Potent and Selective Inhibition of A-to-I RNA Editing with 2'-O-Methyl/Locked Nucleic Acid-containing Antisense Oligoribonucleotides

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Supplementary Table 1: Oligonucleotide Sequences Used for Antisense and RT-PCR

Oligonucleotides Sequence (5' to 3')

Sequence (5 to 5)
CAUAAUCCGAAAGUAUUGAGCAU
CAUAAUCCGAAAGUAUUGAGCAU
ACAGUACAUAAUCCGAAAGUAUU
AC a gu a ca t aa u cc g aa a gu a uu
C <i>t</i> gu <i>g</i> g <i>g</i> a <i>c</i> a <i>g</i> a <i>g</i> g <i>a</i> a <i>c</i> a
GG AAU C AA GT AU AG CU C A
CTGUGGGACAGAGGAACA
CCGCGAATTGAAACGGCTATGCTC
TCCAGACCTGCTGGAGCTAT
TGGCCTTGGATTTCTTTTTG
CCCAAGGAAGTGGTCCAGTTGG
CTGGAACCAGATGGTACGGCC
CCAGAGAGCACAGCGGC
CAAATTGTGCAAGGCTTCCG
TCAGCTCCTGGAGAGTGAGGGTT
TCCAGGAGGAAGCTGAGCAGGTAG

^{*}bold and italicized residues are LNA

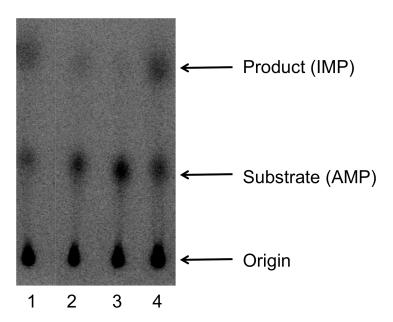


Figure S1. Inhibition of editing of the NEIL1 RNA by the NL18 mixmer. Representative storage phosphor diagram of TLC plate used to separate deamination products arising from the reaction of 50 nM ADAR1 with \leq 18 nM RNA. Lanes 1-3: AON concentrations of 0, 10, 30 nM, respectively. Lane 4: 100 nM of the control AON.

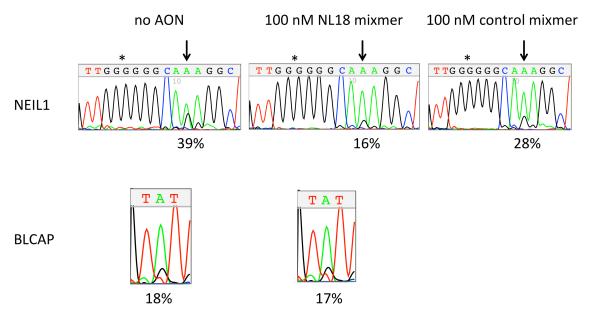


Figure S2. Representative sequencing traces upon isolation of RNA from HeLa cells with treatment of varying concentrations of AON. AON target for inhibition is the NEIL1 pre-mRNA, the BLCAP RNA is used as a control. Arrow indicates recoding site, * indicates exon-exon junction.