

Product Data Sheet

IMPAD1 siRNA (Human)

| Catalog # | Source | Reactivity | Applications | | | |
|-----------------|-----------|---|--|--|--|--|
| CRJ0331 | Synthetic | н | RNAi | | | |
| Description | siRN | A to inhibit IMPAD1 ex | pression using RNA interference | | | |
| Specificity | IMP | AD1 siRNA (Human) is | a target-specific 19-23 nt siRNA oligo duplexes designed | | | |
| | to kr | nock down gene expre | ssion. | | | |
| Form | Lyop | hilized powder | | | | |
| Gene Symbol | IMPA | IMPAD1 | | | | |
| Alternative Na | ames IMP/ | IMPA3; Inositol monophosphatase 3; IMP 3; IMPase 3; Golgi 3-prime | | | | |
| | phos | phoadenosine 5-prim | e phosphate 3-prime phosphatase; Golgi-resident PAP | | | |
| | phos | sphatase; gPAPP; Inosit | ol monophosphatase domain-containing protein 1; | | | |
| | Inos | itol-1(or 4)-monophos | phatase 3; Myo-inositol monophosphatase A3 | | | |
| Entrez Gene | 5492 | 54928 (Human) | | | | |
| SwissProt | Q9N | Q9NX62 (Human) | | | | |
| Purity | > 97 | % | | | | |
| Quality Control | ol Oligo | Oligonucleotide synthesis is monitored base by base through trityl analysis to ensure | | | | |
| | appr | opriate coupling efficie | ency. The oligo is subsequently purified by affinity-solid | | | |
| | phas | e extraction. The anne | aled RNA duplex is further analyzed by mass | | | |
| | spec | trometry to verify the | exact composition of the duplex. Each lot is compared to | | | |
| | the p | previous lot by mass sp | ectrometry to ensure maximum lot-to-lot consistency. | | | |
| Components | Weo | We offers pre-designed sets of 3 different target-specific siRNA oligo duplexes of | | | | |
| | hum | an IMPAD1 gene. Each | vial contains 5 nmol of lyophilized siRNA. The duplexes | | | |
| | can l | pe transfected individu | ally or pooled together to achieve knockdown of the | | | |
| | targe | et gene, which is most | commonly assessed by qPCR or western blot. | | | |

Application key: E- ELISA, WB- Western blot, IH- Immunohistochemistry, IF- Immunofluorescence, FC- Flow cytometry, IC-Immunocytochemistry, IP- Immunoprecipitation, ChIP- Chromatin Immunoprecipitation, EMSA- Electrophoretic Mobility Shift Assay, BL- Blocking, SE- Sandwich ELISA, CBE- Cell-based ELISA, RNAi- RNA interference Species reactivity key: H- Human, M- Mouse, R- Rat, B- Bovine, C- Chicken, D- Dog, G- Goat, Mk- Monkey, P- Pig, Rb-Rabbit, S- Sheep, Z- Zebrafish

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| Component | 15 nmol | 30 nmol |
|--------------------------|--------------|--------------|
| IMPAD1 siRNA (Human) - A | 5 nmol x 1 | 5 nmol x 2 |
| IMPAD1 siRNA (Human) - B | 5 nmol x 1 | 5 nmol x 2 |
| IMPAD1 siRNA (Human) - C | 5 nmol x 1 | 5 nmol x 2 |
| Negative Control | 2.5 nmol x 1 | 2.5 nmol x 2 |
| DEPC Water | 1 ml x 1 | 1 ml x 2 |

Directions for Use

We recommends transfection with 10 nM - 100 nM siRNA 48 to 72 hours prior to cell lysis. Before resuspending, briefly centrifuge the tube to ensure the lyophilized siRNA is at the bottom of the tube. Resuspend the siRNA oligos to an appropriate concentration with DEPC water. For example, resuspend one tube of 5 nmol siRNA oligo in 250 μ l of DEPC water to get a final concentration of 20 μ M.

| Plate | Final volume | Final concentration | siRNA (20 μM) | Lipofectamin |
|---------|--------------|---------------------|---------------|--------------|
| | of medium | of siRNA | | 2000 |
| | | 100 nM | 0.5 μl | 0.25 μl |
| 96-well | 100 μl | 50 nM | 0.25 μl | 0.25 μl |
| | | 10 nM | 0.05 μl | 0.25 μl |
| | | 100 nM | 2.5 μl | 1 µl |
| 24-well | 500 μl | 50 nM | 1.25 μl | 1 μΙ |
| | | 10 nM | 0.25 μl | 1 μΙ |
| | | 100 nM | 5 µl | 2 μΙ |
| 12-well | 1 ml | 50 nM | 2.5 μl | 2 μΙ |
| | | 10 nM | 0.5 μl | 2 μΙ |
| 6-well | 2 ml | 100 nM | 10 µl | 5 µl |
| | | 50 nM | 5 µl | 5 μΙ |

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10 nM

1 µl

5 µl

Storage/Stability

Shipped at 4 °C. Store at -20 °C for one year.

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