SUPPLEMENTARY INFORMATION

Transient Nuclear Lamin A/C Accretion Aids in Recovery from Vapor Nanobubbleinduced Permeabilization of the Plasma Membrane

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Supplementary Table

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Comparison	Common DEGs	
Heating 6h vs VNB 6h	DDX39B, PHLDA2, IGFN1, CARMN, RNY4, UBC, RNY1,BIRC3, LMNA, TRMT9B, TUBA1C	
Heating 24h vs VNB 24h	PNISR, HSPA1A, DDX58, TAGLN, INCENP, TCEA3, SS18L2, LINC00504, FAM228B, MALAT1, CSPP1, RNY1, NDUFS7, PGM5P2, CDC20, CYP3A5, MOB3B, SREK1, PTMS, DZIP1L, GP3135, ZBED6, PITPNM3, LINC01873, ACTA2, LUC7L3, PLEKHN1, TAF3,KMT2B, PRPF38B, MAB21L3, ZNF236, TCIM,	
Heating 48h vs VNB 48h	TAGLN, MYL9	

b

GO biological process (6h)	VNB	Heating
(Striated) muscle cell development	LMNA , PPARA, DDX39B, MYLK3	CSRP2, LMNA , NFATC2, NFATC4, DDX39B, ALPK2
MyD88-independent TLR signaling	BIRC3, UBC, IKBKG	BIRC3, PRKCE, UBC
Response to wounding	CD151, DHFR, HSPB1, LGALS1, PLEC, PPARA, THBS1, TIMP1, C6orf89	EGFR, ETS1, CCN1, DX1, SERPINE1, PRKCE, SDC4, PROCR,TNFRSF12A

Table S1. (a) Significant DEGs that are shared between VNB vs CTR and heating vs CTR are listed per timepoint (6h, 24h and 48h); **(b)** Enrichment analysis in the list of significant DEGs for VNB vs CTR and heating vs CTR pointed to three GO biological processes that are shared 6h after treatment (listed in the left column). The significant DEGs that correspond to the commonly enriched biological processes are listed for VNB vs CTR (middle) and heating vs CTR (right).

Supplementary Figures



Figure S1. (a) FD10 transfection efficiency and relative mean fluorescence intensity for increasing AuNP concentrations as determined by flow cytometry 2 h after VNB treament; **(b)** Cell viability for increasing AuNP concentrations as determined with the metabolic assay Cell Titer Glo 2h or 24h after VNB.



Figure S2. Enriched biological processes from the Gene Ontology (GO) database determined in the list of significant DEGs determined via Metascape for (a) VNB vs CTR at 6h, (b) Heating vs CTR at 6h, (c) VNB vs CTR at 24h, (d) Heating vs CTR at 24h and (e) Heating vs CTR at

48h. More specifically, representatives are shown for clusters of redundant GO biological processes. For each cluster representative the percentage of significant DEGs present in the input list to the total number of genes in that process is plotted as well as the significance (-log corrected p-value) of the enrichment.

Mean chromatin condensation parameter



Figure S3. The chromatin condensation parameter for two different timepoints (6h & 24h) after VNB treatment (VNB) and for the corresponding untreated cells (CTR) in LMNA-WT cells (left) and LMNA-KO cells (right) (***P < 0.001).



Figure S4: (a) The mean nuclear area in LMNA-WT cells at two different timepoints (6h & 24h) after VNB treatment (VNB) and for the corresponding untreated cells (CTR) (*P < 0.05); **(b)** The mean cell area in LMNA-WT cells at two different timepoints (6h & 24h) after VNB treatment (VNB) and for the corresponding untreated cells (CTR) (**P < 0.001)