**Table S1.** Primers used in this study

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| Primer | Sequence (5’ – 3’) | Use |
| C9 Cut M1 | F – GCCGCCGCGGAATTCATGGATAAGAAATACTCAATAGGACTGR - CTTATCCATGAATTCCGCGGCGGCAGATCTCCTCGGTACCGG | To create EcoRI site before Cas9 in pCas-guide-GFP to prepare Cas9-/sgRNA vector |
| C9 Cut M2 | F - CAGCTGGGAGAATTCCCCAAGAAAAAACGCAAGGTGR - TTTCTTGGGGAATTCTCCCAGCTGACTCAAATCAAT | To create EcoRI site after Cas9 in pCas-guide-GFP to prepare Cas9-/sgRNA vector |
| Eco Del 1 C9 | F - GGGCGGCCGGGCATTCGTCGACTGGAACCGGTACCGAGGR - TCGACGAATGCCCGGCCGCCCTATAGTGAGTCGTATTAC | To delete EcoRI site in pCas-guide-GFP  |
| Eco Del 2 C9 | F - GCAGTTAACGCATTCCCCAGTGGAAAGACGCGCAGGCAAR - ACTGGGGAATGCGTTAACTGCCATCCAGCTGATATCCCC | To delete EcoRI site in pCas-guide-GFP |
| sgIQGAP1.1 | F - GATCGCGAAGTGAAAGCCAAATTCAGR - AAAACTGAATTTGGCTTTCACTTCGC | To clone sgIQ 1.1 sequences into pCas-guide-GFP and Cas9-/sgRNA to prepare Cas9/sgIQ 1.1 and Cas9-/sgIQ 1.1 (sgIQ 1.1) |
| sgIQGAP1.2 | F - GATCGATCAGTCCAACAGAAGAAGTGR - AAAACACTTCTTCTGTTGGACTGATC | To clone sgIQ 1.2 sequences into pCas-guide-GFP to prepare Cas9/sgIQ 1.2 |
|  RNA Scaffold | 5’- TAATTTGACTGTAAACACAAA5’- GCACCGACTCGGTGCCACTTT | To test the successful loading and loading efficiency of sgIQ 1.1 after electroporation |
|  IQGAP1  Genome | F - GACATTGCCAGGGATATTCGGR - GGTAACAAATGTCCCATCAG | To amply genomic DNA fragment of IQGAP1 for T7EI Assay |
| LAMP2 RE | F - GGCCCGGGATCCACCATGGTGTGCTTCCGCCTCTTCCR - TTGCTACCATGACC ATCAAATTGCTCATATCCAGCATG | To fuse LAMP2 into pLVX-AcGFP plasmid |
| HN3LAMP2 RE | F- GTGCGGTCTTATGCAATGCAGGTGCAGCTGGTGCAGTCTGR – TTCCAAGCTGCCTCCGCCGCCACTTGAGGAGACGGTGACCAGGGTTC | To fuse HN3LAMP2 into pLVX-LAMP2-AcGFP plasmid |