

## Supplementary

**Table S1:** Correlation of KIR genes and haplotype in child mother and father with the occurrence of FM. Fisher's exact test with risk calculation of odds ratio and 95% confidence analysis was performed. Odds ratios were calculated as follows odds ratio = ((FM-/KIR+)/(FM+/KIR+))/(FM-/KIR-)/(FM+/KIR-). Odds ratio >1 = occurrence or absence of gene affects FM. KIR2DL1, KIR2DL4, KIR3DL2 and KIR2DL3 are present in all donors and therefore were excluded. \*\* Genes are constantly present and therefore were excluded. Significance level  $p < 0.05$ .

KIR	Child			Mother			Father		
	N	p-value	odds ratio (95% CI)	N	p-value	odds ratio (95% CI)	N	p-value	odds ratio (95% CI)
2DL2	51	0.765	1.388 (0.432-4.459)	51	0.202	0.358 (0.085-1.500)	18	1	0.9 (0.133-6.08)
2DL3	51	0.623	1.765 (0.228-13.685)	51	0.04	6.923 (1.23-35.956)	18	1	0.909 (0.754-1.096)
2DL5	50	0.145	0.368 (0.113-1.198)	51	0.065	0.241 (0.058-0.995)	18	0.627	0.500 ( 0.068-3.696)
2DP1	**			51	1	0.969 (0.91-1.031)	**		
2DS1	51	0.08	0.306 (0.094-0.998)	51	0.565	0.642 (0.204-2.017)	18	0.627	0.500 (0.068-3.696)
2DS2	51	0.771	0.75 (0.234-2.404)	51	0.543	0.595 (0.171-2.071)	18	1	1.111 (0.164-7.506)
2DS3	44	0.738	1.447 (0.365-5.735)	44	0.456	2.571 (0.475-13.913)	18	1	0.600 (0.031-11.473)
2DS4	51	0.373	1.056 (0.949-1.174)	51	0.373	1.056 (0.949-1.174)	**		
2DS5	51	0.106	0.317 (0.089-1.132)	51	0.563	0.667 (0.211-2.106)	18	1	2.250 (0.185-27.369)
3DL1	51	0.134	1.118 (0.958-1.304)	51	0.373	1.056 (0.949-1.174)	**		
3DP1	0	0.285	0.906 (0.811-1.013)	51	1	0.969 (0.91-1.031)	18	1	0.909 (0.754-1.096)
3DS1	51	0.07	0.311 (0.091-1.065)	51	0.561	0.616 (0.196-1.933)	18	1	1.333 (0.098-18.192)
Haplotype	51	0.128	6 (0.687-52.383)	51	0.236	4.154 (0.4637-37.513)	18	1	1 (0.072-13.868)

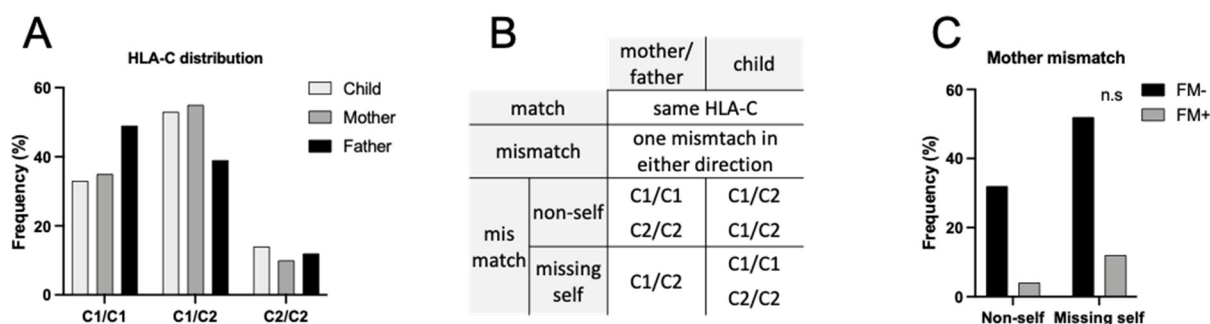
**Table S2:** n values to figures 1 - 6.

Figure	Group	n		
1 A + B	female	23		
	male	28		
1 C + D	age	51		
2 A	FM <sup>+/-</sup>	child	mother	father
	FM <sup>-</sup>	32	32	13
	FM <sup>+</sup>	19	19	7
2 B	child	C1/C1	C1/C2	C2/C2
	mother			
	C1/C1	11	4	0
	C1/C2	10	15	7
	C2/C2	0	4	1
2 C	FM <sup>+/-</sup>	match	mismatch	
	FM <sup>-</sup> mother	11	21	
	FM <sup>+</sup> mother	14	4	
	FM <sup>-</sup> father	7	6	
	FM <sup>+</sup> father	4	3	
2 D	match	26		
	mismatch	25		
2 E + F	C1/C1 match	11		
	C1/C2 match	15		
	non-self	9		
	missing self	16		
3 A	FM <sup>+/-</sup>	2DL3+	2DL3-	
	FM <sup>-</sup>	31	2	
3 B	FM <sup>+</sup>	16	6	
	all	51		
3 C	2DL3 only	8		
	2DL2 and 2DL3	29		
	2DL2 only	14		
3 D + E	FM <sup>+/-</sup>	child	mother	
	FM <sup>-</sup>	32	32	
	FM <sup>+</sup>	19	19	
3 F	KIR gene motif	child	mother	
	A/A cen	13	16	
	A/B cen	29	31	
	B/B cen	9	4	

