Table SI. Information about the procedures and optimized analytical variables for each antibody used in the study and histoscore

Antigen	Staining pattern	Staining evaluation	Clone	Pretreatment; detection platform ^a	Dilution	Source
Ki67 ^b	Nuclear	Nuclear	MIB-1	PT Link Low 95°C 20 min; EnvisionFlex	RTU	Dako (Glostrup, Denmark)
Cyclin D1 ^b	Nuclear and cytoplasmic	Nuclear	EP12	PT Link High 95°C 20 min; EnvisionFlex	RTU	Dako (Glostrup, Denmark)
p16	Nuclear and cytoplasmic	Nuclear and cytoplasmic	E6H4	PT Link Low 95°C 20 min; EnvisionFlex	RTU	Ventana (Tucson, USA)
p53	Nuclear	Nuclear	DO-7	PT Link High 95°C 20 min; EnvisionFlex	RTU	Dako (Glostrup, Denmark)
pRB	Nuclear	Nuclear	1F8	PT Link High 95°C 20 min; EnvisionFlex	1:100	Abcam (Cambridge, UK)
hTERT	Nuclear and cytoplasmic	Nuclear and cytoplasmic	2C4	PT Link Low 95°C 20 min; EnvisionFlex	1:200	Abcam (Cambridge, UK)
PTEN	Nuclear and cytoplasmic	Cytoplasmic	6H2.1	PT Link High 95°C 20 min; EnvisionFlex	1:100	Dako (Glostrup, Denmark)
BRAF/V600E	Cytoplasmic	Cytoplasmic	VE1	Ventana conditions	RTU	Ventana (Tucson, USA)
Cav 3.1	Membranous and cytoplasmic	Membranous and cytoplasmic	Polyclonal	PT Link Low 95°C 20 min; EnvisionFlex	1:150	Antibody BCN (Barcelona, Spain)
Cav 3.2	Membranous, cytoplasmic and nuclear occasionally	Membranous and cytoplasmic	Polyclonal	PT Link High 95°C 20 min; EnvisionFlex	1: 200	Antibody BCN (Barcelona, Spain)

^aEpitope retrieval in the PreTreatment Module, PT-LINK (Dako, Glostrup, Denmark). EnVision FLEX Detection Kit (Dako) using diaminobenzidine chromogen as the substrate. BRAF V600E detection was performed using the Ventana platform (Tucson, AZ, USA). ^bProliferation and cell-cycle markers (Ki67 and cyclin D1) were not evaluated in lentiginous junctural naevi, without clear naevi nest, due to the difficulty of discerning between proliferating naevus cells and proliferating keratinocytes of the epidermal basal layer.

the epidermal basal layer.

Immunohistochemistry staining was graded semiquantitatively by considering the percentage and intensity of the staining. A histological score (*Hsc*) was obtained from each sample and values ranged from 0 (no immunoreactivity) to 300 (maximum immunoreactivity). The score was obtained by applying the following formula, *Hsc*=[1 x (% light staining)] + [2 x (% moderate staining)] + [3 x (% strong staining)].

PTEN: phosphatase and tensin homologue; RTU: ready to use. hTERT: telomerase; cycl D1: cyclin D1.