

Efficient and specific gene knockdown by small interfering RNAs produced in bacteria

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Synthetic small interfering RNAs (siRNAs) are an indispensable tool to investigate gene function in eukaryotic cells^{1,2} and may be used for therapeutic purposes to knock down genes implicated in disease³. Thus far, most synthetic siRNAs have been produced by chemical synthesis. Here we present a method to produce highly potent siRNAs in *Escherichia coli*. This method relies on ectopic expression of p19, an siRNA-binding protein found in a plant RNA virus^{4,5}. When expressed in *E. coli*, p19 stabilizes an ~21-nt siRNA-like species produced by bacterial RNase III. When mammalian cells are transfected by them, siRNAs that were generated in bacteria expressing p19 and a hairpin RNA encoding 200 or more nucleotides of a target gene reproducibly knock down target gene expression by ~90% without immunogenicity or off-target effects. Because bacterially produced siRNAs contain multiple sequences against a target gene, they may be especially useful for suppressing polymorphic cellular or viral genes.

RNA interference (RNAi) by double-stranded (ds) siRNAs induces the degradation of mRNAs bearing complementary sequences^{6,7}. Although most synthetic siRNAs are designed by computer algorithms and produced by chemical synthesis, siRNAs can also be made from transcribed longer dsRNAs that are processed *in vitro* by RNase III family enzymes^{8,9}. In the latter case, the resulting siRNAs contain many sequences against one target (rather than a single sequence as occurs with chemically synthesized siRNA). A pool of several siRNAs can sometimes be more effective and have fewer off-target effects than any one single siRNA^{10,11}. However, thus far, functional siRNAs have not been produced in living cells. Here, we engineer bacterial cells to produce fully processed ready-to-use siRNAs specific for a target gene of interest.

The p19 protein encoded by the plant RNA virus *tombusvirus*⁴ selectively binds to and inhibits the function of siRNAs of ~21 nucleotides, including those containing sequences complementary to virus RNA⁵. A p19 dimer binds to the ~19-nt duplex region of an siRNA in a sequence-independent manner^{12,13}. In agreement with a previous publication, our work shows that endogenous siRNAs in mammalian cells can be isolated using p19 coupled to magnetic beads (Fig. 1a)¹³. We used as a negative control p19 beads incubated with total RNA isolated from *E. coli* (a wild-type strain and a strain transformed with

a pcDNA3.1+ plasmid encoding p19) because *E. coli* lacks canonical RNAi-processing machinery. Surprisingly, p19-coupled beads retrieved ~21-nt dsRNAs from the *E. coli* strain containing the p19 plasmid (Fig. 1a). Although the cytomegalovirus promoter¹⁴ driving expression from this plasmid is mostly used for efficient gene expression in mammalian cells, pcDNA3.1+ plasmids encoding FLAG-tagged p19 or a FLAG-tagged control gene of a similar size (*TREX1*) drove detectable protein expression in *E. coli* (Fig. 1b). We detected small ~21-nt RNAs on SYBR Gold-stained denaturing polyacrylamide gels of total RNA harvested from p19-expressing bacteria, but not on gels of total RNA isolated from bacteria transformed with the empty vector or a vector encoding *TREX1* (Fig. 1b). These data suggest that p19 expression stabilizes a cryptic siRNA-like RNA species in *E. coli*. Similarly sized small RNAs were also detected in p19-expressing, but not wild-type (WT), strains of the Gram-positive bacterium *Listeria monocytogenes* (Supplementary Fig. 1).

To determine if the small RNAs detected in *E. coli* depended on functional p19, we isolated RNA from *E. coli* expressing WT p19 or p19 containing mutations that disrupt siRNA binding^{12,15}. The ~21-nt dsRNA band was more prominent in bacteria expressing WT than mutant p19 (Fig. 1c). Thus siRNA binding to p19 promotes the accumulation of siRNA-like RNAs in *E. coli*.

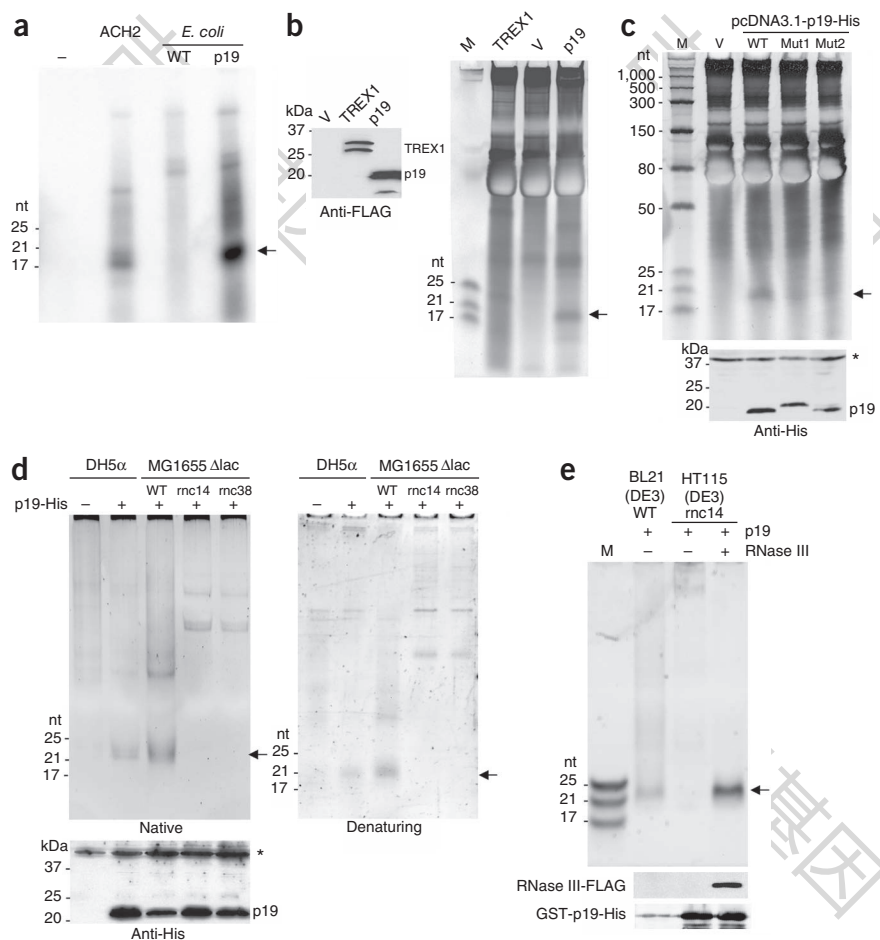
Next we looked for the nuclease responsible for making small RNAs. The most likely candidate was RNase III, an ancestor of eukaryotic Dicer, which is responsible for the final step of siRNA biogenesis¹⁶. *E. coli* RNase III can generate siRNA-sized dsRNAs from longer dsRNAs *in vitro*⁹. We transformed two RNase III mutant strains, *rnc14* (ref. 17) and *rnc38* (ref. 18), with plasmids encoding p19 (Fig. 1d). p19-coupled beads did not pull down any visible small RNAs in either of the RNase III mutant strains. Furthermore, restoration of RNase III expression in HT115(DE3) cells, an *rnc14* strain, restored the production of p19-dependent small RNAs (Fig. 1e). Thus, accumulation of these small RNAs in bacteria depends on ectopic p19 expression and endogenous RNase III expression.

We next asked whether small RNAs generated in p19-expressing *E. coli* exhibit properties similar to those of chemically synthesized siRNAs. We cloned p19 into the pGEX-4T-1 plasmid to express a GST-p19 fusion protein with a C-terminal His tag (Fig. 2a). A T7 promoter driving expression of a hairpin RNA encoding the sequence of the target gene was inserted immediately after the His tag in this plasmid.

