HEPATITIS B VACCINE IN INFANTS: A RANDOMIZED CONTROLLED TRIAL COMPARING GLUTEAL VERSUS ANTEROLATERAL THIGH MUSCLE ADMINISTRATION

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SUMMARY

A significantly diminished antibody response to hepatitis B vaccine has been demonstrated in adults when the buttock is used as the injection site. However, in Brazil, the buttock continues to be recommended as site of injection for intramuscular administration of vaccines in infants. In this age group, there are no controlled studies evaluating the immunogenicity of the hepatitis B vaccine when administered at this site. In the present study, 258 infants were randomized to receive the hepatitis B vaccine either in the buttock (n = 123) or in the anterolateral thigh muscle (n = 135). The immunization schedule consisted of three doses of hepatitis B vaccine (Engerix B®, 10 µg) at 2, 4 and 9 months of age. There were no significant differences in the proportion of seroconversion (99.3% vs 99.2%), or in the geometric mean titer of ELISA anti-HBs (1,862.1 x 1,229.0 mIU/mL) between the two groups. This study demonstrates that a satisfactory serological response can be obtained when the hepatitis B vaccine is administered intramuscularly into the buttock.

KEYWORDS: Hepatitis B; Vaccine efficacy; Injection; Intramuscular; Gluteal injection; Infant.

INTRODUCTION

Early studies on the response to hepatitis B vaccine have demonstrated a 96% seroconversion rate in the target population. However, in 1984 reports appeared of a serological efficacy of only 63 to 68% among vaccinated health workers. In 1985, a surveillance study by the Centers for Disease Control and Prevention (CDC) in 57 hospitals in the United States showed a significantly higher antibody response against the surface antigen of the hepatitis B virus (anti-HBs) in hospitals where the deltoid muscle was used as the injection site compared to the buttocks, 93% and 81% respectively. Based on these findings, administration of hepatitis B vaccine in the buttock is no longer recommended at any age. Administration into the deltoid muscle is recommended for older children and adults, while the anterolateral thigh muscle is preferred for infants and toddlers. In 1998, the hepatitis B vaccine was introduced in the National Immunization Program of the Ministry of Health of Brazil for children under one year of age in all country and for children under 15 years of age in the States of Santa Catarina, Paraná, Distrito Federal and in all the Amazonian region. For infants the anterolateral thigh muscle has been recommended as the injection site. However, in Brazil, the application of intramuscular thigh injections still is not a well-established routine since the buttock continues to be recommended as site of injection for DPT vaccine. The most recent data available from the Ministry of Health of Brazil show that more than 14 million doses of DPT were administered to children under 18 months of age in 1998. A concern has been raised whether turning the anterolateral thigh muscle into the mandatory injection site for hepatitis B administration could result in additional difficulties in implementing children’s vaccination programs due to cultural and technical problems.

All studies reporting reduced serological efficacy of hepatitis B vaccine administered in the buttocks have been conducted on adults. In this age group, an inversely proportional correlation between a high weight and height index and a satisfactory anti-HBs response has been noted. The present study was carried out to evaluate the serological efficacy of hepatitis B vaccination in infants by comparing quantitative anti-HBs levels as a consequence of the injection site used, taking into consideration age, sex and nutritional status.

METHODS

Study design: A randomized controlled trial.

Calculation of sample size: The sample size for the two groups was calculated on the basis of a CDC study on adults that demonstrated a serologic efficacy of 93.9% in vaccinated subjects who received the hepatitis B vaccine in the deltoid, and an efficacy of 81.0% in subjects receiving the vaccine in the buttock. Considering a 5% type I error and 90% statistical power, 157 children were recommended for each group.

Randomization: A total of 314 sealed envelopes containing 156 filling cards for group A (buttock) and 158 filling cards for group B (anterolateral thigh muscle) were mixed by shuffling and then numbering. After the infants filling cards for group A (buttock) and 158 filling cards for group B (anterolateral thigh muscle) were mixed by shuffling and then numbering. After the infants
informing consent, the envelopes were sequentially opened according to admission to the study. Information was then obtained regarding the group to which the infants had been assigned.

**Study population:** The participants were selected from infants aged two to four months whose parents spontaneously sought assistance at one of the basic health units in the City of São Paulo where routine medical visits are performed and the vaccines of the official immunization program are provided. The study was conducted from January 1997 to October 1998. The hepatitis B vaccine was not available to the public health network when the study was performed. Infants whose parents would not be available for follow-up or who did not agree with the randomization of the injection site were not included in the study. Children with a previous vaccination against hepatitis B, a history of blood or immunoglobulin transfusion or of prolonged treatment with corticosteroids, the presence of a clinical situation that would contraindicate routine immunization, or the children of mothers known or suspected to carry HIV or hepatitis B virus infection were also excluded. The parents signed an informed consent form and the study was approved by the Ethical Research Committee of the University of São Paulo School of Medicine.

