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Large recurrent microdeletions associated with schizophrenia

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#	Paper	IF	Citations
1556	Chromosome abnormalities, mental retardation and the search for genes in bipolar disorder and schizophrenia. 2008 , 14, 113-20		18
1555	Recurrent CNVs disrupt three candidate genes in schizophrenia patients. 2008 , 83, 504-10		220
1554	Rare chromosomal deletions and duplications increase risk of schizophrenia. <i>Nature</i> , 2008 , 455, 237-41	50.4	1251
1553	Copy-number variations associated with neuropsychiatric conditions. <i>Nature</i> , 2008 , 455, 919-23	50.4	513
1552	Schizophrenia: Incriminating genomic evidence. <i>Nature</i> , 2008 , 455, 178-9	50.4	39
1551	50 & 100 Years Ago. <i>Nature</i> , 2008 , 455, 179-179	50.4	
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1549	Personal genomes: The case of the missing heritability. <i>Nature</i> , 2008 , 456, 18-21	50.4	1242
1548	What price personal genome exploration?. 2008 , 26, 1105-8		10
1547	Detection of sharing by descent, long-range phasing and haplotype imputation. 2008 , 40, 1068-75		329
1546	Recurrent reciprocal 1q21.1 deletions and duplications associated with microcephaly or macrocephaly and developmental and behavioral abnormalities. 2008 , 40, 1466-71		457
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