Does mouse mammary tumor-like virus cause human breast cancer? Applying Bradford Hill criteria postulates

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Abstract

Background: The role of mouse mammary tumor-like virus (MMTV-like virus) in human breast cancer (BC) has already been widely investigated worldwide with conflicting results. Although the researchers tried to establish the link between MMTV-like virus and BC through the statistical meta-analysis of the previous studies associating MMTV-like virus with BC, they failed to establish a more reliable link due to the shortcomings of the statistical meta-analysis. In the present study, we identified population-wide studies relating MMTV-like virus with BC through the PubMed search engine. Then, we examined the available data of MMTV-like virus prevalence in BC, normal/benign samples, and applied the postulates of Bradford Hill criteria on the available evidence to investigate the association between MMTV-like virus and BC. In addition, to further enhance the reliability of our outcomes, we have also evaluated the methodologies of the previous studies to address the possibility of false-negative and false-positive results.

Results: After a careful evaluation of the extracted data against the postulates of Bradford Hill criteria, it was observed that none of the studies fulfill all the major postulates of Bradford Hill criteria for causation including temporality, consistency, biological gradient, experiment, coherence, specificity, and analogy. Hence, no causal relationship has been suggested between MMTV-like virus and BC patients of the any included population.

Conclusion: The results failed to prove the causal relationship between MMTV-like virus and BC rather suggested it as a co-participant in the pathogenesis of BC.

Keywords: Mouse mammary tumor-like virus (MMTV-like virus), Breast cancer (BC), Bradford Hill criteria

Background

Breast cancer (BC) is a condition in which the breast cells grow abnormally and out of control. A breast is composed of three main parts: lobules, ducts, and connective tissue. The lobules are the glands that produce milk, while ducts are used to carry the milk to the nipple, and connective tissue surrounds and holds everything together (Zucca-Matthes et al. 2016). BC may occur at multiple sites in the breast but ducts and lobules are the major hotspots of the BC. Based on the type of breast cells that turn into cancer, BC has been divided into several types (Sharma et al. 2010). BC is one of the leading causes of mortality worldwide accounting for 14% of cancer-related deaths (Jemal et al. 2011) with a 5-year survival rate of approximately 40–80% (Coleman et al. 2008).

Being female, getting older, consuming alcohol, no pregnancy, obesity, and viral infection are the very common causes of the BC (Lawson 2009). Moreover, the role of human papilloma (HPV) and Epstein–Barr virus (EBV) in the development of BC has already been well acknowledged worldwide (Glenn et al. 2012). There is, however, no clear etiological role of mouse mammary tumor viruses (MMTV) have been identified in the development of BC since the first study which initially identified the presence of MMTV in human BC in 1972 (Axel et al. 1972; Oskouee et al. 2014).
Another virus named MMTV-like virus is 95% homologous to MMTV in terms of sequence homology (Lawson et al. 2006). Controversial evidence has been reported until now regarding the etiologic role of MMTV-like virus in the development of BC. Although the role of MMTV as an etiologic factor in the development of mice memory tumor is well established, however, it is still unclear whether MMTV-like virus contributes to the development of human BC.

The first-ever study documenting the presence of MMTV-like virus was conducted in 1972 by Axel et al. (1972) in the USA. In that study, they detected the presence of MMTV-like virus in a total of 38 BC samples using the molecular hybridization technique and results revealed the presence of MMTV-like virus in approximately 78.9% of the BC samples.

Since then, numerous studies have been carried out worldwide for detecting MMTV-like virus in BC and their outcomes were contradictory because MMTV-like virus was detected in varying detection positivity ratios population-wide, i.e., from 0% (Oskouee et al. 2014; Bindra et al. 2007; Morales-Sanchez et al. 2013; Mota-medifar et al. 2012; Park et al. 2011; Tabriz et al. 2013; Witt et al. 2003) to 100% 78.9% (Axel et al. 1972).

