Association between Epstein-Barr Virus Gene Polymorphism and Breast Cancer Risk among Egyptian Females

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Abstract

Background: Epstein-Barr virus (EBV) has been implicated in the development of breast cancer (BC) since 1995. It is classified into A/B genotypes, C/D subtypes, and F/f variants according to variations in its genome. Aim: To determine the distribution difference of EBV types between BC patients and healthy controls in Egypt and to detect the association between different EBV types and BC characteristics. Methods: Three hundred and sixty-two participants (142 BC patients and 220 controls) were enrolled in this study. All participants were screened for EBV infection by determination of viral-capsid-IgG antibodies in their sera. EBNA-1 gene was detected by PCR in tumor biopsies of seropositive patients and in peripheral blood mononuclear cells of controls. A/B genotyping of EBV was performed by nested-PCR targeting the EBNA-2 gene. C/D subtypes and F/f variants were identified by Restriction fragment length polymorphism at BamHI-I W1/I1 and BamHI-F regions of EBV genome, respectively. Results: Among 362 participants, 300(82.9%) were EBV-seropositive, including 120/142(84.5%) of the BC patients and 180/220(81.8 %) of the controls. EBNA-1 gene was positive in 54(45%) of seropositive BC patients and in 38(21.1%) of seropositive controls. There was a significant association of EBNA-1 gene with breast cancer (OR=3.05, 95%CI=1.84-5.07). Moreover, EBNA-1 gene positivity was significantly associated with the more aggressive tumors. Genotype-A and prototype-F were predominant among patients (90.4%, 100%, respectively) as well as among controls (91.7%, 100%, respectively) with no statistical significant association with BC risk. However, subtype-D was significantly more frequent in patients (95.6%) than in controls (64.7%) and was significantly associated with a higher BC risk as compared to subtype-C (OR=11.7, 95%CI=2.4-57.08). Subtype-D was significantly associated with higher grades tumors (100% among grade III), with progesteron receptor-negative tumors and with HER2-positive tumors (100% for each). The combined genotypes that significantly associated with BC risk were ADF (OR=4.9) and BDF (OR=5.5). Conclusions: Subtype-D of EBV could be the only EBV type implicated in BC development among Egyptian females and associated more with poor prognosis.

Keywords: Epstein-Barr virus- breast- cancer- RFLP- genotyping- Egypt

Introduction

Epstein-Barr virus (EBV), the etiological agent of infectious mononucleosis, is a Herpes virus that belongs to the Herpesviridae family. It is highly widespread, affecting more than 90% of the world’s population (Dowd et al., 2013).

EBV genome is 172,000 base pairs long and although it can code for more than 85 proteins, the well-known ones are few; Six EBV nuclear antigens (EBNAs 1, 2, 3A, 3B, and 3C, and EBNA-LP); three latent membrane proteins (LMPs 1, 2A, 2B), also known as latent genes , and small noncoding EBV-encoded RNAs (EBERs 1 and 2) (Kalla and Hammerschmidt, 2012).

The presence of polymorphisms in several regions of the EBV genome define different EBV types and variants, which have been shown to have a distinctive geographical distribution (Ayadi et al., 2006).

EBV is classified into two genotypes: genotype-A and genotype -B (sometimes known as type-1 and type-2). EBV Genotypes-A and-B were initially distinguished based on changes in the EBNA2 sequence ,which only shows 70% of gene homology and 54% of protein homology between genotype-A and genotype -B (Farrell, 2015). Although Palser and his colleagues observed that differences between genotype-A and genotype -B could be caused by variations in the sequences of EBNA2 and EBNA3A, EBNA3B, and EBNA3C genes, they found
that the most frequent variance between genotype-A and genotype-B is the differences in the EBNAA2 gene (Palser et al., 2015).

EBV BamHI-F and BamHI-W/ Bam HI-I boundary regions in the EBV genome have been identified as BamHI endonuclease restriction sites. Therefore, restriction fragment polymorphism (REFLP) analysis would help to further classify each of genotype-A and genotype-B into C/D subtypes and into F/f variants based on the presence or lack of BamHI restriction sites in the previously mentioned regions. Subtype-C is distinguished by the absence of a BamHI restriction site between the BamHI-W and BamHI-I fragments, while subtype-D has an extra BamHI site in the same region. In the BamHI-F fragments, prototype-F has no BamHI site, whereas variant-F has an extra BamHI site (Ayadi et al., 2007).

