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700,000 ORESTES

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Now that we have the whole human genome sequence, the challenge remains to identify all the genes and transcripts hidden within it. In the October 9 [Proceedings of the National Academy of Sciences](#), Anamaria Camargo and colleagues, from the [Ludwig Institute for Cancer Research](#) in São Paulo, Brazil, report the results from a Brazilian project aimed at defining the human transcriptome (*Proc Natl Acad Sci USA* 2001, **98**:12103-12108). The approach being used exploits open reading frame expressed sequence tags [ORESTES](#). This differs from the conventional expressed sequence tag (EST) strategy in that it provides sequence information along the whole length of each transcript, rather than just the ends. The method involves low-stringency PCR to produce cDNA libraries, samples of which are then sequenced. Camargo *et al.* generated almost 700,000 ORESTES from 24 types of normal or malignant tissue using 3,540 mini-libraries. They predict that their ORESTES dataset may represent as many as 60% of all human genes (including abundant and rare transcripts). The ORESTES approach generates a larger coverage and a greater number of contigs per gene than to standard EST methods, offering the possibility to complete the closure of most sequences using RT-PCR.

References

1. *Proceedings of the National Academy of Sciences*, [<http://www.pnas.org>]
2. Ludwig Institute for Cancer Research , [<http://www.ludwig.org.br>]
3. Shotgun sequencing of the human transcriptome with ORF expressed sequence tags.