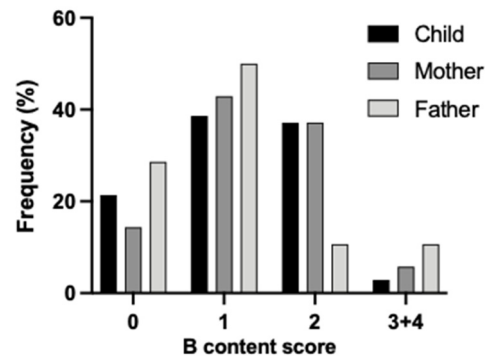
Figure	Group	n			
3 F	A/A tel	25	27		
	A/B tel	24	23		
	B/B tel	2	1		
4	KIR	FM <sup>-</sup>	FM <sup>+</sup>	FM <sup>-</sup> act	FM <sup>+</sup> act
	all act	-	-	42	10
	2DL1/S1	42	10	42	10
	2DL1/S5	30	8	30	8
	2DL1	39	9	39	9
	2DL2/3/S2(DX27)	36	10	36	10
	2DL3	42	10	42	10
	2DL2/3/S2(CH-L)	18	3	18	3
	2DL5	31	9	31	9
	2DS4	41	10	41	10
	2DS3/S4	31	9	31	9
5 A	NKG2A	26	7	26	7
	FM <sup>-</sup>	23			
5 B	FM <sup>+</sup>	10			
	FM <sup>-</sup>	23			
5 B	FM <sup>+</sup>	13			
	father	16			
6 A	match	mother		father	
	mismatch	25		12	
6 B		20		9	
	HLA-C1/C1	19		15	
	HLA-C1/C2	16		15	
6 C	HLA-C2/C2	4		4	
	2DL2+	4			
	2DL2/L3+	19			
6 D	2DL3+	12			
	cen A/A	16			
	cen A/B	23			
	cen B/B	5			
	tel A/A	23			
	tel A/B	21			
	tel B/B	0			

**Table S3:** n-values to supplementary figure 1 -4.

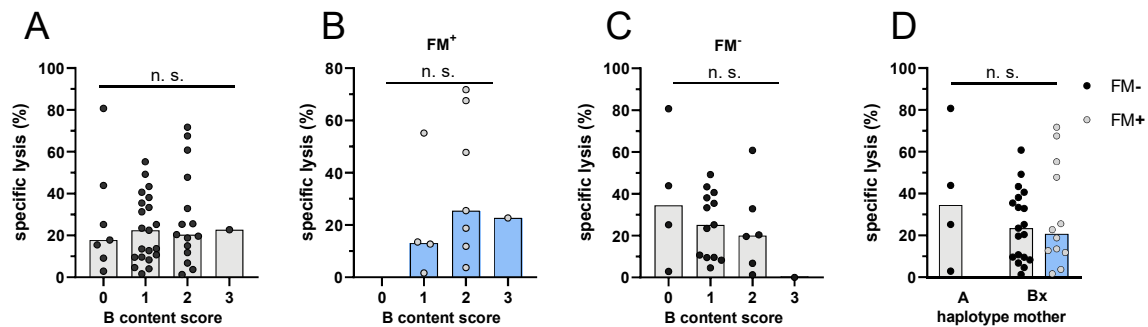
Figure	Group	N		
S1 A	HLA-C	Child	Mother	Father
	C1/C1	23	24	16
	C1/C2	37	38	13
	C2/C2	10	7	4
S1 C	FM+/-	non-self	missing-self	
	FM-	8	13	
	FM+	1	3	
S2	B content score	Child	Mother	Father
	0	15	10	8
	1	27	30	14
	2	26	26	3
	3+4	2	4	3
S3 A	0	7		
	1	21		
	2	15		
	3+4	1		
S3 B	0	0		
	1	4		
	2	7		
	3+4	1		
S3 C	0	4		
	1	13		
	2	6		
	3+4	1		
S3 D	Haplotype	FM-	FM+	
	A	4	0	
	Bx	19	12	



**Figure S1:** Child, mother and father were HLA-C genotyped and grouped according to their HLA-C genotype into HLA-C1/C1 (C1), HLA-C1/C2 (C1/2) or HLA-C2/C2 (C2). (A) HLA-C distribution of the analyzed cohort. (B) Criteria for grouping depending on HLA match or mismatch of mother and child. (C) Mismatched mothers were grouped into mismatch non-self or missing self. Statistical analysis of frequencies was performed using Fisher's exact test for two groups or Chi2 for more than two. Statistical significance  $p < 0.05$ .



**Figure S2:** B content score frequencies in the analyzed cohort. B content score was determined after KIR genotyping in child, mother and father. Subsequently, they were grouped depending on their B content score.



**Figure S3:** Parental cytotoxicity against filial leukemic blasts. Parental NK cells were isolated and IL-2/IL-15 pre-activated overnight. The next day their cytotoxicity was analyzed against filial blasts and the control cell line K562 in a FACS-based killing assay. (A) All mothers were grouped depending on their B content score and the lysis of leukemic blasts was compared between the groups. Mothers were further grouped into (B) FM+ and (C) FM- and the specific lysis was compared between the groups. (D) Mothers were grouped depending on their KIR haplotype and specific lysis of leukemic blasts was compared. Each dot resembles one sample, bars show mean of the group. Statistical analysis was performed using Kruskal-Wallis test followed by Dunn's multiple comparison test for more than two groups. Significance  $p < 0.05$ , n. s. not significant.