients harbor the sole *TP53* mutation without 17p13 deletion, whereas 17p13 deletion in the absence of *TP53* mutation is less frequent. The direct clinical implications of such a complex and heterogeneous picture of *TP53* defects in CLL are that both FISH cytogenetics and sequencing should be done to have a comprehensive assessment of the *TP53* gene status.⁷³ Because leukemic clones may evolve, 17p13 deletion and *TP53* mutation analyses should be repeated at each disease progression requiring treatment if it was normal before. Sanger sequencing is the currently recommended approach for *TP53* mutation analysis.

Prognostic scores allow very-low-risk patients to be sorted out; these patients have a life expectancy similar to that expected in the normal population and show a low likelihood of progression. On the other hand, prognostic scores also allow for identification of very-high-risk patients who generally progress and shortly succumb to the disease. However, prognostic scores are not 100% accurate (the outcome discrimination capacity is 73% for the CLL-IPI and 68% for the integrated mutational-cytogenetic model) because they all comprise intermediate-risk categories, which are a case mix containing both low- and high-risk patients whose outcome is not captured by the scoring system.^{27,31} Although scoring systems may be useful for patient counseling and for the definition of the timing of follow-up, they can turn a “watch-and-wait” strategy into a “watch-and-worry”

situation in patients without treatment indication. Therefore, at the time of diagnosis, in asymptomatic patients who do not require treatment, current guidelines do not support the need for a full prognostic assessment also incorporating molecular biomarkers such as gene mutations, FISH, and *IGHV* analysis.

Perspectives

Conventional Sanger sequencing has limited sensitivity and allows for the identification of *TP53* mutations represented in at least 10% to 20% of the alleles. Next-generation sequencing is more sensitive than Sanger sequencing for the identification of small *TP53*-mutated subclones (down to 1%), which occur in a significant fraction of CLL, have the same unfavorable prognostic impact as clonal *TP53* defects, and anticipate the development of a chemorefractory phenotype among CLL patients requiring treatment.^{26,28} However, given the risk of false-positive calls when dealing with the identification of small subclonal mutations by next-generation sequencing, methodologies and analyses should be standardized and harmonized at the international level before incorporating this approach in the clinical setting.

The robustness of *NOTCH1* mutations as a biomarker of resistance to rituximab and ofatumumab in CLL, as well as the possibility that obinutuzumab may overcome such resistance, requires independent clinical validation.⁶⁶⁻⁶⁸ Thus, it is premature to recommend *NOTCH1* mutation testing routinely in clinical practice.

Treatment-emergent mutations of *BTK* and *PLCG2* are found in 85% of ibrutinib-resistant CLLs and these mutations are detected up to 15 months before clinical relapse.⁶⁹ Although these observations support *BTK* and *PLCG2* mutations as a sensitive biomarker of ibrutinib resistance, the notion that such mutations can be detected even in CLL patients without clues of clinical relapse may suggest a limited specificity for ibrutinib resistance. Prospective investigations are thus needed before translating *BTK* and *PLCG2* mutations as biomarkers to inform the decision to switch patients from ibrutinib treatment toward alternative options.

Conclusions

The genome of CLL has a strong impact on the clinical outcome of the disease and has revealed important molecular biomarkers. In clinical practice, predictive biomarkers heavily impact on clinician’s choice because they can inform CLL treatment decisions. Currently, the best example is represented by *TP53* disruption by gene mutation and/or deletion that associates with chemoimmunotherapy failure and mandates treatment with innovative drugs, including ibrutinib, idelalisib, or venetoclax. The mutation status of *IGHV* genes represents an additional predictive biomarker, which helps identify patients who may benefit the most from chemoimmunotherapy with FCR. Assessment of these biomarkers at the time of treatment requirement is recommended by most current guidelines for the management of CLL. Other molecular predictors are under investigation, but their application in clinical practice is premature.

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