Neonatal Visual Evoked Potentials in Infants Born to Mothers Prescribed Methadone

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KEY WORDS
flash visual evoked potential, VEP, methadone, infant

ABBREVIATIONS
CI—confidence interval
FAEE—fatty acid ethyl ester
IUGR—intrauterine growth restriction
NAS—neonatal abstinence syndrome
VEP—visual evoked potential

Dr McGlone contributed to study design, recruited all patients, collected and analyzed data, and drafted the manuscript; Drs Hamilton and Bradnam developed the original idea; Drs Hamilton and McCulloch made substantial contribution to study design and analysis of data; Mr Boulton contributed to study design and collection and analysis of data; Dr Bradnam made substantial contribution to study design; Dr Weaver contributed to study design; Dr Mactier made substantial contribution to study design and execution of study; Drs Hamilton, McCulloch, Bradnam, Weaver, and Mactier and Mr Boulton revised the manuscript critically for important intellectual content; and all authors approved the final version of the manuscript.

www.pediatrics.org/cgi/doi/10.1542/peds.2012-2113
doi:10.1542/peds.2012-2113

Accepted for publication Nov 2, 2012

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PEDIATRICS (ISSN Numbers: Print, 0031-4005, Online, 1098-4275)

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FINANCIAL DISCLOSURE: The authors have indicated they have no financial relationships relevant to this article to disclose.

FUNDING: Funding was provided by Yorkhill Children’s Foundation and Tenovus Scotland.

WHAT’S KNOWN ON THIS SUBJECT: Impaired visual development has been reported in infants born to mothers prescribed methadone in pregnancy. Immature visual evoked potentials have been reported in this population, but data were confounded by gestation, growth restriction, and illicit drug use.

WHAT THIS STUDY ADDS: Visual evoked potentials are small and immature in infants exposed to methadone and other drugs of misuse in utero. These changes are independently associated with methadone exposure and persist after controlling for gestation, socioeconomic deprivation, alcohol consumption, and cigarette smoking.

abstract

OBJECTIVE: Drug misuse in pregnancy is associated with impaired infant visual development. Pilot data showed abnormal flash visual evoked potentials (VEPs) in neonates exposed to methadone in utero, but results were confounded by intrauterine growth restriction, gestation, and ongoing drug misuse. This large cohort study aimed to clarify the effects on neonatal flash VEPs of maternal drug misuse in pregnancy, including prescription of substitute methadone and subsequent development of neonatal abstinence syndrome.

METHODS: This was a prospective cohort study. Flash VEPs were recorded within 3 days of birth from 100 healthy infants of drug-misusing mothers prescribed substitute methadone during pregnancy and 50 comparison infants matched for birth weight, gestation, and socioeconomic deprivation. VEP morphology was classified as mature, typical, or immature, and amplitudes and implicit times of the major waveform components measured. Drug exposure was determined by maternal history, maternal and infant urine, and meconium toxicology.

RESULTS: VEPs from maternal drug-exposed infants were more likely to be of immature waveform (P < .001) and were smaller in overall amplitude (median 27 μV vs 39 μV, P < .001) compared with non-drug-exposed infants. Most infants were exposed to illicit drugs in addition to prescribed methadone; differences in VEP parameters were independently associated with maternal prescribed methadone and persisted after correcting for birth weight, cigarette smoking, and excess in utero alcohol exposure.

CONCLUSIONS: In utero exposure to prescribed substitute methadone is associated with altered flash VEPs in the newborn period and these infants may warrant early clinical visual assessment. Pediatrics 2013;131:e857–e863
Maternal drug misuse is a significant social problem affecting an estimated 250,000 to 350,000 children in the United Kingdom. Management of maternal opiate misuse in pregnancy includes substitute prescribing of methadone, a synthetic opioid that stabilizes lifestyle, lessens risk-taking behavior, and reduces the incidence of preterm birth and intrauterine growth restriction (IUGR). Methadone use in pregnancy is associated with improved compliance with antenatal care, but most mothers also misuse other substances, particularly opiates and benzodiazepines. There is increasing evidence to suggest that maternal drug misuse in pregnancy has adverse effects on infant visual development, with reported abnormalities, including reduced visual acuity, nystagmus, delayed visual maturation, strabismus, and refractive errors. Flash visual evoked potentials (VEPs) in infancy reflect the integrity and maturity of the visual system and we have demonstrated abnormal VEPs in newborn infants exposed to methadone in utero. That pilot study was, however, limited by small numbers and confounded by poly-drug exposure and IUGR. Abnormal infant flash VEPs in other high-risk neonatal groups have been linked with long-term visual, motor, and learning disabilities. The aims of the current study were to investigate the effects of maternal drug misuse on neonatal flash VEPs and to assess any relationship between early visual electrophysiology and subsequent neonatal abstinence syndrome (NAS).