Diagnosis of EBV infection depends on a variety of tests, including nonspecific tests like the monospot test for heterophile antibodies, serological tests for detection of EBV-specific antibodies and molecular assays for detection of nucleic acid (Hess, 2004). Viral capsid antigen (VCA)-IgG, VCA-IgM, and EBNA-1 IgG are frequently required for the detection of EBV antibodies. VCA-IgG antibodies are superior to VCA-IgM in diagnosing past EBV infections. Furthermore, approximately 5–10% of healthy people infected with EBV never produce EBNA-IgG and this percentage is higher in immunodeficient people. Therefore, VCA-IgG antibodies are the most accurate single test for detection of past infection with EBV (Smatti et al., 2018). EBNA1 gene is constantly present in all EBV-infected cells. As a result, it is used for molecular detection of EBV infection (Sun et al., 2015).

Breast cancer (BC) is the most frequently diagnosed malignancy as well as the leading cause of cancer death among women, worldwide. In accordance to WHO estimates, BC would be diagnosed in 2.1 million women every year. According to reports in 2018, over 627,000 women died from the disease, accounting for almost 15% of all cancer-related fatalities among women, worldwide (Farahmand et al., 2019). BC is a complex disease with an aetiology that is poorly understood. Many traditional factors were identified as risk factors for BC; family history, early menarche, late menopause, obesity and nulliparity (Kaminska et al., 2015).

In 1995, Labrecque and his colleagues were the first to identify the association between EBV and BC, and since then, many studies have confirmed the role of EBV in the development of BC (Labrecque et al., 1995). Recently, there has been an increasing interest to specify the association between different types of EBV and malignant diseases (Neves et al., 2017). However, the majority of these studies were focused on EBV-associated nasopharyngeal carcinoma (NPC) and EBV-associated gastric carcinoma, and they proposed that the pathological effects of EBV might vary according to the type of EBV strain (Corvalan et al., 2006; Chen et al., 2010).

To date, most of the researches that studied the relationship between EBV and breast carcinoma were interested in investigating whether or not there is an association between EBV infection and the development of BC, and none of these studies determined the association between different types of EBV and BC. Therefore, we investigated the variations in the distribution of different types, subtypes, and variants of EBV between BC women and healthy controls in Egypt and to clarify the relationship between specific EBV types and the histopathological features of BC.

Materials and Methods

Study subjects

This case-control study included 362 female participants; 142 breast cancer (BC) patients and 220 healthy controls. All patients and controls were recruited from the Oncology Center, Faculty of Medicine, Mansoura University (OCMU) in Egypt through the period from January 2019 to December 2020. The ages of all patients and controls were matched (p >0.05). Patients’ ages ranged from 37 to 79 years old with a median age of 54 years. The diagnosis of BC in all enrolled patients was based on clinical examination, radiological findings, and histopathological examination of tissue biopsy. Control subjects were selected from women who attended OCMU for annual mammography as part of a breast checkup and we excluded those with a history of BC and those with fibroadenomas.

Sample collection

Fresh tumor tissue and blood specimen were obtained from BC patients, whereas only blood specimen was obtained from the control group. The tissue sample from each tumor was divided into two parts; one part was fixed in formaldehyde and delivered to OCMU’s pathology laboratory for histopathological examination, while the other part was stored at -80°C to be used for PCR. The blood specimen was obtained via venipuncture and divided into two tubes; a plain tube for separation of serum and an EDTA tube for separation of peripheral blood mononuclear cells (PBMCs), both of which were kept at –80°C until use.

Study design

A Flowchart showing the workflow of this study is presented in Figure 1. All included BC patients and controls were screened for EBV infection by the detection of anti-viral capsid antigen (VCA) IgG antibodies in their sera. Then EBV seropositive subjects were investigated for the presence of EBV DNA by PCR amplification of the EBNA-1 gene; either in tumor biopsy of BC patients or in PBMCs of controls. Thereafter, A/B genotypes, C/D subtypes and F/f variants of EBV were identified in all EBV-DNA positive specimens by analysis of three regions in the EBV genome (EBNA-2 gene, BamHI-I W1/I1 and BamHI-F regions, respectively).

