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 2 **FIG S3** Overexpression of σ NS complements siRNA knockdown of σ NS during
 3 infection. Cells that constitutively express siRNAs directed against σ NS or GFP were
 4 transfected with expression plasmids encoding GFP, WT σ NS (σ NS), or WT σ NS with
 5 σ NS-siRNA-resistant sequences (σ NS MM) and incubated for 24 h. Cells were
 6 adsorbed with reovirus strain T3D at an MOI of 5 PFU/cell and incubated for 24 h. Cell
 7 lysates were collected for (A) immunoblotting and (B) infectious virus quantification by
 8 plaque assay. (A) Immunoblot analysis of proteins expressed following
 9 complementation. Protein expression was evaluated using monoclonal antibodies
 10 specific for GFP or alpha-tubulin (α -tub) and guinea-pig sera specific for σ NS. (B)
 11 Infectious virus quantification following complementation. Titer values that differ

12 significantly from those obtained from cells expressing siRNAs against σ NS
13 complemented with GFP by one-way analysis of variance (ANOVA) with Dunnett's
14 multiple-comparison test are shown (****, $P < 0.0001$).

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