

Supplementary material and methods

Epigenetic changes in the early stage of silica-induced cell transformation

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Table S1: Physical characteristics of the silica samples.

Sample ^a	Origin	Main phase	Primary particle size ^b	Main impurities (wt %) ^c	BET SSA (m ² /g) ^d	DLS ^e	
						Z-average diameter in water (nm)	Polydispersity index
NM-203	Pyrogenic	Amorphous	24.7 ± 17.7 nm	Al (0.43)	204	318.7 ± 4.9	0.420 ± 0.026
Min-U-Sil 5	Natural; US Silica Co. (Berkeley)	Crystalline	1.33 ± 0.85 µm	Al (0.16) ^d	5	1385.7 ± 377.0	0.380 ± 0.130

^a In-depth physical and chemical characterizations of the samples (except DLS data) are available in Rasmussen *et al*, 2013 (NM-203) and Elias *et al*, 2006 (Min-U-Sil);

^b Determined by transmission electron microscopy;

^c Impurities were determined by energy dispersive spectrometry for NM-203 and by inductively coupled plasma spectrometry for Min-U-Sil;

^d The BET (Brunauer-Emmet-Teller) specific surface area (SSA) was determined by nitrogen adsorption;

^e Dynamic light scattering (DLS) was performed using a Nanozetasizer (Malvern). Samples were dispersed in water as described in the Materials and methods section. Values represent the mean ± standard deviation of three DLS analyses.

Table S2: Antibodies used for Western Blot and ChIP analyses.

Target	Supplier	Reference	Molecular weight (kDa)
DNMT1	Cell Signaling	5119	200
DNMT3a	Cell Signaling	2160	130
DNMT3b	Abcam	ab2851	100
HDAC1	Cell Signaling	2062	63
HDAC2	Cell Signaling	2540	60
HDAC3	Cell Signaling	2632	49
HDAC6	Cell Signaling	7612	140
Total histone H3	Millipore	06-755	17
Acetylated histone H3	Millipore	06-599	17
Total histone H4	Millipore	05-858	10
Acetylated histone H4	Millipore	06-866	10
β-actin	Cell Signaling	4970	42
c-Myc	Cell Signaling	13987	57
RNA polymerase II	Abcam	ab210527	
Trimethylated histone H3 lysine 4	Abcam	ab8580	
Acetylated histone H3 lysine 4	Abcam	ab113672	
Acetylated histone H3 lysine 9	Millipore	06-942	
Acetylated histone H3 lysine 27	Abcam	ab4729	
5-methylcytosine	Diagenode	C15200081	

Table S3: Primers used for QRT-PCR and ChIP analyses.

Target gene	Forward and reverse sequences	Amplicon size
<i>β-actin</i>	F: GGCCAACCGTGAAAAGATGA R: TACGACCAGAGGCATACAGG	104
<i>α-tubulin</i>	F: CACTACACCATTGGCAAGGA R: GCTGTGGAAAACCAAGAAGC	102
<i>Gapdh</i>	F: CTGCACCACCAACTGCTTAG R: GTCTTCTGGTGGCAGTGAT	108
<i>Tbp</i>	F: ACGGACAAC TGCGTTGATT TT R: ACTTAGCTGGGAAGGCCAAC	128
<i>c-myc</i>	F : CCCTAGTGCTGCATGAGGA R : TGCCTCTTCTCCACAGACAC	103
<i>CollA2</i>	F: TTGCAATCGGGATCAGTACG R : TGGGACCATCAACACCATCT	103
P1	F: CGCCCTTATATTCCGGGG R: GAGCTGAGTGAGGCGAGTC	96
P2	F: AGGGATCCTGAGTCGCAGT R: CGCTCACTCCCTGTCTCT	92
e2	F: GAGGAGACATGGTGAACCAGA R: AACCGCTCCACATACAGTCC	93
neg	F: CAGTCTCACCCGACCATCT R: CTGACAGAGCTTCCCCAAA	92