

**CASE REPORT**

**Case Report: III° atrioventricular block due to fulminant myocarditis managed with non-invasive transcutaneous pacing** [version 2; referees: 2 approved]

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**Abstract**

Fulminant myocarditis is a life-threatening clinical condition. It is the inflammation of myocardium leading to acute heart failure, cardiogenic shock and cardiac arrhythmias. Incidence of fulminant myocarditis is low and mortality is high. Most grievous complications of fulminant myocarditis is mainly cardiac arrhythmias; if there is delay on active management of the patient, it may be fatal. Here, we describe a case of III° atrioventricular block due to fulminant myocarditis that was managed with non-invasive transcutaneous cardiac pacing in the absence of ECMO. The non-invasive transcutaneous pacemaker is a safe, effective and convenient device to revert arrhythmias.
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Competing interests: No competing interests were disclosed.

How to cite this article: Devkota K, Wang YH, Liu MY et al. Case Report: III° atrioventricular block due to fulminant myocarditis managed with non-invasive transcutaneous pacing [version 2; referees: 2 approved] F1000Research 2018, 7:239 (doi: 10.12688/f1000research.14000.2)

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Grant information: The author(s) declared that no grants were involved in supporting this work.

First published: 28 Feb 2018, 7:239 (doi: 10.12688/f1000research.14000.1)
Figure 2

Laboratory measurements. Her lips were cyanosed and she had 90/41mmHg and SPO2 80%.

At the hospital, she again had convulsion for 1–2 minutes and was paced at a rate of 120 bpm. She again had tachycardia and dyspnoea. Both pupils were round, 4mm and reactive. She had no other significant external examination findings.

A lateral chest radiograph was normal. Chest examination showed b/l crackles and irregular heart rate with no murmurs. The abdomen was soft and non-tender. Electrocardiogram (ECG) showed- III° atrio-ventricular (A-V) block; left anterior fascicular block, ST-T changes (Figure 1). She had cardiac arrest three times in the emergency department at 15–20 minutes interval. She was resuscitated with chest compression along with Isoproterenol and adrenaline. Subsequently, her heart rate was maintained in between 70–90 beats/min. Provisional diagnosis was acute fulminant myocarditis with bronchial pneumonia.

The patient was admitted to hospital after explaining the disease condition and prognosis to parents. She was on continuous oxygen, dopamine, diazepam, Immunoglobulin, ceftriaxone, IV fluids on a maintenance dose and nebulization with Ipratropium bromide (250mcg nebulization) along with a high dose of vitamin C (to reduce the risk of myocardial injury), Coenzyme Q 10 (for myocardial protection), fructose diphosphate (to improve cardiac metabolism) and mannitol (to reduce cerebral edema). However, she again had a cardiac arrest. In addition, ECG showed sinus P wave and no QRS with heart rate dropped from 70 to 20bpm. The patient was resuscitated with chest compression, atropine, and adrenaline. Isoproterenol was started at 1.5mcg/kg/min and increased up to 2mcg/kg/min. Subsequently, her heart rate was maintained at 60–70 bpm. Her heart rate again decreased to 30bpm when isoproterenol was discontinued. As there was no extracorporeal membrane oxygenation (ECMO) machine in our hospital and transfer was not possible, the patient was prepped for non-invasive transcutaneous cardiac pacing.

Cardiac pacing was adjusted to 16 mA and rate 90 bpm. The patient’s heart rate was controlled at 80–100 bpm. Her complexion gradually became reddish, cyanosis gradually improved but she had developed eyelid edema. She had passed urine about 130ml twice in 12 hours. Dopamine was increased to 7mcg/kg/min and she was started on furosemide.

At first 24 hours after cardiac pacing, the patient was conscious. She had passed urine 4 times about 700ml. But facial puffiness was still present. Her heart rate was maintained at the rate of 110–130bpm and SPO2 was 96% with oxygen. Chest pacing was reduced to 14 mA, and the frequency was changed to 70 bpm. After 48 hours of pacing, the heart rate was improved to 100–110bpm with few ventricular premature beats. Then pacing was reduced to 12 mA, frequency changed to 60bpm. The pacing current and frequency were gradually slowed down and discontinued. Then, sinus rhythm was established with the heart rate of 100–110bpm with ECG monitoring. The heart rate fluctuated at 80–100bpm with frequent ventricular premature beats. Echocardiography showed left ventricular myocardial wall thickening and thickening of endocardium with left ventricular ejection fraction (LVEF 50%), suggestive of endocardial fibroelastosis (EF). Chest radiograph showed increased lung texture and enlarged cardiac shadow (Figure 2). Captopril, hydrochlorothiazide, and spironolactone were started to reduce cardiac remodeling and to protect heart function. Furosemide was continued.

