Pulse oximetry screening for critical congenital heart disease in term and late pre-term newborn

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Abstract

Introduction: Neonatal screening for critical congenital heart disease (CCHD) aids in identifying infants for whom early diagnosis and therapy are beneficial.

Objective: To study utilisation of pulse oximetry in term and late preterm neonates for early diagnosis of CCHD.

Method: This prospective study was conducted in newborns in the Kanuni Sultan Suleyman Education and Research Hospital, Turkey from December 2012 to August 2013. Pre-ductal and post-ductal pulse oximetry was carried out on all neonates between 24-72 hours of age. Screening was considered to be positive when the oxygen saturation level was ≤ 95% or if the pre-ductal and post-ductal saturations, taken after 3 hours from the initial measurements, differed by more than 3%.

Results: There were 971 (97%) babies with negative screening and 29 (3%) babies with positive screening. The sensitivity was 93%, specificity 99%, positive predictive value 81% and negative predictive value 99%.

Conclusions: Pulse oximetry screening in term and late preterm neonates is effective for early diagnosis of CCHD.

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(Key words: Critical congenital heart disease, early detection, pulse oximetry)

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Introduction

Reported prevalence of congenital heart disease (CHD) at birth is 9 per 1000 live births with around 25% being considered critical1. Neonates with such unrecognized critical congenital heart disease (CCHD) can manifest with growth retardation, heart failure, cardiovascular collapse, hypoxic-ischemic encephalopathy and even death. Early detection has led to a decline in poor postoperative outcome due to CCHD2-3.

Though antenatal ultrasonography usually detects major CHD, they can be missed, especially in developing countries, because of the absence of routine antenatal visits4. Pulse oximetry is a non-invasive tool used to detect CCHD5,6. American Academy of Pediatrics recommends pulse oximetry screening of newborns in the case of potential CHD7. Neonatal screening for CCHD aids in identifying infants for whom early diagnosis and therapy are beneficial.

Objective

To study utilisation of pulse oximetry in term and late preterm neonates for early diagnosis of CCHD.

Method

This prospective study was conducted on newborns in the Kanuni Sultan Suleyman Education and Research Hospital, Turkey from December 2012 to August 2013. Demographic data was obtained from medical records. The study was approved by the local ethics committee. Written informed consent was received from all parents/guardians. Patients were included in the study if they were full-term or late preterm newborns. Exclusion criteria included neonates with prenatal diagnosis of CHD and co-morbidities such as preterm delivery, respiratory distress syndrome, dysmorphic syndromes, multiple congenital anomalies, and sepsis.

Pre-ductal and post-ductal pulse oximetry was done in all neonates between 24-72 hours of age by placing sensor on right wrist and right foot. Saturations were recorded one minute after a stable reading. Values greater than 95% were considered normal. Screening was considered to be positive when the oxygen saturation level was ≤95% or if the pre-ductal and post-ductal saturations, taken after 3 hours from the initial measurements, differed by more than 3%. In such instances an echocardiogram was done, without regard to any
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clinical signs. The authors phoned the parents with regard to the status of their child at 3 months after birth if the newborn had negative screening.

CCHD was defined as CHD needing surgery or catheter intervention during infancy, including D-transposition of great arteries, coarctation of aorta, Fallot’s tetralogy, hypoplastic left heart syndrome, pulmonary atresia, hypoplastic or interrupted aortic arch, double outlet right ventricle and truncus arteriosus. The monitoring was conducted using Nellcor Puritan Bennett N-560 pulse oximeters.

Number Cruncher Statistical System 2007 Statistical Software (Utah, USA) was used to calculate frequencies and percentages. Median and interquartile range was used to describe patient age. Chi-square test and Fisher’s exact test were used to compare independent groups of categorical data. A p value <0.05 was taken as statistically significant. Sensitivity, specificity, positive and negative predictive values of pulse oximetry screening to detect CCHDs were done.

Results
One thousand neonates were enrolled in the study of whom 482 (48%) were females and 518 (52%) were males. The gestational ages ranged from 34 to 40 weeks (mean 37.7 ± 1.3 weeks). Birth weights ranged from 1800 to 4850g (mean 3224.1 ± 397.1g). Six hundred and twenty one (62%) babies were born by Caesarean section which was the commonest mode of delivery. A comparison of the demographic data between neonates with and without CCHD is shown in Table 1. CCHD was significantly higher in male newborns (p<0.05).

