Seroprevalence of Anti-EBV IgG among Various Age Groups from Khon Kaen Province, Thailand

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Abstract

Epstein-Barr virus (EBV) is an extremely common herpesvirus that may cause asymptomatic infection or various diseases, including infectious mononucleosis, certain lymphoproliferative diseases and several types of neoplasms. Vaccine development is an important strategy to reduce the burden of EBV-associated diseases and the timing of vaccinations should be before primary infection occurs. In the past, more than 90% of Thai children were infected with EBV in early childhood. Now, due to the improved healthcare system in Thailand, we aim to determine current prevalence of EBV infection among people in different age groups. A total of 538 sera were collected from residents of Khon Kaen province in northeastern Thailand for detecting anti-EBV IgG. Sera of infants under 2-years-old were also tested for anti-EBV IgM and EBV-DNA. The prevalence of anti-EBV IgG was 47.1% (95% CI: 23.3-70.8) in infants aged 0-6 months, 34.9% (95% CI: 23.1-46.7) in those aged 6-24 months, 87.9% (95% CI: 79.5-96.3) in children aged 3-5 years and then maintained at above 95% through adulthood. These seropositivity rates among Thai children remain similar to those found in a previous study conducted 20 years ago. Thai children are still exposed to EBV from an early age. Therefore, a prophylactic vaccine should be given within the first two years of life.

Keywords: Seroprevalence - anti-EBV IgG - children - adult - Thailand

Introduction

Epstein-Barr virus (EBV), a member of the Herpesviridae family, is a major cause of infectious mononucleosis in young children and adolescents (De Paschale and Clerici, 2012). More than 90% of the adult population worldwide has antibodies against EBV (Hsu and Glaser, 2000). Once infected, lifelong immunity usually persists. Nevertheless, immune responses are unable to eliminate the virus, resulting in EBV latency in memory B lymphocytes. Clinical manifestations of EBV infections vary according to age, ranging from silent infections in infants to infectious mononucleosis in adolescents and adults. In addition, many serological and molecular studies have provided evidence that EBV is associated with various types of malignancies such as lymphoma, nasopharyngeal carcinoma (NPC), Hodgkin disease and gastric carcinoma (Cohen et al., 2001; Hjalgrim et al., 2000; Mitarnun et al., 2004; Piriou et al., 2012; Tiwawech et al., 2005).

The seroprevalence of EBV antibodies increases with age but varies across populations according to geographic region, ethnicity and socioeconomic status (Balfour et al., 2013; Condon et al., 2014; Dowd et al., 2013; Kagro et al., 1994). In the United States, a seroprevalence survey from 2009 found that around 50% of children between 6-8 years had antibodies against EBV (Balfour et al., 2013). It was estimated that about half of the infections occurred in late childhood, causing high numbers of infectious mononucleosis cases.

In contrast, an EBV seroepidemiology survey conducted in Thailand from 1997-1998 found that more than 90% of Thai children between 6-8 years had immunity to the EBV viral capsid antigen (Pancharoen et al., 2001; Poovorawan et al., 1997; Pancharoen et al., 2001). With respect to the improvement of the healthcare system in Thailand during the past decade, we estimated a higher seronegativity rate among infants and young children and designed this study to determine the current status of EBV seroprevalence across age groups. The results of this study will help to identify a target age group for future prophylactic vaccines.

Materials and Methods

The proposal of this study was approved by the Institutional Review Board of the Faculty of Medicine, Chulalongkorn University (IRB No 526/57).
The sera analyzed in this study were collected in 2014 during research on the impact of universal hepatitis B immunization on newborns as part of the Expanded Program on Immunization (EPI). The age of the subjects ranged from birth to 57 years. All subjects lived in Khon Kaen province in the northeast of Thailand. A total of 538 samples were randomly selected to be used in this study. The subjects had previously been informed by the further study of their sera via written informed-consent.

