

Real-world data on chronic myelomonocytic leukemia: clinical and molecular characteristics, treatment, emerging drugs, and patient outcomes

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Table S1: NGS panels used for gene mutation analysis.

NGS panel	Genes and regions studied
Ion Ampliseq™ AML Research Panel (ThermoFisher Scientific)	Coding regions: <i>CEBPA, DNMT3A, GATA2, TET2, TP53</i> Hotspot regions: <i>ASXL1, BRAF, CBL, FLT3, IDH1, IDH2, JAK2, KIT, KRAS, NPM1, NRAS, PTPN11, RUNX1, WT1</i>
	Coding regions: <i>ASXL1, BCOR, CALR, CEBPA, ETV6, EZH2, IKZF1, NF1, PHF6, PRPF8, RB1, RUNX1, SH2B3, STAG2, TET2, TP53, ZRSR2</i>
OncoPrint™ Myeloid Research Panel (ThermoFisher Scientific)	Hotspot regions: <i>ABL1, BRAF, CBL, CSF3R, DNMT3A, FLT3, GATA2, HRAS, IDH1, IDH2, JAK2, KIT, KRAS, MPL, MYD88, NPM1, NRAS, PTPN11, SETBP1, SF3B1, SRSF2, U2AF1, WT1</i> Genes involved in gene fusions: <i>ABL1, ALK, BCL2, BRAF, CCND1, CREBBP, EGFR, ETV6, FGFR1, FGFR2, FUS, HMGA2, JAK2, KMT2A, MECOM, MET, MLLT10, MLLT3, MYBL1, MYH11, NTRK3, NUP214, PDGFRA, PDGFRB, RARA, RBM15, RUNX1, TCF3, TFE3</i>

Table S2: Cytogenetic abnormalities observed in 50 of 156 CMML patients.

Karyotype	<i>n</i> = 156 <i>n</i> (%)
Normal	106 (67.9)
Abnormal	50 (32.1)
+8 isolated	11
-7/del(7q) isolated	7
-Y	7
Complex karyotype	6
Others	
47,XX,+8[5]/48,idem,+21[2]	1
46,XY,t(5;6)(q31;q22);t(12;20)(p13;q11)[20]	1
46,XX,del(17)(p12)[20]	1
47,XY,+21[7]/46,XY[20]	1
46,XX,t(5;16)[3]/46,XX[17]	1
46,XY,t(3;7)(q26;q21)[10]/46,XY,idem,+13[4]	1
47,XY,+mar[2]/46,XX[30]	1
47,XY,+21[12]/46,XY[8]	1
46,XX,del(11)(q22q24)[14]/46,idem,del(12)(p12p13)[5]	1
46,XY, t(2;3) (p21;q27)[26]	1
46XY,del(11)(q14.3) [20]	1
46,XX,del(5)(q33)[5]/46,XX[15]	1
46,XY,add(6)(q23)[4]/46 XY[16]	1
46,XX,del-5(q13q33)[18]	1
46,XY,del(12)(p12)[8]/46, XY[12]	1
46,XY,del(20)(q11q13)[7]/46,XY[13]	1
47,XY,+11[12]/46,XY[8]	1
46,XY,t(14;15)(q32;q22)[14]	1
46,XY,del(20)(q11q13)[8]/46,XY[8]	1

