

Thin and thick primary cutaneous melanomas reveal distinct patterns of somatic copy number alterations

Supplementary Material

SCNA > 0.45 Chromosome Arm

M2

Chr	% Alteration/Arm	Gain/Loss
6p	47%	Gain
6q	86%	Loss
7p	53%	Gain
9p	69%	Loss
11p	87%	Gain
18q	55%	Gain
8q	complete (98%)	Gain

M11

Chr	% Alteration/Arm	Gain/Loss
6p	47%	High Gain
11p	63%	Gain

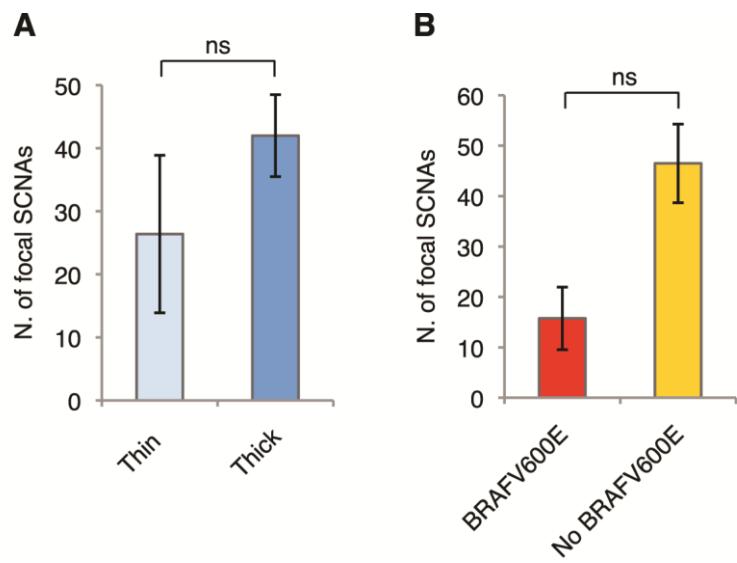
M14

Chr	% Alteration/Arm	Gain/Loss
4q	55%	Gain
6p	70%	High Gain
7p	61%	Gain
8p	48%	Gain
8p+q	p half (50%)	Gain
	q complete (99%)	Gain
9p	49%	Loss
10q	75%	Loss
10p	p complete (99%)	Loss
11p	76%	Gain
13q	q complete (96%)	Gain
14q	45%	Loss
18p+q	p complete (99%)	Gain
	q complete (96%)	Gain
20p	70%	Gain

