Gene EBNA3C: Type of Infection by EBV (EBV1 and EBV2) Correlation With Clinical and Biochemical Parameters (AST, ALT and GGT) in Individuals With Infectious Mononucleosis in the Metropolitan Area of Belém, Pará, 2005-2016

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Abstract

Background: Two types of EBV (EBV1 and 2) have been shown to infect humans. This study aimed to detect the types of EBV that cause infectious mononucleosis and correlate these viral types with biochemical parameters (AST, ALT and GGT) in the metropolitan region of Belém from 2005 to 2016. Methods: A total of 76 cases of infectious mononucleosis were processed at the Instituto Evandro Chagas, Ananindeua, Brazil. PCR was used to analyze the EBNA3C region for the recognition of EBV types. Biochemical testing (AST, ALT and GGT) was performed by the COBAS INTEGRA clinical biochemistry PLUS 400 / ROCHE automatic analyzer. The data were evaluated using the Statistical Package for Social Science - SPSS 17.0 and GraphPadPrism 7.0 for Windows (GraphPad software, San Diego, CA, USA). Results: EBV1 infection was observed in 71.1% (54/76) of individuals, among whom those > 14 years constituted 66.7% (36/54); the average age was 23 years, and the number of women infected was higher (61.1% 33/54) than that of men 38.9%21/54). The symptoms/clinical signs observed in infection by EBV1 were cervical lymphadenopathy in 64.8% (35/54), fever in 63% (34/54), headache and arthralgia in 20.3% (11/54), and exanthema in 18.5% (10/54). Infection by EBV2 was observed in only 17.1% (13/76) of cases. Coinfection by EBV1 and EBV2, most frequently showing symptoms of fever and cervical lymphadenopathy, occurred in 66.7% (6/9) and 55.6% (5/9) of individuals.

Conclusion: EBV1 was predominant in 71% of clinical cases of infectious mononucleosis. The correlation of biochemical parameters in infection by EBV1, EBV2, and coinfection by EBV1/2 revealed a statistically significant difference in mean changes of EBV1 in individuals older than 14 years.

Background

Epstein-Barr virus (EBV) belongs to the order *Herpesvirales*, family *Herpesviridae,*
subfamily Gammaherpesvirinae, genus Lymphocryptovirus and species Human
gammaherpesvirus 4. EBV was the first oncogenic virus found to infect humans. The
hexagonal nucleocapsid viral particles were formed with linear, double stranded,
enveloped DNA, with a diameter of 180 to 200 nm 4.
Although symptomatic infections with these viruses occur in benign form, EBV has been
implicated in the genesis of a variety of lymphoproliferative disorders and severe
epithelial neoplasms, such as African Burkitt's lymphoma, carcinoma and
nasopharyngeal 6.
In Brazil, several studies have recorded the high frequency of antibodies in the studied
populations. Studies conducted by Monteiro et al. (1998) found that at least 70% of the
serum samples analyzed in the city of Belém, state of Pará, contain IgG antibodies to EBV,
at the outpatient clinic level ranging from 53.8% to 95.6%, or in the community (81.1% to
100%) 7.
Positive indexes were expressive, even in the youngest lowest age groups. These results
suggest that active and recent infection (infectious mononucleosis) was detected in 10.6%
(25/234) of children and adolescents in northern Brazil 8.
Data from Young and Murray (2003) demonstrated that EBV is present in approximately
90% of individuals and is controlled by the immune system, mainly by cellular immunity,
which may make the subject more susceptible to virus proliferation and may trigger
lymphoproliferative disorders 9,10.
It is increasingly important to identify the epidemiological characteristics associated with
the risk of EBV infection in populations to reduce the clinical conditions associated with
possible morbidity and mortality 11.
The difference between the sequences encoding EBV nuclear antigens (EBNA2, 3A, 3B, and
3C) allows for the identification of different genotypes with distinct epidemiological
characteristics 12.

According to Young et al. (2000), human and geographic assemblages can influence the distribution of EBV genotypes 1 and 2, which change with rates of detection of these viral genotypes in diseases associated with EBV 13. Type 1 EBV has a higher incidence in western regions, and type 2 is found more frequently in sub-Saharan Africa and New Guinea than in other parts of the world 14.

The phenotypic difference between EBV1 and 2 is more evident during immortalization of B cells in EBV1 lymphoblastoid cell lines (LCLs) compared to EBV2. This fact reinforces the biological and functional difference between the two viral types, where B cell immortalization in vitro was shown to be more effective by type 1 EBV 15. According to Mandell et al. (2001), differentiation of genotypes can clarify the different immune responses during viral persistence 16.