Laboratory tests

Serological testing: The sera were stored at -20°C at the Center of Excellence in Clinical Virology, Department of Pediatrics, Faculty of Medicine, Chulalongkorn University until used for testing. The presence of anti-EBV-CA (Epstein-Barr virus capsid antigen) IgG was detected by quantitative analysis using commercially-available ELISA kits (EUROIMMUN, Luebeck, Germany) according to the provided instructions. Results were also interpreted according to those instructions. Samples were recorded as seropositive, seronegative or borderline according to levels of anti-EBV-CA IgG being above 22 RU/ml, below 16 RU/ml or between 16 and 22 RU/ml, respectively. All samples from children under 2 years of age were further tested for anti-EBV IgM using the same, commercially-available ELISA kits. Results of the IgM test were interpreted as positive, negative or borderline according to the included instructions.

Molecular study

In addition to IgM detection, sera from children under 2 years of age were subjected to viral DNA extraction using an Exogene Viral DNA/RNA kit (GeneAll, Seoul, Korea) according to the manufacturer’s instructions. Human herpesvirus DNA was amplified using nested consensus primers for herpes viruses as previously published (Sakulwira et al., 2004). The polymerase chain reaction (PCR) conditions were the same as in the previous study (VanDevanter et al., 1996) and the final PCR products were identified by electrophoresis in 2% agarose gel (FMC Bioproducts, Rockland, ME). The positive control was a tissue biopsy (nasopharyngeal carcinoma tissue) positive for EBV. The sensitivity of this method to detect EBV DNA is 100 copies per 100 ng of DNA (VanDevanter et al., 1996).

Data analysis

Analysis involved dividing sera samples into ten age groups in order to depict the age characteristics of EBV seroprevalence, especially during early childhood. Demographic data were presented in terms of the median, range and percentages. Statistical significance was determined by a $P = 0.05$. In addition, we reviewed the previous literature reporting the EBV seroprevalence of various countries in recent years to demonstrate trends of seropositivity across different age groups.

Results

Of the 538 participants, 267 (49.6%) were male and 271 (50.4%) were female. The median age was 13 years and ages ranged from 2 days to 57 years. The overall seropositivity rate, defined by an anti-EBV-CA IgG equal to or above 22 RU/ml, was 87.9% (95% CI 85.2-90.7). We interpreted those with borderline (between 16 and 22 RU/ml) or negative (below 16 RU/ml) ELISA results as seronegative. After organizing the subjects into age groups, the seropositivity rate of anti-EBV IgG in infants under 6 months was 47.1% (95% CI 23.3-70.8). Then the rate slightly decreased to 34.9% (95% CI 23.1-46.7) among children aged from 6 months to two years. In children 3-5 years of age, the seropositivity rate was significantly increased to 87.9% (95% CI 79.5-96.3). Afterward, the rate was maintained at above 95% through adulthood, as shown in Table 1.

Further tests to detect anti-EBV IgM and EBV DNA in the 80 samples from children less than 2 years of age found that 3 samples (3.8%) were positive for IgM and none were positive for EBV DNA. The 3 samples positive for anti-EBV IgM were from the infants aged 11, 13 and 18 months. The 11-month-old infant was also positive for anti-EBV IgG, whereas only IgM was detected in the 13-month-old and 18-month-old infants.

Table 1. Anti-EBV IgG Seropositivity Status by Commercially Available ELISA kit (n=538)

<table>
<thead>
<tr>
<th>Age groups</th>
<th>Number</th>
<th>Number</th>
<th>Percent</th>
<th>95% CI</th>
</tr>
</thead>
<tbody>
<tr>
<td>0-6 months</td>
<td>17</td>
<td>8</td>
<td>47.1</td>
<td>23.3-70.8</td>
</tr>
<tr>
<td>6 months-2 years</td>
<td>63</td>
<td>22</td>
<td>34.9</td>
<td>23.1-46.7</td>
</tr>
<tr>
<td>3-5 years</td>
<td>58</td>
<td>51</td>
<td>87.9</td>
<td>79.5-96.3</td>
</tr>
<tr>
<td>6-8 years</td>
<td>53</td>
<td>53</td>
<td>100</td>
<td></td>
</tr>
<tr>
<td>9-11 years</td>
<td>58</td>
<td>56</td>
<td>96.6</td>
<td></td>
</tr>
<tr>
<td>12-14 years</td>
<td>55</td>
<td>53</td>
<td>96.4</td>
<td></td>
</tr>
<tr>
<td>15-20 years</td>
<td>60</td>
<td>59</td>
<td>98.3</td>
<td></td>
</tr>
<tr>
<td>21-26 years</td>
<td>57</td>
<td>55</td>
<td>96.5</td>
<td></td>
</tr>
<tr>
<td>27-40 years</td>
<td>59</td>
<td>58</td>
<td>98.3</td>
<td></td>
</tr>
<tr>
<td>&gt;40 years</td>
<td>58</td>
<td>58</td>
<td>100</td>
<td></td>
</tr>
<tr>
<td>Total</td>
<td>538</td>
<td>473</td>
<td>87.9</td>
<td>85.2-90.7</td>
</tr>
</tbody>
</table>