Histopathological examination

Diagnosis of BC and determination of its histological type was achieved in OCMU’s pathology laboratory by examination of hematoxylin-eosin (H-E) stained slides prepared from tumor biopsies. Only tumor biopsies with cancerous cells of more than 60% were included in the study. Grading of the tumors was performed according to the Scarff Bloom Richardson classification (Bloom...
Detection of EBV-viral capsid antigen (VCA) IgG antibodies

Screening of all included BC patients and controls for EBV infection was performed by detection of anti-VCA IgG antibodies by ELISA (Bio-Rad Medical Diagnostics, GmbH, Germany) following the manufacturer guidelines.

Polymerase chain reaction (PCR) amplification of EBNA-1 gene

Detection of EBV DNA was performed by PCR amplification of EBNA-1 gene. DNA was extracted from all specimens with serologically confirmed EBV infection (120 tumor biopsies from BC patients and 180 PMNCs from controls) using the QIAamp DNA Mini Kit (Qiagen, GmbH, Hilden, Germany), following the manufacturer’s tissue and blood protocols. Then EBV DNA was detected by PCR amplification of the EBNA-1 gene using specific primers listed in Table 1 according to the previously described PCR protocol (Hashimoto et al., 1995).

Detection of A/B genotypes of EBV

Nested PCR that targets EBNA-2 gene was used for genotyping of EBV into genotype-A and genotype-B (Ayee et al., 2020). In the first round of PCR amplification, a common region of EBNA-2 was amplified using EBNA-2A and EBNA-2B as forward and reverse primers, respectively. A PCR reaction was performed in a total volume of 12.5 µl and the final concentrations of each component were as follows: 1X One TaqR Quick-Load 2X master mix with standard buffer (Bio Labs, UK), 0.5 µM from each of the forward and reverse primers, and 0.5 µg/µL of the extracted DNA. Amplification was achieved in a thermal cycler under the following conditions: 2 min at 94°C for initial denaturation, then followed by 35 cycles of amplification (each cycle consisted of 60 s at 94°C, 90 s at 52°C and 4 min at 72°C), followed by a cycle of 10 min at 72°C for final extension.

In the 2nd round of nested PCR, the reaction volume was 12.5 µl and the components were the same as in the 1st round except that 0.5 µl of the amplified product from the 1st round was used as a template and three primers were used; EBNA-2C as a forward primer (common to genotype-A and genotype-B), EBNA-2D and EBNA-2E as reverse primers specific for genotype-A and genotype-B, respectively. The cycling conditions were as follows: An initial cycle of heat denaturation for 2 min at 94°C, 35 amplification cycles (30 s at 94°C, 60 s at 52°C, and 2 min at 72°C), and a final extension cycle for 10 min at 72°C. The resultant amplicon was visualized by electrophoresis on an agarose gel with 2% ethidium bromide. Samples were identified by the presence of their respective bands (250 bp for genotype-A and 300 bp for genotype-B).

Statistical analysis

The Statistical Package for Social Sciences (SPSS) version 22 was used to analyze the data. The qualitative data were expressed as frequencies and percentages. The differences in distribution of EBV types between tissue biopsies of BC patients and sera of the control group were compared by a univarite binary logistic regression test with the calculation of odds ratio (OR) and 95% confidence intervals (CIs). The difference between groups was considered statistically significant if the p-value was less than 0.05.

Results

Detection of anti-VCA IgG of EBV by ELISA

Screening of 142 BC patients and 220 healthy females for past exposure to EBV infection revealed that 120/142 (84.5%) of the cases and 180/220 (81.8%) of the controls were seropositive for anti-VCA IgG with no statistically significant difference between them (p=0.5), Table 2.