The metropolitan area of Belém still lacks studies to characterize the different types of circulating EBV (EBV1 and EBV2) associated with clinical and demographic characteristics (gender, age and origin) and molecular findings that describe the epidemiology of the infection caused by these viral agents due to its characteristics of viral persistence that can induce chronic infections and reactivations in human populations with genetic competence to possible oncogenic events.

The objective of this study was to detect the types of Epstein-Barr virus (EBV1 and EBV2) that cause infectious mononucleosis (IgMVCA/EBV+) and correlate them to biochemical parameters (ASL, ALT and GGT) from the metropolitan area of Belém between 2005 and 2016.

Methods

Type of study
The current study was descriptive and retrospective. Samples from 76 clinical cases of infectious mononucleosis that were reactive to EBV/VCA IgM, were tested by the RIDASCREEN® (R-Biopharm, Darmstadt, Germany) enzyme immunoassay kit and processed in the Virology section and the biochemical dosages (AST, ALT and GGT) were processed in the pathology section of the Evandro Chagas Institute, Ananindeua, Brazil, from the metropolitan area of Belém. The samples were collected between 2005 and 2016 and later evaluated by PCR for the EBNA3C gene for recognition of EBV types.

Identification of the EBNA3C gene through PCR

Nucleic acid extraction

Samples were extracted using the QIAamp DNA Mini Kit (Qiagen, Germantown, MD) according to the manufacturer’s protocol.

Identification of the EBV EBNA3C gene

For the identification of EBV type, we used the EBNA–3C primer 17. Due to the flanking regions of the type-specific variation primary sites, the resulting PCR products were of two different sizes for EBV1 (153 bp) and EBV2 (246 bp). Five microliters of the eluted ADN were used for PCR amplification with a primer concentration of 0.05 μM (5’ GCCAGAGGTAAGTGGACTTT 3’ and 5’ TGGAGAGGTCAGGTTACTTA 3’, respectively). PCR was performed in 0.5 mL microcentrifuge tubes in a final 25 μL mixture containing 0.125 μL (5 U / μL) Platinum Taq DNA Polymerase (Invitrogen, Brazil), 1.5 mM MgCl 2 (Invitrogen™, Brazil), 0.2 mM dNTPs (Invitrogen™, Brazil), 5 μL of 10X buffer (Invitrogen™, Brazil) and 2 μL (20 μM / μL) of the abovementioned primers. After denaturation of the DNA mold at 94°C / 1 min, PCR cycle conditions (PTC 100 / Peltier Effect Cycle, Thermostable Controller) included 40 cycles of denaturation at 94°C / 30 sec, annealing at 58°C / 30 sec, extension at 72°C / 1 min, after 1 final extension cycle at 72°C / 7 min. Water
(negative control) and B958 and P3HR1 cell lines (positive controls) were used for the amplification of EBV1 and EBV2 in each amplification series.

Ethical Considerations

The research study was approved by the research ethics committees of the Evandro Chagas Institute (CAAE n° 65332717.2.0000.0019, with legal opinion n° 2098453, dated June 4, 2017).

Data Analysis

The results were organized and stored in a database in Microsoft Office Access 2016, *Statistical Package for Social Science* - SPSS 17.0 and GraphPadPrism 7.0 for Windows (GraphPad software, San Diego, CA, USA) 18.

Results

Detection of EBV 3C Gene of the EBV

EBV DNA was detected in 76 patients with infectious mononucleosis by PCR for the EBNA3C gene. In total, 55.3% (42/76) were female, and 44.7% (34/76) were male. The patients were from the cities of Belém with 76.3% (58/76), Ananindeua with 22.4% (17/76) and Marituba with 1.3% (1/76).

As for the clinical picture, a multiplicity of signs and symptoms were noticed, such as fever in 65.8% (50/76), cervical lymphadenomegaly in 60.5% (46/76), pharyngitis in 19.7% (15/76), arthralgia in 17.0% (13/76), and headache in 9.2% (7/76). For fever days, 22% (11/50) reported their permanence for up to 5 days, 16% (8/50) for 10 days, 20% (10/50) for 15 days, 16% (8/50) for 20 days, 16% (8/50) for more than 20 days, and 10% (5/50) had no information in the epidemiological record.

EBV1 infection was observed in 71.1% (54/76); the most frequent age group was >14 years old, with 66.7% (36/54) of the individuals. Since the age range was 23 years, the
number of women was 61.1% (33/54), which was higher than in men, with 38.9% (21/54).

