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RNAi to RNAi

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RNA interference (RNAi) is a powerful technique for gene silencing but the mechanisms by which double-stranded RNA (dsRNA) affects target gene activity is still poorly understood. In the Early Edition of the [Proceedings of the National Academy of Sciences](#), Nathaniel Dudley and colleagues from the [University of North Carolina](#) at Chapel Hill describe a method for isolating dsRNA molecules that prevent RNAi and give insights into the [mechanisms involved](#). They co-injected pools of dsRNAs into *Caenorhabditis elegans* embryos and screened for inhibition of RNAi-induced embryonic lethality. This led them to isolate *gfl-1*, a homolog of the human *GAS41* gene, a predicted DNA-binding protein identified by virtue of its amplification in glioblastomas. The authors used their 'RNAi-to-RNAi' assay to test *polycomb*-group genes and found that *polycomb*-like genes *mes-3*, *mes-4* and *mes-6* were also required for RNAi. Furthermore, mutants null for these genes were also RNAi-deficient. Further work will be required to understand the role of these chromatin-binding factors in the mechanisms of RNAi.

References

1. RNA-triggered gene silencing
2. *Proceedings of the National Academy of Sciences*, [<http://www.pnas.org>]
3. University of North Carolina, [<http://www.unc.edu>]
4. Using RNA interference to identify genes required for RNA interference, [<http://www.pnas.org/cgi/doi/10.1073/pnas.062605199>]