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Mammalian **circadian clocks** are regulated by a series of interacting positive and negative transcription-translation feedback loops in the hypothalamic suprachiasmatic nucleus (SCN). In the April 13 **Science**, Yagita *et al.* describe an *in vitro* model system that resembles the molecular oscillator in the SCN (*Science* 2001, **292**:278-281). Cultured fibroblasts exhibit cycling circadian gene expression when stimulated with the vasoconstricting peptide endothelin-1. Endothelin-1 induced cycling phases of expression of the mammalian clock genes *per1* and *per2*, the cryptochrome gene *Cry1*, and the transcriptional activator *Bmal1*. Experiments with embryonic fibroblasts derived from mice lacking both **mCry1** and **mCry2** genes showed that the *mCry* genes are essential for generating molecular rhythm in fibroblasts, as they are in the SCN clock. Thus, the periodic mRNA expression profiles of clock genes in fibroblasts display key features of an SCN-like timekeeper, providing a model system for exploring circadian rhythms *in vitro*.

References

1. Life's 24-hour clock: molecular control of circadian rhythms in animal cells.
2. *Science*, [<http://www.sciencemag.org>]
3. Mammalian Cry1 and Cry2 are essential for maintenance of circadian rhythms.