Regarding the origin of the individuals with EBV genotype 1, the frequencies were as follows: city of Belém with 76.0% (41/54), Ananindeua with 22.2% (12/54) and Marituba with 1.8% (1/54).

The symptoms/clinical signs observed in infection by EBV1 were cervical lymphadenopathy in 64.8% (35/54), fever in 63% (34/54), headache and arthralgia in 20.3% (11/54), and exanthema in 18.5% (10/54) (Fig 1).

In EBV2 infection, 17.1% (13/76) cases were observed, where the most frequent symptom was fever in 76.9% (10/13). The individuals were distributed in only two ranges, being > 14 years old, with 53.8% (7/13) individuals, compared to 6 to 14 years old, in 46.2% (6/13) individuals; the average age was 24 years, with the males being more frequent 76.9% (10/13) than females 23.1% (3/13).

Confection with EBV1 and EBV2 is more frequent in children and cervical lymphadenopathy in 66.7 (6/9) and 55.6% (5/9), respectively. The most frequent age group was > 14 years, with 44.4% (4/9) of the individuals. Since the age group of 2 to 5 years presented 22.2% (2/9) individuals and 6 to 14 years with 33.3% (3/9) individuals, with an average age of 21 years, the number of women of 66.7% (6/9) was higher than that of men with 33.3% (3/9).

In the assessment of hepatic function, alterations to AST in EBV1 infection were confirmed in 14.8% (8/54) of cases with results above the reference values (5–40 U/L) and 7.7% (1/13) for EBV2. It is worth mentioning that the reference values are the same in the three age groups (Fig 2). When analyzing the mean values by age group, the range of >14 years was outside the normality limits with a higher frequency of iterations, thus presenting a statistically significant difference (p-value < 0.05).

Fig 2. Evaluation of ALS of individuals with EBV 1, EBV 2 and EBV 1 and 2 infection.
Values above the reference values (2–41 U/L) were also observed for ALT (Fig 3). The mean value was outside the limits of normality in the age groups of 6 to 14 years and >14 years, presenting a significant difference (p-value < 0.05) by the *Wilcoxon* test.

Fig 3. Evaluation of AST of individuals with EBV 1, EBV 2 and EBV 1 and 2 infection.

For the evaluation of GGT, some altered results were observed, but no changes were found in the age group of 2 to 5 years, given the reference values: (5 - 55 U/L). Changes in the age group from 6 to 14 years were all above normal, given the same reference range of the previous age group, and the greatest number of alterations was observed in the range of >14 years, above or below, given reference values: (12 - 43 U/L) (Fig 4). The mean value was outside the limits of normality in the age group of > 14 years, and the mean values presented a significant difference (p-value <0.05) by the *Wilcoxon* test.

Fig 4. Evaluation of GGT activity in individuals with EBV 1, EBV 2 and EBV 1 and 2 infection.

**Discussion**

Primary EBV infection is usually asymptomatic and may progress to benign lymphoproliferative disease called infectious mononucleosis (IM), especially in late childhood or early adulthood in developing countries 4.

Infectious mononucleosis is characterized by significant clinical polymorphisms in which factors such as age, immune status and comorbidities have been described as parameters in clinical evolution from asymptomatic infections to more severe conditions evidenced by acute complications, such as multiple organ failure, disseminated intravascular coagulation, ulcer/perforation of digestive tract, coronary artery aneurysm, lymphomas and lymphohistiocytes and EBV-associated hemophagocytosis 19, 20.

Mendoza et al. (2008) confirmed that EBV infection has an incubation period ranging from 4–6 weeks with prodromal symptoms of asthenia, anorexia, headache and chills, which
often precede the signs and symptoms of mononucleosis: fever (that can reach 39–40°C) accompanied by pharyngotonsillitis and lymphadenopathy 21, 22, 23. In agreement, in our study, fever was the clinical finding in 65.8% (50/76), cervical lymphadenomegaly in 60.5% (46/76), pharyngitis in 19.7% (15/76), arthralgia in 17.0% (13/76), and headache in 9.2% (7/76) of the 76 patients analyzed.

Regarding the type of EBV infection, there are two different types of EBV 24. EBV types are related to variation in the EBNA2 and EBNA3 gene sequences, commonly known as types 1 and 2 17, 25.

Studies conducted in other countries have demonstrated the predominance of EBV1 infection in China with rates of 76.3%, Argentina in 75.9%, Sweden in 67% and Hong Kong 57% 26, 27, 28, 29.

