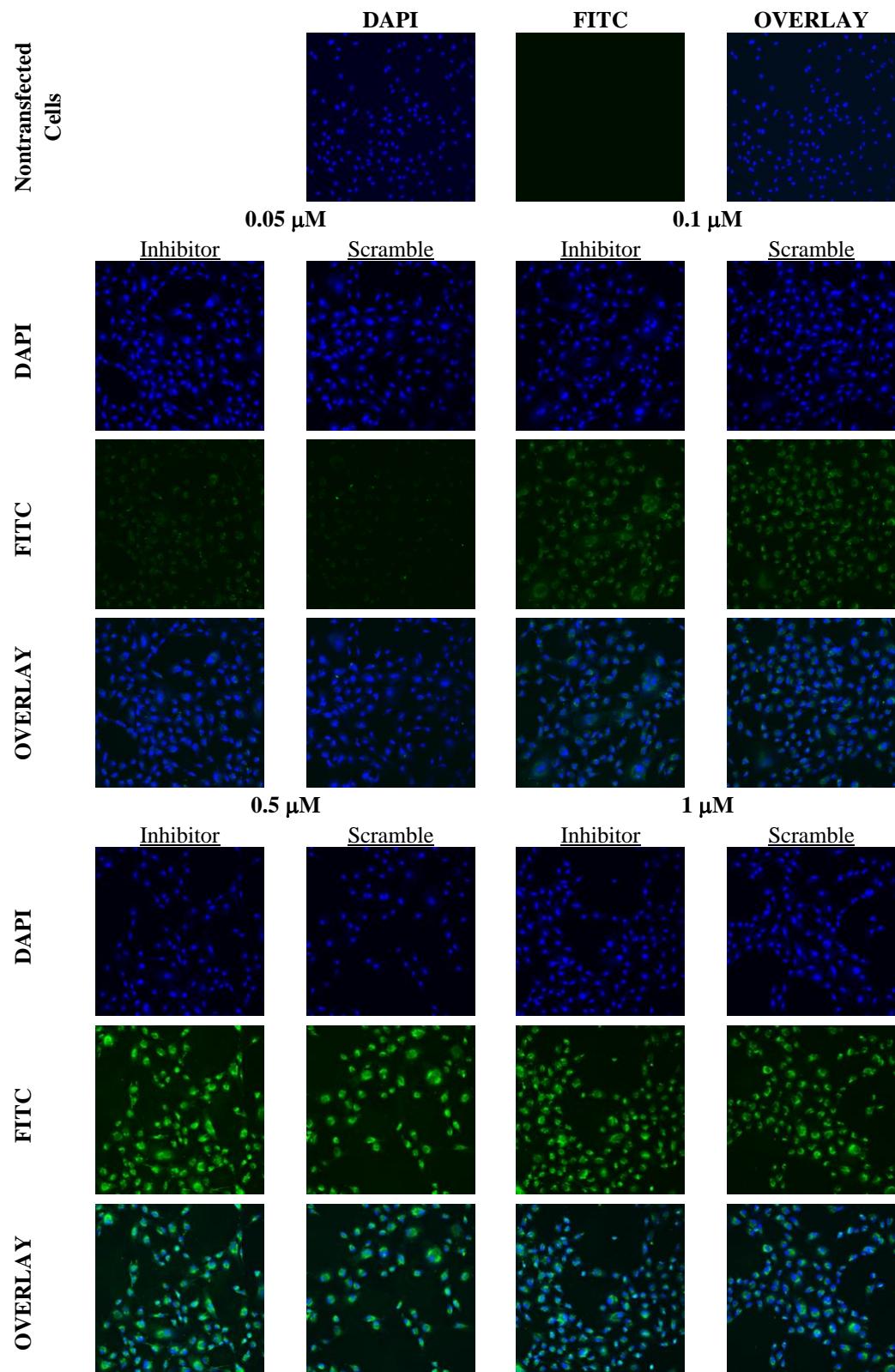