Detection of EBV DNA by PCR

EBV-DNA was detected in 54/120 (45%) of BC tissue biopsies and in 38/180 (21.1%) of control PBMCs. There was a significant association between BC and EBV-DNA positivity (OR=3.05, 95%CI: 1.84-5.07, p<0.001), Table 2.
Genotyping of EBV

Analysis of A/B types of EBV

Successful amplification of the EBNA-2 gene was observed in 52/54 (96.3%) of BC patients and in 36/38 (94.7%) of controls. Nested PCR of the EBNA-2 gene revealed that genotype-A EBV was more frequently detected than genotype-B EBV in both BC patients (90.4% vs. 9.6%) and healthy control groups (91.7% vs. 8.3%). There was no statistically significant difference in the distribution of genotypes-A and -B between BC patients and the control group (p=0.8), Table 3.

Analysis of C/D subtypes of EBV

Amplification of the BamHI W1/I1 boundary region was successful in 45/54 (83.3%) of BC patients and in 34/38 (89.5%) of controls. Among 45 BC patients with successfully amplified BamHI W1/I1 region, subtype-D was detected more frequently than subtype-C (95.6% vs. 4.4%). Similarly, in the control group, subtype-D was more frequent than subtype-C (64.7% vs. 35.3%). There was a statistically significant difference in the distribution of subtypes-D and-C between BC patients and the control group (p=0.002). Subtype-D EBV was significantly more frequently associated with tissue biopsies from breast tumors (95.6%) than with PMNCs from controls (64.7%) (OR=11.7, 95% CI=2.4-57.1), Table 3.

Analysis of F/f variants of EBV

The BamHI-F region was amplified successfully in 48/54 (88.9%) of BC tumor biopsies and in36/38 (94.7%) of controls. The successfully amplified EBV from all BC patients and controls was of F-variant, Table 3.

Combined Genotypes of EBV

A comparison of the combined genotypes of EBV between BC patients and controls revealed that the most predominant combination of EBV genotypes among BC patients as well as among the control group was ADF (84.6% vs. 62.5%, respectively). In BC patients, BDF was the second most frequent combined genotype (10.3%), followed by ACF and BCF (6.2% for each). Whereas in the control group, the second most frequent combined genotype was ACF (28.1%), followed by BDF (6.2%) and BCF (3.1%). The distribution of the EBV genotype combination between BC patients and controls was statistically significant (p=0.022). The combined genotypes that significantly associated with BC risk were ADF (OR=4.9, 95% CI=1.61-14.99) and BDF (OR=5.5, 95% CI=1.25-24.41), Table 3.

Association between EBV DNA positivity and tumor characteristics

Among 120 seropositive BC patients, there was a

Table 1. Target Genes and Their Primer Sequence

| Target gene | Primer sequence | Product size (bp) |
|-------------|-----------------|------------------|
| EBNA-1      | F: 5’CACTTTAGAGCTCTGGAGGA-3’  R: 5’TAAAGATAGCGACGAGCAG-3’  | 112 |
| EBNA-2      | First round primers |                |
|            | EBNA-2A  F: 5’TGGAAACCCGTCGACTCT-3’  | 801 |
|            | EBNA-2B  R: 5’TAAAGATAGCGACGAGCAG-3’  | |
|            | Second round primers |                |
|            | EBNA-2C  F: 5’AGGGATGCGCTGGACACAAGA-3’  | |
|            | EBNA-2D  R: 5’GCCTGGGTTGTGACAGAG-3’  | Genotype-A: 250 |
|            | EBNA-2E  R: 5’TGAAGAGATATGCTCTAGGAG-3’  | Genotype-B: 300 |
| BamHI-I W1/I1 region | F: 5’ACCTGATCGACTCTTGCGAAAC-3’  R: 5’TCTGCTCAACACCTCACTGT-3’  | Subtype-C: 205 |
|            | BamHI-F region  F: 5’TCCCCAGCTGTTACCACATTTC-3’  R: 5’GGCAATGCGGACGTCTTGGTA-3’  | Prototype-F: 198 |

Table 2. Frequency of EBV Infection in Breast Cancer Patients and Controls

| anti-VCA IgG (n) | Control group No. (%) | BC patients No. (%) | P-value | OR (95% CI) |
|------------------|-----------------------|---------------------|----------|-------------|
| Anti-               | 220 (21.1)            | 54 (45)             | <0.001   | 3.05 (1.84-5.07) |
| VCA                 | Seronegative          | 40 (18.2)           | 22 (15.5) | Reference   |
| Seropositive        | 180 (81.8)            | 120 (84.5)          | 0.507    | 1.2 (0.68-2.14) |
| EBNA-1 gene (n)     | 180                   | 120                 |          |             |
| Negative            | 142 (78.9)            | 66 (55)             |          | Reference   |
| Positive            | 38 (21.1)             |                     |          |             |