Table 1: Comparison of demographic data between neonates with and without CCHD (n=1000)

<table>
<thead>
<tr>
<th>Characteristic</th>
<th>Critical congenital heart disease (CCHD)</th>
<th>p value</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Yes (n=28)</td>
<td>No (n=972)</td>
</tr>
<tr>
<td>Sex</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Male n (%)</td>
<td>20 (71)</td>
<td>498 (51)</td>
</tr>
<tr>
<td>Female n (%)</td>
<td>08 (29)</td>
<td>474 (49)</td>
</tr>
<tr>
<td>Gestational age (weeks)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>34-37 n (%)</td>
<td>20 (71)</td>
<td>826 (85)</td>
</tr>
<tr>
<td>38-40 n (%)</td>
<td>08 (29)</td>
<td>146 (15)</td>
</tr>
<tr>
<td>Birth weight (g)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>≥ 2500 n (%)</td>
<td>25 (89)</td>
<td>860 (88)</td>
</tr>
<tr>
<td>&lt;2500 n (%)</td>
<td>03 (11)</td>
<td>11 (12)</td>
</tr>
<tr>
<td>Mode of delivery</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Caesarean n (%)</td>
<td>13 (46)</td>
<td>608 (63)</td>
</tr>
<tr>
<td>Vaginal n (%)</td>
<td>15 (54)</td>
<td>364 (37)</td>
</tr>
</tbody>
</table>

There were 971 (97%) babies with negative screening and 29 (3%) babies with positive screening. Of the 971 neonates with negative screening, 6 had oxygen saturation levels ≤ 95 in the first measurement and showed normal saturation when repeated after 3 hours of initial measurement. Echocardiography was done in all 29 newborns with positive screening. Of the 29 neonates, three had normal cardiac anatomy and function and twenty-six had a pathologic echocardiogram. Of the 971 negative screening patients who had normal oxygen saturation in both measurements, two had a postnatal CCHD diagnosis (Figure 1).
Pulse oximetry CCHD screening had a sensitivity of 93%, a specificity of 99%, a positive predictive value of 81% and a negative predictive value of 99%. There was no difference among the predictive performances of pulse oximetry in detecting CCHD between pre-ductal and post-ductal measurements (Table 2).

Table 2: Predictive performance of pulse oximetry in detecting critical congenital heart disease

<table>
<thead>
<tr>
<th>Measurement</th>
<th>Sensitivity</th>
<th>Specificity</th>
<th>Positive predictive value</th>
<th>Negative predictive value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Pre-ductal</td>
<td>93%</td>
<td>99%</td>
<td>81%</td>
<td>99%</td>
</tr>
<tr>
<td>Post-ductal</td>
<td>93%</td>
<td>99%</td>
<td>81%</td>
<td>99%</td>
</tr>
</tbody>
</table>

Discussion

In our study, using pulse oximetry screening we were able to diagnose 26 neonates with CCHD. The prevalence of CCHD in our study was higher than reported prevalence of CHD. This may be due to high-risk pregnancies being referred to our tertiary hospital.

Though early diagnosis and treatment of CCHD is crucial, there is no consensus about the recommended screening protocol for infants with a positive screen. In our study, the oxygen saturation threshold was ≤95%. We made a second measurement separated by 3 hours from the first measurement if the first neonate saturation was ≤95%. Of 971 neonates with negative screening, 6 neonates had oxygen saturation levels ≤ 95 in first measurement and normal saturation after 3 hours. Therefore two repeated measurements have been shown to be accurate in detecting CCHD.

In our study, sensitivity of pulse oximetry screening was found to be 93% for detection of CCHD. Several studies have shown that when pulse oximetry is done in the first 24 hours of life, there can be more false positives. In our study, pulse oximetry was done between 24-72 hours age. Moreover, newborns with co-morbidities such as preterm delivery, respiratory distress syndrome, dysmorphic syndromes, multiple congenital anomalies and sepsis were excluded from the study. This may account for the lower false positivity rate in our study.

Routine use of pulse oximetry screening may detect CCHD in asymptomatic newborns before hospital discharge. It prevents previously well infants from acute collapse at the time of ductal closure and lead to improve morbidity with worse surgical and neurodevelopmental outcomes.

Conclusions

Pulse oximetry screening in term and late preterm neonates is effective for early diagnosis of CCHD.

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