Abbreviations
A-V block- Atrioventricular block; ECG- Electrocardiogram; ECMO- Extracorporeal Membrane Oxygenation; LVEF- Left ventricular ejection fraction

Introduction
Fulminant myocarditis is a life-threatening clinical condition. It is inflammation of myocardium leading to acute heart failure and cardiac conduction abnormalities with rapid deterioration. There are about 10–38% cases of fulminant myocarditis among all cases of acute myocarditis. Causes of fulminant myocarditis may be of viral, bacterial or non-infectious origin. Diagnosis of fulminant myocarditis is very difficult because of non-specific symptoms and diagnostic tools. There may be signs of acute heart failure, cardiogenic shock, or life-threatening cardiac arrhythmias. Cardiac arrhythmias are varied in presentation, ranging from Sinus arrest, AV block, Ventricular tachycardia and Ventricular fibrillation during acute phase. Here we present a case of fulminant myocarditis presenting with different clinical features and III° A-V block, which was successfully managed with non-invasive transcutaneous pacing.

Case report
A 3 ½ year old female child having a productive cough and 5–6 episodes of the passage of loose stool for 2 days was taken to local hospital after she had sudden convulsion for about 2 minutes. At the hospital, she again had convulsion for 1–2 minutes and her heart rate dropped to 30 bpm. She was given atropine, dexamethasone, dextrose and per rectal choral hydrate at the local hospital (doses not known) and immediately referred to Renmin Hospital. She had no significant past medical history, drug sensitivity or allergies. On arrival she was conscious but lethargic and dyspnoic. Both pupils were round, 4mm and reactive to light. Her vitals were T36.4 °C, HR 62 bpm, RR 65/min, BP 90/41mmHg and SPO2 80%. Table 1 shows the results of routine laboratory measurements. Her lips were cyanosed and she had pale and cold extremities. Neck and throat examination was normal. Chest examination showed b/l crackles and irregular heart rate with no murmurs. The abdomen was soft and non-tender. Electrocardiogram (ECG) showed- III° atrio-ventricular (A-V) block; left anterior fascicular block, ST-T changes (Figure 1). She had cardiac arrest three times in the emergency department at 15–20 minutes interval. She was resuscitated with chest compression along with Isoproterenol and adrenaline. Subsequently, her heart rate was maintained in between 70–90 beats/min. Provisional diagnosis was acute fulminant myocarditis with bronchial pneumonia.

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Table 1. Laboratory investigations from day of admission to discharge.