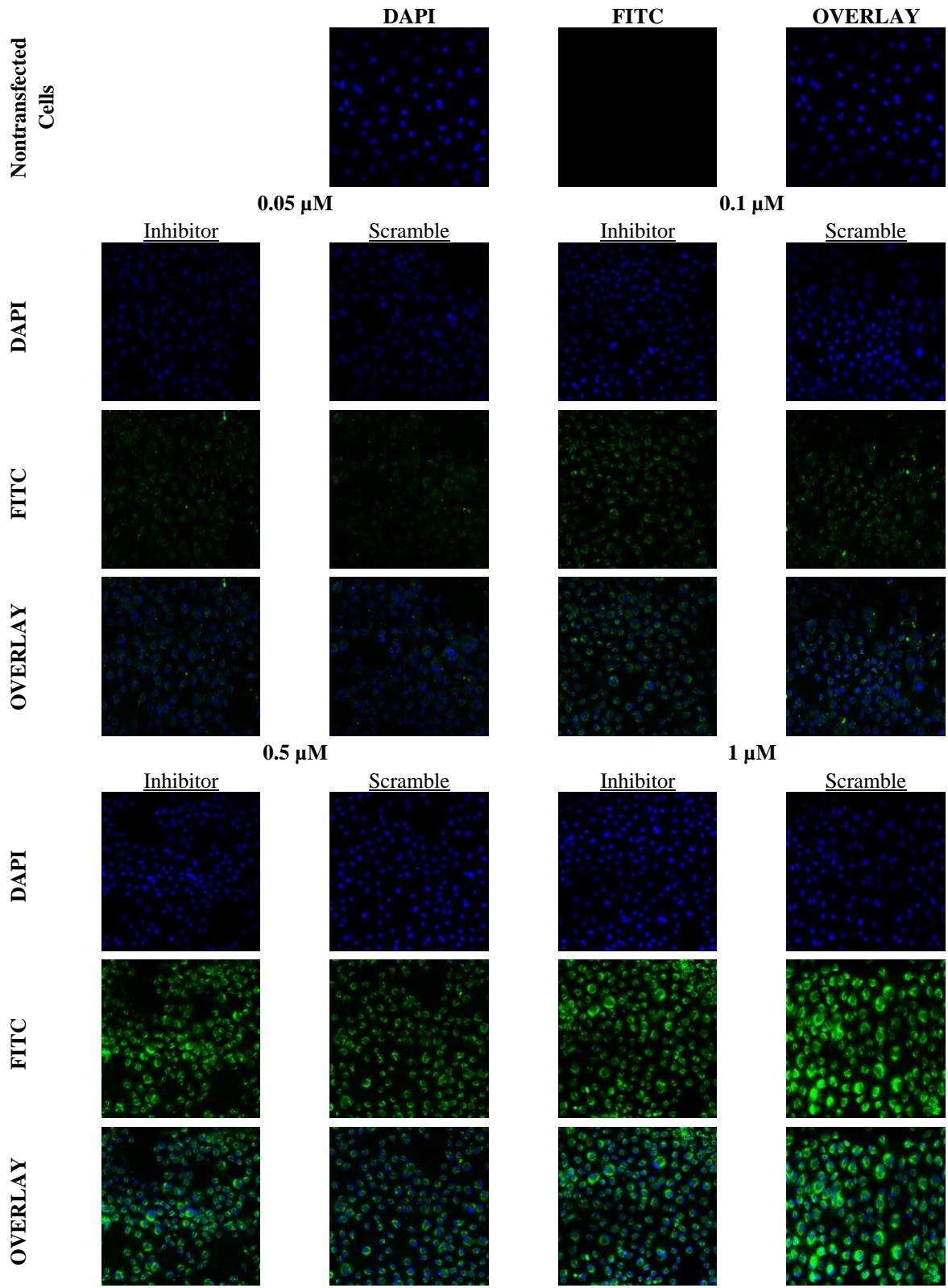