**Immunization schedule:** The immunization schedule consisted of three applications of hepatitis B vaccine (Engerix B<sup>®</sup>, Smith Kline Beecham, Rixensart, Belgium) as a 10 µg dose was injected IM into the upper, outer right quadrant of the buttock (Group A) or into the anterolateral surface of the right thigh (Group B). The nurses in charge of vaccine administration were trained to apply IM injections into the anterolateral thigh muscle. Needles measuring 30 or 25 mm in length by 0.7 mm in diameter were used for the buttock and anterolateral thigh muscle injections, respectively. The vaccine was stored at 2-8 °C.

**Intervention plan:** The Engerix B<sup>®</sup> vaccine (Smith Kline Beecham, Rixensart, Belgium) as a 10 µg dose was injected IM into the upper, outer right quadrant of the buttock (Group A) or into the anterolateral surface of the right thigh (Group B). The nurses in charge of vaccine administration were trained to apply IM injections into the anterolateral thigh muscle. Needles measuring 30 or 25 mm in length by 0.7 mm in diameter were used for the buttock and anterolateral thigh muscle injections, respectively. The vaccine was stored at 2-8 °C.

**Statistical analysis:** First descriptive analysis (frequency distribution, arithmetic and geometric means, medians and standard deviation [SD]) were calculated. The Student t-test was calculated to compare means of normally distributed samples (age, weight and height); when not applicable, Wilcoxon-Mann-Whitney U test was used<sup>25</sup>. Quantitative variables were dichotomized into two categories, the cut-off point lying close to the median. The efficacy of the vaccine in each group was assessed from the proportion of non-responders according to the injection site studied. To analyze the variable anti-HBs > 10 mIU/mL according to the group studied, we also calculated the geometric mean for antibody levels, which is not affected by extreme values. The group of infants with anti-HBs levels < 100 mIU/mL was analyzed separately by univariate analysis to detect typical characteristics, and later by unconditional logistic regression to control for possible confounding variables<sup>17</sup>. The strength of the association between variables was estimated by calculating the odds ratio (OR) and 95% confidence interval (95% CI). Statistical significance was evaluated by the likelihood ratio test (LR).

**RESULTS**

Of 314 infants, 258 (82.2%) completed all stages of the study. No significant differences were observed between the group that concluded the study and the infants lost to follow-up with respect to age, sex, weight, height, and vaccine application site. Of those who remained in the study (n = 258), the mean interval (SD) between the 1<sup>st</sup> and the 2<sup>nd</sup> dose applications was 67.0 days (15.8 days), that between the 2<sup>nd</sup> and 3<sup>rd</sup> doses was 175.8 days (40.7 days), and that between the 3<sup>rd</sup> dose and blood collection was 144.5 days (50.5 days). No significant differences were seen in the characteristics of the infants who received the vaccine in the anterolateral thigh muscle group and an efficacy of 99.2% (122/123).
Table 1

<table>
<thead>
<tr>
<th>Variable</th>
<th>Anterolateral thigh muscle</th>
<th>Buttock</th>
</tr>
</thead>
<tbody>
<tr>
<td>Number</td>
<td>135</td>
<td>123</td>
</tr>
<tr>
<td>Sex (female)</td>
<td>63 (46.7%)</td>
<td>63 (51.2%)</td>
</tr>
<tr>
<td>Age (days)</td>
<td>74.1 ± 15.2</td>
<td>73.6 ± 14.9</td>
</tr>
<tr>
<td>Weight (Kg)</td>
<td>5.5 ± 0.8</td>
<td>5.5 ± 0.9</td>
</tr>
<tr>
<td>Length (cm)</td>
<td>56.7 ± 3.1</td>
<td>56.8 ± 3.2</td>
</tr>
<tr>
<td>Dose interval (1st and 2nd)</td>
<td>67.9 ± 17.1</td>
<td>66.4 ± 14.1</td>
</tr>
<tr>
<td>Dose interval (2nd and 3rd)</td>
<td>173.4 ± 36.0</td>
<td>175.8 ± 45.3</td>
</tr>
<tr>
<td>Interval between 3rd dose and blood collection</td>
<td>144.6 ± 51.1</td>
<td>144.4 ± 50.2</td>
</tr>
</tbody>
</table>

* The differences are not statistically significant

Twelve of the 258 infants (8.5%) had anti-HBs levels < 100 mIU/mL. Of these, seven belonged to the 135 infants who received the vaccine in the anterolateral thigh muscle group (p = 0.06). The median anti-HBs titer was 2.051 mIU/mL for the anterolateral thigh muscle group and 1.485 mIU/mL for the buttock group (p = 0.14). Twenty-two of the 258 infants (8.5%) had anti-HBs levels < 100 mIU/mL. Comparison of the risk of vaccine failure (anti-HBs < 10 mIU/mL) with respect to the nutritional profile of each study group was not performed due to the small number of infants presenting this response (one child in each group).

