

## **Dilated Cardiomyopathy Case Reports**

Dilated cardiomyopathy (DCM) is a significant myocardial disorder characterized by **left ventricular dilation and impaired systolic function**, often leading to heart failure, arrhythmias, thromboembolic events, and sudden cardiac death. **Case Reports and Case Series in Cardiology Journal** welcomes the submission of **dilated cardiomyopathy case reports and case series** that present rare clinical scenarios, diagnostic challenges, imaging findings, genetic associations, and therapeutic outcomes.

This dedicated section focuses on clinically meaningful reports involving **idiopathic, familial, viral, ischemic, and secondary dilated cardiomyopathy**, offering valuable educational insights for cardiologists, physicians, and clinical researchers.

We invite manuscripts covering:

- Idiopathic dilated cardiomyopathy
- Familial and genetic DCM cases
- Viral myocarditis progressing to DCM
- Postpartum dilated cardiomyopathy
- Alcohol-induced cardiomyopathy
- Chemotherapy-related cardiomyopathy
- Pediatric dilated cardiomyopathy
- Arrhythmia-associated DCM
- Advanced heart failure due to DCM
- Echocardiography and cardiac MRI findings
- Device therapy (ICD, CRT) case reports
- Long-term outcome and follow-up case series

At **Case Reports and Case Series in Cardiology Journal**, we encourage authors to submit rare, educational, and clinically impactful DCM cases that improve diagnosis, treatment planning, and patient outcomes.

## **Why Publish With Us?**

- Rapid peer-review process
- DOI allocation for every article
- International editorial support
- Global visibility
- Ideal platform for cardiomyopathy case reports

## **Manuscript Submission**

Authors are requested to submit their manuscript by using Online Manuscript Submission

Portal: <https://www.casereportsincardiology.org/submit.html> (or) also invited to submit through the Journal

E-mail Id: [editor@casereportsincardiology.org](mailto:editor@casereportsincardiology.org).