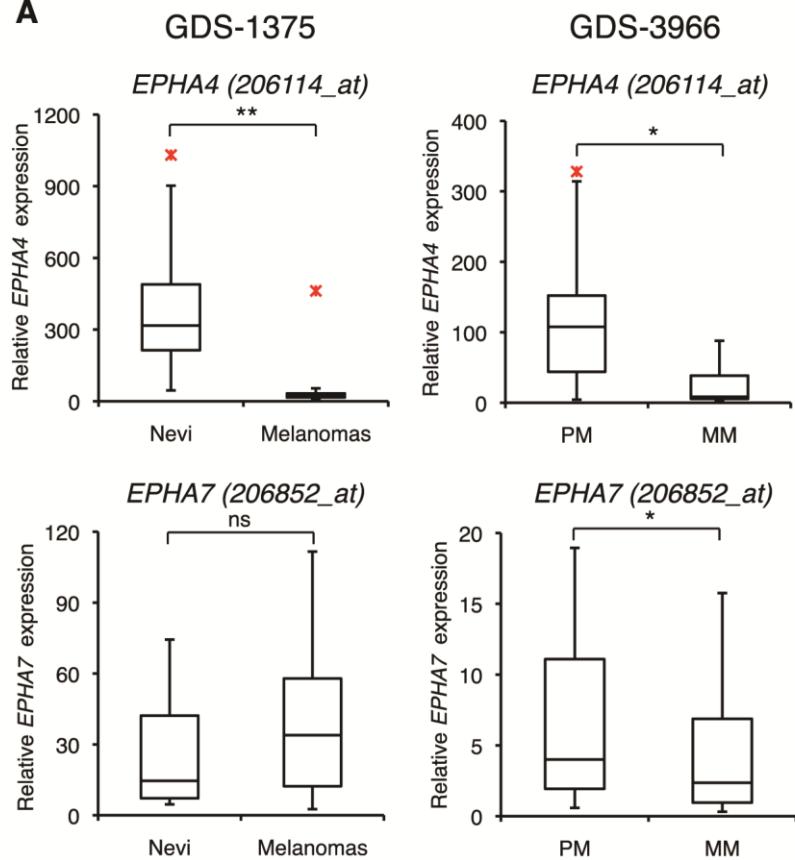
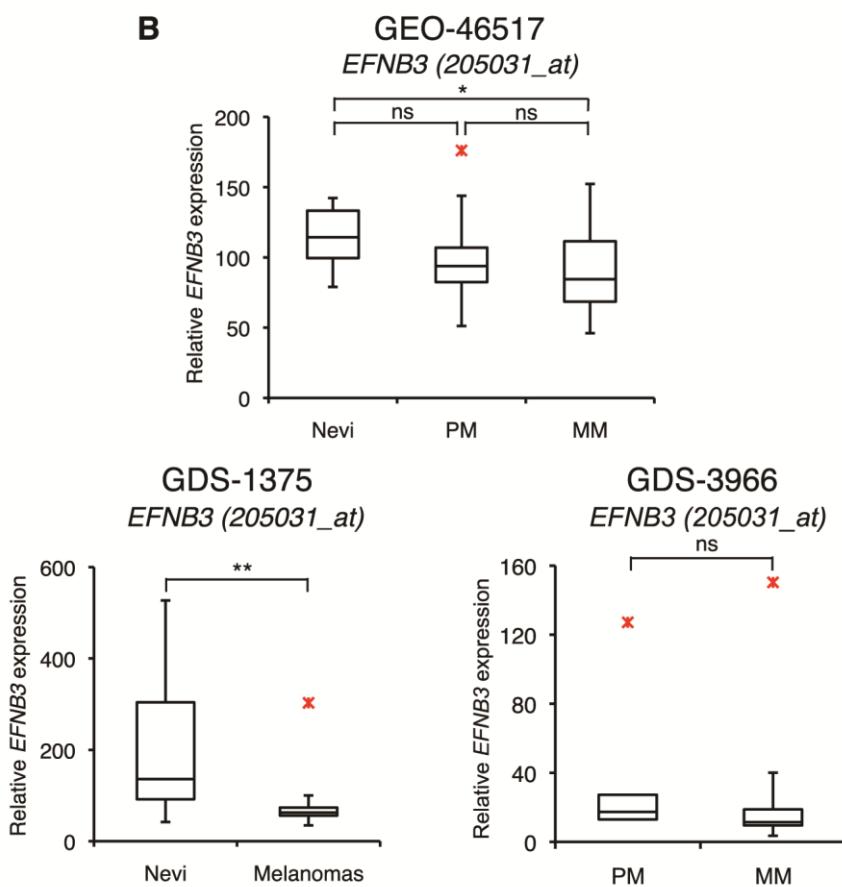
Common aberrations

Chr	% Alteration/Arm	Gain/Loss	Samples
6p	47%	Gain	M2, M11 M9, M14
6q	at least 85%	Loss	M2, M9
7p	at least 53%	Gain	M2, M14
8p	at least 48%	Gain	M9, M11
9p	at least 50%	Loss	M2, M14
11p	at least 76%	Gain	M2, M11, M14
18q	at least 50%	Gain	M2, M11

Supplementary Figure 1. Chromosome aberrations in four thick melanomas.