Supplementary Figure S1. mi-21 levels upon plating, throughout passaging, and after serum restriction.



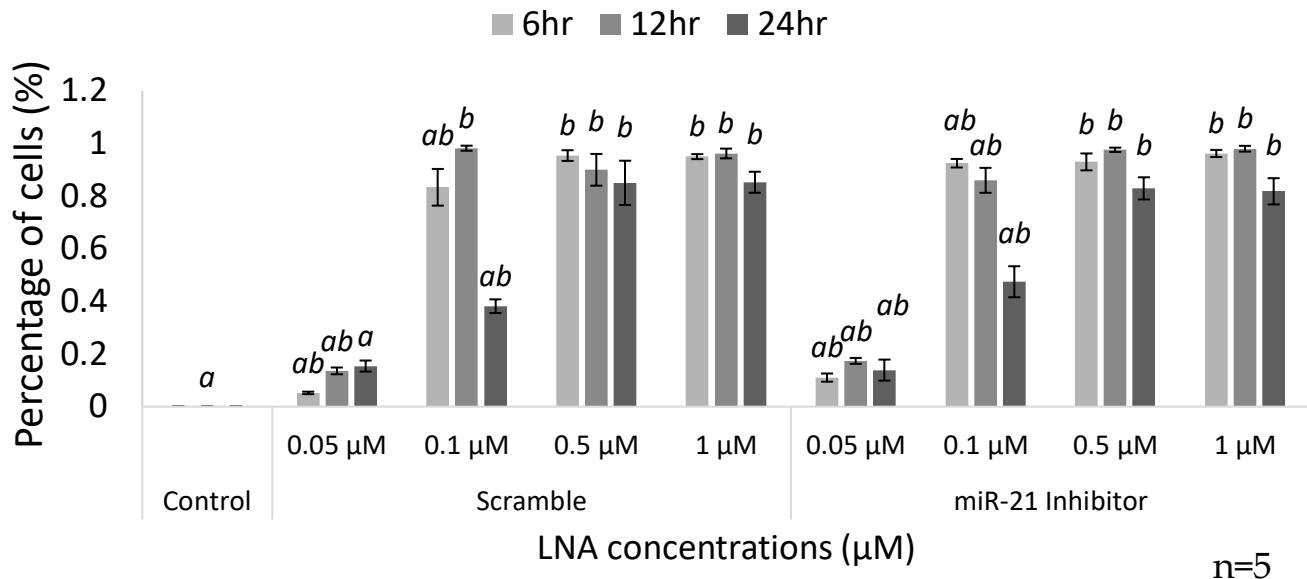
(A)



(B)

(C)

Number of Fluorescent Cells

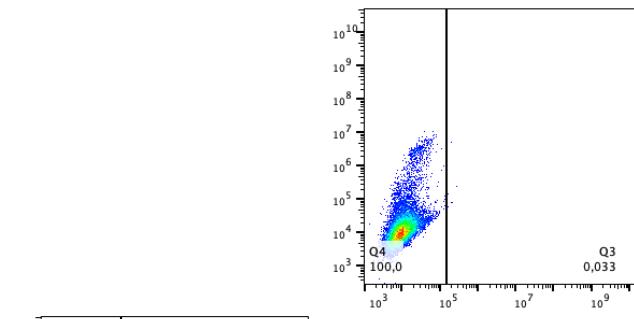


Supplementary Figure S2. Additional Confocal microscopy of LNAs at (A) 6 and (B) 24hrs with (C) number of fluorescent cells