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Figure 1 Ectopic p19 expression captures small RNAs in *E. coli*. (a) p19-coupled magnetic beads were incubated with total RNA isolated from mammalian ACH2 cells, or from *E. coli* cells that were either WT or transformed with a pcDNA3.1-p19 expression plasmid. Captured RNAs were 5' 32 P-labeled, separated on a native polyacrylamide gel and detected by autoradiography. (b) FLAG-tagged p19 or TREX1, or empty vector (V), were expressed in *E. coli*. Left, anti-FLAG immunoblot. Right, total RNAs were separated on a denaturing polyacrylamide gel and stained with SYBR Gold. (c) Top, total RNAs purified from *E. coli* expressing an empty vector (V), or WT or mutant (Mut1 (ref. 12): W39G, W42G; and Mut2 (ref. 15): K71A, R72G; mutants were defective in siRNA binding) His-tagged p19 proteins were separated on a denaturing polyacrylamide gel and stained with SYBR Gold. Bottom, anti-His immunoblot. (d) p19-coupled magnetic beads were incubated with total RNA extracted from WT *E. coli* (DH5 α or MG1655 Δ lac) or RNase III mutant strains (*rnc14* and *rnc38* in MG1655 Δ lac background) expressing or not His-tagged p19. p19-captured RNAs were separated on native (left) or denaturing (right) gels and stained with SYBR Gold. Bottom, anti-His immunoblot. The asterisks (*) in (c,d) indicate equal loading of a background band. (e) p19-coupled magnetic beads were incubated with total RNA extracted from *E. coli* WT BL21(DE3) cells or *rnc14* mutant HT115(DE3) cells that were co-transfected with p19 and a vector encoding Flag-tagged *E. coli* RNase III (or empty vector). p19-bound RNAs were separated on a native polyacrylamide gel and stained with SYBR Gold. Bottom, anti-FLAG and anti-His immunoblots. Arrows indicate the ~21-nt small RNA band. M, markers. Data are representative of at least two independent experiments.



We first used a hairpin encoding full-length *EGFP* (*EGFPFL*). The expression of the GST-p19-His fusion protein and hairpin RNA were both induced by isopropyl β -D-thiogalactoside (IPTG). We used nickel (Ni) affinity chromatography to capture the GST-p19-His protein, and 0.5% SDS to selectively elute p19-bound RNAs, which were predominantly ~21 nt long (Fig. 2b and Supplementary Fig. 2). Small RNAs were further purified from other longer RNAs by anion exchange high-performance liquid chromatography (HPLC). To verify that these bacterial small RNAs, which we refer to as pro-siRNAs for prokaryotic siRNAs, are double stranded, we treated them with a variety of nucleases. Like chemically synthesized siRNAs, bacterial small RNAs were sensitive to RNase A, but were insensitive to enzymes that digest single-stranded (ss)RNA or DNA (Xrn1, RNase T1, exonuclease T (Exo T), exonuclease I (Exo I) or DNase Turbo) (Fig. 2c). Next, we tested whether transfection of HeLa cells stably expressing *d1EGFP* (HeLa-d1EGFP) with bacterial small RNAs that were purified from *E. coli* expressing p19 and an *EGFPFL* hairpin would result in incorporation of the RNAs into Argonaute (Ago), the central component of the RNA-induced silencing complex (RISC). To do this we performed immunoprecipitation with a pan-Ago antibody and analyzed the ability of the associated RNAs to hybridize to an *EGFP* probe (northern blot analysis) (Fig. 2d). RNAs that immunoprecipitated with anti-Ago were ~21 nt long and hybridized to the *EGFP* probe only in cells transfected with the *EGFPFL* pro-siRNA; in contrast, no small RNA immunoprecipitated with control mouse IgG.

Thus bacterial small RNAs were similar to synthetic siRNAs in chemical composition and were incorporated into the RISC.

Because pro-siRNAs exhibit properties of siRNAs, we next tested whether pro-siRNAs can suppress expression of specific target genes. We transfected HeLa-d1EGFP cells with a chemically synthesized *EGFP* siRNA or with pro-siRNAs purified from *E. coli* expressing p19 and an *EGFPFL* hairpin or a hairpin encoding a 100-nt fragment of *EGFP* that overlapped with the chemically synthesized siRNA sequence (*EGFP100*). As measured by quantitative RT-PCR (qRT-PCR) and flow cytometry, both *EGFPFL* and *EGFP100* pro-siRNAs knocked down *EGFP* expression more effectively than equimolar concentrations of chemically synthesized siRNA (Fig. 2e and Supplementary Fig. 3a). Pro-siRNAs made from the p19-expressing plasmid without any *EGFP* sequence or expressing only the antisense half of the *EGFP* hairpin did not effectively knockdown *EGFP* (Supplementary Fig. 3b). As expected, silencing by pro-siRNA was not dependent on Dicer because *EGFPFL* pro-siRNA still functioned in Dicer-deficient HCT116 cells¹⁹ and recombinant Dicer protein did not further process pro-siRNAs *in vitro* (Supplementary Fig. 4).

To test the effectiveness of pro-siRNA-mediated knockdown of endogenous and viral genes, we used convenient restriction sites to clone and express hairpins from the coding regions of *LMNA* (which encodes two splice variant products, lamin A and lamin C), *PLK1*, *TP53* and HIV *vif* (viral infectivity factor) and *gag* (capsid antigen). These hairpins contained 200–579 nt of each sense and antisense