| Blood Investigations | 0 DOA | 1 DOA | 2 DOA | 3 DOA | 4 DOA | 7 DOA | 12 DOA | 19 DOA | 27 DOA |
|----------------------|-------|-------|-------|-------|-------|-------|--------|--------|--------|
| WBC (4-10×10^9/L)   | 19.86 | ↑     | 20.02 | 17.16 | ↑   | 18.42 | ↑     | 15.61  | ↑     | 12.51  | ↑     | 10.52 | ↑     | 10.01 | ↑     | 8.43 |
| Neutrophils (50–75%)| 68.7  |     | 70.8  | 85.4  | ↑   | 80.47 | ↑     | 47.6   | ↓     | 55.8   | 69.3  | 57.6  | 57.8  | 5    | 5     |
| Lymphocytes (20–40%)| 23.9  |     | 22.4  | 10.4  | ↓   | 14.4  | ↓     | 45.9   | ↑     | 37     | 27    | 35.9  | 35    |     |      |
| Monocytes (3–8%)    | 7.3   |     | 6.2   | 4.1   |     | 4.1   |       | 5.6    |       | 5      | 3.1   | 5.6   | 5     |      |      |
| Eosinophils (0.5–5%)| 0     |     | 0.5   | 0     |     | 0     |       | 0.6    |       | 1.6    | 0.4   | 0.6   | 1.6   |      |      |
| Basophils (0–1%)    | 0.1   |     | 0.1   | 0.1   |     | 0.1   |       | 0.1    |       | 0.6    | 0.2   | 0.3   | 0.6   |      |      |
| ANC (2–7.5×10^9/L)  | 13.65 | ↑   | 14.1  | 14.67 | ↑   | 13.67 | ↑     | 7.42   | ↑     | 3.89   | 7.83  | 1.68  | 3.16  | 2.86 |      |
| ALC (0.8–4×10^9/L)  | 4.74  | ↑   | 2.22  | 1.78  |     | 1.88  |       | 7.16   | ↑     | 5.86   | 1.68  | 3.16  | 2.86  |      |      |
| Hemoglobin (110–170 g/L) | 110 | ↓   | 107   | 95    | ↓   | 97    |       | 108    | ↓     | 110    | 116   | 122   | 129   |      |      |
| Platelets count (100–300×10^9/L) | 203  | 200  | 143   | 132   | 403  | 257   | 311    | 403    | 257    |      |      |      |      |      |
| ESR (0–15 mm in 1 hr)| 1     |     |       |       |     |       |       |        |       |        |       |      |      |      |      |
| Blood Glucose (3.89–6.11 mmol/L) | 11.2 | ↑   | 6.1   | 3.8   | ↓   | 4.6   |       | 4.2    |       | 7.2    | 5.3   |      |      |      |      |
| Potassium (K) (3.5–5.4 mmol/L) | 4.3  | ↓   | 5.18  | 4.46  |     | 3.5   |       | 3.52   |       | 4.13   | 4.4   |      |      |      |      |
| Sodium (Na) (135–148 mmol/L) | 131  | ↓   | 138   | 129   | ↓   | 133.8 | ↓     | 134.6  | ↓     | 141    | 142   |      |      |      |      |
| Calcium (Ca) (2.05–2.55 mmol/L) | 1.58 | ↓   | 1.6   | 1.62  | ↓   | 2.27  |       | 2.23   |       | 2.38   |      |      |      |      |      |
| Blood Urea (1.8–7.1 mmol/L) | 21.35| ↑   | 21.45 | 21.79 | ↑   | 6.88  |       | 4.98   |       | 5.29   | 4.1   |      |      |      |      |
| Creatinine (44–106 umol/L) | 158.5| ↑   | 167   | 211.6 | ↑   | 63.5  |       | 57.5   |       | 49.6   | 50    |      |      |      |      |
| Uric Acid (129–417 umol/L) | 977  | ↑   | 980   | 935   | ↑   | 274   |       | 253    |       | 169    |      |      |      |      |      |
| ALT (8–40 U/L)       | 8526  | ↑   | 6589  | 1406  | ↑   | 552   | ↑     | 52     | ↑     | 27     |      |      |      |      |      |
| AST (5–40 U/L)       | 4724  | ↑   | 3245  | 2319  | ↑   | 309   | ↑     | 309    | ↑     | 31     |      |      |      |      |      |
| αHBDH (72–182 IU/L)  | 3895  | ↑   | 2145  | 1061  | ↑   | 631   | ↑     | 489    | ↑     | 192    | ↑     |      |      |      |      |
| ALP (40–150 IU/L)    | 151   | ↑   | 144   | 140   | ↑   | 127   |       | 126    |       | 141    |      |      |      |      |      |
| γGTP (5–54 U/L)      | 26    |     | 28    | 28    |     | 58    | ↑     | 53     |       | 48     |      |      |      |      |      |
| LDH (100–300 IU/L)   | 10140 | ↑   | 9876  | 2155  | ↑   | 746   | ↑     | 456    | ↑     | 223    | 145   |      |      |      |      |
| Total Protein (60–85 g/L) | 56.8 | ↓   | 60.4  | 63.6  |     | 66.8  |       | 72.8   |       | 72.7   |      |      |      |      |      |
| Albumin (35–55 g/L)  | 33.5  | ↓   | 33    | 32    | ↓   | 39    |       | 43     |       | 41.1   |      |      |      |      |      |
| Globulin (20–35 g/L) | 23.3  |     | 26.9  | 31.6  |     | 34    |       | 34     |       | 31.6   |      |      |      |      |      |
| Creatine Kinase (25–200 IU/L) | 1170 | ↑   | 1245  | 1679  | ↑   | 109   | 89    | 39     |       | 35     |      |      |      |      |      |
| CKMB (0–25 U/L)      | 247   | ↑   | 187   | 104   | ↑   | 48    | ↑     | 48     | ↑     | 10     | 13    |      |      |      |      |
| Troponin T (0–0.08 ng/ml) | 0.361| ↑   | 0.024 | 0.018 |     | 0.024 |       | 0.018  |       |      |      |      |      |      |      |
| ASO Titre (0–166 IU/ml) | 7    |     |       |       |     |       |       | 24     |       |      |      |      |      |      |      |
| CRP (0–10 mg/L)      | 0.9   |     |       |       |     |       |       | 0.1    |       |      |      |      |      |      |      |