In general, a statistical meta-analysis is usually preferred when establishing a correlation between the virus and the disease as compared to the single study. This choice is based on the multiple advantages of the meta-analysis such as increased number of objects, greater diversity among the objects, and conclusion with a high level of evidence over the individual single study which has disadvantages like a small cohort of patients and conclusions with a low level of evidence. By keeping in view the inconsistencies in the MMTV-like virus detection ratios in worldwide published studies, recently, researchers have analyzed the previously published studies by the means of statistical meta-analysis to yield more useful pieces of information.

Previously, a statistical meta-analysis was performed to find out the causal relationship between MMTV-like virus and BC by Wang et al. (2014) through the available literature of MMTV-like virus and BC in various authentic research engines including MEDLINE (PubMed), Embase, EBSCO (ASP/BSP), and China National Knowledge Infrastructure (CNKI). They obtained more than 22 studies from different populations such as Western, Asian population and African populations and their revealed no significant association between MMTV-like virus and BC.

Although evaluating the results of previous studies documenting the role of MMTV-like virus in the development of BC through statistical meta-analysis was a better choice than generalizing the results of an individual study, however, we did not consider statistical meta-analysis reliable to establish a causal-relationship between MMTV-like virus and the BC development because of some serious limitations such as its inability to analyze the methodologies of the previous studies, so there is no way to evaluate the possibility of false-negative and false-positive results, nor does statistical meta-analysis provide any information regarding the effect of heterogeneity-specific nature of the understudied populations on MMTV-like virus detection. In addition, statistical meta-analysis results in publication biasness, where meta-analysis does not select studies with no results even though they contain valuable information.

By looking at the discrepancy in the outcomes of the previously published studies and significant shortcomings of the statistical meta-analysis, we performed the population-wise valuation of the results of previous studies detecting MMTV-like virus in BC using the Bradford Hill criteria. These criteria are widely used and accepted worldwide over many years for establishing a causal relationship between a presumed cause and an observed effect of public health research (Fedak et al. 2015).

In the course of evaluation, we analyzed, whether or not these studies fulfill all the postulates of Bradford Hill criteria to declare a causal relationship between MMTV-like virus and BC. In addition, we also evaluated the methodologies used by the previous studies to address the possibility of false-positive and false-negative results for better outcomes. The outcomes of the present study will help to establish a more reliable population-wise causal relationship between MMTV-like virus and BC and determine the more appropriate treatment strategies for BC patients.

Methods
In the present study, we implemented a two-phase methodology (Fig. 1).

Literature search
All the relevant articles associating MMTV-like virus with BC were identified through the PubMed search database using the keywords: “Breast Cancer” AND “MMTV-like virus.” We also defined “Retroviridae” AND “Breast neoplasia” as medical subject headings (MeSH) terms. Mesh terms and keywords were combined during the search process. All the literature was searched available until March 2020, with the “Original Article” filter. In total, 1428 original articles were identified through the PubMed search engine.

Relevant data extraction
From 1429 original articles, the 40 relevant articles were identified having the desired information by initially
reading the title, abstract, and then the complete article. Furthermore, a comprehensive table was constructed having all the required information from the selected relevant studies.

Evaluation of the results using the postulates of Bradford Hill criteria
Based on the extracted data, all the identified studies were carefully evaluated against the following Bradford Hill criteria postulates:

1. Strength: larger the association, more probability of the causal relationship, 2. Temporality: cause must lead to the induction of an effect. If the delay is expected between the cause and effect, then the effect has to occur after the delay, 3. Consistency: different studies conducted by different researchers at different places with different sample sizes and reporting the similar results increase the chances of the causal relation between the cause and effect, 4. Plausibility: there should be plausible mechanism between the cause and effect, 5. Biological gradient: greater response is produced by the causative agent in response to the greater exposure. However, in some cases, effect can be triggered by the mere presence of the factor, while in other cases, greater exposure can lead lower effect as well, 6. Experiment: the relationship between the cause and effect should be explained by the experiments and experiment should results in the reduction of effect when the causative agent is removed, 7. Coherence: causal relationship should not conflict with already known literature about the disease or exposure, 8. Specificity: causality is more likely if the effect has only one cause, 9. Analogy: previous evidence of the association between the cause and effect should support the current statement for the causal relationship.