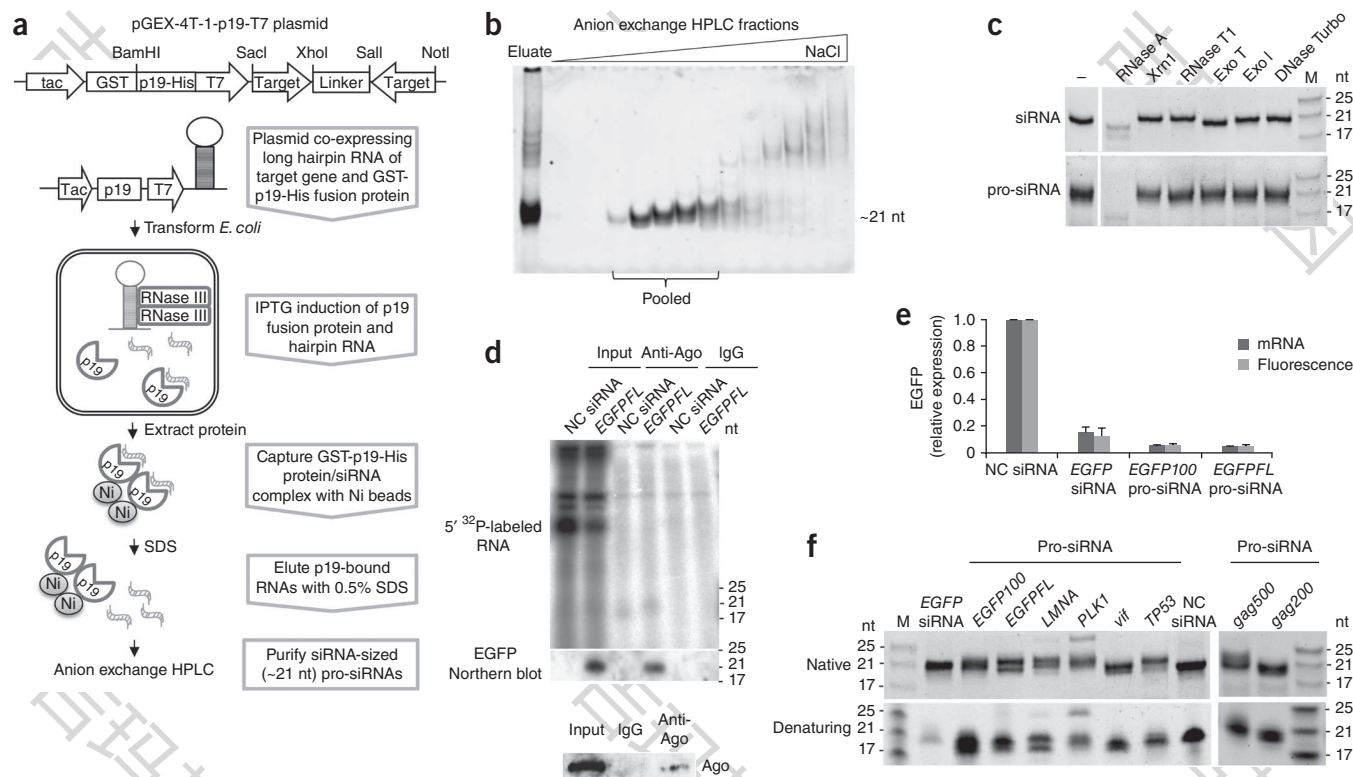


Figure 2 pro-siRNAs knock down target gene expression. **(a)** Schematic of pGEX-4T-1-p19-T7 plasmid and the method to produce pro-siRNAs from *E. coli*. **(b)** Anion exchange HPLC fractions of SDS-eluted RNAs (isolated from *E. coli* engineered as in **a** to express pro-siRNAs) were separated on a native polyacrylamide gel and stained with SYBR Gold. HPLC fractions containing only ~21-nt small RNAs were combined (pooled) for use in subsequent experiments. **(c)** Synthetic siRNAs (top) and HPLC-purified pro-siRNAs (bottom) were incubated with indicated nucleases and separated on a native polyacrylamide gel stained with SYBR Gold. –, no nuclease. **(d)** Anti-Ago mAb 2A8 or mouse total IgG was used to immunoprecipitate RNAs from HeLa-d1EGFP cells transfected with negative control (NC) siRNA or EGFPFL pro-siRNAs. Immunoprecipitated RNAs were analyzed by 5' ³²P end-labeling (top) or by northern blot analysis using a probe complementary to the full-length EGFP coding sequence (middle). Bottom, anti-Ago immunoblot. **(e)** HeLa-d1EGFP cells were transfected with 4 nM of synthetic EGFP siRNA or with pro-siRNAs. EGFP expression was measured by qRT-PCR (mRNA) and flow cytometry (fluorescence). Data are normalized to cells treated with negative control (NC) siRNA. mRNA level is relative to GAPDH. Data are mean \pm s.d. of two (qRT-PCR) and three (EGFP fluorescence) independent experiments. **(f)** HPLC-purified pro-siRNAs or synthetic siRNAs (EGFP and NC) were separated on native (top) or denaturing (bottom) polyacrylamide gels stained with SYBR Gold. M, markers. Data are representative of at least two independent experiments.

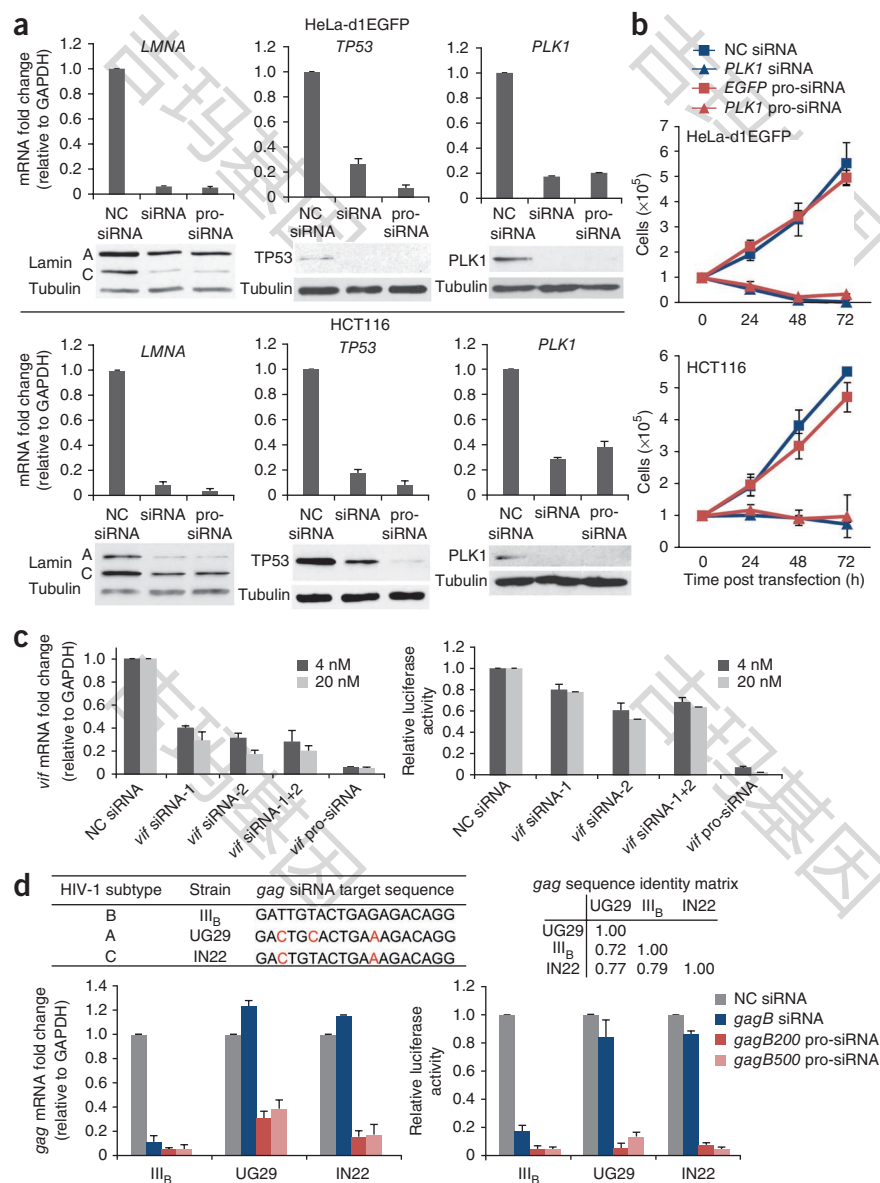
sequence (523 nt for LMNA, 299 nt for PLK1, 300 nt for TP53, 579 nt for vif, and 200 and 500 nt for gag). The HPLC-purified pro-siRNAs for each gene contained a few different-sized species that migrated close to the 21-nt marker on both native and denaturing polyacrylamide gels (Fig. 2f). For LMNA and PLK1 pro-siRNAs, a minor RNA band migrated at ~25 nt. We transfected HeLa-d1EGFP and HCT116 with pro-siRNAs and commercially available chemically synthesized siRNAs (LMNA and TP53 siRNAs were from a single sequence; PLK1 siRNAs were a pool of four siRNAs and were chemically modified by proprietary methods for enhanced stability and reduced off-target effects²⁰). The extent of gene knockdown was similar when cells were transfected with siRNAs and pro-siRNAs at 4 nM (Fig. 3a). To more closely evaluate the potency of pro-siRNAs, we did dose-response experiments comparing transfection with pro-siRNAs (0.2, 2, 20 nM) targeting LMNA, TP53 and PLK1 to five commercial siRNAs for each gene (four siRNAs from Dharmacon, of which the PLK1 siRNAs were chemically modified²⁰) (Supplementary Fig. 5). The potency of the commercial siRNAs varied, as best evaluated at the lowest concentration. The pro-siRNAs, whose sequences were not optimized, achieved similar gene knockdown as the commercially optimized siRNAs. At a concentration of 2 nM, each pro-siRNA achieved a knockdown of ~90%. Because siRNA design algorithms are imperfect, identifying

potent siRNAs often requires testing several sequences, which can be time consuming and costly. pro-siRNAs might circumvent the need to test multiple sequences to identify a single potent siRNA.