| Atrial Blood Gas | PH (7.35 – 7.45) | 7.34 | ↓ | 7.39 | ANC: Absolute Neutrophil count |
|                 | PaO2 (80–100 mmHg) | 82.1 | 88 |       | ALC: Absolute Lymphocytes cont |
|                 | PaCO2 (35–45 mmHg) | 23  | 36 |       |                                        |
|                 | HCO3 (22–26 mEq/L) | 13.3  | 24.2 |       |                                        |
|                 | Anion Gap (10–15 mEq/L) | 31  | 15.2 |       |                                        |

| Stool Routine       | Normal |                                        |
| Urine routine       | Normal |                                        |
| Mycoplasma titer    | Negative |                                   |
Figure 1. (A) Electrocardiogram (ECG) at emergency showing III atrio-ventricular block; left anterior fascicular block, ST-T changes; (B) ECG recording during transcutaneous pacing; (C) ECG at the time of discharge, which is normal.

Figure 2. (A) Patient on non-invasive transcutaneous pacing; (B) Echocardiography after 48 hours of admission showing left ventricular myocardial wall thickening and thickening of endocardium; (C) Chest X ray showing increased lung texture and enlarged cardiac shadow.
Mannitol was stopped after the patient’s MRI scan revealed normal findings.

The patient’s HR was in between 80–100 bpm, with blood pressure was increasing gradually. Dopamine was tapered and stopped at 72 hours, after her BP reached 110/78 mmHg. The chest became gradually clear and her heart sounds were also normal. ECG monitoring also showed improvement with decreased numbers of premature beats and gradual change of S-T segments towards normal.

The patient was discharged on the 28th day after admission after her all routine investigations returned to normal (Table 1). ECG showed sinus rhythm with heart rate 102 bpm.

Echocardiography showed normal cardiac chambers, normal wall motion with EF 60%. Her final diagnosis was fulminant myocarditis with III° A-V block and bronchial pneumonia. On discharge, the patient was advised to continue captopril 6.25mg bid, metoprolol succinate 6.25mg bid, prednisone 1mg / kg orally for six months.

At six month follow up the patient’s echocardiography had returned to normal with LVEF 65%, and prednisone was reduced to 0.5mg/kg orally for 15 days and with a tapering dose for the next 15 days. After one year follow-up, she had no complaints and no significant abnormalities noted on echocardiography.

All doses of medications can be seen in Table 2.

