

Supplementary information

Supplementary figure S1. a) size distributions and b) Zeta-potentials of PDA/PEI NPs, PDA/PEI NPs/pGL-3, and PDA/PEI NPs/ NC complexes. The mass ratio of PDA/PEI NPs and nucleic acids in those complexes was 10. Data were presented as mean \pm standard deviation, student t-test, * indicates p<0.05, ns denotes non-significance (p>0.05). c) Gel separation of PDA/PEI NPs-nucleic acid complexes. The ratios of PDA/PEI NPs-to-nucleic acid were a: 0; b: 0.1; c: 0.2; d: 0.5; e: 1; f: 2; g: 5 and h: 8.



Supplementary figure S2. Cell viability of a) corneal endothelial cells and b) epithelial cells after treated with PDA/PEI NPs/NC complexes at various concentrations. Data were presented as mean \pm standard deviation, one-way ANOVA, * indicates p<0.05.



Supplementary figure S3. Stability test of PDA/PEI NPs/nucleic acid complexes in various days after complexes preparation DLS.



Supplementary figure S4. Immunocytochemistry staining of f-actin skeleton. AAP cells were pretreated PDA/PEI NPs/miR-21-5p or PDA/PEI NPs/NC for 24 hours. Shown were representative images of 5 replicates.



Supplementary figure S5. Hematoxylin-eosin (HE) staining of eyes that injected with sterile PBS or PDA/PEI NPs/NC at 24 hours. The gross morphology could be observed without obvious inflammation infiltration. The representative images of three replicates were shown. Scale bar 500 μ m (upper panel), 100 μ m (lower panel).

Target genes	Representative transcript	Cumulative weighted context++ score	Previous TargetScan publication(s)
SMAD7	ENST00000262158.2	-0.46	2005, 2007, 2009, 2011
TIMP3	ENST00000266085.6	-0.46	2005, 2007, 2009, 2011
RHOB	ENST00000272233.4	-0.12	2005, 2007, 2009, 2011

Supplementary table S6. Predicted target genes of miR-21-5p from TargetScan. After searching for the publications of glaucoma pathological researches, RhoB, SMAD7, and TIMP3 were selected and presented from 384 target genes of miR-21-5p. The cumulative weighted context++ score, which is a scoring method of predicted efficacy of targeting of the sites (Agarwal et al., 2015), was listed for reference.



Supplementary figure S7. Dual-luciferase reporter assay verification of target genes of miR-21-5p. The relative luciferase activity was significantly decreased in the presence

of wild type (WT) 3' untranslated region, and then restored in mutant type (MT) constructs. (n=3, *** p<0.001, student t-test)



Supplementary figure S8. The proteins interaction network of the miR-21-5p targets. The network of protein-protein interaction for the three genes (RhoB, SMAD7, and TIMP3) protein products was constructed according to the STRING database (https://string-db.org/).



Supplementary figure S9. The expression of MMP9 in outflow tissue. The protein level of MMP9 was significantly increased in outflow tissue.