Figure 1. The seroprevalence of EBV Infection in the Asian Countries: Hong Kong (Kangro et al., 1994), Bangladesh (Haque et al., 1996), Thailand in 1997 (Poovorawan et al., 1997), Thailand in 2014 (Current Study), Japan (Takeuchi et al., 2006), China (Xiong et al., 2014) and Taiwan (Chen et al., 2015). The data was Potted by the Mean age in Each Specific Age-Group in the Studies
Interestingly, the seropositivity rate of infants below six months of age was higher than in infants between 6 months and two years: 47.1% vs. 34.9%. This pattern has also been observed in other studies and is due to preexisting maternal antibodies active in infants under six-months-old (Leogrande et al., 1993). Longitudinal serological data have shown that maternal anti-EBV protection usually lasts for 3-4 months (Biggar et al., 1978; Chan et al., 2001). Once the level of maternal antibodies declines, infants begin to seroconvert. In our study, we observed that the seropositivity rate increased significantly to 87.9% in children aged 3-5 years. Thus, the majority of children in our study seroconverted during preschool age.

Among the 80 sera of children below two years of age, IgM antibodies were detected in only three samples (3.8%), collected from children aged 11, 13 and 18 months. The presence of anti-EBV IgM signifies an acute infection or a reactivation of the virus (De Paschale and Clerici, 2012; Nystad and Myrrem, 2007). Since we tested for IgM in young infants, it was very likely that a positive IgM was the result of an acute infection. Thus, our study showed that the earliest age to become infected was 11 months. However, anti-EBV IgM is difficult to detect because its levels decline quickly (Xiong et al., 2014) and it may persist at a very low concentration (Yuan et al., 2013). A previous study reported that the positive rate of EBV-IgM in the acute phase of the disease was only 25% in infants (Dohno et al., 2010). Therefore, it is possible that some Thai children were infected even before 11 months; we cannot demonstrate the earliest age of primary infection due to the lack of longitudinal data.

The molecular technique is another tool to help diagnose EBV infections with equivocal serological test results. During infectious mononucleosis, EBV virions and naked EBV DNA pass into the peripheral blood and can be detected in serum or plasma samples within 14 days after the onset of symptoms (De Paschale and Clerici, 2012; Kimura et al., 2008). However, EBV DNA can also be detected from reactivated EBV infections (Dohno et al., 2010) and little is known about the EBV DNA prevalence rate in asymptomatic infections. We performed PCR on the 80 sera samples from children under 2 years of age in order to detect acute infections where EBV DNA can be found in peripheral blood before anti-EBV IgG or IgM can be detected. However, no EBV DNA was found in our sera samples. This could have been due to the low viral load of asymptomatic infections, because our small sample size was too small to detect EBV DNA positive patients or because our test was not sensitive enough to detect EBV DNA in the sera samples from infectious mononucleosis (IM) patients. Previous studies have shown that the EBV viral load in EBV-associated IM was approximately 10^5 copies/ugDNA (Cheng et al., 2007). The lower limit of DNA detection in our amplification method was also around 10^3 copies/ugDNA. Therefore, another method, such as real-time PCR, may be more sensitive in detecting EBV DNA from low viral load samples.

While the seroprevalence of anti-EBV IgG in Thailand remained relatively the same during the past 20 years, changes in the prevalence of EBV antibodies in the United States have been observed comparing between 2003-2004

![Figure 2. The seroprevalence of EBV Infection in America and Europe Countries: USA in 2003-2004 and 2009-2010 (Balfour et al., 2013), South Italy (Leogrande et al., 1993) and UK (Kangro et al., 1994). The Data was Plotted by the Mean age in Each Specific Age-Group in the Studies](image)


Seroprevalence of Anti-EBV in Children and Adults of Different Ages in Thailand