EBV, Epstein-Barr virus; VCA, Viral capsid antigen; EBNA, Epstein Barr virus nuclear antigen; BC, breast cancer.
significant association between detection of EBV-DNA (EBNA-1 gene) and each of the large tumor size and higher tumor stage (P = 0.001 for each). Also, positivity of EBV-DNA was significantly associated with carcinomas of infiltrating ductal type (75%), and with tumors of high histological grade (71.4%, 50%, and 14.3% for grades III, II, and I, respectively). Moreover, the proportion of EBV DNA positive tumors was significantly higher in metastatic tumors (p=0.001). On the other hand, there was no observed association between EBV-DNA detection in BC tissues and steroid hormone receptors. However, EBV-DNA was detected significantly more frequently in HER2-positive tumors than in HER2-negative tumors (75% vs. 25%, p= 0.001), Table 4.

Association between EBV genotypes and tumor characteristics

A/B genotypes of EBV did not show significant associations with any of tumor characteristics. Similarly, no association of statistical significance was observed between the C/D subtypes of EBV and either tumor size, tumor stage, tumor histology, presence of metastasis, or oestrogen receptor. However, the C/D subtypes of EBV from BC biopsies showed significant associations with tumor grade, progesterone receptor, and HER2- positive
tumors. The proportion of subtype D positive BC biopsies increased significantly with tumour grade, from 86.7% for grade II to 100% for grade III. Although the D-subtype of EBV was detected more frequently in progesterone receptor-negative tumors (100%) than in progesterone receptor-positive tumors (86.7%), it was significantly more frequent in HER2-positive tumors than HER2-negative tumors (100% vs. 83.3%, p= 0.01), Table 5.

Discussion

The presence of different polymorphisms in certain regions of the EBV genome resulted in the presence
of different genotypes, subtypes, and variants of EBV (Ayadi et al., 2007). However, little is known about how EBV genetic diversity may affect EBV-associated disease (Palser et al., 2015). Moreover, up to date, there is a controversy whether these EBV variants are geographically related, or disease-associated. In 1995, the association of EBV with BC was reported but no study has been conducted in Egypt since then to investigate if there is a specific association of BC with certain types of EBV or if it is simply a geographical distribution.

In this study, the seroprevalence of EBV was 84.5% in BC patients and was 81.8% in the control group with no statistical significant difference. This is consistent with the high prevalence of EBV in the world. It is estimated that more than 90% of the world’s population is EBV-seropositive (Tzellos and Farrell, 2012). Smatti et al., (2017) reported that the EBV seroprevalence rate among healthy blood donors was 100% in Egyptians and Pakistanis, 98.3% in Indians, 96.8% in Syrians, and 97.8% in Qataris.

Although this study detected a high seroprevalence of EBV among all study participants, the latent infection with EBV (as detected by molecular detection of the EBNA-1 gene) was significantly higher in BC tumor biopsies (45%) as compared to PMNCs of the control group (21.1%). This is in line with two previous Egyptian studies that detected EBV infection in 35.3% and 25% of breast carcinomas (Mohamed et al., 2007; Fawzy et al., 2008). Additionally, using PCR as a method of EBV detection, the prevalence of EBV infection among BC patients in various regions of the world was about 10–50% (Perkins et al., 2006; Lorenzetti et al., 2010). Other studies, on the other hand, failed to detect EBV among BC patients (Lespagnard et al., 1995; Kijima et al., 2001). This discrepancy in the results could be attributed to a variety of factors, such as geographical variation, differences in EBV infection detection methods, specimen type, use of different genes in the EBV genome for PCR, and the inclusion of different histological types of BC. However, studies that used biopsy specimens and used nested PCR as a method for detection showed less heterogeneity (Farahmand et al., 2019).