The assessment of each postulate was qualitative/descriptive, as there was an element of subjectivity in applying quantitative scoring. Evidence collected for each postulate is presented in Table 1 and results section with a final judgment as to whether the viewpoint was fulfilled or not.
| Studied Population | Technique used for the viral genome detection | Target gene | No. of normal samples screened | No. of normal samples positive for MMTV-LIKE VIRUS | Percentage positivity of MMTV-LIKE VIRUS in normal samples | No. of adjacent/benign samples screened | No. of adjacent/benign samples positive for MMTV-LIKE VIRUS | Percentage positivity of MMTV-LIKE VIRUS in adjacent/benign samples | No. of breast cancer samples screened | No. of breast cancer samples positive for MMTV-LIKE VIRUS | Percentage positivity of MMTV-LIKE VIRUS in breast cancer samples | References |
|-------------------|-----------------------------------------------|-------------|---------------------------------|---------------------------------------------------|----------------------------------------------------------|----------------------------------------|------------------------------------------------------------|---------------------------------------------------------------------|-----------------------------------------|-------------------------------------------------------------|---------------------------------------------------------------------|------------------|
| Australia         | PCR                                           | Env         | 0                               | 0                                                 | 25                                                      | 6                                      | 24                                                         | 25                                                                  | 9                                       | 36                                                          | Naru et al. (2017)                                              |                  |
|                   | PCR                                           | Env         | 40                              | 13                                               | 32                                                      | 0                                      | 0                                                         | 50                                                                  | 39                                      | 78                                                          | Glenn et al. (2012)                                                |                  |
|                   | Nested PCR                                    | Env         | 0                               | 0                                                 | 0                                                       | 0                                      | 0                                                         | 42                                                                  | 0                                       | 0                                                           | Park et al. (2011)                                                 |                  |
|                   | PCR, Immunohistochemistry                      | Env, gp52   | 29                              | 0                                                 | 0                                                       | 0                                      | 0                                                         | 74                                                                  | 33                                      | 45                                                          | Lawson et al. (2010)                                               |                  |
|                   | PCR                                           | Env         | 0                               | 0                                                 | 0                                                       | 0                                      | 0                                                         | 51                                                                  | 28                                      | 56                                                          | Mok et al. (2008)                                                  |                  |
|                   | Semi-nested PCR                               | Env         | 111                             | 2                                                 | 1.8                                                     | 25                                     | 5                                                         | 20                                                                  | 288                                             | 90                                                          | Ford et al. (2004)                                                 |                  |
|                   | Semi-nested PCR                               | Env         | 0                               | 0                                                 | 0                                                       | 0                                      | 0                                                         | 128                                                                 | 50                                      | 3906                                                     | Faedo et al. (2004)                                                |                  |
|                   | PCR                                           | Env         | 0                               | 0                                                 | 0                                                       | 0                                      | 0                                                         | 44                                                                  | 20                                      | 4545                                                     | Lawson et al. (2004)                                               |                  |
|                   | PCR                                           | Env         | 0                               | 0                                                 | 0                                                       | 0                                      | 0                                                         | 59                                                                  | 22                                      | 373                                                       | Lawson et al. (2006)                                               |                  |
| Pakistan          | PCR                                           | LTR         | 0                               | 0                                                 | 0                                                       | 0                                      | 0                                                         | 55                                                                  | 19                                      | 345                                                       | Naushad et al. (2017a)                                              |                  |
|                   | PCR                                           | LTR, Env    | 0                               | 0                                                 | 0                                                       | 0                                      | 0                                                         | 250                                                                 | 29                                      | 116                                                      | Naushad et al. (2017b)                                             |                  |
