

Repping *et al.*, Polymorphism for a 1.6-Mb deletion of the human Y chromosome persists through balance between recurrent mutation and haploid selection

Supplementary Table 2. Initial screen, prevalence of the gr/gr deletion by Y haplotype. The gr/gr deletion was found in five Y haplotypes present in both groups of men (spermatogenic failure and unknown spermatogenic phenotype). These five haplotypes are shaded. In all five haplotypes the frequency of the gr/gr deletion allele is higher among men with spermatogenic failure.

Haplotype ^a	Spermatogenic failure		Unknown spermatogenic phenotype	
	# men tested	% with gr/gr deletion	# men tested	% with gr/gr deletion
A	0	-	8	0
B	0	-	3	0
DE*(xD2b)	49	4.1	14	0
D2b	0	-	2	100
C	2	0	1	0
F(xJ,xK)	82	4.9	36	0
J	62	6.5	17	5.9
K(xP)	21	4.8	37	2.7
P(xR1a)	218 ^b	1.8	83 ^c	0
R1a	37	0	14	0
Total	<u>471^b</u>		<u>215^c</u>	

^aSee **Supplementary Table 7**.

^bTwo of the original 473 men screened for the gr/gr deletion were excluded (WHT3107 for a 47,XY Y karyotype; WHT3173 for a b2/b4 duplication, see **Figure 3** main text and **Supplementary Table 1**).

^cOne of the 216 men screened for the gr/gr deletion was excluded for a b2/b4 duplication (PD0267; **Supplementary Table 1**).