| Table 2. List of medications, including doses and duration, given to the patient during hospital admission. |
| --- |
| **Medicine** | **Doses** | **Route/duration** |
| Atropine | 0.25mg | IV When required |
| Adrenaline | 0.2mg | IV When required |
| Isoproterenol | 0.2mcg | IV- bolus at ER |
| 0.15mcg/kg/min | IV in 50ml of 5% glucose |
| 0.2mcg/kg/min | IV in 50ml of 5% glucose |
| Dopamine | 3–5mcg/kg/min | IV in 50ml of 5% glucose |
| Diazepam | 0.5mg/kg | IV when required |
| Phenobarbital | 2mg/kg | IV when required |
| Mannitol | 42 ml | IV 6 hourly for 2 days from DOA |
| 42 ml | IV 8 hourly for next 2 days |
| 42ml | IV 12 hourly for next 2 days then stop |
| Fructose diphosphate | 3.4g /OD | IV for 10 days |
| Ceftriaxone | 100mg/kg/day | IV 12 hourly from DOA for 10 days |
| Piperacillin tazobactam | 1.125gm/day | IV 12 hourly from 3 DOA for 10 days |
| Immunoglobulin | 5 gm | IV daily for 5 days |
| Methylprednisolone | 1.5mg/kg/day | IV for 5 days |
| prednisone | 10 mg / OD | PO from 6 DOA and on discharge also |
| Captopril | 6.25mg / BID | PO from 3 DOA and on discharge also |
| Spironolactone | 10mg/OD | PO from 5 DOA and on discharge also |
| Furosemide | 10mg | IV 12hourly from 2nd DOA to 5DOA |
| Hydrochlorothiazide | 10mg | PO 12 hourly from 5 DOA till discharge |
| Coenzyme Q10 | 5 mg | PO 8hourly from 2nd DOA till discharge |
| Vitamin C | 3 gm | IV 12 hourly from 2nd DOA till discharge |
Discussion
In children, sometimes myocarditis is self-limiting. However, if it progresses there is the risk of acute cardiac failure, hemodynamic disturbances, and arrhythmias leading to significant morbidity and mortality. Mortality due to myocarditis for infants is more than 75%, whereas for children it is more than 25%. There is no any specific clinical course and investigations to diagnose fulminant myocarditis. Initially, they present with flu-like symptoms and later develop sudden onset of cardiac symptoms that rapidly deteriorate. Neonates may present with fever, poor feeding, and listlessness and sometimes with danger signs like apnea, episodic cyanosis, and diaphoresis. Older children present with respiratory or gastrointestinal symptoms. Among them only a few present with chest pain. Diagnosis is mainly done on the basis of: Clinical presentation, blood profile, including CBC, electrolytes, creatinine kinase, creatine kinase MB isoenzyme, C-reactive protein, Troponin T, Troponin I, antistreptolysin O titer, polymerase chain reaction to detect viral antigens, autoantibodies marker, liver enzymes, ECG, Echocardiography, ultrasonography and even Cardiac MRI which are mostly supportive. If echocardiography shows low LVEF in children with fulminant myocarditis, the prognosis is poor. Mortality in fulminant myocarditis is mainly due to cardiac arrhythmias among which structural changes, parameters of ventricular dynamics and vascular changes are responsible for the increased incidence. Acute fulminant myocarditis, if properly and aggressively treated, has excellent long-term survival even if the patient may present with severe hemodynamic compromise. Complete heart block on initial ECG may also have an excellent prognosis, although mechanical assistance may be warranted as shown in the study by Lee E Y et al. This can be managed with percutaneous cardiopulmonary support, ECMO, intra-aortic balloon pumping, or the ventricular assisted device. ECMO remains an effective approach in children for the management of acute fulminant myocarditis. In addition, intravenous immunoglobulin and high dose steroids help to reduce inflammation. Patients managed with immunoglobulin, steroids or mechanical support for fulminant myocarditis may have higher survival rate compared to those not receiving these therapies.

In the present case, we tried to manage initially with Isoproterenol but were unsuccessful. So we applied the non-invasive transcutaneous pacemaker to revert the A-V block. Pads or electrodes detachment, patient non-cooperation and skin-burn due to high voltage electric current are its limitations. In contrast, intraventricular cardiac pacing is time-consuming; much more risky and surgical site wound infection is common. Beland et al, Kelly et al in their articles note that non-invasive transcutaneous pacemaker is the safe, effective and suitable equipment for children.

Conclusion
Acute fulminant myocarditis is a grievous condition with high morbidity and mortality. No delay should be had on starting immunoglobulin and steroids if suspected. If there is the arrhythmia, the patient should be immediately started on ECMO, percutaneous cardiopulmonary support or ventricular assisted device. If these are not available, then non-invasive transcutaneous cardiac pacing must be started, which is a safe, convenient and cost-effective device to revert arrhythmias caused by myocarditis.

Consent
We have taken written informed consent from the child’s legal guardian (her father) to use and publish his child’s medical case history and any accompanying images.

Data availability
All data underlying the results are available as part of the article and no additional source data are required.

Competing interests
No competing interests were disclosed.

Grant information
The author(s) declared that no grants were involved in supporting this work.

Acknowledgements
We wish to thank Dr. Wu Bing, (Resident Cardiology Department II, Renmin Hospital), entire doctor and nursing staffs Department of Pediatrics I, Renmin Hospital and to the child parents for the consent.