To examine the potential toxicity of pro-siRNAs, we analyzed the growth of HeLa-d1EGFP and HCT116 (human colon carcinoma) cells after they were transfected with either a negative control siRNA or EGFP pro-siRNA (Fig. 3b). Their growth curves were not significantly different. To compare the effectiveness of gene knockdown by pro-siRNAs and siRNAs, we examined cell proliferation after knocking down PLK1, which kills dividing cells²¹. PLK1 siRNAs and pro-siRNAs markedly reduced viability with indistinguishable kinetics (Fig. 3b).

We next used pro-siRNAs to knockdown the HIV gene vif, which targets the host restriction factor APOBEC3G for ubiquitylation and degradation, thereby preventing APOBEC3G packaging into virions. Vif is therefore dispensable for the initial round of HIV replication but essential for the spread of the infection to new cells. We compared the efficacy of vif pro-siRNAs with two validated chemically synthesized siRNAs^{22,23}. Neither siRNAs nor pro-siRNAs targeting vif altered the percentage of initially infected HeLa-CD4 cells (data not shown), but both suppressed vif gene expression and inhibited subsequent rounds of infection, as assessed in the TZM-bl luciferase reporter cell line (Fig. 3c).

Figure 3 pro-siRNA-mediated knockdown of endogenous and viral gene expression in human cells. **(a)** Synthetic siRNAs or pro-siRNAs specific for the indicated target genes or negative control (NC) siRNA were transfected (4 nM) into HeLa-d1EGFP (top) or HCT116 (bottom) cells. Target gene expression was measured by qRT-PCR (bar graphs) or immunoblots (tubulin is loading control). **(b)** HeLa-d1EGFP (top) or HCT116 (bottom) cells were transfected with *PLK1* siRNA or pro-siRNA (4 nM) or NC siRNA or *EGFP* pro-siRNA as nontargeting controls, respectively. Cells were counted at indicated time points. **(c)** HeLa-CD4 cells were transfected with *vif* siRNAs (*vif* siRNA-1 and *vif* siRNA-2, alone or together) or *vif* pro-siRNA 24 h before HIV-1 infection. Left, *vif* mRNA was measured by qRT-PCR 36 h after infection. Right, infectivity of culture supernatants from transfected HeLa-CD4 cells was measured by HIV-1 *tat*-driven expression of a luciferase reporter gene in TZM-bl assay. **(d)** Top, target sequence of the *gag* siRNA (HIV-1 clade B) used here; corresponding sequences in clade A UG29 and clade C IN22 strains are shown together (left) with the proportion of identical bases in the entire *gag* coding region for the three isolates tested (right). Bottom, HeLa-CD4 cells were infected with HIV-1 clade B (III_B), U87.CD4.CXCR4 (ref. 33) cells were infected with HIV-1 clade A (UG29) and U87.CD4.CCR5 cells³³ were infected with HIV-1 clade C (IN22). Cells were transfected with 20 nM of the indicated synthetic or pro-siRNA 12 h before HIV-1 infection. Left, *gag* mRNA was measured by qRT-PCR 36 h after infection. Right, infectivity of culture supernatants was measured by TZM-bl assay. Data are mean \pm s.d. of three (**a–c**) and two (**d**) independent experiments. mRNA expression and TZM-bl luciferase data are normalized to cells transfected with NC siRNA.



vif pro-siRNAs were more potent than chemically synthesized siRNAs.

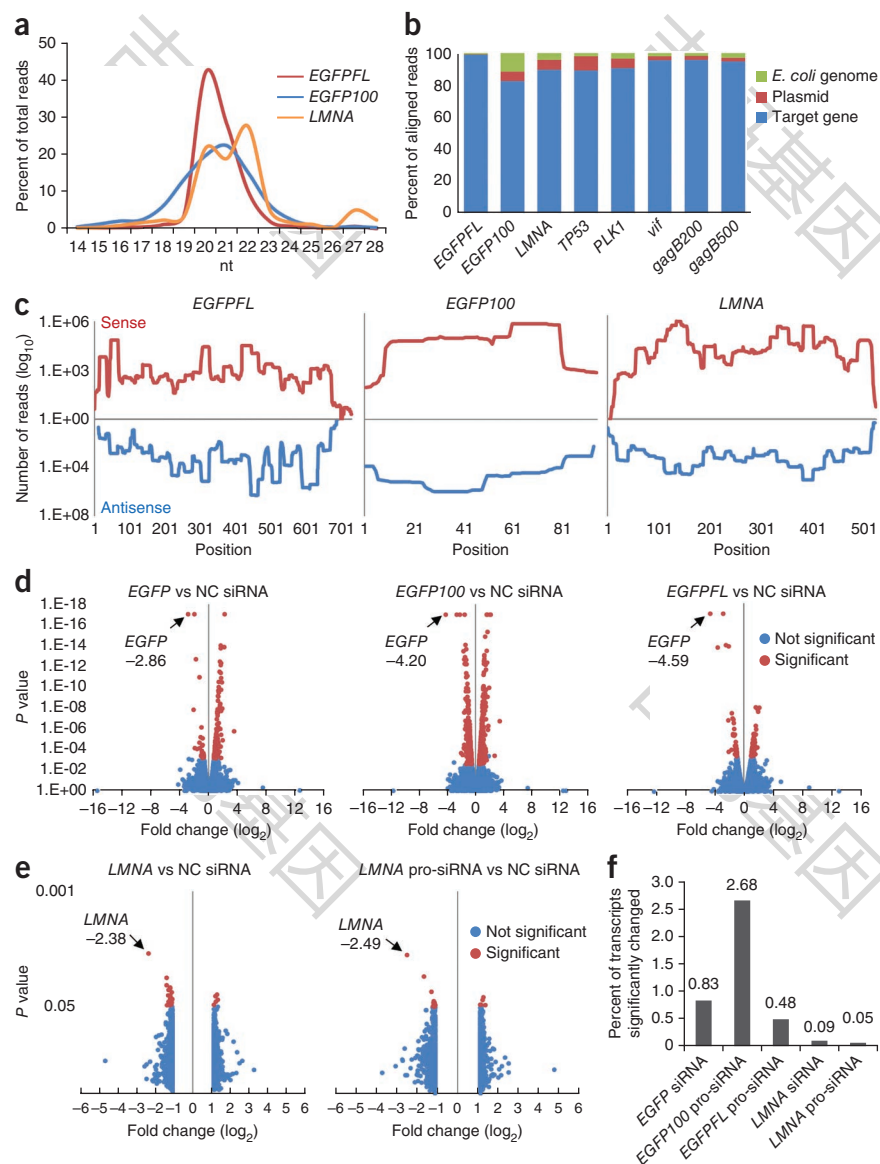
One major obstacle to using RNAi to suppress HIV or other viruses is virus sequence diversity. Because pro-siRNAs target many sequences within a gene (see below), compared to chemically synthesized siRNAs, pro-siRNAs directed against a viral gene might have broader activity against diverse viral strains and might be less likely to incite generation of siRNA-resistant mutants. We previously tried unsuccessfully to identify an siRNA against HIV-1 clade B *gag* that could also inhibit viral isolates from other clades²². To investigate whether pro-siRNAs might have broader activity, we used hairpins encoding 200 and 500 nt from the *gag*-coding region of clade B HIV-III_B virus. Compared to a chemically synthesized *gag* siRNA, the *gagB200* and *gagB500* pro-siRNAs more potently suppressed HIV-III_B (Fig. 3d). More importantly, both *gag* pro-siRNAs, but not the chemically synthesized *gag* siRNA, knocked down expression of *gag* mRNA and inhibited *in vitro* spread of viruses of different clades (UG29, clade A; IN22, clade C). However, not surprisingly, the *gag* pro-siRNAs more efficiently inhibited the III_B virus than the UG29 or IN22 viruses. These data suggest that pro-siRNAs could be particularly useful for targeting heterogeneous and rapidly evolving viral genes.