In this case-control study, EBV-DNA positivity increased the risk of BC development by about threefold (OR = 3.05). This is similar to two recent meta-analysis that reported that the risk of BC association with EBV

| Tumor characteristics | A/B genotypes No. (%) (n=52) | D/C subtypes No. (%) (n=45) | P-value |
|-----------------------|-----------------------------|-----------------------------|---------|
|                       | Genotype- A                  | Genotype- B                  |         |
|                       |                             |                             |         |
|                       | Total No. (%)                | Subtype-D                    | Subtype-C |         |
|                       |                             |                             | P-value  |         |

Table 5. Association between Specific EBV Genotypes and Tumor Features

IDC, Infiltrating ductal carcinoma; ILC, Invasive lobular carcinoma; PR, Progesterone receptor; ER, Estrogen receptor; HER2, Human epidermal growth
infection is 3.84 and 4.74 folds higher as compared to controls (Bae and Kim, 2016; Farahmand et al., 2019).

In the current study, in an attempt to clarify if there is an association between BC and specific types of EBV in Egypt, we detected the distribution frequency of different EBV types obtained from tumor tissues of BC women and compared them with those obtained from PMNCs of healthy non-BC females.

Regarding the distribution of A/B genotypes of EBV among the studied BC patients and healthy controls, we found that most of the EBV detected in both BC tumor biopsies and PMNCs of the controls were of genotype-A (90.9% and 91.7%, respectively, p=0.8). The high prevalence of genotype-A among the participants of this study is consistent with its geographical distribution. Genotype-A is the most prevalent genotype of EBV in the world, predominantly in the United States, France, Germany, North Africa, Turkey, and Asia. Whereas, genotype of EBV predominates more in Central Africa and New Guinea (Ibrahim et al., 2010). In this study, the distribution difference of genotype-A and genotype-B was statistically insignificant between BC patients and the control group. As a result, it appears that neither genotype-A nor genotype-B has a preferential association with BC in Egypt. In the same context, the association of A/B genotypes of EBV with other malignant diseases such as NPC seems to be variable worldwide. Tamura et al. (1993) reported no association of A/B genotypes of EBV with NPC in Japan, as they were predominant in patients as well as in general populations. Klemenc et al., (2006) and Ayadi et al., (2007), reported a high incidence of genotype-A in Slovenian and Tunisian patients with NPC, respectively. Ayee et al., (2020) identified genotype-B as the virulent genotype in Ghana and associated it with the likelihood of NPC development in Ghanaian patients.

Regarding the polymorphism in the BamHI-I W1/I1 region. This study demonstrated that the frequency of subtype-D EBV is significantly higher in BC tumor biopsies (95.6%) as compared to controls (64.7%) (OR =11.7, 95%CI = 2.4-57.1, P =0.002). Similarly, Zhang et al., (2017) found a significant association between subtype-D and the risk of BC among Chinese populations (OR = 2.86), suggesting that subtype-D may play a role in the pathogenesis of BC. It is worth noting that the association of EBV subtype-D with other malignant diseases has previously been demonstrated; Abdel Hamid et al., (1992) discovered that subtype-D is detected more frequently in Egyptian NPC than subtype-C. Similarly, it was reported that EBV of subtype-D was associated with NPC in each of Slovenian patients (62.5%), Tunisian population (98%) and Northern China (32.2%) (Cui et al., 2011). Moreover, subtype-D was found to be associated not only with NPC but also with gastric carcinoma in Iranian population (90%), in Southern Tunisian population (100%), in Latin American population (85%) (Corvalan et al., 2006; Abdirad et al., 2007; BenAyed-Guerfali et al., 2011).

The genetic evidence that C/D subtypes of EBV could be related to the development of tumors is that some EBV genes, such as BamHI-A Rightward Frame-1 (BARF1) and Latent Membrane Protein-2A are present near the C/D locus. These genes are involved in transformation and immortalization during the tumorigenic process (Corvalan et al., 2006). However, the precise mechanisms need more exploration.