Our findings revealed that EBV1 was frequent in 71.1% (54/76) of cases of infectious mononucleosis from the metropolitan area of Belém, Brazil. A total of 68 samples of Chinese individuals were studied with the aim of identifying circulating types according to the EBNA 3C gene by PCR. In total, 76.3% (45/59) of samples were EBV-1, 20.3% (12/59) were EBV-2, and 3.4% (2/59) EBV-1 and EBV-2 (coinfected), and 13.2% (9/68) of the samples did not amplify.

A study conducted by Deng et al. (2014) with samples of 209 Japanese patients obtained the following results: 146/209 (69.9%) samples had EBV, 107/146 (73.3%) were EBV-1, 27/146 (18.5%) were EBV-2 and 12/146 (8.2%) were coinfectected (EBV-1 and EBV-2), and 63/209 (30.1%) did not amplify the EBNA3C gene 30.

A study conducted in Qatar revealed similar frequencies with a predominance of EBV1 (72.5%, 37/51) compared to genotype 2 (3.5%), and mixed infections were detected in 4% of the samples 31. The determination of the types of EBV in the present study made it possible to distinguish the molecular epidemiology and circulation of these viral agents.
Hepatic Evaluation

The most clinically relevant transferases are aminotransferase (aspartate amino transferase - AST, alanine amino transferase - ALT and γ-glutamyl transferase - GGT), which express the main indexes of liver function 32, where small changes may occur in normal individuals (less than twice the reference value). In patients with infectious mononucleosis caused by EBV, values up to 5 to 10 times higher than the reference values have been reported and may even progress to fulminant hepatitis, which is not present with bilirubin abnormalities 33, 34.

Ninety-five patients with infectious mononucleosis and 95 healthy controls were analyzed for AST, ALT and GGT; alterations were elevated in patients with infectious mononucleosis compared to the controls 34.

When compared to the EBV types (EBV1, EBV2 and EBV1/2), our results were statistically significantly correlated with the age group and AST and ALT values (p <0.005). Similar data were cited by Zhang et al. (2018), who reported that ALT, AST and GGT levels were significantly increased in cases of infectious mononucleosis compared to controls, indicating that transferase levels can be used to diagnose and treat as a risk alert for the infection caused by MI34.

Conclusion

This pioneering study identified EBV1 in 71.1% of clinical cases of infectious mononucleosis in the metropolitan area of Belém, Pará, from 2005 to 2016. In 11.8% of the cases, coinfections with EBV1 and EBV2 occurred. Cervical lymphadenopathy and fever were the most relevant clinical findings and signs in the EBV types. The correlation of biochemical parameters (AST, ALT and GGT) with type of infection by EBV1, EBV2, and by EBV1/2 revealed a statistically significant difference in mean changes of EBV1 in
individuals older than 14 years.

Abbreviations

EBV: Epstein-Barr virus; LCLs: lymphoblastic cell lines; AST: aspartate amino transferase; ALT: alanine amino transferase; GGT: y-glutamyl transferase; MI: infectious mononucleosis.

Declarations

Competing interests
There authors declare that there are no competing interests.

Authors’ contributions
The work presented here was carried out as a collaboration between all authors. TAFM, IBC, IBC, TLSC, BMRC, AAP, AESS, FLPR, and AJMF carried out most experiments. TAFM and JLFM made contributions to design, data analysis and data interpretation. TAFM and RCMS drafted the manuscript. JLFM and TAFM provided the most financial support. TAFM, AAP and AESS collected and assembled the data. All the authors have given final approval to publish the manuscript.

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Availability of data and materials
All data generated or analyzed during this study are included in this published study. No other data are available for supplementary materials.

Ethics approval and consent to participate
All participants provided informed written consent for all study procedures and for the use
of their data for scientific evaluation and publication in a blinded form. This study was conducted in accordance with the Declaration of Helsinki, and it was approved by the Evandro Chagas Institute with legal opinion no 2098453, dated June 4, 2017).

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Figures
Figure 1

Signs and symptoms of infectious mononucleosis in patients infected with EBV-1, 2005 to 2016.

Figure 2

Evaluation of ALS of individuals with EBV 1, EBV 2 and EBV 1 and 2 infection.
Figure 3
Evaluation of AST of individuals with EBV 1, EBV 2 and EBV 1 and 2 infection.

Figure 4
Evaluation of GGT activity in individuals with EBV 1, EBV 2 and EBV 1 and 2 infection.