References
1. Lee EY, Lee HL, Kim HT, et al: Clinical features and short-term outcomes of pediatric acute fulminant myocarditis in a single center. Korean J Pediatr. 2014; 57(11): 489–495. PubMed Abstract | Publisher Full Text | Free Full Text
2. Lobo ML, Taguchi Â, Gaspar HA, et al: Fulminant myocarditis associated with the H1N1 influenza virus: case report and literature review. Rev Bras Ter Intensiva. 2014; 26(3): 321–326. PubMed Abstract | Publisher Full Text | Free Full Text
3. Vashist S, Singh GK: Acute myocarditis in children: current concepts and management. Curr Treat Options Cardiovasc Med. 2009; 11(3): 383–391. PubMed Abstract | Publisher Full Text
4. Ichikawa R, Sumitomo N, Komori A, et al: The follow-up evaluation of electrocardiogram and arrhythmias in children with fulminant myocarditis. Circ J. 2011; 75(4): 932–8. PubMed Abstract | Publisher Full Text
5. Forcada P, Beigelman R, Milei J: Inapparent myocarditis and sudden death in pediatrics. Diagnosis by immunohistochemical staining. Int J Cardiol. 1996; 56(1): 53–7. PubMed Abstract | Publisher Full Text
6. Maron BJ, Gohman TE, Aeppli D: Prevalence of sudden cardiac death during competitive sports activities in Minnesota high school athletes. J Am Coll Cardiol. 1998; 32(7): 1881–4. PubMed Abstract | Publisher Full Text
7. Neuspiel DR, Kuller LH: Sudden and unexpected natural death in childhood and
adolescence. JAMA. 1985; 254(10): 1321–5.

8. Topaz O, Edwards JE: Pathologic features of sudden death in children, adolescents, and young adults. Chest. 1985; 87(4): 476–82.

9. Niimura I, Maki T: Sudden cardiac death in childhood. Jpn Circ J. 1989; 53(12): 1571–80.

10. Levine MC, Klugman D, Teach SJ: Update on myocarditis in children. Curr Opin Pediatr. 2010; 22(3): 278–83.

11. Pei L, Yang N, Yang YH, et al.: Clinical features and prognostic factors in children with fulminant myocarditis. Zhongguo Dang Dai Er Ke Za Zhi. 2015; 17(11): 1232–6.

12. Klein RM, Vester EG, Brehm MU, et al.: [Inflammation of the myocardium as an arrhythmia trigger]. Z Kardiol. 2000; 89 Suppl 3: 24–30.

13. McCarthy RE 3rd, Boehmer JP, Huban RH, et al.: Long-term outcome of fulminant myocarditis as compared with acute (nonfulminant) myocarditis. N Engl J Med. 2000; 342(10): 690–5.

14. Saji T, Matsuura H, Hasegawa K, et al.: Comparison of the Clinical Presentation, Treatment, and Outcome of Fulminant and Acute Myocarditis in Children. Circ J. 2012; 76(5): 1222–1228.

15. Taoka M, Shiono M, Hata M, et al.: Child with fulminant myocarditis survived by ECMO Support--report of a child case. Ann Thorac Cardiovasc Surg. 2007; 13(1): 60–4.

16. Ning B, Zhang C, Lin R, et al.: Local experience with extracorporeal membrane oxygenation in children with acute fulminant myocarditis. PLoS One. 2013; 8(12): e82258.

17. Tucker CE, Fernandez MJ, Morrison CC: Successful Treatment of Fulminant Myocarditis. J Clin Case Rep. 2013; 3: 256.

18. Beland MJ, Heslefin PS, Finlay CD, et al.: Noninvasive transcutaneous cardiac pacing in children. Pacing Clin Electrophysiol. 1987; 10(6): 1262–1270.

19. Kelly JS, Royster RL, Angert KC, et al.: Efficacy of noninvasive transcutaneous cardiac pacing patients undergoing cardiac surgery. Anesthesiology. 1989; 70(5): 747–51.
Open Peer Review

Current Referee Status:  

Version 2

Referee Report 26 March 2018

doi:10.5256/f1000research.15664.r32382

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Authors have answered the questions raised by the reviewer in an appropriate way

Competing Interests: No competing interests were disclosed.

Referee Expertise: Myocarditis, cardiomyopathies, heart failure

I have read this submission. I believe that I have an appropriate level of expertise to confirm that it is of an acceptable scientific standard.

