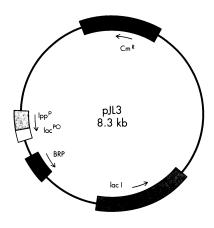


BRP-vectors

for controlled protein release into cell culture medium



The BRP vectors harbor the gene coding for the Bacterial Release Protein (BRP) which makes the inner and outer membrane of *E. coli* permeable. BRP induction with either IPTG (pJL3) or mitomycin C (pSW1) results in the release of cellular proteins into the cell culture medium.

Advantages:

- inducible expression of BRP
- adjustable permeabilization of the E. coli membranes
- controlled protein secretion
- protein release into the cell culture medium from intact cells
- complete system with description and protocols.

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USA: USB Tel: 800-321-9322; 216-765-5000 Japan: Funakoshi Tel: 03-5684-1620, Fax: 03-5684-1620 Circle No. 11 on Readers' Service Card 5) Genic exchange. There has been a tacit assumption that while Africa has been the source of genic exchange of all kinds to other areas, it has somehow remained genetically pristine and isolated. Yet genic exchange involving northern Africa, certainly in Holocene times, across the Straits of Gibraltar and from the Levant, has modified gene pools over much of northern and eastern Africa. To the southeast, Madagascar was settled by sea from Indonesia some 1000 years ago, and there is evidence that this migratory process was reflected in landings on the east African coast, yet another influence on African genomes.

For all these reasons we expect that Africa will have great, even the greatest, diversity at any time in the human past. As both the "out of Africa" and the multiregional hypotheses of modern human origins place the source of those origins in Africa, greater diversity there does not solely support either concept. Only hominid fossil and archaeological data can do that.

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References

- A. G. Thorne and M. H. Wolpoff, Am. J. Phys. Anthropol. 55, 337 (1981); M. H. Wolpoff, W. Xinzhi, A. G. Thorne, in The Origins of Modern Humans: A World Survey of the Fossil Evidence, F. H. Smith and F. Spencer, Eds. (Liss, New York, 1984), pp. 411– 483
- A. C. Wilson and R. L. Cann, Sci. Am. 266, 68 (April 1992).
- A. G. Thorne, in Proceedings of the 8th Panafrican Congress of Prehistory and Quaternary Studies, Nairobi, September 1977, R. E. Leakey and B. A. Ogot, Eds. (TILLMIAP, Nairobi, 1981), pp. 180–181.

Response: Thorne et al. discuss genetic diversity in Africa and the reasons for it, a subject we only touched upon in our article. We do expect greater diversity in Africa than in the rest of the world because this continent has been occupied by modern humans and their immediate ancestors for a longer time. Thorne et al. give time depth of human occupation as the third of five reasons for this expectation, but give four other reasons, of which the first (size), second (environmental diversity), and fifth (genetic exchange) are unconvincing: Asia tends to come before, or to be comparable with, Africa for these three reasons; comparison with New Guinea is not taken from our article. Reason four (central place) is really another facet of reason three (time depth). The real issue, however, is the relevance of our observations to the multiregional evolution hypothesis, which these authors espouse.

This hypothesis was inspired by claims of regional continuity of cranial morphology, which are, however, far from being accepted unanimously by paleoanthropologists. Several articles in (1) have reexamined them independently, and there is a roughly one-to-one split among "pro" and "con" discussants. A recent, thorough reanalysis (2) uses a new morphological approach, reaching conclusions that disagree with the multiregional hypothesis.

What is the relevance of our observations to this hypothesis? Can one, should one, integrate it with that of demic expansions? This would involve Homo sapiens sapiens expanding from Africa and admixing with descendants of H. erectus or archaic H. sapiens in other parts of the world. The gene frequency data we have presented do not indicate a need to postulate such admixtures; but at the same time, they do not exclude them, as we mentioned in our article and explain in more detail (3). Because of the absence of recombination, mitochondrial DNA data are more informative on this issue and have so far not given evidence in favor of the participation of older local inhabitants in the European, East-Asian, or Oceanian gene pools. One cannot, however, say that this did not happen infrequently (4).

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References

- F. H. Smith and F. Spencer, Eds., The Origins of Modern Humans: A World Survey of the Fossil Evidence (Liss, New York, 1984).
- M. Mirazon Lahr, thesis, Cambridge University (1993).
- L. L. Ćavalli-Sforza, P. Menozzi, A. Piazza, History and Geography of Human Genes (Princeton Univ. Press, Princeton, NJ, in press).
- R. L. Cann, in *The Human Revolution*, P. Mellows and C. Stringer, Eds. (Princeton Univ. Press, Princeton, NJ, 1989), pp. 17–30.

Corrections and Clarifications

Reference 4 in the article "Demic expansions and human evolution" by L. L. Cavalli-Sforza et al. (29 Jan., p. 639) was incorrect. It should have read as follows: "F. Weidenreich, Evolution 1, 221 (1947); C. Coon, The Living Races of Man (Knopf, New York, 1965); M. H. Wolpoff, in (2), pp. 62–108."