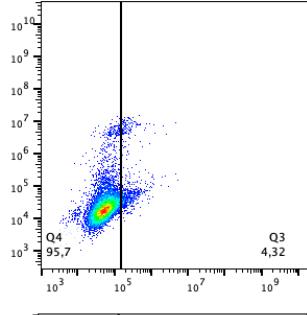
Nontransfected Control

0.05 μ M

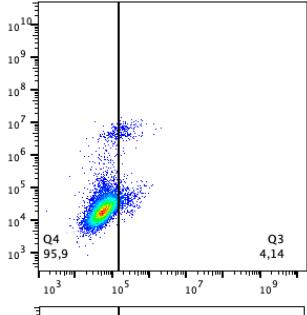
Inhibitor



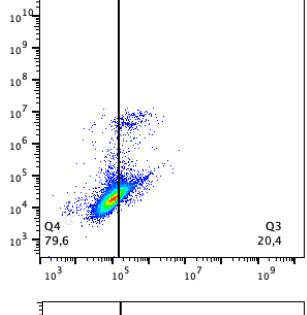
Scramble



Inhibitor

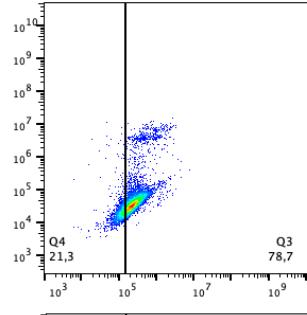


Scramble