METHODS

This prospective cohort study was conducted at the Princess Royal Maternity in Glasgow, UK. Eligible infants were born to drug-misusing mothers prescribed substitute methadone during pregnancy. Exclusion criteria were birth before 36 completed weeks’ gestation, congenital ocular abnormality, and significant neonatal illness. Infants <36 weeks’ gestation were excluded to remove the potential confounding effect of preterm birth on the neonatal VEP waveform. For comparative purposes, healthy, non-drug-exposed infants were recruited within the same maternity unit. To correct for potential confounding factors, comparison infants were matched for completed week of gestation, birth weight ± 250 g, and Carstairs deprivation index score ± 1 (deprivation category based on the presence of overcrowding, unemployment, social class, and car ownership). Ethics approval was granted by the West of Scotland Research Ethics Committee, and written informed parental consent was obtained for all participants.

Sample Size

A pilot study demonstrated differences in flash VEP morphology, amplitude, and implicit times of the major components between 21 methadone-exposed infants and 20 control infants. Local audit suggested that ~20% of pregnant women prescribed substitute methadone used no additional illicit substances and so 100 maternal drug-exposed infants were expected to include a subgroup of 20 who had been exposed to methadone alone. Approximately 50% of infants were predicted to develop significant NAS. A group of this size (n = 50) was likely to have 95% confidence intervals (CIs) of the mean for flash VEP P2 amplitude of ±2 μV and for P2 implicit time of ±11 ms, which is adequately narrow for clinical purposes.

Clinical Data Collection

Maternal and infant demographic characteristics were collected at study enrollment and the severity of NAS recorded from the case notes. The latter was classified by using a modified Lipsitz score: 1 = no NAS, 2 = mild NAS (no pharmacological treatment required), 3 = moderate NAS (standard oral morphine treatment), and 4 = severe NAS (escalated treatment with phenobarbital).

Drug Exposure and Toxicology

After study recruitment, a confidential interview was conducted with all mothers regarding drug and alcohol use during pregnancy. Maternal urine samples were collected routinely at the hospital booking visit and specimens of the infant’s urine and meconium were obtained as soon as possible after delivery. Urine samples were analyzed by the regional toxicology laboratory by using the enzyme multiplied immunosassay technique assays run to Substance Abuse and Mental Health Services Administration guidelines on an Abbott Architect Analyzer (Abbott, Abbott Park, IL). Assays included methadone, opiates, benzodiazepines, amphetamines, cannabinoids, and cocaine metabolites. Meconium samples were screened for the same drugs by using enzyme linked immunosorbent assay; selected positive samples were further analyzed by using solid phase and liquid-liquid extraction followed by gas chromatography–mass spectrometry or liquid chromatography–mass spectrometry. History and urine and meconium toxicology were combined to provide an overall drug exposure pattern for each study infant. Meconium samples were also analyzed for fatty acid ethyl esters (FAEEs) to identify infants exposed to excess alcohol in utero by using liquid chromatography–mass spectrometry. A cutoff value of >10,000 ng/g of meconium was used to signify excess alcohol consumption in pregnancy. Meconium samples were collected from control infants.
and analyzed for drugs of misuse and FAEEs.

**Neonatal VEP Recording**

Flash VEPs were recorded within the first 72 hours of life from an active midline occipital electrode over the visual cortex at Oz, with a midline frontal reference electrode at Fz, according to international standards. Stimulus generation, recording, and data storage were carried out by using an Espion evoked potential system (Diagnosys LLC, Lowell, MA) and flashes delivered by using a hand-held light-emitting diode stimulator to deliver a white pulse flash with time-integrated luminance of 28 cd.s.m\(^{-2}\) at a frequency of 1 Hz. Impedance was <10 kohms, recorded at the start and end of each recording. A minimum of 30 trials per average were recorded and the procedure repeated to check reproducibility. Each VEP recording session took ~30 minutes to complete. All VEPs were recorded before commencement of any pharmacological treatment of NAS.