Regarding the polymorphism at the BamHI-F region, we detected that EBV from all BC tumor biopsies and control PMNCs was of prototype-F. Therefore, the association of prototype-F with Egyptian BC is not likely. It was proposed that f-variant is only a geographic polymorphism because the EBV that was detected in Egyptian, Algerian, and Tunisian NPCs was of prototype-F, whereas the EBV that was detected in Asian NPCs was of variant-f (Cui et al., 2011). However, Zhou et al., (2001) reported that variant-f is found only in Chinese NPCs and not in Chinese HD patients. Therefore, the hypothesis that the f-variant is a geographically restricted polymorphism needs further investigation.

In the present study, ADF was the most prevalent combination of EBV genotypes among BC patients as well as among control subjects. Likewise, Klemenc et al., (2006) found that ADF is the most predominant combination of EBV genotypes in Slovenian patients with NPC as well as in the healthy population. Additionally, the risk of BC was significantly higher with each of ADF (OR=4.92) and BDF (OR=5.54) combinations and not significantly increased with BCF (P=0.217). Therefore, our findings could point to the implication of specific combinations of EBV genotypes in the development of BC in Egypt. Consequently, our results are not consistent with the hypothesis suggested by Khanim et al (1996) that EBV strains are geographically determined rather than disease-associated.

As regards the association between EBV and the characteristics of BC. EBV was detected more frequently in BC of larger size, higher stage, and higher grade. These findings are similar to that made previously by Murray et al., (2003) and Mohamed et al., (2007) who found that EBV is significantly associated with the more aggressive tumors. Other studies, however, debate our findings as they did not find any statistically significant association between EBV positivity and either grade, stage, or metastasis of the tumor (Richardson et al., 2015; Ahmed and Yussif, 2016; EI-Naby et al., 2017).

In this study, there was no demonstrable significant association between EBV positivity and the steroid-hormone receptor. This agrees with many previous studies (Tiwawech et al., 2008; Ahmed and Yussif, 2016; Bae and Kim, 2016; Preciado et al., 2005). On the contrary, other investigators have shown that EBV is significantly associated with steroid hormone receptor-negative tumors (Bonnet et al., 1999; Murray et al., 2003; EI-Naby et al., 2017).

We found that 75% of EBV-DNA positive BC biopsies are also reactive to HER2 expression. This is close to the result reported by Mohamed et al., (2007) who reported that 55.9% of DNA positive cases were HER2 positive. This finding correlates with the recognition that EBV could transform epithelial cells of the breast into malignancy via stimulating the HER2/HER3 signalling pathways (Szostek et al., 2009). Furthermore, our findings point to some extent that subtype-D of EBV could be the only EBV type...
that might be implicated in the development of BC with poor prognosis because it was the only type that showed more association with tumor of higher grade and with progesterone receptor-free tumors (p = 0.04).

In Conclusion, to the best of our knowledge, this is the first study in Egypt that provides baseline data on the prevalence of different EBV types in BC patients as well as in healthy controls. Overall, this study found that the D-subtype of EBV is BC-associated. Whereas, A/B genotypes and F/f variants of EBV are not associated with BC. Additionally, specific combinations of EBV genotypes (ADF and BDF) were BC-associated. Moreover, our findings indicated that EBV could have an impact on the histological criteria of BC and thus could have a role in prognosis and in optimizing the therapeutic strategies for BC. Nevertheless, our results are considered a platform for any future studies with a larger number of patients with BC in Egypt as well as in other parts of the world to investigate whether variations in the distribution of different types of EBV are geographically related or BC related.

Author Contribution Statement

Mashaly M: Contributes to the concept and design of the study, performed the laboratory work concerned with EBV diagnosis (ELISA, PCR, REFLP), wrote the original draft of the manuscript, revised statistical analysis and interpreted the obtained results. Ghorab D: performed the histopathological part of the study for breast cancer diagnosis. Hegazy M, Abdelkhalek M and Gaballah K: Contributed to the clinical examination of all participants and collection of the study samples and data. Elzehery R: Contributes to the concept and design of the study, performed the laboratory work concerned with EBV diagnosis (ELISA, PCR, REFLP). All authors read and revised the manuscript.

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Ethical approval

The present study received an ethical approval from Ethical Committee of Faculty of Medicine – Mansoura University (MFM-Institutional Research Board).

Availability of data

All data of this study are available on request in SPSS file.

Conflict of interest

The authors declare that they have no known competing financial interests or personal relationships that could have appeared to influence the work reported in this paper.

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