Magnetocardiography-Guided Management of an Unusual Case of Isoimmune Complete Atrioventricular Block Complicated by Ventricular Tachycardia

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Cardiac manifestations of immune-mediated damage include transient incomplete AV block [7], junctional ectopic tachycardia [8], endocardial fibroelastosis [9, 10], and cardiomyopathy [11, 12]. However, even in large series of isoimmune AV block, ventricular tachycardia (VT) is rarely diagnosed [13]. Whether this is because VT is not a common manifestation of disease, or VT has been incorrectly diagnosed by M-Mode/Doppler technology, is not known. We had the opportunity to use fMCG to diagnose VT in a fetus with isoimmune AV block, and followed this infant after delivery.

Key Words
Arrhythmia · AV block · Magnetocardiography

Abstract
A fetus who was diagnosed at 25 weeks of gestation with isoimmune AV block presented at 34 weeks with a precipitous fall in ventricular rate and periods of tachycardia. Magnetocardiography revealed the tachycardia to be ventricular. After delivery, nonsustained ventricular tachycardia continued. The baby then successfully paced, and at higher ventricular rates the tachycardia resolved. Five years later the child has normal ventricular function and is doing well.

The technique of fetal magnetocardiography (fMCG) has elucidated important features of fetal arrhythmias and has become an important tool in guiding in utero management of these fetuses [1–5]. Complete atrioventricular (AV) block is the most common finding in affected fetuses exposed to maternal SSA/Ro and SSB/La ribonuclear proteins (isoimmune AV block) [6], but other

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This research was supported by NIH grants R01HL63174 (R.T.W.) and R21HL094922 (J.F.S.).
hours after the ultrasound revealed third-degree AV block with a ventricular rate of 63 and an atrial rate of 120 bpm. The fetal heart was structurally normal with good biventricular systolic function and no signs of heart failure or endocardial fibroelastosis. The mother was immediately started on Dexamethasone 4 mg orally once a day.

At 28 weeks, an fMCG was performed in the Department of Medical Physics at the University of Wisconsin in Madison using a 37-channel biomagnetometer in a specially constructed low-permeability room, as previously described \[2\]. Atrial and ventricular rates, QRS duration and QT and PR intervals were measured from the signal-averaged fMCG. 5–10 tracings each of 10–15 min duration were recorded from multiple probe positions, then digitized and band-pass filtered (1–80 Hz). Fetal heart rate accelerations were correlated with fetal movement by simultaneous actocardiography. This initial fMCG showed a reactive heart rate pattern (fig. 1A) and no other arrhythmia (table 1).

Weekly sonograms from 26–33 weeks gestation showed no change in the heart rate or cardiac function. At 34 weeks, the fetal ventricular rate dropped to 48 bpm. A second fMCG showed a non-reactive heart rate pattern (fig. 1B) and several episodes of VT at rates of 85–180 bpm (fig. 2, table 1). Because of the findings of VT and the poor heart rate variability, the infant was delivered that same day by cesarean section. The postnatal 12-lead electrocardiogram recorded at 2 h of age revealed a ventricular rate of 45 bpm, an atrial rate of 150 bpm and third-degree AV block (QTc of 442 ms). During continuous monitoring in the nursery, episodes of non-sustained VT were noted. Isoproterenol failed to increase the ventricular rate >50 bpm, so a temporary transvenous pacing catheter was placed. Despite an increased heart rate, over the next several hours the infant developed hypotension and decreased urine output. Adrenal insufficiency was suspected and later confirmed by low serum cortisol levels. The infant stabilized after solumedrol infusion. On day 10, an epicardial dual chamber pacemaker was implanted. The baby was discharged home in good condition on day 17. She was treated with hydrocortisone for 2 months. Five years later, the child is healthy, with normal ventricular function. Based on numerous ambulatory ECGs, VT has not recurred.

### Table 1. Magnetocardiography results at 28 and 34 weeks in a fetus with isoimmune complete AV block

<table>
<thead>
<tr>
<th>GA weeks</th>
<th>Atrial rate, bpm mean (range)</th>
<th>Ventricular rate, bpm mean (range)</th>
<th>Reactive HR pattern</th>
<th>ectopy %</th>
<th>VT %</th>
<th>QTc ms</th>
</tr>
</thead>
<tbody>
<tr>
<td>28</td>
<td>130 (120–155)</td>
<td>63 (60–96)</td>
<td>yes</td>
<td>6.7</td>
<td>none</td>
<td>484</td>
</tr>
<tr>
<td>34</td>
<td>130 (118–160)</td>
<td>50 (49–59)</td>
<td>no</td>
<td>3.1</td>
<td>0.1</td>
<td>574</td>
</tr>
</tbody>
</table>

**Fig. 1.** Fetal heart rate patterns of acceleration. Each individual black line is an R-R interval. The graph shows the heart rate over 5 min. **A** At 28 weeks, the tracing shows good variability over a wide ventricular rate range. **B** At 34 weeks, not only has the ventricular rate dropped precipitously, but there is no variability whatsoever in the ventricular rate range. The tall lines represent ventricular ectopy.
Discussion

We report the unusual occurrence of isoimmune complete AV block and VT diagnosed by fMCG after a dramatic decrease in the ventricular rate in a 34 week fetus. Other unusual features are that the occurrence late in gestation after previous fMCG showed only third-degree AV block and that the VT occurred 9 weeks after the initiation of transplacental anti-inflammatory treatment. These findings led to an immediate operative delivery of a non-hydropic infant who was paced shortly thereafter and has survived with normal ventricular function.

The most common technique for the diagnosis of fetal arrhythmias evaluates the AV relationship and atrial and ventricular rates by M-Mode or Doppler echocardiography. However, echocardiography measures the mechanical effect of an electrical event rather than direct recordings of cardiac electrical impulses. In addition, diagnosis is difficult if the arrhythmia is transient and since measurements of conduction intervals are necessary for diagnosis (such as long QT syndrome or bundle branch block). Recently, the use of tissue velocity imaging, which allows precise timing of atrial and ventricular events, has increased the sophistication of echocardiography arrhythmia diagnosis [14], but conduction intervals other than...
the P-R interval can only be measured by fMCG or fetal electrocardiography.

Fetal electrocardiogram, which has the potential to be more easily available and less expensive than fMCG, remains limited in arrhythmia diagnosis because the maternal electrocardiogram voltage potential is 10–100 times stronger than the fetal signal resulting in significant artifact. Additionally, useful signals from the fetus are difficult to obtain after 27 weeks, probably due to insulating properties of vernix caseosa [15]. Such problems are not encountered with fMCG, which reliably measures signals from 20–40 weeks of gestation, provided that the recordings are made within a magnetically shielded room.

In the fetus reported here, it is unknown why the ventricular rate dropped abruptly in only 1 week’s time. It may be that despite the anti-inflammatory treatment there was on-going damage to the conduction system, resulting in a slower ventricular pacemaker more distal from the AV node. The finding that the QRS duration was narrow at this lower rate suggests that the new ventricular focus was proximal to the bundle of His.

The occurrence of VT accompanying isoimmune AV block in the fetus is unusual. VT may be an escape rhythm due to the low ventricular rate, or it may be due to myocardial ischemia secondary to low cardiac output from the abrupt fall in heart rate. Both may further be affected by progressive autoantibody-mediated myocardial injury. We further speculate that the findings of VT with severe bradycardia may be one of the mechanisms of sudden death in the fetus with isoimmune disease. These findings underscore the importance of vigilant monitoring in the third trimester of fetuses with isoimmune disease and no heart failure, even if they seem clinically stable.

References

