PublisherInfo				
PublisherName	:	BioMed Central		
PublisherLocation		London		
PublisherImprintName	:	BioMed Central		

Creating protein folds

ArticleInfo		
ArticleID	:	3756
ArticleDOI	:	10.1186/gb-spotlight-20000901-02
ArticleCitationID	:	spotlight-20000901-02
ArticleSequenceNumber	:	193
ArticleCategory	:	Research news
ArticleFirstPage	:	1
ArticleLastPage	:	2
ArticleHistory	:	RegistrationDate: 2000-09-01OnlineDate: 2000-09-01
ArticleCopyright	:	BioMed Central Ltd2000
ArticleGrants	:	
ArticleContext	:	130591111

William Wells Email: wells@biotext.com

An exon, the basic unit of DNA that gets shuffled around during evolution, has an average coding capacity of 40 amino acids, or roughly half of a small folded protein domain. Exon exchange between homologous proteins can lead to slightly altered proteins, but in the August 29 Proceedings of the National Academy of Sciences Riechmann and Winter ask whether shuffling between unrelated sequences can generate new folds (*Proc Natl Acad Sci USA* 2000, **97**:10068-10073). Their starting material is DNA encoding half of a beta-barrel domain, plus fragmented genomic DNA from *Escherichia coli*. The fusion products that can fold are selected by their resistance to proteolysis; one of these proteins is significantly more stable than the original intact protein. The genomic segments that survive the selection do not share sequence homology with the starting beta-barrel sequence, although beta structure predominates in the final protein products.

References

- 1. DNA shuffling of a family of genes from diverse species accelerates directed evolution.
- 2. Proceedings of the National Academy of Sciences, [http://www.pnas.org/]