

Table S1: Cancer predisposition syndromes (CPSs) detected in 193 patients according to age at time of neoplasm diagnosis. BWS: Beckwith-Wiedemann Syndrome, CMMRD: Constitutional mismatch repair deficiency Syndrome, FA: Fanconi anemia, feat.: features, HSCR: Hirschsprung disease, HLH: Hemophagocytic lymphohistiocytosis, IHH: Isolated hemihypertrophy, LFS: Li-Fraumeni Syndrome, MEN: Multiple endocrine neoplasia Syndrome, mut.: gene mutation, NBS: Nijmegen breakage Syndrome, NF1: Neurofibromatosis type I, RTPS: Rhabdoid tumor predisposition Syndrome, S.: Syndrome, VHL: Hippel-Lindau Syndrome

Age at time of cancer diagnosis [m: month(s) or y: year(s)]	No. of cases	Cancer predisposition syndrome (CPS)																				
		BWS	BRCA1 mut.	CMMRD	DICER1 mut.	Down S.	Edwards' S.	FA + BRCA2 mut.	Feat. of dysmorphia	HSCR	HLH	IHH	IMAGE S.	LFS	LFS + NBS	MEN S.	NBS	NF1	Noonan S.	RTPS	VHL	WAGR S.
0 to < 1 y	49	5			2	3	1		1			6		5				23		2	1	
0 to < 1 m	11	1				2												7		1		
1 to < 2 m	6					1								1				4				
2 to < 3 m	3						1							1						1		
3 to < 4 m	3																	3				
4 to < 5 m	1								1													
5 to < 6 m	5	2										1		1				1				
6 to < 7 m	4	1										2		1								
7 to < 8 m	3				2													1				
8 to < 9 m	3																	2			1	
9 to < 10 m	5	1										3						1				
10 to < 11 m	2													1				1				
11 to < 12 m	3																	3				
1 to < 2 y	12					3		1			2			4				2				
2 to < 3 y	25	3	1			4				1		1		4				10				1
3 to < 4 y	25	2		2		3					2	4		6			1	5				
4 to < 5 y	14					3						4	1	2				4				
5 to < 6 y	8											2		1				5				
6 to < 7 y	4													1				2	1			
7 to < 8 y	5											1		1		1		2				
8 to < 9 y	2																	2				
9 to < 10 y	7					1								1		1		4				
10 to < 11 y	9								1					2				6				
11 to < 12 y	5					1								2				2				
12 to < 13 y	7					1									1	1		4				
13 to < 14 y	3																	3				
14 to < 15 y	2										1							1				
15 to < 16 y	3															1		2				
16 to < 17 y	7													3			1	3				
17 to < 18 y	6								1					2				3				
No. in total	193	10	1	2	2	19	1	1	3	1	5	18	1	34	1	4	2	83	1	2	1	1

Table S2: Cancer predisposition syndromes (CPSs) detected in 193 patients according to diagnosed neoplasm defined by ICD-10 classification. In two cases two genetic disorders were detected (FA with BRCA2 gene mutation, LFS with NBS), in three cases multiplied cancers were diagnosed (C49 with C64 in one case, C49 with C92 in two cases). BWS: Beckwith-Wiedemann Syndrome, CMMRD: Constitutional mismatch repair deficiency Syndrome, FA: Fanconi anemia, HSCR: Hirschsprung disease, HLH: Hemophagocytic lymphohistiocytosis, IHH: Isolated hemihypertrophy, LFS: Li-Fraumeni Syndrome, MEN: Multiple endocrine neoplasia Syndrome, NBS: Nijmegen breakage Syndrome, NF1: Neurofibromatosis type I, RTPS: Rhabdoid tumor predisposition Syndrome, VHL: Hippel-Lindau Syndrome

Cancer predisposition syndrome (CPS)	No. of cases	Neoplasms diagnosed in CPS-positive patients defined by ICD-10 classification																							
		C22	C30	C37	C38	C40	C41	C47	C48	C49	C49 C64	C49 C92	C56	C64	C71	C72	C73	C74	C75	C80	C81	C83	C85	C91	C92
BWS	10	2								1				7											
BRCA1 gene mutation	1																							1	
CMMRD	2																							2	
DICER1 gene mutation	2									1				1											
Down Syndrome	19																						11	8	
Edwards' Syndrome	1	1																							
FA + BRCA2 mutation	1										1														
Features of dysmorphia	3									1															2
HSCR	1							1																	
HLH	5																					1	4		
IHH	18													18											
IMAGE Syndrome	1												1												
LFS	34	5	1	1		1		7				2		13						1	1			2	
LFS + NBS	1											1													
MEN	4					1											3								
NBS	2																		1					1	
NF1	83	2			1			39	1	5				31	1	1	1	1							
Noonan Syndrome	1													1											
RTPS	2						1			1															
VHL	1													1											
WAGR Syndrome	1												1												
No. in total	193	10	1	1	1	2	1	47	1	9	1	2	1	28	46	1	1	4	1	1	1	1	1	21	10