

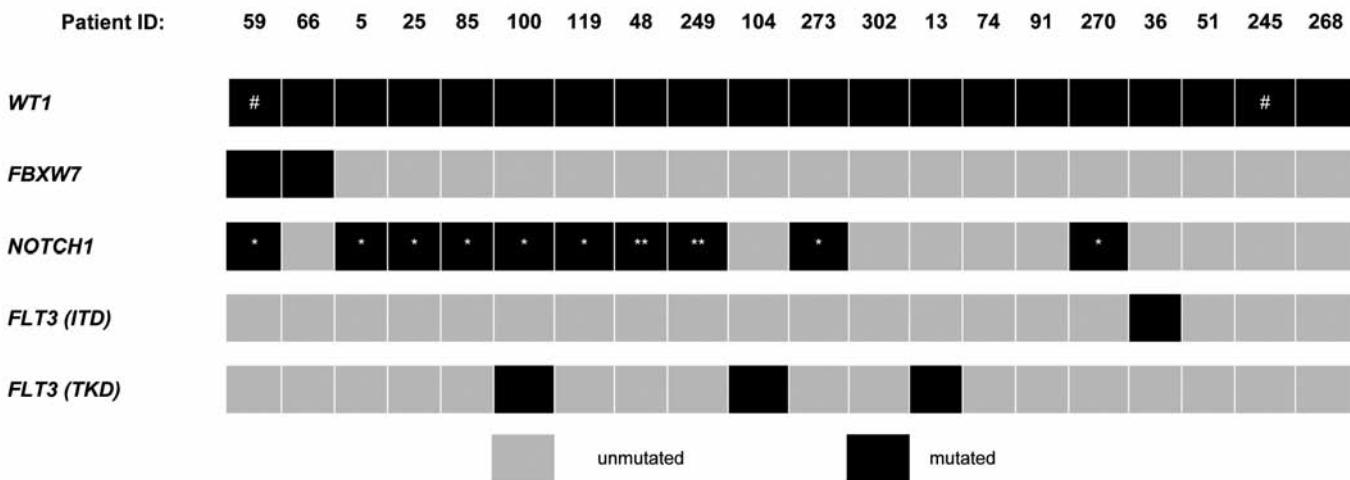
Prognostic implications of mutations and expression of the Wilms tumor 1 (WT1) gene in adult acute T-lymphoblastic leukemia

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Online Supplementary Figure S1. Coexistence of WT1 mutations with mutations in the FBXW7, NOTCH1, and FLT3 genes. T-ALL patients with WT1 mutations (n=20) are listed and coexistence of mutation in the FBXW7, NOTCH1, and FLT3 genes are indicated by black boxes. # WT1 mutations in exons 7 and 9; * NOTCH1 mutations in the HD domain; ** NOTCH1 mutations in the PEST domain; ITD: internal tandem duplication; TKD: D835 mutation in the tyrosine kinase domain.

Online Supplementary Table S1. T-ALL patients with *WT1* mutations.

Patient ID	Age	Sex	DNA change [#]	Clones*	Protein change ^{††}	Outcome	T-ALL subtype	
<i>WT1</i> ^{mut7}								
36	20	M	clone a: 1305_1306insTTGAAAT clone b: [1304_1305insT; 1305_1306insACGATAC]	[5/10] [5/10]	Val371X Arg370LeufsX8	no clinical data available	early T-ALL	
48	27	M	1334_1338dupCGGTC	[5/10]	Ala382Glyfs§	no clinical data available	mature T-ALL	
51	22	F	[1300_1301insA; 1302dupG]	homozygote	Arg369ThrfsX7	failure/death	early T-ALL	
249	18	M	1337_1340delinsAGATGAAGCACCGATACTT	[11/11]	Ser381ArgfsX2	relapse/death	thymic T-ALL	
268	63	F	[1295-5_1309del20; 1312T>C; 1315_1316delAG]	[8/9]	[Asp367_Val371; Val347 SerfsX10]	CCR/alive	early T-ALL	
270	28	M	1305delinsCCC	[8/10]	Arg370ProfsX6	failure/death	mature T-ALL	
273	30	F	clone a: 1334delinsAA clone b: 1333_1334insG	[3/6] [3/6]	Arg380AlafsX5 Ser381ValfsX4	relapse/death	thymic T-ALL	
302	31	F	1319_1320insCC	[6/6]	Thr377Argfs§	failure/death	mature T-ALL	
5	24	M	1334delinsTA	[5/10]	Arg380X	CCR/alive	thymic T-ALL	
13	32	M	clone a: [1304_1305insT; 1305_1306insACGATAC] clone b: 1305_1306insTTGAAAT	[4/10] [5/10]	Arg370LeufsX8 Val371X	SCT/alive	early T-ALL	
25	21	F	clone a: 1305_1306insTAGCGCGGGGGCGGA TTCAGGGCGGCC clone b: 1305delinsCCCCC	[4/10] [5/10]	Val371SerfsX23 Arg370ProfsX16	relapse/alive	thymic T-ALL	
66	40	M	1333dupA	[11/14]	Arg380ThrfsX5	death in CR	thymic T-ALL	
74	32	F	1333dupA	[14/14]	Arg380ThrfsX5	relapse/death	thymic T-ALL	
85	15	M	1305_1306insGTAACCCG	[5/5]	Val371X	SCT/alive	early T-ALL	
91	26	M	1333_1334insCGGAGGGA	[4/5]	Ser381Argfs§	SCT/alive	mature T-ALL	
100	40	M	1334delinsAA	[7/11]	Arg380LysfsX5	SCT/alive	early T-ALL	
104	40	M	1301delinsGG	[5/5]	Arg369GlyfsX16	SCT/death	early T-ALL	
119	23	F	1337_1338insGGGGTTGGCATGGTAGGACGGT	[11/12]	Arg380TrpfsX12	SCT/alive	early T-ALL	
<i>WT1</i> ^{mut7} and <i>WT1</i> ^{mut9}								
				Exon 7	Exon 9			
245	23	M	[1422_1423insGCGGGCC(+) =, 1581G>C*]	[4/6]	[homozygote]	[Met410Glnfs§ (+) Arg462Pro]	SCT/death	early T-ALL
59	36	F	[1334delinsGGG(+) =, 1581G>C*]	[5/10]	[12/13]	[Arg380Glyfs§ (+) Arg462Pro]	death in CR	thymic T-ALL

[#]The stop codons created by these *WT1*^{mut7} do not occur until the 3' part of exon 7. ^{*}Nucleotide sequence variations numbering is according to Genbank accession number NM_024426. ^{††}The sequence variations are designated according to the current recommendations of the Human Genome Variation Society (<http://www.hgvs.org/mutnomen/>). [§]The protein changes are theoretically deduced. *1581G>C mutation previously reported in AML (see ref. 5 in main text). ^{*}Number of mutated clones/total number of analyzed clones. CCR: continuous complete remission; CR: complete remission; SCT: stem cell transplantation in first CR.