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the prnp gene provides instructions for making a protein called prion protein (prp), which is active in the brain and several other tissues. although the precise ...

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nature protocols vol 7 pages p1397 - p1409 ... associated publication: sensitive detection of pathological prion protein by cyclic amplification of protein misfolding.

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protocols. in order to provide effective surveillance for cjd and other prion diseases, we strongly recommend that an effort be made to have an autopsy performed in ...

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prion diseases denote a distinct form of infectivity that is based in the misfolding of a self-protein (prp c) into a pathological, infectious conformation (prp sc).

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synthetic human prion protein is a synthetic peptide from the human prion protein. cd230. also known as the prion protein (prp) exists in two alternate forms; a ...

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prion diseases are transmissible, fatal, neurodegenerative diseases in human and animals. the molecular basis of neurodegeneration in prion diseases is largely unclear.

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