The fact that the two groups showed a similar proportion of seroconversion may be of practical importance since not all public health units in developing countries are staffed by workers trained in administering IM injections into the anterolateral thigh muscle. If the buttock is considered a possible alternative for hepatitis B vaccine administration, many operational and cultural problems may be avoided, eliminating possible delays in the implementation of infant immunization against hepatitis B.

No significant difference in anti-HBs levels > 5,000 mIU/mL was detected between groups. However, when the number of infants with anti-HBs levels < 100 mIU/mL was analyzed, a significant association was obtained between this parameter and the vaccine administration site (p = 0.04). In the multivariate analysis, the only variable associated with anti-HBs levels < 100 mIU/mL was the vaccine application site (OR = 2.54, 95% CI = 1.02-6.76). However, the prevalence of infants with post-vaccination levels < 100 mIU/mL was low, both in the study population as a whole (8.5%) and in each group specifically (5.2 % for the group vaccinated in the anterolateral thigh muscle and 12.2% for the group vaccinated in the buttock).

The clinical significance of this response pattern to hepatitis B vaccine is not well defined. Some investigators believe that anti-HBs levels after vaccination should exceed 100 mIU/mL since there are reports of individuals with anti-HBs levels < 100mIU/mL who present clinical or subclinical hepatitis B virus infection with transiently positive antigenemia and hepatitis B virus DNA25. Other investigators prefer the use of terms such as “seroconverted” for subjects who exhibit an anti-HBs response > 10 mIU/mL, as opposed to the term “seroprotected”, which is reserved for those whose anti-HBs levels exceed 100 mIU/mL.43,14

Studies involving the clinical and laboratory monitoring of populations at a high risk of acquiring the hepatitis B virus have demonstrated that the hepatitis B vaccine induces a long-lasting protective immune response. Natural boosters occur frequently and, due to an anamnestic type response, even individuals considered to be low
responders (anti-HBs < 100 mIU/mL) continue to be protected against chronic hepatitis B virus infection for years. Additional booster doses of hepatitis B vaccine are not recommended by the World Health Organization. Recent studies have shown that vaccine-dependent immunity is of long duration, suggesting that there is no need to administer booster doses of the vaccine to prevent a fall in or disappearance of anti-HBs levels over the years.

The administration of IM injections into the buttock is believed to involve greater risk than that into the anterolateral thigh muscle because of possible damage to the sciatic nerve. However, in most reported cases of sciatic nerve palsies in infants and children after gluteal injections there was evidence of direct injection of neurotoxic material, such as antibiotics, bismuth and quinine, into or near the nerve. Otherwise, there are also anecdotal reports of complications following intramuscular injections into the anterolateral thigh muscle, including muscle necrosis deep to the fascia lata, acute chemical synovitis, and fibrosis and contracture of the quadriceps muscle resulting in loss of knee flexion. There are no controlled, prospective studies comparing the risks involved in the use of these two injection sites for immunization in infants. In Brazil, approximately 15 million doses of vaccines are administered each year to infants using the buttock as site of injection. In the present study a satisfactory serological response was observed in infants when the hepatitis B vaccine was administered into the buttock suggesting that this procedure may be considered in environments where, for cultural or operational reasons, it can facilitate the implementation of vaccination programs.

RESUMO

Vacina da hepatite B em lactentes: um ensaio clínico controlado, aleatorizado, comparando a administração no glúteo com a administração no vasto lateral da coxa

Diminuição significativa da resposta sorológica à vacina da hepatite B foi documentada em adultos quando a mesma é aplicada no glúteo. Entretanto, em alguns países, o glúteo ainda é bastante utilizado para aplicação intramuscular de vacinas em lactentes. Nesta faixa etária, não há estudos controlados que tenham avaliado a imunogenicidade da vacina da hepatite B quando administrada em diferentes locais. No presente estudo, 258 lactentes foram sorteados de forma aleatória para receber a vacina da hepatite B no glúteo (n = 123) ou no músculo vasto lateral da coxa (n = 135). O esquema vacinal consistiu em 3 doses da vacina da hepatite B (Engerix B®, 10 μg) aos 2, 4 e 9 meses de idade. Não se observaram diferenças significativas nas proporções de soroconversão (99,3% X 99,2%), ou na média geométrica dos títulos de anti-HBs (1.862 X 1.229 mUI/mL) entre os dois grupos. Este estudo demonstra uma resposta sorológica satisfatória em lactentes quando a vacina da hepatite B é aplicada no glúteo.

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REFERENCES


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