

Supplemental Materials

Supplemental Tables

Supplemental Table S1. Gene targets of SM CLL/lymphoma panel

Transcription regulators	<i>ARID1B, ARID2, BCORL1, CREBBP, KMT2A, KMT2D, KDM6A, TET2</i>
TLR signaling pathway	<i>MYD88</i>
NF-κB signaling pathway	<i>BIRC3, CARD11, MAP2K1, TNFAIP3, TRAF3</i>
BCR signaling pathway	<i>CD28, CD79A, CD79B, TNFRSF14</i>
MAPK signaling pathway	<i>BRAF, KRAS, NF1, NRAS</i>
DNA damage and repair	<i>ATM, TP53</i>
RNA splicing and processing	<i>SF3B1</i>
Others	<i>APC, CCND1, CCND2, CHD8, DIS3, DNMT3A, EZH2, FOXO1, GNA13, ID3, IDH2, NOTCH1, NOTCH2, RHOA, STAT3, STAT5B, TCF3, ZAP70</i>

Abbreviations: TLR, Toll-like receptor; MAPK, mitogen-activated protein kinase; **SM CLL**, **St. Mary's chronic lymphocytic leukemia**

Supplemental Table S2. List of detected mutations

See the file "CLL Supplementary Table 2.xlsx"

Supplemental Table S3. Univariate and multivariate **Cox proportional hazards** regression analyses of genetic factors for time to treatment

Variables	Univariate analysis			Multivariate analysis		
	HR	(95% CI)	<i>p</i>	HR	(95% CI)	<i>p</i>
Somatic hypermutation	0.40	(0.19-0.84)	0.016	0.57	(0.23-1.42)	0.226
Isolated del(13q)	1.93	(0.74-5.01)	0.178			
Trisomy 12	1.67	(0.62-4.50)	0.313			
Complex karyotype	1.74	(0.70-4.33)	0.237			
<i>ATM</i> abnormality	1.86	(0.72-4.80)	0.203			
<i>TP53</i> abnormality	2.95	(1.22-7.12)	0.016	2.86	(1.17-7.00)	0.021
<i>NOTCH1</i> mutation	4.51	(1.87-10.91)	<0.001	4.99	(1.49-16.69)	0.009
<i>MYD88</i> mutation	0.68	(0.34-1.39)	0.292			

Abbreviations: HR, hazard ratio; CI, confidence interval

Supplemental Table S4. Univariate and multivariate **Cox proportional hazards** regression analyses of genetic factors for overall survival

Variables	Univariate analysis			Multivariate analysis		
	HR	(95% CI)	<i>p</i>	HR	(95% CI)	<i>p</i>
Somatic hypermutation	0.16	(0.04-0.60)	0.006	0.50	(0.10-2.51)	0.403
Isolated del(13q)	1.39	(0.14-13.50)	0.774			
Trisomy 12	1.42	(0.17-11.64)	0.741			
Complex karyotype	1.45	(0.16-13.05)	0.738			
<i>ATM</i> abnormality	1.69	(0.20-14.15)	0.630			
<i>TP53</i> abnormality	7.26	(1.73-30.58)	0.007	7.07	(1.56-32.1)	0.001
<i>NOTCH1</i> mutation	7.21	(1.45-35.88)	0.016	3.47	(0.44-27.6)	0.239
<i>MYD88</i> mutation	0.56	(0.12-2.64)	0.463			

Abbreviations: HR, hazard ratio; CI, confidence interval