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In the July issue of [Nature Medicine](#) Carl Novina and colleagues at the [Massachusetts Institute of Technology](#) describe the use of [short interfering RNA](#) (siRNA) strategies to prevent infection by the AIDS-causing virus, HIV-1 (*Nature Medicine*, online 3 June 2002, DOI:10.1038/nm725). Novina *et al.* designed siRNA against the human CD4 gene, encoding a major HIV surface receptor. Cell transfections of CD4-siRNA reduced CD4 expression by about 75%, and blocked HIV-1 viral infection. Novina *et al.* also developed siRNA directed against the viral *gag* gene to block expression of the p24 antigen, a major structural protein of the virus. The p24-siRNA effectively silenced viral expression and inhibited viral infections. Novina *et al.* demonstrated in tissue culture models (HeLa cells or human T-cell lines) that siRNA can regulate viral entry and syncytium formation, alter free viral titers and affect HIV infections. This proof-of-principle study hints at the therapeutic potential of siRNA technology.

References

1. *Nature Medicine*, [<http://medicine.nature.com>]
2. Massachusetts Institute of Technology , [<http://www.mit.edu>]
3. RNA interference - 2001.