**Data Analysis**

Flash VEPs were categorized as present or absent. When present, the amplitude and implicit times of peaks and troughs were measured. Peaks and troughs were defined as follows: P1 = any positive component before P2; P2 = the major positive component between 126 and 300 ms and preceding N3 if present; N3 = negative component between 200 and 400 ms; P3 = positive component after the N3. The total summed amplitude of all peaks and troughs was calculated. Each flash VEP was also categorized by waveform morphology, based on normal maturation of VEP morphology in infants from preterm to early postterm. Descriptive categories were: typical (predominant negativity near 300 ms [N3] with either no detectable P2 or with P2 amplitude less than one-third of the N3 amplitude), immature (predominant negativity near 300 ms [N3] with either no detectable P2 or with P2 amplitude less than one-third of the N3 amplitude), and atypical (reproducible VEP with an unusual waveform that did not meet criteria of other categories) (Fig 1). All VEPs were assessed by 2 observers,

![Neonatal VEP waveform morphology](image)
1 of whom was blinded to the infant’s group and clinical course.

Statistics

VEP implicit times and amplitudes were tested for normality by using Anderson-Darling tests; as the data were of skewed distribution, comparisons between groups were done by using Mann-Whitney tests and Kruskal-Wallis tests. VEP morphology was compared between groups by using χ² tests. Linear and logistic regression models were used to correct for potential confounders and to assess the effects of independent drugs of misuse on neonatal flash VEPs.

RESULTS

One hundred maternal methadone-exposed infants (98% of eligible infants approached) and 50 matched comparison infants were recruited. Sixteen methadone-exposed infants were excluded because of preterm birth (n = 11) or significant illness (n = 5). Maternal and infant demographic characteristics are shown in Table 1. Although birth weights were comparable, maternal methadone-exposed infants had smaller occipitofrontal head circumferences (33.5 cm vs 34.1 cm, 2-sample t test P = .015). Substitute methadone-prescribed mothers were more likely to report cigarette smoking (95% vs 60%, χ² test P < .001) (Table 1). The most commonly misused drugs in addition to methadone were opiates (74%), benzodiazepines (66%), and cannabis (62%) (Table 2, Fig 2). Only 9 infants were exposed to methadone alone (Fig 2); a further 8 were additionally exposed to other opiates giving a subgroup of 17 exposed to opiates alone. Meconium toxicology analysis was performed on 30 of the comparison infants, 3 of whom tested positive for cannabinoids. Of those infants whose meconium sample was sufficient to test for FAEEs, 26 drug-exposed (26/63, 41%) and 5 comparison infants (5/21, 23%) had elevated levels, indicative of excess alcohol exposure in utero. Fourteen drug misusing mothers were taking prescribed antidepressant and/or antipsychotic medication during pregnancy, compared with none of the comparison mothers. To investigate any potential confounding effect of these maternal prescribed drugs on the newborn infant, VEP and VEP parameters were compared between drug-exposed infants whose mothers were taking prescribed antidepressant medication and those who were not. There was no difference in VEP amplitude, implicit time, or morphology between these groups.

Comparison of Flash VEPs

Methadone-exposed infants were less likely to demonstrate P1 and N2 components than comparison infants (P1: 21% vs 48%, χ² = 11.6, P = .001; N2: 38% vs 60%, χ² = 6.5, P = .011) and had VEPs of smaller total amplitude: 27 μV (interquartile range 17–42) vs 39 μV (interquartile range 28–67) (Mann-Whitney test P < .001, 95% CI –20 to –6) (Table 3). Overall, maternal methadone-exposed infants had more

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### Table 1: Demographic Characteristics of Infants and Mothers