Version 1

Referee Report 14 March 2018

doi:10.5256/f1000research.15664.r32381

Mani Prasad Gautam
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The Authors have discussed an interesting case of fulminant myocarditis in a child which have been managed with transcutaneous pacing for the third degree AV block along with other supportive care including intravenous methylprednisolone and immunoglobulin. In overall the article is at satisfactory level in its scientific contents and usefulness to other practitioners. Some improvement has been seen after the first review and have adequate standard of scientific publication.

Competing Interests: No competing interests were disclosed.

I have read this submission. I believe that I have an appropriate level of expertise to confirm that it is of an acceptable scientific standard.
Sabine Pankuweit  
Department of Cardiology, Angiology and Intensive Care, Philipps University of Marburg, Marburg, Germany

The authors described a case of III° atrioventricular block due to fulminant myocarditis that was managed with non-invasive transcutaneous cardiac pacing in the absence of ECMO. In this case the non-invasive transcutaneous pacemaker was a safe, effective and convenient device to revert arrhythmias.

Nevertheless, as the authors stated, if there is the arrhythmia, the patient should be immediately started on ECMO, percutaneous cardiopulmonary support or ventricular assisted device. Authors should specify and discuss, what the most important arrhythmias with negative prognosis to what extent are.

Are there any efforts made in this case, to get information on the etiology of the disease (e.g. viral, autoimmune), was there any discussion on an endomyocardial biopsy.

Data in children with regard to the prognosis and outcome in fulminant myocarditis should be added.

Is the background of the case’s history and progression described in sufficient detail?  
Yes

Are enough details provided of any physical examination and diagnostic tests, treatment given and outcomes?  
Partly

Is sufficient discussion included of the importance of the findings and their relevance to future understanding of disease processes, diagnosis or treatment?  
Partly

Is the case presented with sufficient detail to be useful for other practitioners?  
Partly

**Competing Interests:** No competing interests were disclosed.

**Referee Expertise:** Myocarditis, cardiomyopathies, heart failure

I have read this submission. I believe that I have an appropriate level of expertise to confirm that it is of an acceptable scientific standard.

Referee Report 05 March 2018

do:10.5256/f1000research.15216.r31290

Mani Prasad Gautam  
Department of Medicine, Bharatpur Hospital, Chitwan, Nepal
The Authors have discussed an interesting case of fulminant myocarditis in a child which have been managed with transcutaneous pacing for the third degree AV block along with other supportive care including intravenous methylprednisolone and immunoglobulin. In overall the article is at satisfactory level in its scientific contents and usefulness to other practitioners. But there is some space to improve its rating. The background information provided here is an overall information, it would have been better if the author has also focused on various arrhythmias associated with fulminant myocarditis as the myocarditis tends to have various refractory arrhythmias including ventricular tachycardia. It would have been better if the authors were focused on particular theme, such as on various arrhythmias and their medical management or overall medical management. The case description is somehow lacking smoothness in the flow of information. That could be because lack of cohesion in sentences and writing in the form of clinical vignette can improve its content. Similarly there are a lot of English grammar related errors and rampant use of short form that should be corrected. In addition, it is advised to use scientific words rather than layman terms in academic writings. Taking help from an English language expert might also improve its contents. The case report section should be made little bit short. The discussion section should be more descriptive by discussing clinical features of myocarditis including various arrhythmias and medical management of myocarditis. The use of Vitamin C, Coenzyme Q10, Fructose diphosphate and Mannitol can not be understood from the discussion section which I was expecting there. I hope, the revision after considering above advice will improve the scientific content of the article.

References
1. Gautam M, Sogunuru G, Subramanyam G, Viswanath R: Tuberculous myocarditis presenting as a refractory ventricular tachycardia of biventricular origin. Journal of College of Medical Sciences-Nepal. 2012; 7 (2). Publisher Full Text

Is the background of the case's history and progression described in sufficient detail?
Partly

Are enough details provided of any physical examination and diagnostic tests, treatment given and outcomes?
Yes

Is sufficient discussion included of the importance of the findings and their relevance to future understanding of disease processes, diagnosis or treatment?
Partly

Is the case presented with sufficient detail to be useful for other practitioners?
Yes

Competing Interests: No competing interests were disclosed.

I have read this submission. I believe that I have an appropriate level of expertise to confirm that it is of an acceptable scientific standard, however I have significant reservations, as outlined above.
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