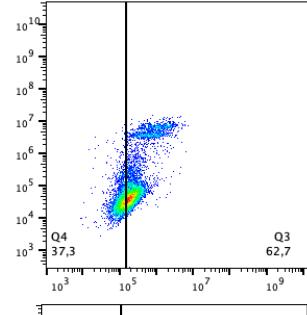


0.5 μ M

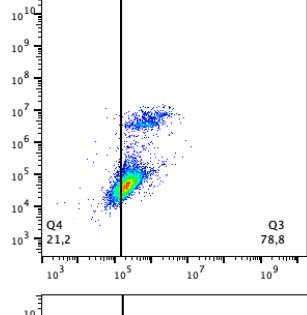
Inhibitor



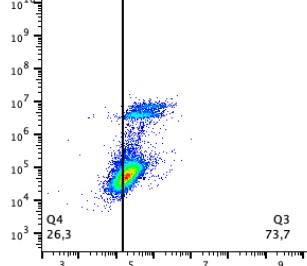
Scramble



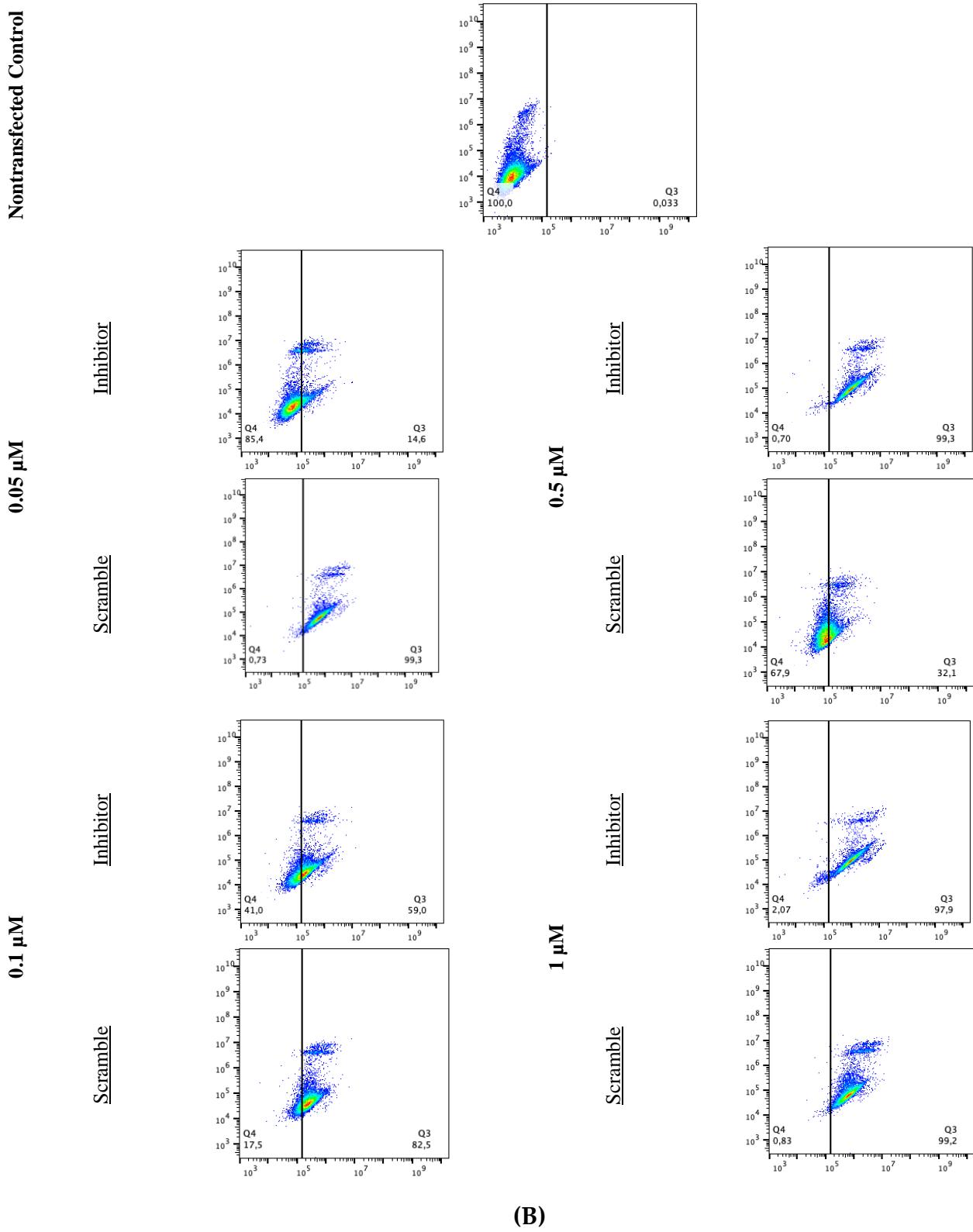
Inhibitor



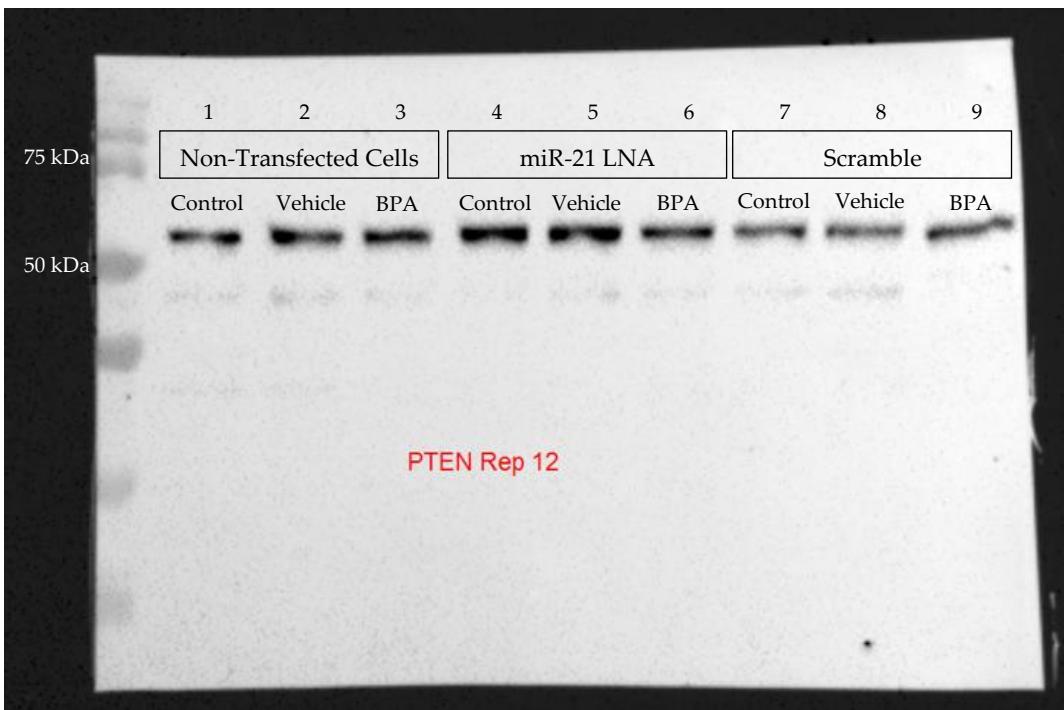
Scramble



(A)