|                   | PCR                                           | LTR, Env    | 6                               | 0                                                 | 0                                                       | 0                                      | 0                                                         | 80                                                                  | 37                                      | 4625                                                     | Naushad et al. (2014)                                               |                  |
| Studied Population | Technique used for the viral genome detection | Target gene | No. of normal samples screened | No. of normal samples positive for MMTV-LIKE VIRUS | No. of adjacent/benign samples screened | No. of adjacent/benign samples positive for MMTV-LIKE VIRUS | No. of normal samples screened | No. of normal samples positive for MMTV-LIKE VIRUS | No. of adjacent/benign samples screened | No. of adjacent/benign samples positive for MMTV-LIKE VIRUS | No. of breast cancer samples screened | Percentage positivity of MMTV-LIKE VIRUS in breast cancer samples | References |
|-------------------|-----------------------------------------------|-------------|-------------------------------|-----------------------------------------------|----------------------------------------|-------------------------------------------------|-------------------------------|-----------------------------------------------|----------------------------------------|-------------------------------------------------|-------------------------------|-------------------------------------------------|-----------|
| Iran              | Nested PCR                                    | Env         | 65                            | 0                                             | 0                                      | 0                                              | 0                             | 0                                             | 0                                      | 65                                              | 0                             | 0                                              | Oskouee et al. (2014) |
|                   | Nested PCR                                    | Env         | 0                             | 0                                             | 0                                      | 59                                             | 3                             | 5                                             | 59                                     | 19                                              | 32.2                          | Shariatpanahi et al. (2017) |
|                   | Real-time PCR                                 | Env         | 0                             | 0                                             | 0                                      | 0                                              | 0                             | 0                                             | 40                                     | 0                                               | 0                             | Tabriz et al. (2013) |
|                   | Nested PCR                                    | Env         | 300                           | 0                                             | 0                                      | 0                                              | 0                             | 0                                             | 50                                     | 0                                               | 0                             | Motamedifar et al. (2012) |
| Morocco            | PCR                                           | Env         | 100                           | 0                                             | 0                                      | 0                                              | 0                             | 0                                             | 100                                    | 12                                              | 12                            | Reza et al. (2015) |
| Austria            | PCR, Nested PCR                               | Env         | 0                             | 0                                             | 0                                      | 0                                              | 0                             | 0                                             | 50                                     | 0                                               | 0                             | Witt et al. (2003) |
| Egypt              | Nested PCR                                    | Env         | 0                             | 0                                             | 0                                      | 50                                             | 2                             | 4                                             | 100                                    | 36                                              | 36                            | Hafez et al. (2013) |
| Vietnam            | PCR                                           | Env         | 60                            | 0                                             | 0                                      | 0                                              | 0                             | 0                                             | 120                                    | 1                                               | 0.83                          | Ford et al. (2003) |
| Argentina          | PCR                                           | Env         | 74                            | 1                                             | 1.35                                   | 74                                             | 0                             | 0                                             | 74                                     | 23                                              | 31                            | Melana et al. (2002) |
| Sweden             | RT-PCR                                        | Env         | 0                             | 0                                             | 0                                      | 11                                             | 0                             | 0                                             | 35                                     | 0                                               | 0                             | Bindra et al. (2007) |
| Saudi Arabia       | PCR                                           | Env         | 51                            | 0                                             | 0                                      | 93                                             | 9                             | 9.7                                           | 101                                    | 6                                               | 5.9                           | Al Dossary et al. (2018) |
| Tunisia            | Semi-nested PCR                               | Env         | 0                             | 0                                             | 0                                      | 0                                              | 0                             | 0                                             | 122                                    | 17                                              | 3.9                           | Hachana et al. (2008) |