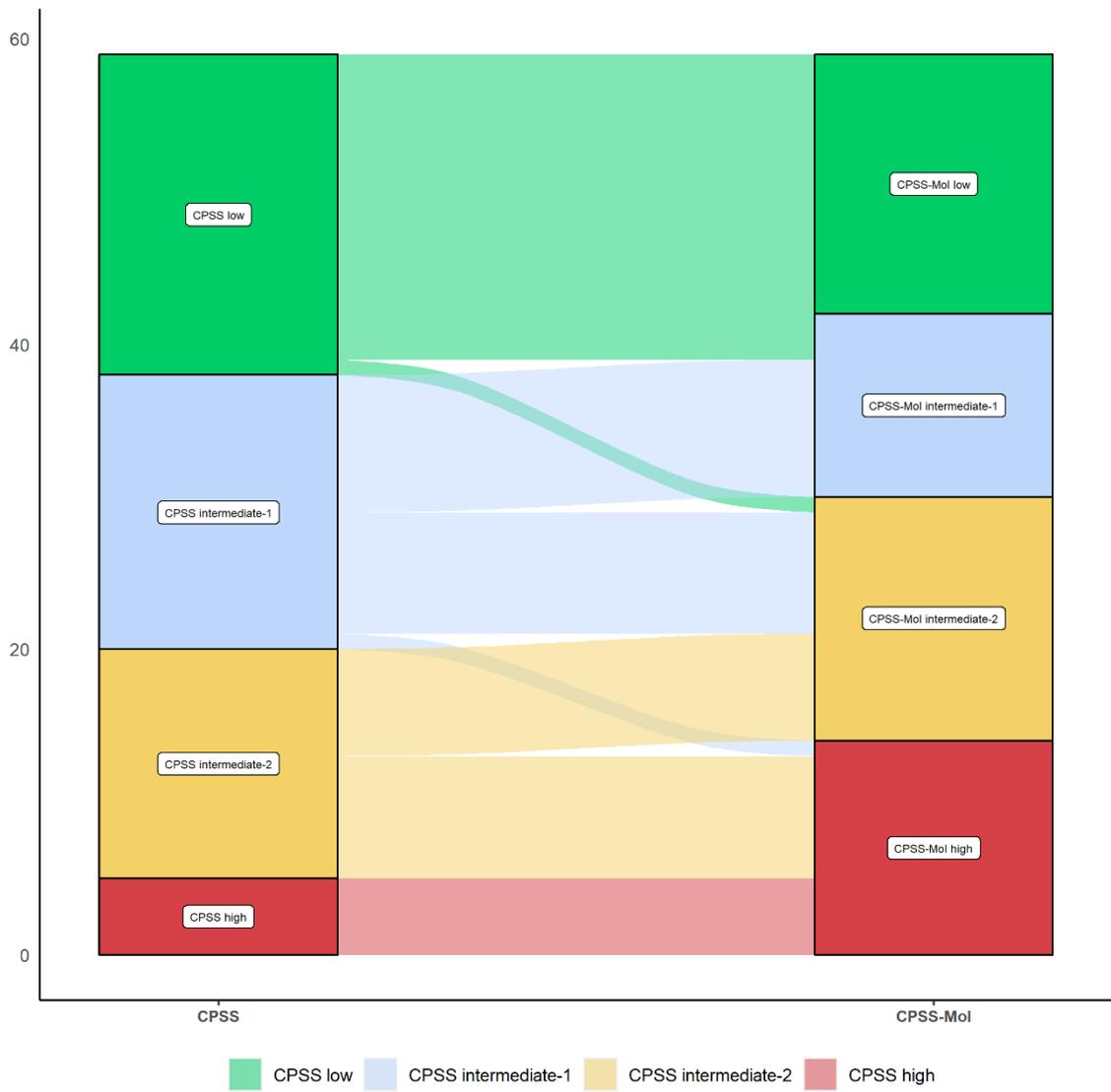


Figure S1: Alluvial plot of the patients included in the study according to CPSS and CPSS-Mol scores.

