



PublisherInfo		
PublisherName	:	BioMed Central
PublisherLocation	:	London
PublisherImprintName	:	BioMed Central

Analysing Xist

ArticleInfo		
ArticleID	:	4378
ArticleDOI	:	10.1186/gb-spotlight-20020116-01
ArticleCitationID	:	spotlight-20020116-01
ArticleSequenceNumber	:	44
ArticleCategory	:	Research news
ArticleFirstPage	:	1
ArticleLastPage	:	2
ArticleHistory	:	RegistrationDate : 2002-01-16 OnlineDate : 2002-01-16
ArticleCopyright	:	BioMed Central Ltd2002
ArticleGrants	:	
ArticleContext	:	130593311

Jonathan B Weitzman

Email: jonathanweitzman@hotmail.com

Inactivation of the X chromosome requires the *Xist* gene, whose product, a noncoding RNA, associates with chromatin on the inactive X chromosome and causes transcriptional silencing. In an Advanced Online Publication from [Nature Genetics](#), Anton Wutz and colleagues at the [Whitehead Institute for Biomedical Research](#) report their analysis of functional domains within the mouse *Xist* RNA (DOI:10.1038/ng820). They used mouse embryonic stem (ES) cells expressing different *Xist* transgenes under the control of a tetracycline-inducible promoter. Deletion of the 5' end completely abolished silencing function, whereas large deletions in the middle and 3' region of *Xist* did not affect *Xist* function. Localization studies indicated that chromosome localization and gene silencing functions are independent and reside in distinct domains of the *Xist* RNA. A conserved sequence in the 5' region is required for silencing, while the localization sequences show little homology and are scattered throughout the gene.

References

1. X-chromosome inactivation: counting, choice and initiation.
2. Nature Genetics , [<http://www.nature.com/ng>]
3. Whitehead Institute for Biomedical Research , [<http://wi.mit.edu>]