Suppl. Table S4: Correlation of TMB with molecular subtypes and genomic alterations

Parameter	Category (n)	Median TMB	P-value §
			(Wilcoxon test)
Molecular subtype (AIMS) †	Basal-like (82)	1.49	
	HER2-enriched (50)	1.60	0.299
	other (4)	1.25	0.512
ATM	mut (6)	1.44	
	wt (143)	1.52	0.931
ARID1A	mut (5)	2.46	
	wt (144)	1.47	0.040
BRCA1	mut (20)	1.85	
	wt (129)	1.43	0.091
BRCA2	mut (9)	2.70	
	wt (140)	1.43	0.004
CCNE1	amp (11)	1.30	
	wt (138)	1.53	0.569
NOTCH1	mut (5)	2.50	
	wt (144)	1.50	0.063
MYC	amp (39)	1.43	
	wt (110)	1.56	0.269
PIK3CA	mut/amp (17)	1.39	
	wt (132)	1.56	0.900
PTEN	mut/del (17)	1.87	
	wt (132)	1.47	0.099
TP53	mut (103)	1.65	
	wt (46)	1.14	0.011
HRD-panel #	mut/del (45)	1.89	
_	wt (104)	1.37	< 0.001

<sup>§</sup> P-values not corrected for multiple testing

<sup>&</sup>lt;sup>†</sup> Molecular subtype from RNA-seq was available only for 136 of the 149 samples with WES (see Supplementary Figure S1).

<sup>\*\*</sup> The predefined HRD-panel encompassed: BRCA1, BRCA2, ATM, PALB2, BARD1, DRIP1, RAD51B, RAD51C, RAD51D, FAAP20, CHECK2, FAN1, FANCE, FANCM, POLQ, NBN