<table>
<thead>
<tr>
<th>Category</th>
<th>Cases (n = 100)</th>
<th>Comparisons (n = 50)</th>
<th>P Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Gender: % male</td>
<td>46</td>
<td>44</td>
<td>NS</td>
</tr>
<tr>
<td>Mode of delivery: %</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>SVD</td>
<td>72</td>
<td>70</td>
<td>NS</td>
</tr>
<tr>
<td>LUSCS</td>
<td>21</td>
<td>20</td>
<td></td>
</tr>
<tr>
<td>Instrumental</td>
<td>7</td>
<td>10</td>
<td></td>
</tr>
<tr>
<td>Gestation, wkc</td>
<td>39.3 (38.2–40.1)</td>
<td>39.7 (38.1–41.6)</td>
<td>NS</td>
</tr>
<tr>
<td>5-min Apgar</td>
<td>9 (9–10)</td>
<td>9 (9–10)</td>
<td></td>
</tr>
<tr>
<td>Birth weight, g</td>
<td>2892 (±505)</td>
<td>3005 (±539)</td>
<td>NS</td>
</tr>
<tr>
<td>SGA, %</td>
<td>18</td>
<td>20</td>
<td>NS</td>
</tr>
<tr>
<td>LBMI, %</td>
<td>20</td>
<td>18</td>
<td>NS</td>
</tr>
<tr>
<td>OFC, cm</td>
<td>33.5 (±1.6)</td>
<td>34.1 (±1.8)</td>
<td>.015</td>
</tr>
<tr>
<td>Microcephaly, %</td>
<td>8</td>
<td>8</td>
<td>NS</td>
</tr>
<tr>
<td>Feeding at D/C, %</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Bottle</td>
<td>87</td>
<td>90</td>
<td></td>
</tr>
<tr>
<td>Breast</td>
<td>7</td>
<td>8</td>
<td>NS</td>
</tr>
<tr>
<td>Mixed</td>
<td>6</td>
<td>2</td>
<td></td>
</tr>
<tr>
<td>Maternal smoking, %</td>
<td>95</td>
<td>60</td>
<td>&lt;.001</td>
</tr>
<tr>
<td>Maternal BMI, %</td>
<td>23 (21–26)</td>
<td>23.5 (21–30)</td>
<td>NS</td>
</tr>
<tr>
<td>Maternal DEPCAT</td>
<td>7 (5–7)</td>
<td>6 (4–7)</td>
<td>NS</td>
</tr>
</tbody>
</table>

Microcephaly was defined as OFC < second centile. D/C, discharge; DEPCAT, Carstairs deprivation index score; LBMI, low birth weight; LUSCS, lower uterine segment caesarean delivery; NS, not significant; OFC, occipitofrontal head circumference; SGA, small for gestational age; SVD, spontaneous vertex delivery.

1 Categories were compared by using χ² tests; birth weight and OFC by using 2-sample t tests; and gestation, Apgar scores, maternal BMI, and DEPCAT by using Mann-Whitney tests.

2 Categorical data (gender, delivery, SGA, LBMI, microcephaly, feeding, and smoking) are given as percentage of each cohort.

3 Gestation, Apgar scores, maternal BMI, and DEPCAT scores are medians (interquartile range).

4 Birth weight and OFC are means (± SD).

### Table 2: Drug Exposure: Comparison of History and Toxicology (Cases)

<table>
<thead>
<tr>
<th>Drug</th>
<th>History (n = 100)</th>
<th>Maternal Urine (n = 84)</th>
<th>Infant Urine (n = 70)</th>
<th>Meconium (n = 74)</th>
<th>Combined (n = 100)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Methadone, %</td>
<td>100</td>
<td>92</td>
<td>61</td>
<td>96</td>
<td>100</td>
</tr>
<tr>
<td>Other opiates, %</td>
<td>54</td>
<td>56</td>
<td>36</td>
<td>81</td>
<td>74</td>
</tr>
<tr>
<td>BDZ, %</td>
<td>51</td>
<td>58</td>
<td>33</td>
<td>53</td>
<td>66</td>
</tr>
<tr>
<td>Amphetamine, %</td>
<td>2</td>
<td>1</td>
<td>0</td>
<td>14</td>
<td>13</td>
</tr>
<tr>
<td>Cannabis, %</td>
<td>19</td>
<td>39</td>
<td>9</td>
<td>65</td>
<td>62</td>
</tr>
<tr>
<td>Cocaine, %</td>
<td>5</td>
<td>5</td>
<td>3</td>
<td>15</td>
<td>14</td>
</tr>
</tbody>
</table>

Data are the percentage of positive results for each technique. The combined column combines the history and toxicology results to give a pattern of overall drug exposure for each infant. BDZ, benzodiazepine.
immature/atypical VEPs and fewer mature responses ($\chi^2 = 13.6, P = .001$) (Fig 3). All of these differences persisted after correcting for occipitofrontal head circumference, maternal cigarette smoking, and excess maternal alcohol intake during pregnancy with a linear regression model for VEP amplitude and a binary logistic regression model for the prevalence of VEP components.