Because mammalian cells are sensitive to bacterial endotoxin, which elicits production of inflammatory mediators by stimulating

Toll-like receptor 4, we assessed whether purified pro-siRNAs are contaminated with endotoxin. Although SDS-eluted pro-siRNAs contained substantial amounts of endotoxin as measured by Limulus amoebocyte lysate (LAL) assay, HPLC-purified pro-siRNAs, even at concentrations as high as 320 nM, contained no detectable endotoxin (Supplementary Table 1). We also tested for endotoxin contamination by measuring expression of proinflammatory cytokine and interferon-stimulated genes in monocyte-derived human macrophages incubated with (Supplementary Fig. 6a) or transfected with (Supplementary Fig. 6b) HPLC-purified pro-siRNAs. Neither pro-inflammatory nor interferon-stimulated genes were induced by pro-siRNAs in these highly endotoxin-sensitive cells.

To ascertain the sequence composition of pro-siRNAs, we cloned and deep sequenced pro-siRNAs using a method established for eukaryotic siRNAs (sequencing summary in Supplementary Table 2). Most reads were 20–22 nt in length (Fig. 4a and Supplementary Fig. 7). The majority of reads (on average ~75%) aligned to the target gene sequence, plasmid backbone or the *E. coli* genome and, consistent with the efficiency of target gene knockdown, the majority of

Figure 4 pro-siRNA sequences and assessment of off-target effects. (a) Length distribution of *EGFPFL*, *EGFP100* and *LMNA* pro-siRNAs assessed by deep sequencing. (b) Percentage of deep sequencing reads aligning to the target gene hairpin, the *E. coli* genome or the plasmid backbone. (c) pro-siRNA deep sequencing read alignment to indicated positions of the *EGFPFL*, *EGFP100* and *LMNA* target genes (sense and antisense). (d) Volcano plots of expression changes versus *P* value of all annotated transcripts detected by RNA deep sequencing in HeLa-d1EGFP cells transfected with *EGFP* synthetic siRNA or *EGFP100* or *EGFPFL* pro-siRNAs relative to expression in cells transfected with a negative control (NC) siRNA. Arrows indicate *EGFP* and the numbers indicate its fold change (\log_2). Cut-off for significance is *q* value < 0.05 (the default in Cufflinks). (e) Volcano plots of expression changes (1.2-fold less or more) versus *P* value detected by microarray in HeLa-d1EGFP cells transfected with *LMNA* synthetic siRNA or pro-siRNAs relative to expression in cells transfected with a NC siRNA. Arrows indicate *LMNA* and the numbers indicate its fold change (\log_2). Cut-off for significance is *P* < 0.05 (paired *t*-test by dChip). (f) Percent of significantly changed transcripts in e and f. Numbers on top of bars indicate exact percentages. Significantly changed genes are listed in **Supplementary Table 4**.



aligned sequences (82–99%) originated from the target gene sequence (Fig. 4b). Reads spanned the entire target gene sequence, but some reads concentrated at specific sites ('hot spots') (Fig. 4c, Supplementary Figs. 7 and 8). We detected some sequence strand bias for most of the hot spots (Supplementary Fig. 8a). Because our data (Fig. 2c,f) strongly suggested that pro-siRNAs are double stranded, we suspected that strand bias may have been due to differences in ligation efficiency during cloning, a well-known problem²⁴, rather than to the presence of many ssRNAs. To evaluate this, we designed forward and reverse DNA oligonucleotide probes (26–27 nt) aligning to three *EGFPFL* pro-siRNA hot spots and performed solution hybridization and native gel electrophoresis (Supplementary Table 3 and Supplementary Fig. 8b). The relative intensity of hybridized bands was approximately equal for sense and antisense probes for each hot spot and loosely correlated with the number of reads from each hot spot (Supplementary Fig. 8c–e). Thus, the strand bias in the deep sequencing data likely reflects ligation bias introduced during cloning.

To further investigate the hot spot pattern, we compared two independent preparations of *EGFPFL* pro-siRNAs cloned using different sets of adaptors. Their potency, size profile and sequence content were similar, but not identical. The most abundant hot spots were consistent in the two samples, but the strand bias changed with the adaptors, consistent with cloning bias (Supplementary Fig. 9a–d). Hot spots might be due to intrinsic sequence preferences for RNase III cleavage, or to differences in RNA stability or RNA binding to p19 after cleavage. To determine whether 'hot spots' are determined by sequence differences at or close to the hot spot, we constructed hairpins of equal sizes from

the 5' and 3' ends of the full-length *EGFP* sequence. The pro-siRNAs generated from the two half *EGFP* hairpins contained hot spots mostly identical to those generated from the full-length *EGFPFL* hairpin (Supplementary Fig. 9e). Thus hot spots seem to be determined by local sequence differences. However, a basic bioinformatic analysis searching for preferred sequence motifs or bases in the hot spots was inconclusive (data not shown). *E. coli* RNase III might process dsRNA into siRNA-sized small RNAs *in vivo* through a mechanism that differs from Dicer²⁵, whose cleavage of a long dsRNA results in phased and evenly distributed sequences along a target gene.

Because pro-siRNAs contained nontargeting sequences derived from the plasmid or *E. coli* genome, we were concerned about possible off-target effects²⁶. To evaluate off-target effects, we compared by RNA deep sequencing the RNA expression profile of HeLa-d1EGFP cells transfected with 4 nM of negative control siRNA, chemically synthesized *EGFP* siRNA or *EGFPFL* or *EGFP100* pro-siRNAs (sequencing summary in Supplementary Table 2). We used Tophat and Cufflinks to analyze the data and generated volcano plots of all annotated transcripts²⁷ (fold change versus *P* value, Fig. 4d). Compared to chemically synthesized *EGFP* siRNA, *EGFPFL* and *EGFP100* pro-siRNAs induced

notable changes in expression of a smaller and larger number of genes, respectively (Fig. 4f, Supplementary Fig. 10a and Supplementary Table 4). *EGFPFL* pro-siRNAs also produced the fewest changes in long noncoding RNA levels (Supplementary Fig. 10b,c). *EGFP100* pro-siRNAs, made from a shorter hairpin (100 bp), contained a higher proportion of sequences mapping to plasmid and *E. coli* genomic sequences, compared to pro-siRNAs made from longer hairpins (200–720 bp, Fig. 4b). Thus, longer target gene hairpins will likely generate pro-siRNA preparations that will have fewer off-target effects. We also compared by microarray the gene expression profiles of cells transfected with *LMNA* chemically synthesized siRNAs and pro-siRNAs. Consistent with the *EGFPFL* data, *LMNA* pro-siRNAs made from a longer hairpin (523 bp) led to fewer significantly changed ($P < 0.05$) genes compared with the chemically synthesized *LMNA* siRNA (Fig. 4e,f, Supplementary Fig. 10d and Supplementary Table 4). For both *EGFP* and *LMNA*, the intended target was always the most downregulated gene, and pro-siRNAs consistently induced greater suppression than the siRNA. The significantly changed genes in each of these experiments were not enriched for innate immune genes²⁸ (Supplementary Table 4), confirming that the pro-siRNAs did not stimulate an innate immune response. Thus pro-siRNAs offer highly specific knockdown that is at least as good as synthetic siRNAs without the need to test multiple sequences.

