

Journal Information

Journal Title	PSYCHIATRIC GENETICS
Publisher	LIPPINCOTT WILLIAMS & WILKINS
ISSN	0955-8829
E-ISSN	1473-5873
Database	Clarivate
Domain	Health Sciences, Life Sciences, Social Sciences

Classification History

Year	A+	A	B	C
2026		✓	✓	
2025		✓	✓	
2024		✓	✓	
2023		✓	✓	
2021/2022		✓	✓	
2020		✓	✓	
2019		✓	✓	
2018		✓		

Classification: The table above shows all journal classifications recorded for each classification year in the Green Journals database. A journal may have more than one classification in the same year.

Recent Articles

UK-based mental health professionals' perceptions of genetic and environmental risk for psychiatric conditions 2026-10-9 Authors: Helena L. Davies, Jessica Mundy, Karla Mohoric et al. DOI: 10.1097/ypg.0000000000000426
Causal associations between chronic hepatitis B and major depressive disorder in East Asian populations: a bidirectional two-sample Mendelian randomization study 2026-10-9 Authors: Ya-juan Le, Hai-Ying Jin, Li-Li Cai et al. DOI: 10.1097/ypg.0000000000000424
Can neurodevelopmental disorders be part of brain-lung-thyroid syndrome? A case report 2026-10-5 Authors: Selman Yildirim, Cagil Ozyilmaz, Esra Hosoglu et al. DOI: 10.1097/ypg.0000000000000428
The role of PRKRA mutation in antipsychotic-induced extrapyramidal symptoms: a case report and implications for precision psychiatry 2026-10-5 Authors: Mude Jeevan Naik, Simran Arora, Likhita Thippeswamy et al. DOI: 10.1097/ypg.0000000000000430
CLCN6 mutation in an Indian patient with rapidly progressive dementia 2026-10-5 Authors: Manik Inder Singh Sethi, Mude Jeevan Naik, Guru S. Gowda et al. DOI: 10.1097/ypg.0000000000000433

Female-specific associations of glyoxalase 1 polymorphisms with schizophrenia in a Han Chinese case-control study

2026-9-24

Authors: Amy Jing-Wen Yin, Wei-Wei Wang, Wenge Zhang et al.

DOI: 10.1097/ypg.0000000000000425

Disruption of GAD1 protein architecture by a novel missense variant in a consanguineous family with autosomal recessive intellectual disability

2026-9-15

Authors: Eesha Sajjad, Shagufta Naz, Haiba Kaul et al.

DOI: 10.1097/ypg.0000000000000427

Prenatal diagnosis and genetic counseling of a de novo 10q11.22q11.23 duplication associated with a normal development at 12 months of age

2026-9-1

Authors: Xingxing Wang, Lijun Liu, Kai Jin

DOI: 10.1097/ypg.0000000000000432

A de novo 1.62 Mb deletion at 2q34 with nonpenetrant neurodevelopmental phenotype at 12 months

2026-9-1

Authors: Qiu Guo, Xinfang He, Xu Liu et al.

DOI: 10.1097/ypg.0000000000000431

Contrasting outcomes of 16p11.2 microdeletion and microduplication in prenatal diagnosis: phenotypic variability and genetic counseling strategies

2026-7-29 Vol. 36 Issue 3 Pages 151-155

Authors: Qiu Guo, Li Zhang, Yi Zuo et al.

DOI: 10.1097/ypg.0000000000000421