**Table 3**

<table>
<thead>
<tr>
<th>Parameter</th>
<th>Cases (n = 100)</th>
<th>Comparisons (n = 50)</th>
<th>P Value</th>
<th>Adjusted P Valuea</th>
</tr>
</thead>
<tbody>
<tr>
<td>P1, %b</td>
<td>21</td>
<td>48</td>
<td>.001</td>
<td>.003</td>
</tr>
<tr>
<td>N2, %b</td>
<td>38</td>
<td>80</td>
<td>.011</td>
<td>.011</td>
</tr>
<tr>
<td>P2, %b</td>
<td>89</td>
<td>98</td>
<td>.106</td>
<td>—</td>
</tr>
<tr>
<td>N3, %b</td>
<td>87</td>
<td>84</td>
<td>.021</td>
<td>—</td>
</tr>
<tr>
<td>P1 IT, msb</td>
<td>133 (118–175)</td>
<td>137 (114–157)</td>
<td>.036</td>
<td>—</td>
</tr>
<tr>
<td>P2 IT, msb</td>
<td>207 (191–221)</td>
<td>206 (191–228)</td>
<td>.890</td>
<td>—</td>
</tr>
<tr>
<td>N3 IT, msb</td>
<td>286 (247–328)</td>
<td>321 (250–357)</td>
<td>.262</td>
<td>—</td>
</tr>
<tr>
<td>Total amplitude, $\mu V$c</td>
<td>27 (17–42)</td>
<td>39 (28–67)</td>
<td>&lt;.001</td>
<td>.001</td>
</tr>
</tbody>
</table>

Percentage responses were compared by using $\chi^2$ tests; implicit times and amplitudes were compared by using Mann-Whitney tests. IT, implicit time; —, not significant.

a Adjusted P value is after correcting for occipitofrontal head circumference, maternal smoking, and excess maternal alcohol intake during pregnancy with a linear regression model for VEP amplitude and a binary logistic regression model for the prevalence of VEP components.

b Percentage of VEPs containing the individual component.

c Median (interquartile range) for implicit times and amplitude.

**Neonatal VEPs and In Utero Drug Exposure**

Regression analysis confirmed that differences in VEP parameters persisted after correcting for other illicit drug use: VEP amplitude, $P = .012$; P1 response, odds ratio 0.02, 95% CI 0.00–0.16, $P = .001$; N2 response, odds ratio 0.27, 95% CI 0.09–0.84, $P = .024$. We found no associations between the prescribed maternal dose of methadone and amplitude, latency, or morphology of infant flash VEPs.

**DISCUSSION**

Substitute methadone is the currently recommended treatment of pregnant opioid-dependent women.23 Several observational studies have reported abnormalities of visual and neurologic development in infants of drug-misusing mothers prescribed substitute methadone in pregnancy,5–12,17 but to date there has been no prospective study of such infants, and so the prevalence of visual dysfunction in this population remains unknown.

We have previously described abnormal flash VEPs in newborn infants exposed to methadone in utero17 but these data were confounded by IUGR and gestation, and numbers were too small to investigate the individual effects of methadone and other illicit drugs. This larger cohort study has confirmed substantial differences between neonatal flash VEPs of infants exposed to methadone in utero and those of non–drug-exposed comparison infants and suggests that prescribed substitute methadone, rather than other illicit drugs, may be to blame. Reduced VEP amplitude with immature waveform is consistent with the clinical finding of delayed visual maturation previously reported in similar but older populations.10–12

In utero methadone exposure in animal studies causes depletion of both acetylcholine and nerve growth factor in the brain as well as alterations in

**FIGURE 2**

Euler diagram illustrating pattern of combined drug exposure for all 100 cases. Stimulants: cocaine and/or amphetamines.

**FIGURE 3**

VEP waveform morphology for cases and comparisons. Drug-exposed infants had a greater proportion of immature or atypical VEPs (cases 28%, controls 10%) and fewer mature responses (cases 21%, controls 48%) compared with comparison infants. $\chi^2 = 13.6, P = .001$.  

- **TABLE 3**

VEP Parameters: Case and Comparison Infants

- **FIGURE 3**

VEP waveform morphology for cases and comparisons. Drug-exposed infants had a greater proportion of immature or atypical VEPs (cases 28%, controls 10%) and fewer mature responses (cases 21%, controls 48%) compared with comparison infants. $\chi^2 = 13.6, P = .001$.  

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Pediatrics 2013;131;e857
DOI: 10.1542/peds.2012-2113 originally published online February 18, 2013;

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