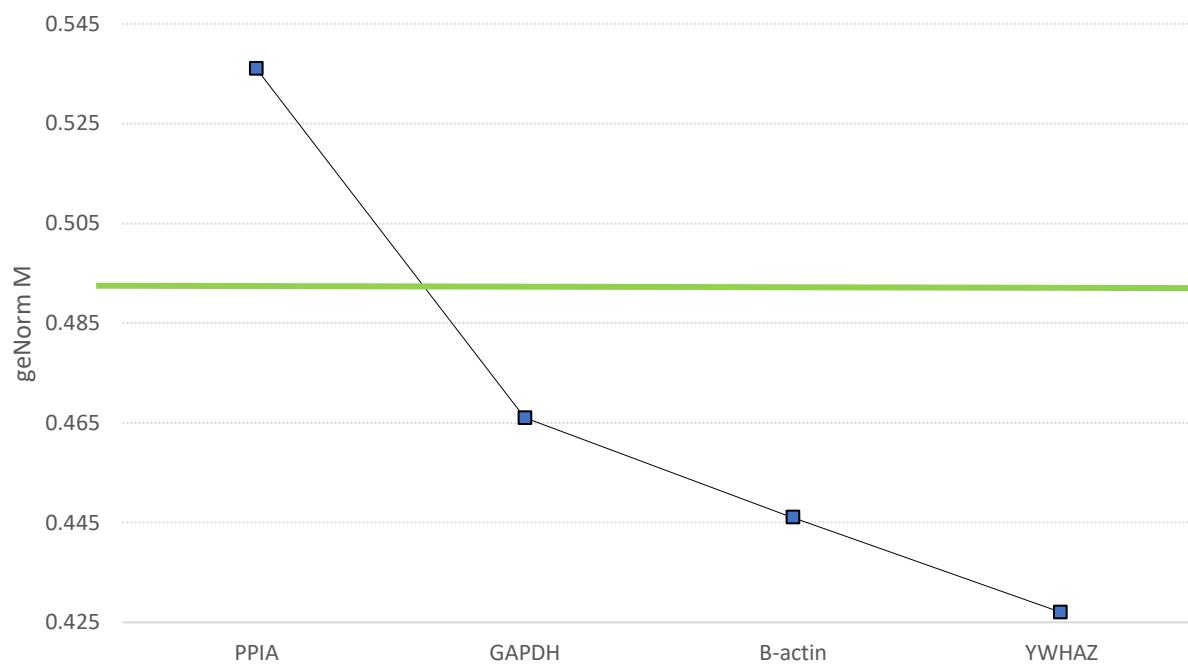
Supplementary Figure S3. Additional Flow Cytometry of LNAs at (A) 6 and (B) 24hrs.



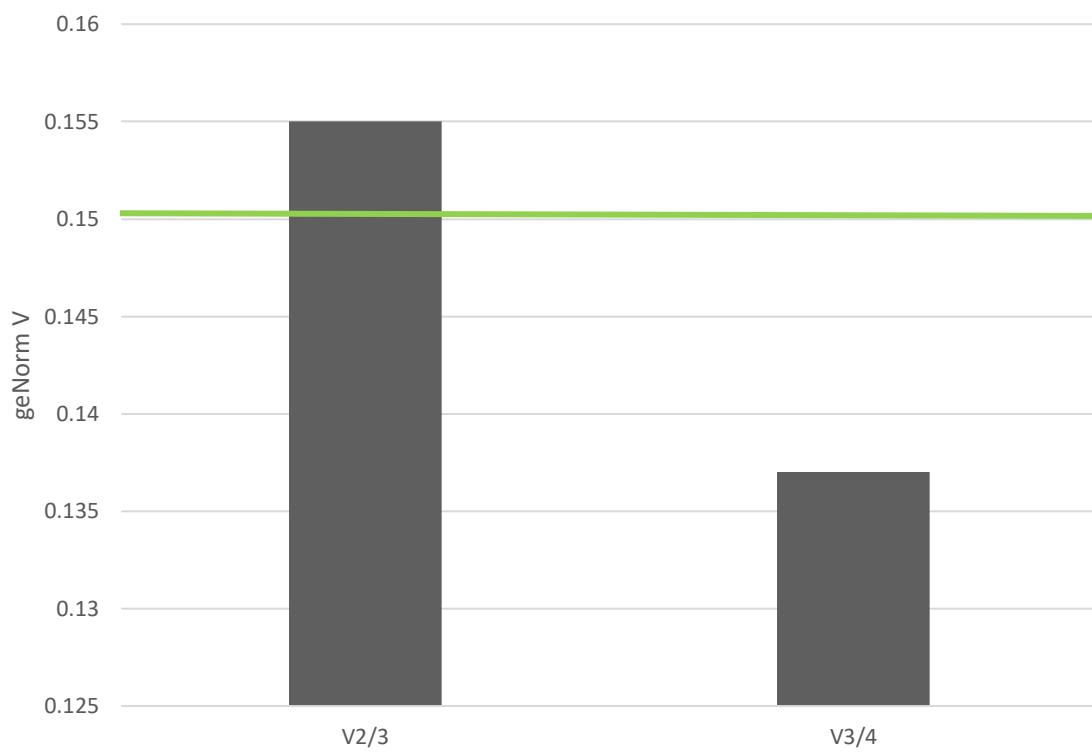
Supplementary Figure S4. Original PTEN blot after miR-21 LNA and BPA treatment.

(A)

Average expression stability of remaining reference targets

**(B)**

Determination of the optimal number of reference targets



Supplementary Figure S5. Optimal reference gene selection (A) and number of references (B) across treatments using geNorm.