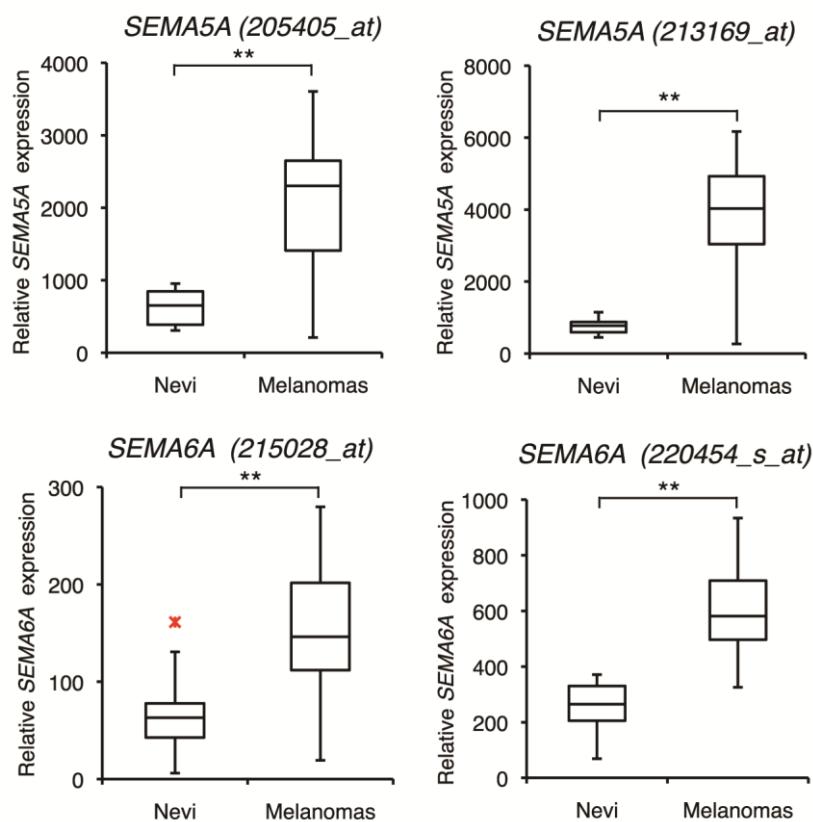


Supplementary Figure 2. (A) Number of focal SCNAs in thin (n=5) and thick melanomas (n=5) ($p=0.27$, Mann-Whitney U test). (B) Number of focal SCNAs in melanomas with BRAFV600E (n=4) and with no BRAFV600E mutation (n=6) ($p=0.08$, Mann-Whitney U test).

A**B**

Supplementary Figure 3. (A) Expression of *EPHA4* and *EPHA7* mRNA determined by different publicly available microarray data sets (GDS-1375 and GDS-3966). (B) Expression of *EFNB3* mRNA determined by different publicly available microarray data sets (GEO-46517, GDS-1375 and GDS-3966). In GEO-46517 nevi (n=9), primary (n=31) and metastatic (n=73) melanomas. In GDS-1375 nevi (n=17) and melanomas (n=45). In GDS-3966 primary melanomas (n=31) and metastatic melanomas (n=52). PM, primary melanomas; MM, metastatic melanomas. Red crosses indicate outlier values. *, p<0.05; **, p<0.001.

GDS-1375



Supplementary Figure 4. Expression of *SEMA5A* and *SEMA6A* mRNA in nevi (n=17) and melanomas (n=45) determined by a publicly available microarray data set (GDS-1375). Red crosses indicate outliner values. **, p<0.001.

Table S1:	Summary of genomic data
Table S2:	Summary of somatic mutations
Table S3:	Non-synonymous and stopgain SNVs
Table S4:	Small indels
Table S5:	Splice variants
Table S6:	SCNA gain
Table S7:	SCNA loss
Table S8:	Mutations found in TCGA and Cosmic databases