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Journal Information

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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

Evaluation of the risk of renal cell carcinoma and other cancers in individuals with MITF p.E318K in a large single-institution cohort
2026-10-30 Pages jmg-2026-111922
Authors: Leigh Anne Stout, Cindy Hunter, Courtney Schroeder et al.
DOI: 10.1136/jmg-2026-111922

Founder derivative chromosome causing a recessive-like recurrent disorder: der(17)-associated 19p13.3 duplication in the Upper Tigris Basin
2026-10-26 Pages jmg-2026-111825
Authors: Murat Ozturk, Esra Habiloğlu, Burçin Gönüllü Polat et al.
DOI: 10.1136/jmg-2026-111825

Deletion size and background genetic variation shape congenital heart disease phenotypes in 3016 individuals with 22q11.2 deletion syndrome
2026-10-14 Pages jmg-2026-111761
Authors: Jhih-Rong Lin, Daniella Miller, Dana Luong et al.
DOI: 10.1136/jmg-2026-111761

UK consensus recommendations for onward testing for Lynch syndrome in patients with deficient mismatch repair (dMMR) non-colorectal/non-endometrial cancers
2026-10-14 Pages jmg-2026-111623
Authors: Terri Patricia McVeigh, Ruth Armstrong, John Burn et al.
DOI: 10.1136/jmg-2026-111623

Dual impact of uniparental disomy in imprinting and recessive disorders: a case of UPD(15) with unexpected inheritance and literature review
2026-10-14 Pages jmg-2026-111582
Authors: Solène Doppler, Benjamin Durand, Corentin Schwarz et al.
DOI: 10.1136/jmg-2026-111582

Refinement of the UKCGG/CanVIG framework for triage of putative germline variants identified via tumour and ctDNA sequencing: maximising clinical utility and harmonising gene and variant-restricted reporting across UK clinical practice
2026-10-14 Pages jmg-2026-111648
Authors: Terri Patricia McVeigh, Helen Hanson, Katie Snape et al.
DOI: 10.1136/jmg-2026-111648

Clinicopathological, imaging and genetic spectrum of inherited rimmed vacuolar myopathies in a Chinese cohort
2026-10-12 Pages jmg-2026-111585
Authors: Ze-Ling Zheng, Yu-Hua Lin, Qi-Fang He et al.
DOI: 10.1136/jmg-2026-111585

Recurrent helical COL1A1 E678K variant in three families with an EDS-predominant phenotype expanding the clinical and molecular spectrum of COL1-related overlap disorder
2026-10-12 Pages jmg-2026-111532
Authors: Shaimaa Helal, Enas Nasr, Rafael Pacheco et al.
DOI: 10.1136/jmg-2026-111532

Integrating deep-exome sequencing and droplet digital PCR for genetic diagnosis of 142 patients with somatic overgrowth and vascular malformation
2026-10-12 Pages jmg-2026-111671
Authors: Karin Sollander, Anna Hammarsjö, Andrés Metz Erlendsson et al.
DOI: 10.1136/jmg-2026-111671

Somatic panel testing reveals no common alternative oncogenic driver in BRCA1/2 -mutant high-grade serous ovarian cancer with discordant low genomic instability score
2026-10-8 Pages jmg-2026-111819
Authors: Aanum Aanum, Antonio Pereira, Bailey Andrew et al.
DOI: 10.1136/jmg-2026-111819