Here we showed that bacteria can produce siRNAs that are not toxic and efficiently suppress expression of exogenous genes (*EGFP*), viral genes (*vif* and *gag*) and endogenous genes (*PLK1*, *TP53*, *LMNA*) in mammalian cells. Because pro-siRNAs are natural products of RNase III, they likely have favorable ends (e.g., 5'-phosphate, 3'-hydroxyl and 3' overhangs) for efficient loading by Ago into the RISC, and they do not activate cytosolic innate immune RNA sensors. Although we used one plasmid to express p19 and the target gene hairpin, it is possible to use two separate plasmids to express p19 and the sense and antisense strands of the target sequence (Supplementary Fig. 11). Like chemically synthesized siRNAs, but unlike short hairpin RNA-mediated stable gene knockdown, the pro-siRNA-mediated gene silencing is temporary.

Without much optimization we achieved an average yield of ~4 nmol (~42 µg) pro-siRNA per liter of *E. coli* culture, which, albeit modest, is more than enough for most laboratory experiments, given the activity of pro-siRNAs at low nanomolar concentrations. The engineered plasmid or *E. coli* genome could potentially be further optimized to maximize yield and improve effectiveness and/or specificity. For example, we doubled the yield of *EGFPFL* pro-siRNA by overexpressing *E. coli* RNase III (Supplementary Fig. 12).

Previous studies described the use of *E. coli* to induce RNAi. For example, RNase III-deficient *E. coli* expressing dsRNAs can be fed to *Caenorhabditis elegans* to knockdown expression of worm genes¹⁷, and bacteria-derived dsRNAs can be applied to plants to induce specific gene knockdown²⁹. In worms and plants, unlike mammalian cells, gene silencing is enhanced by RNA-dependent RNA polymerases that amplify small amounts of RNA. More recently, *E. coli* genetically engineered to express an invasin (to induce bacterial uptake) and listeriolysin (to allow bacterial RNAs to escape from phagolysosomes) delivered dsRNAs into the cytoplasm of human cells through “trans-kingdom RNAi” technology^{30,31}. In all these scenarios, target gene silencing requires host cell processing of long siRNA precursors by Dicer.

pro-siRNAs could become a valuable addition to existing RNAi techniques for both research and therapeutics. Use of pro-siRNAs would eliminate the need to purchase and test multiple individual chemically synthesized siRNAs. When generated from longer hairpins, pro-siRNA preparations containing multiple sequences might

trigger fewer off-target effects than individual siRNAs and, in the cases of virus infection or cancer, might more effectively prevent target gene escape by mutation. The method we developed for producing pro-siRNAs was adapted from well-established techniques for making recombinant proteins from *E. coli* and could easily be adapted and scaled-up in an industrial setting. In addition, mammalian cDNA libraries might be used to generate pro-siRNA libraries for siRNA screening purposes. That said, chemical synthesis provides the opportunity for chemical modifications to increase potency, enhance stability and reduce off-target effects or to couple fluorophores or targeting moieties. More work is needed to determine if such modifications might also be possible for pro-siRNAs, for example, by adding modified ribonucleotides to bacterial cultures during IPTG induction or by performing the same coupling reactions with purified pro-siRNAs as are used to modify chemically synthesized siRNAs.

A recent study found that yeast Ago protein expressed in *E. coli* binds small RNAs that are predominantly a mixture of sense and antisense strands derived from sequences of the expression plasmid³². That study confirms our finding that siRNA-like small RNAs are generated in bacteria and can be stabilized by binding to proteins like Ago or p19. Future studies should evaluate whether siRNAs generated in bacteria might have functional significance, for example, in bacterial defense against foreign genetic elements and pathogens or in regulation of endogenous bacterial genes.

METHODS

Methods and any associated references are available in the [online version of the paper](#).

Accession code. Small RNA deep sequencing data, SRA: [SRP018503](#). Microarray data, GEO: [GSE44105](#). mRNA deep sequencing data, GEO: [GSE44256](#).

Note: Supplementary information is available in the [online version of the paper](#).

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AUTHOR CONTRIBUTIONS

L.H. and J.L. designed the experiments with advice from J.J., L.M., and P.D. J.J. and L.M. prepared p19 beads and RNA deep sequencing libraries. P.D. constructed *E. coli* mutant strains. E.K. performed siRNA comparison and macrophage transfection experiments. L.H. performed all other experiments. L.H. and J.L. wrote the paper.

COMPETING FINANCIAL INTERESTS

The authors declare competing financial interests: details are available in the [online version of the paper](#).

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ONLINE METHODS

Bacterial strains and culture conditions. All *E. coli* strains used in this study are listed in **Supplementary Table 5**. *E. coli* strain DH5 α was used for cloning and for initial characterization of the siRNA-like RNA species. For recombinant protein expression and pro-siRNA production, we used T7 Express Iq (New England Biolabs (NEB)), a BL21-derived *E. coli* strain. We used two mutants of RNase III, *rnc-14::Dtn10* (Tet^R) and *Drnc-38* (Kan^R). These were moved by P1 transduction from parent strains HT115(DE3)¹⁷ and SK7622 (ref. 18) into *E. coli* strain MG1655 Δ lacZYA (also referred to as MG1655 Δ lac). All *E. coli* strains were cultured in LB broth, Lennox (BD) at 37 °C with shaking at 250 r.p.m. and antibiotics when required were used at the following concentrations: carbenicillin (100 μ g/ml), kanamycin (50 μ g/ml), spectinomycin (50 μ g/ml), tetracycline (12.5 μ g/ml).

Listeria monocytogenes strain 10403S was cultured in brain-heart infusion medium (BD Biosciences) at 30 °C. Transformation of bacterial cells was done as previously described³⁴.

Genes and plasmids. The p19 gene used in this study was cloned from *Tomato bushy stunt virus* (gift of James Carrington, Donald Danforth Plant Science Center). All plasmids are listed in **Supplementary Table 6** and will be made available through Addgene. To produce p19 in *E. coli*, we used pcDNA3.1⁺ (Invitrogen) to express the p19 protein with a C-terminal FLAG tag (pcDNA3.1-p19-FLAG) or an N-terminal His tag (pcDNA3.1-His-p19). Plasmid pcDNA3.1-TREX1-FLAG encodes a C-terminal FLAG-tagged TREX1 protein. To express p19 in *L. monocytogenes*, pLIV-1-His-p19 plasmid was used, which encodes p19 with an N-terminal His tag in the pLIV-1 plasmid (gift of Darren Higgins, Harvard Medical School). *E. coli* RNase III with an N-terminal FLAG was cloned into pcDNA3.1⁺ and pCDF-1b (Novagen) plasmids.

We used two strategies for pro-siRNA production in *E. coli*. In one approach p19-His was fused to GST in pGEX-4T-1 (to express GST-p19-His fusion protein). On the same plasmid we cloned a hairpin RNA-expressing cassette consisting of an inverted repeat separated by a 32-bp linker downstream of a T7 promoter (**Fig. 2a**). The hairpin RNA sequences were: *EGFPFL*, the entire 720-bp *EGFP* coding sequence (from pEGFP-N1, Clontech); *EGFP100*, 100 bp from nt 219 to 318; *EGFP Hotspot-1* 360 bp from nt 1 to 360; *EGFP Hotspot-2* 360 bp from nt 361 to 720; *LMNA* (NM_005572.3), 523 bp from nt 267 to 789; *TP53* (NM_000546.5), 301 bp from nt 376 to 676; *PLK1* (NM_005030.3), 299 bp from nt 92 to 390; *vif* (K03455), the entire 579-bp; *gag* (K03455), *gagB200*: 200 bp from nt 1183 to 1382, *gagB500*: 500 bp from nt 1004 to 1503. (GenBank entries listed; numbers refer to position with respect to the translation start site).

In another approach we used two compatible plasmids for pro-siRNA production. The GST-p19-His protein was cloned under the control of the T7 promoter in pRSF-1b (Novagen) or pCDF-1b to generate pRSF-GST-p19-His and pCDF-GST-p19-His. The second plasmid is a L4440 plasmid encoding the entire *EGFP* coding sequence (L4440-EGFP).

