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Fast forward to a sensitive test for prion diseases

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Prion diseases such as scrapie, bovine spongiform encephalopathy (BSE) and variant Creutzfeldt-Jakob disease can currently be diagnosed only by post-mortem examination of brain tissue. The trace amounts of prions in the blood of infected individuals remain beyond detection. In the June 14 *Nature*, Gabriela Saborio and colleagues at the [Serono Pharmaceutical Research Institute](http://www.spri.serono.com/) in Geneva, Switzerland, report that minute amounts of prion protein (PrP^{Sc}) can convert large amounts of normal protein (PrP^C) into its faulty state *in vitro* and this could be applied to the detection of the prions in spinal fluid or blood.

The procedure that Saborio *et al.* have developed is conceptually analogous to polymerase chain reaction cycling. Aggregates formed when PrP^{Sc} is incubated with PrP^C are disrupted by sonication to generate multiple smaller units for the continued formation of new PrP^{Sc}. After cyclic amplification, they found that 97% of the protease-resistant PrP present in the sample corresponded to newly converted protein (*Nature* 2001, **411**:810-813).

"Accelerating the prion natural conversion process in the laboratory can compress years of [real] time into a few hours," said Claudio Soto, the senior author of the paper. The method may also provide an opportunity to determine whether PrP^{Sc} replication results in the generation of infectivity *in vitro*.

References

1. Saborio GP, Permanne B, Soto C: Sensitive detection of pathological prion protein by cyclic amplification of protein misfolding. *Nature* 2001, 411:810-813., [<http://www.nature.com/nature>]
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