SUPPLEMENTARY APPENDIX

Detection of ABL1 kinase domain mutations in therapy-naïve BCR-ABL1-positive acute lymphoblastic leukemia

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Supplementary Table 1

gender	age at ID	date ID	mutation at ID	GMALL ¹	alloSCT	2nd gen TKI ²	Pona ³	therapy data incomplete ⁴	molecular follow- up ⁵	outcom e 6
F	75	Apr-07							no indication for SQ	remission
F	66	Jan-08						х		deceased (< 6 m.)
F	58	Jan-08						х		deceased (< 6 m.)
F	48	Apr-08		Х						deceased (< 6 m.)
F	63	Jul-08		Х	Х					NA
F	71	Aug-08								remission
F	67	Oct-08	c.827A>G, D276G, 3%		х			х		NA
М	44	Nov-08						х		NA
М	40	Jan-09			х	х		х	acquired mutation	relapse
F	46	Jan-09						х		deceased (< 6 m.)
М	18	Mar-09						Х		NA
M	66	Apr-09						х		relapse, deceased (< 6m)
М	39	May-09		Х	х				no indication for SQ	NA
F	74	Jul-09						х	no indication for SQ	remission
М	84	Dec-09						х		NA
М	47	Apr-10						Х		NA
F	51	Apr-10		Х	х				no indication for SQ	remission
М	61	May-10		Х	х				no indication for SQ	remission
M	72	Jun-10	c.756G>T, Q252H, 46% c.756G>C, Q252H, 8%	х		х			mutation present at relapse	remission, relapse
F	20	Jul-10		х						deceased (< 6 m.)
М	63	Aug-10		х						deceased (< 6 m.)

F	62	Nov-10				х		relapse
F	71	Dec-10	х					remission
М	67	Dec-10	х		Х		acquired mutation	remission, relapse
М	62	Mar-11	х					remission
F	21	Apr-11	х	Х			no indication for SQ	remission
F	34	Apr-11		х				remission
М	56	Jun-11	х					deceased (< 6 m.)
F	61	Jul-11		х			no indication for SQ	NA
М	27	Aug-11				х		NA
F	53	Sep-11	х					deceased (< 6 m.)
М	44	Nov-11		х	Х	х	no indication for SQ	remission, relapse
М	47	Dec-11	х	Х		х		NA
М	39	Mar-12	х	Х		х		NA
М	69	Apr-12	х					relapse, deceased (< 6m)
M	47	May-12				Х		NA
М	66	May-12	х		Х			relapse
F	73	Jul-12				х		NA
F	80	Aug-12				Х		NA
F	68	Sep-12	х					remission
М	47	Oct-12	х			х		deceased (< 6 m.)
F	75	Nov-12				х	no indication for SQ	remission
М	55	Jan-13	х				no acquired mutation	relapse
М	76	Feb-13	х				acquired mutation	relapse
M	33	Feb-13	Х					deceased (< 6 m.)
M	69	Mar-13				х		NA
М	18	Mar-13		х		х		relapse
F	59	Mar-13				х	acquired mutation	relapse
F	65	Mar-13	х					remission
F	81	May-13				х		NA
F	74	Jun-13				х		NA
F	44	Aug-13				х		NA
M	32	Aug-13				х		NA
F	38	Sep-13			Х	х	acquired mutation	relapse

М	70	Sep-13		х					no acquired mutation	remission, relapse
M	70	Nov-13		X				Х	mutation	NA NA
М	60	Nov-13		Х						NA NA
F	61	Feb-14						Х		NA
М	77	Feb-14				X	Х		acquired mutation	remission, relapse
М	40	Feb-14		Х					acquired mutation	relapse
М	61	Feb-14						Х		NA NA
F	84	Mar-14						Х		NA
М	71	May-14		Х		х	Х		acquired mutation	remission, relapse
F	52	Jun-14	c.944C>T T315l, 14%	х	х		х			remission
М	49	Jun-14			Х	х	Х	Х	no indication for SQ	relapse
F	70	Aug-14		х		х		х	no acquired mutation	remission, relapse
М	68	Aug-14				Х		Х		NA
М	62	Sep-14				х	Х		acquired mutation	relapse
М	50	Oct-14		х	Х				no indication for SQ	remission
F	76	Nov-14		х					no indication for SQ	NA
М	64	Apr-15		Х			Х		acquired mutation	remission, relapse
F	42	Apr-15		х						deceased (< 6 m.)
М	48	May-15		х						NA
М	28	May-15					Х		acquired mutation	relapse
F	66	Jul-15			х			х	no indication for SQ	NA
М	74	Jul-15						Х		deceased (< 6 m.)
М	49	Jul-15			х			Х	acquired mutation	relapse
М	52	Jul-15		x	x	x		х	no acquired mutation	remission, relapse
F	82	Aug-15		х		х			acquired mutation	relapse
М	79	Oct-15	c.730A>G M244V, 2%					х		NA
М	47	Nov-15			Х		Х		acquired mutation	relapse
М	48	Jan-16		Х	Х	Х			no indication for SQ	remission
М	76	Feb-16	c.763G>A E255K, 1%					х		NA
F	42	Feb-16						х		NA
F	60	Mar-16						х		NA

F	41	Mar-16				х		NA
M	68	Mar-16				х		NA
M	64	Sep-16		Х	Х		no indication for SQ	remission
М	46	Oct-16	Х	х			no indication for SQ	remission
M	62	Oct-16				х		NA
F	45	Nov-16						NA

Abbre viations: F: female, M: male; ID: initial diagnosis; allo SCT: allogeneic stem cell transplantation, SQ: sequencing of the ABL1 kinase domain, NA: not available

- 1) Patient treated as part of the GMALL study or according to the study protocol at the respective time.
- 2) Patient received dasatinib or nilotinib at any point during treatment as single agent or part of a combination therapy.
- 3) Patient received ponatinib at any point during treatment.
- 4) The information on the therapy of this patient could be potentially incomplete.
- 5) For those patients, samples for mutation testing over a follow-up period of at least 6 months, were available. This excludes patients who deceased early (within the first six months) or for whom no material for *BCR-ABL1* expression and mutation testing was available. We defined the indication for mutation testing as a *BCR-ABL1/ABL1* ratio of least ~1% (0.962%).
- 6) Outcome is broadly classified as (a) remission, (b) relapse and (c) deceased within the first 6 moths (m.). The latter patients were excluded from the longitudinal mutation testing.