All cloning was done using PCR and standard techniques. All primers (with information for restriction enzyme sites) are listed in **Supplementary Table 7**.

Cells. Human HeLa-d1EGFP, HCT116, HCT116 *Dicer*^{-/-}, HeLa-CD4, TZM-bl, U87.CD4.CXCR4 and U87.CD4.CCR5 (ref. 33) were cultured in DMEM medium (Invitrogen) supplemented with 10% heat-inactivated FBS (FBS). ACH2 cells (human leukemia T-cell line CEM latently infected with HIV-1) were cultured in RPMI medium (Invitrogen) supplemented with 10% heat-inactivated FBS. For assays using primary monocyte-derived human macrophages, monocytes were isolated from the blood of a healthy donor by Ficoll-Paque Plus (GE Healthcare) density separation. Human samples were obtained with approval by the Boston Children's Hospital Investigation Review Board. Monocytes were plated on PRIMARIA plates (FALCON) in RPMI medium (Invitrogen) supplemented with 10% heat-inactivated human serum and adherent cells were cultured for 5 d to allow differentiation into macrophages.

RNA isolation and qRT-PCR. Total RNA was isolated from 3 ml of *E. coli* stationary phase culture with 1 ml Trizol reagent (Invitrogen) following the manufacturer's protocol. RNA from human cells was collected in Trizol and extracted according to the manufacturer's protocol. Total RNA (1 μ g) was converted to cDNA using SuperScript III Reverse Transcriptase (Invitrogen). For qRT-PCR, 10 μ l reaction, containing SsoFast EvaGreen mastermix (Bio-Rad),

appropriate primers (**Supplementary Table 5**), and template cDNAs made from 10 ng RNA, was amplified on a Bio-Rad CFX 96 Thermal Cycler. All qRT-PCR data were normalized to the human *GAPDH* gene. qRT-PCR primers for human genes (**Supplementary Table 7**) were selected from PrimerBank (<http://pga.mgh.harvard.edu/primerbank/>).

siRNA isolation from total RNA using p19 magnetic beads. p19 magnetic beads were prepared at NEB as previously described¹³. To pull down siRNAs, 50 μ g of total RNA (isolated from human or *E. coli* cells) was used following the manufacturer's protocol¹³.

His-tag purification of GST-p19-His and bound pro-siRNA. GST-p19-His was purified as follows. A fresh single transformant of T7 Express Iq containing pGEX-4T-1-p19-T7 was used to inoculate 300 ml LB medium in a 1.5-l flask. When the A₆₀₀ reached 0.3–0.6, protein and pro-siRNA expression were induced by adding 0.5 mM IPTG for 1 h. Cells were centrifuged and lysed in 10 ml lysis buffer (50 mM phosphate buffer pH 7.0, 300 mM NaCl, 10 mM imidazole, 1% Triton X-100, 1 mg/ml lysozyme) at 4 °C for ~30 min followed by sonication (Misonix S-4000) until the lysate was not viscous. After centrifugation the lysate was incubated with rotation with 1 ml Ni-NTA resin (Thermo Scientific) overnight at 4 °C. The resin was washed with lysis buffer four times, each time for 10 min at 4 °C with rotation. Bound GST-p19-His was eluted in lysis buffer containing 300 mM imidazole at room temperature.

To purify p19-bound pro-siRNA the procedure was as above until the final elution step when 500 μ l 0.5% SDS was added for 10 min at room temperature with rotation. This step was repeated and both SDS eluates were combined and passed through a 0.22- μ m centrifuge filter (Corning) before HPLC purification on a Bio WAX NP5 anion exchange column (Agilent Technologies). The HPLC buffers were: buffer A, 25 mM Tris-HCl, 2 mM EDTA; buffer B, 25 mM Tris-HCl, 2 mM EDTA, 5 M NaCl. HPLC was initiated with a flow rate of 1 ml/min at 25 °C. Elution was done using a linear gradient of 0–10% buffer B over 4 min, followed by 10% buffer B for 6 min, and a second linear gradient of 10–25% buffer B over 15 min at a reduced flow rate of 0.5 ml/min. pro-siRNA eluted in the second gradient was collected by isopropanol precipitation.

Polyacrylamide gel electrophoresis (PAGE) of RNA. For denaturing electrophoresis of RNA, mini-sized, precast 15% polyacrylamide TBE-Urea gels (Invitrogen) were used. RNA samples were heated to 95 °C for 5 min in Gel Loading Buffer II (Ambion) and then immediately placed on ice until gel loading. Electrophoresis was performed in a 70 °C water bath (to ensure complete denaturation of siRNA) and gels were stained with SYBR Gold (Invitrogen). For analysis of *E. coli* total RNA, 20- μ g samples of Trizol-isolated RNA were loaded. RNA size standards (microRNA marker, siRNA marker and Low Range ssRNA Ladder) were from NEB.

For native electrophoresis of RNA, mini-sized homemade 10–20% polyacrylamide TBE gels were used with the Bio-Rad Mini-PROTEAN Tetra Cell. RNA samples were prepared in Gel Loading Buffer II (Ambion) without heat denaturation and electrophoresis was done at room temperature.

Nuclease sensitivity assay. The nucleases tested were: RNase A, RNase T1 and Turbo DNase (all from Ambion), Xrn1, exonuclease T and exonuclease I (all from NEB). For each assay, 200 ng of an unmodified synthetic negative control siRNA (GenePharma) and *vif* pro-siRNA were used and assays were incubated in a 20- μ l reaction volume using standard amounts of enzymes at 37 °C for 1 h. Treated RNAs were purified by phenol/chloroform extraction followed by isopropanol precipitation.

Tests for endotoxin activity and immune activation of primary human monocyte-derived macrophages. RNA samples diluted in ddH₂O to the indicated concentration were analyzed by the single vial Gel Clot LAL assay (detection limit 0.25 EU/ml, Lonza) following the manufacturer's protocol. Lipopolysaccharide (LPS) from *E. coli* O111:B4 (Sigma-Aldrich) was used as a positive control.

To test for cytokine gene activation, monocyte-derived macrophages plated in 24-well plates (1 \times 10⁵ cells/well) were incubated with medium containing RNA or LPS at the indicated concentration for 4 h before harvesting RNA. siRNAs and pro-siRNAs (20 nM) were also transfected into cells using Lipofectamine 2000 (Invitrogen) and total RNA was harvested 24 h after transfection.

5' ^{32}P labeling of RNA. RNA samples were dephosphorylated by Antarctic Phosphatase (NEB) for 30 min at 37 °C in the presence of Murine RNase Inhibitor (NEB). The Antarctic Phosphatase was deactivated by incubation at 65 °C for 5 min and the RNA was end-labeled with γ - ^{32}P ATP (PerkinElmer) and T4 Polynucleotide Kinase (NEB). Gels were exposed using a phosphorimager screen and visualized using a FLA-9000 Image Scanner (Fujifilm).

