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| 1. Record Nr. | UNISA996395868203316 |
| Autore | Cawdrey Daniel <1588-1664.> |
| Titolo | The inconsistencie of the independent way [[electronic resource]] : with scripture, and it self. Manifested in a threefold discourse, I. Vindiciæ vindiciarum, with M. Cotton. II. A review of M. Hookers Survey of church-discipline. The first part. III. A diatribe with the same M. Hooker concerning baptism of infants of non-confederate parents, cap 2. of his third part. / / By Daniel Cawdrey, a member of the assembly, and late preacher at Martins in the Fields |
| Pubbl/distr/stampa | London, : Printed by A. Miller for Christopher Meredith at the sign of the Crane in Pauls Church-yard, MDCLI. [1651] |
| Descrizione fisica | [24], 219, [1] p |
| Soggetti | Congregational churches Baptism Church polity Church discipline |
| Lingua di pubblicazione | Inglese |
| Formato | Materiale a stampa |
| Livello bibliografico | Monografia |
| Note generali | In three parts, each with separate dated title page. Errata: p. [24]. Annotation on Thomason copy: "may: 15". Reproduction of the original in the British Library. |
| Sommario/riassunto | eebo-0018 |

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| 2. Record Nr. | UNINA9910137089803321 |
| Autore | Jean-Luc Vilotte |
| Titolo | Promiscuous functions of the prion protein gene family |
| Pubbl/distr/stampa | Frontiers Media SA, 2015 |
| Descrizione fisica | 1 online resource (113 p.) |
| Collana | Frontiers Research Topics |
| Soggetti | Biology, life sciences |
| Lingua di pubblicazione | Inglese |
| Formato | Materiale a stampa |
| Livello bibliografico | Monografia |
| Sommario/riassunto | <p>The cellular prion protein PrPC is a ubiquitous GPI-anchored protein. While PrPC has been the focus of intense research for its involvement in a group of neurodegenerative disorders known as transmissible spongiform encephalopathies (TSE), much less attention has been devoted to its physiological function. This notably relates to the lack of obvious abnormalities of mice, goat or cattle lacking PrPC. This apparently normal phenotype in these PrPC-deficient animals however contrasts with the very high degree of conservation of the prion protein gene (Prnp) in mammalian species (over 80%), and the presence of genes with similarities to Prnp in birds, reptiles, amphibians and fish. This high conservation together with its ubiquitous expression, - albeit at highest levels in the brain-, suggest that PrPC has major physiological functions. Dissecting PrPC function is further complicated by the occurrence, in mammals, of two potentially partially redundant homologues, Doppel, and Shadoo. The biological overlaps between members of the prion protein family are still under investigation and much debated. Similarly, although in vitro analyses have suggested various functions for PrPC, notably in cell death and survival processes, some have yielded conflicting results and/or discrepancies with in vivo studies. This Research Topic brings together the accumulated knowledge regarding the biological roles of the prion protein family, from the animal to the molecular scale.</p> |