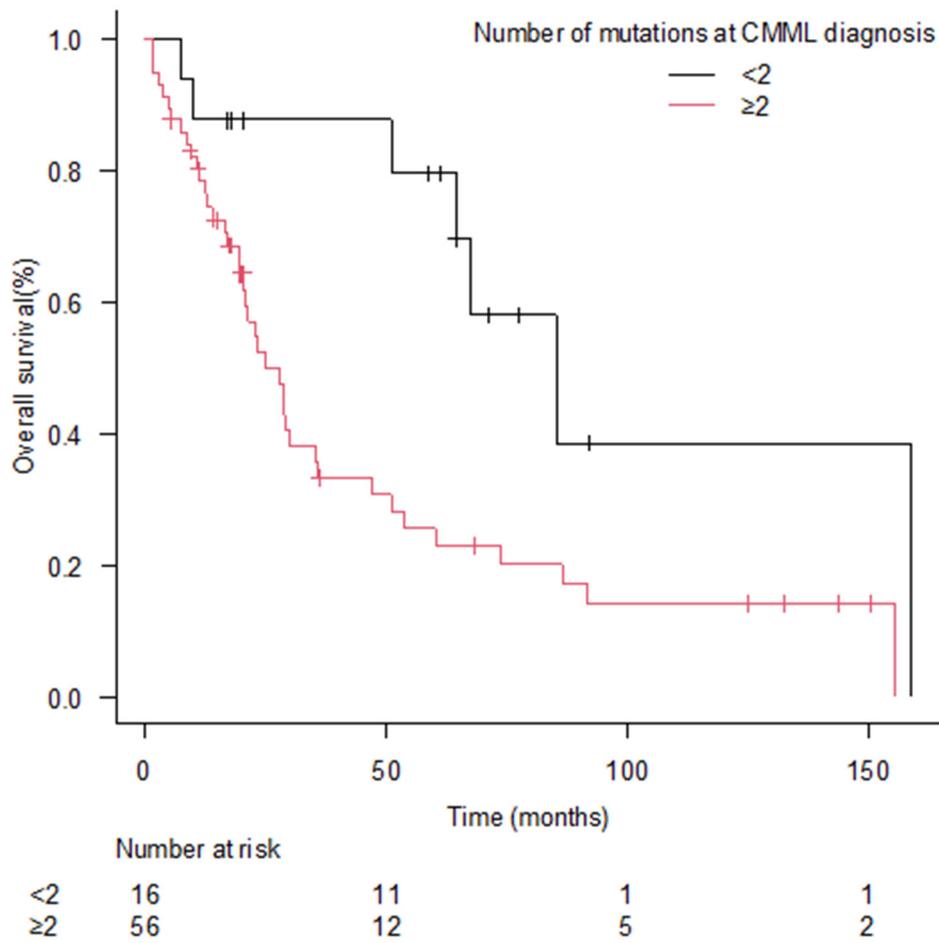


Figure S2: Overall survival according to the number of mutations detected at CMML diagnosis.

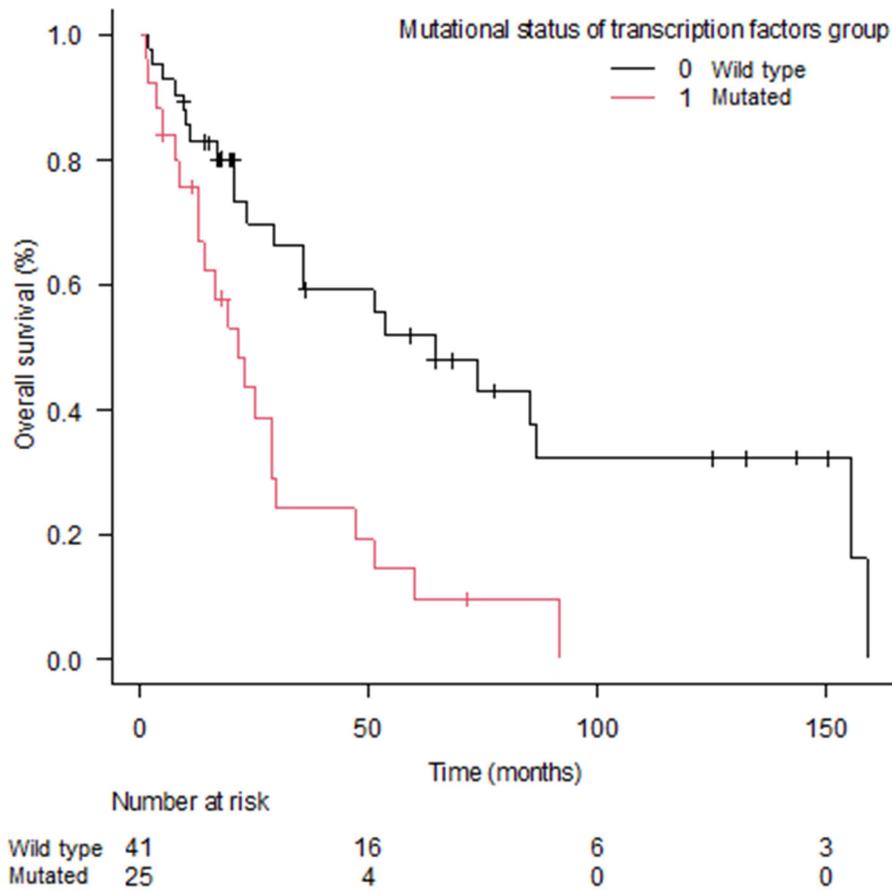


Figure S3: Overall survival according to the mutation status of transcription factors.