Small RNA northern blot analysis. Northern blot analysis for small RNAs was done as previously described³⁵. The *EGFP*-specific sense probe was a ^{32}P -UTP-internally labeled RNA prepared by *in vitro* transcription using T7 RNA polymerase (NEB) and a PCR-generated DNA template of the full-length *EGFP* gene that incorporated a T7 promoter.

siRNA transfection for testing RNA silencing efficiency. All siRNA transfections were done using Lipofectamine 2000 following the manufacturer's protocol. Briefly, cells were plated in 24-well plates (1×10^5 per well) and the transfection complex (containing 1.0 μl Lipofectamine 2000 and siRNAs) was added directly to the medium. RNA and protein samples were isolated from cells 24 h after transfection. For the PLK1 cell killing experiment, cells were counted using a TC-10 automatic cell counter (Bio-Rad). The following siRNAs were used: ON-TARGETplus Non-targeting siRNA #4 (D-001810-04-05, Dharmacon); siGENOME Lamin A/C Control siRNA (D-001050-01-20, Dharmacon); Set of 4: siGENOME LMNA siRNAs (MQ-004978-01-0002, Dharmacon); ON-TARGETplus SMARTpool - Human PLK1 (L-003290-00-0005, Dharmacon); Set of 4 Upgrade: ON-TARGETplus PLK1 siRNA (LU-003290-00-0002, Dharmacon); Set of 4: siGENOME TP53 siRNA (MQ-003329-03-0002, Dharmacon); negative-control siRNA (NC siRNA, B01001, GenePharma); positive-control siRNA TP53 (B03001, GenePharma); custom *EGFP* siRNA (sense, GGCUCAGUCCAGGAGCGCACC; antisense, UGCGCUCUGGACGUAGCCUU); custom *vif* siRNA-1 (ref. 22) (sense, GUUCAGAGUACACAUCCCT; antisense, GGGUUGUACUUCU GAACCTT); custom *vif* siRNA-2 (ref. 23) (sense, CAGAUGGCAGGUGA UGAUUGT; antisense, AAUCAGCACCUGCCAUCUGTT); and custom *gag* siRNA (ref. 24): (sense, GAUUGUACUGAGAGACAGGCU; antisense, CCUGUCUCUCAGUACAAUCUU).

RISC immunoprecipitation. Cells (3×10^6) were transfected with 4 nM NC siRNA or *EGFPFL* pro-siRNAs. After 24 h cells were scraped from the plate in 2 ml lysis buffer (150 mM KCl, 25 mM Tris-HCl pH 7.5, 2 mM EDTA, 0.5 mM DTT, 1% NP-40 and Roche Complete Protease Inhibitor Cocktail). Cells were then mechanically disrupted for 1 min using a micro-MiniBeadbeater (BioSpec). The cell lysate was incubated at 4 °C with rotation for 1 h to ensure complete lysis. Immunoprecipitation was performed by adding anti-Ago (2A8) antibody (Millipore, MABE56) or mouse total IgG (Jackson Labs) at 1:100 dilution together with 30 μl protein G Dynabeads (Invitrogen) and samples were rotated at 4 °C overnight. After washing four times in lysis buffer, precipitated RNAs were isolated using Trizol reagent from 90% of the reaction mix, and 10% was saved for immunoblot input.

Western immunoblot. Protein samples were prepared by heating cells to 95 °C for 5 min in 1 \times SDS loading buffer before SDS-PAGE. Immunoblot was done using SNAP i.d. Protein Detection System (Millipore) following the manufacturer's protocol. Antibodies and their dilutions were: anti-FLAG (M2) 1:1,000 (Sigma-Aldrich, F1804); anti-His tag 1:500 (Covance, MMS-156P); anti-PLK1 1:100 (Santa Cruz, sc-17783); anti-Lamin A/C 1:1,000 (Santa Cruz, sc-7292); anti-p53 (DO-1) 1:500, (Santa Cruz, sc-126); anti-beta-Tubulin 1:10,000 (Sigma-Aldrich, T5168); anti-Ago (2A8) 1:1,000 (Millipore, MABE56). Horseradish peroxidase-conjugated anti-mouse or anti-rabbit IgG secondary antibodies were used at 1:5,000 dilution followed by incubating the membranes in SuperSignal West Pico Chemiluminescent Substrate (Thermo Scientific).

Solution hybridization and native gel electrophoresis assay. DNA oligonucleotides purchased from IDT were PAGE purified. Purified DNA oligonucleotides (10 pmol) were end-labeled with γ - ^{32}P ATP by T4 polynucleotide kinase (NEB) and 2 pmol was then mixed with 5 ng of pro-siRNAs in buffer containing 20 mM Tris-HCl pH 7.9, 100 mM NaCl and 2 mM EDTA. Samples

were heated to 80 °C for 10 min and allowed to cool to room temperature. A fraction of the sample was separated on a native 15% polyacrylamide gel. The gel was directly exposed to a phosphorimager screen. Multi-gauge software (Fujifilm) was used for image quantification.

siRNA library preparation, deep sequencing and data analysis. siRNAs were cloned according to the Illumina small RNA sample preparation guide v1.5 with the following exceptions. Custom 5' RNA ligation adaptors were synthesized with a 4-nt nucleotide barcode sequence (Supplementary Table 8). Small RNA libraries were pooled and sequenced on one sequencing lane of an Illumina GAII sequencer (Genome Technology Core, Whitehead Institute or NEB). Novocraft software was used for sequence alignment. Reference genome was *E. coli* K12 substr. MG1655. We wrote Perl software scripts for data analysis.

mRNA profiling by microarray and deep sequencing. siRNAs and pro-siRNAs (4 nM) were transfected into HeLa-d1EGFP cells and RNA was isolated 24 h after transfection. Nontargeting siRNA #4 (Dharmacon) was used as negative control siRNA. Data from biological duplicates were analyzed at the Microarray Core, Dana-Farber Cancer Institute for microarray analysis using GeneChip 1.0 ST (Affymetrix). Microarray data were analyzed using dChip software and *P* values of gene expression changes were calculated using paired *t*-test method³⁶.

For RNA deep sequencing, a Ribo-Zero rRNA Removal Kit (Epicentre) was used to remove large ribosomal RNAs from total RNA following the manufacturer's protocol. For each condition, two libraries were constructed and sequenced independently from duplicate samples. rRNA-depleted RNA (from 500 ng total RNA) was used to construct a deep sequencing library using NEBNext Ultra RNA Library Prep Kit for Illumina (NEB #E7530) according to the manufacturer's protocol. Illumina GAII was used for sequencing (NEB). Tophat and Cufflinks software suites were used to analyze the RNA deep sequencing data from biological duplicates according to ref. 27. Reference genome was Human genome GRCh37/hg19 and annotations of lincRNA transcripts were downloaded from UCSD genome browser.

Flow cytometry. For EGFP, cells were removed from plates by trypsin digestion and re-suspended in FACS buffer, DPBS (Invitrogen) containing 2% heat-inactivated FBS. Intracellular staining of p24 antigen was performed using an Intracellular Staining Kit (Invitrogen) according to the manufacturer's protocol and fluorescein-labeled p24 antibody (1:200, Beckman Coulter, cat no. KC57-FITC). Fluorescence was analyzed on a FACSCalibur (BD) using FlowJo software (Tree Star).

HIV infection and TZM-bl assay. HeLa-CD4 cells were transfected with 4 nM siRNA or pro-siRNA in 24 well plates (1×10^5 cells/well). Cells were infected 12 h post-transfection with HIV_{IIIIB} (~400 ng/ml p24) and culture medium was changed 12 h post-infection. For HIV_{UG29} U87.CD4.CXCR4 cells were used and for HIV_{IN22} U87.CD4.CCR5 cells were used. Culture medium was collected for TZM-bl assay and RNA was extracted for qRT-PCR 36 h post-infection. TZM-bl cells, plated in 24 well plates (1×10^5 cells/well) 12 h before, were analyzed 24 h after adding culture supernatants by luciferase assay performed using a Luciferase Assay System kit (Promega) following the manufacturer's protocol.

RNase A digestion assay for *E. coli* total RNA. ~2 μg of total *E. coli* RNA were incubated with 1.0 unit of RNase A for 15 min at 37 °C in 1 \times DNase I reaction buffer (NBE) supplemented with 400 mM NaCl. The resulting products were analyzed on a 0.8% agarose gel containing ethidium bromide.

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Erratum: Efficient and specific gene knockdown by small interfering RNAs produced in bacteria

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In the version of this article initially published online, in Figure 4d, the label *EGFPFL* vs NC siRNA has been corrected to read *EGFPFL* vs NC siRNA. The error has been corrected for the print, PDF and HTML versions of this article.