|                   | Antigen detection                             | gp52        | 0                             | 0                                             | 0                                      | 17                                             | 0                             | 0                                             | 33                                     | 23                                              | 70                            | Levine et al. (1984) |
|                   | PCR                                           | Env         | 140                           | 2                                             | 1.4                                    | 0                                              | 0                             | 0                                             | 38                                     | 28                                              | 7.37                          | Pogo et al. (2010) |
| Studied Population | Technique used for the viral genome detection | Target gene | No. of normal samples screened | No. of normal samples positive for MMTV-LIKE VIRUS | Percentage positivity of MMTV-LIKE VIRUS in normal samples | No. of adjacent/benign samples screened | No. of adjacent/benign samples positive for MMTV-LIKE VIRUS | Percentage positivity of MMTV-LIKE VIRUS in adjacent/benign samples | No. of breast cancer samples screened | No. of breast cancer samples positive for MMTV-LIKE VIRUS | Percentage positivity of MMTV-LIKE VIRUS in breast cancer samples | References |
|-------------------|---------------------------------------------|-------------|-------------------------------|-----------------------------------------------|-----------------------------------------------------|--------------------------------------|---------------------------------------------------------------|---------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|---------------------------------------------|---------------------------------------------------------------|-----------------------------------------------------------------------------------------------------------------------------|--------------------------|
| USA               | Nested PCR                                  | Env         | 35                            | 0                                             | 0                                                  | 0                                    | 0                                           | 0                                               | 73                                         | 27                                           | 369                                                                 | Etkind et al. (2000) |
| USA               | PCR                                         | Env         | 140                           | 2                                             | 1.4                                                | 0                                    | 0                                           | 0                                               | 709                                        | 323                                          | 455                                                                 | Pogo et al. (2010)    |
| USA               | PCR                                         | Env         | 107                           | 2                                             | 1.8                                                | 29                                   | 2                                           | 6.9                                             | 314                                        | 121                                          | 38.5                                                                 | Wang et al. (1995)    |
| USA               | Molecular hybridization                      | –           | 0                             | 0                                             | 0                                                  | 0                                    | 0                                           | 0                                               | 38                                         | 30                                           | 789                                                                 | Axel et al. (1972)    |
| Mexico            | PCR                                         | Env         | 0                             | 0                                             | 0                                                  | 65                                   | 0                                           | 0                                               | 86                                         | 0                                            | 0                                                                 | Morales-Sanchez et al. (2013) |
| Mexico            | PCR                                         | Env         | 0                             | 0                                             | 0                                                  | 0                                    | 0                                           | 0                                               | 119                                        | 5                                            | 4.2                                                                 | Zapata-Benavides et al. (2007) |
| Myanmar           | Nested PCR                                  | Env         | 0                             | 0                                             | 0                                                  | 458                                  | 72                                          | 16                                              | 458                                        | 57                                           | 12.4                                                                | Cedro-Tanda et al. (2014) |
| Myanmar           | Nested PCR                                  | Env         | 0                             | 0                                             | 0                                                  | 0                                    | 0                                           | 0                                               | 58                                         | 1                                            | 1.7                                                                 | San et al. (2017)     |
### Table 1 (continued)

| Studied Population | Technique used for the viral genome detection | Target gene | No. of normal samples screened | No. of normal samples positive for MMTV-LIKE VIRUS | Percentage positivity of MMTV-LIKE VIRUS in normal samples | No. of adjacent/benign samples screened | No. of adjacent/benign samples positive for MMTV-LIKE VIRUS | Percentage positivity of MMTV-LIKE VIRUS in adjacent/benign samples | No. of breast cancer samples screened | No. of breast cancer samples positive for MMTV-LIKE VIRUS | Percentage positivity of MMTV-LIKE VIRUS in breast cancer samples | References |
|--------------------|-----------------------------------------------|-------------|-------------------------------|-----------------------------------------------|----------------------------------------------------------|----------------------------------------|-------------------------------------------------|----------------------------------------------------------|----------------------------------------|-------------------------------------------------|----------------------------------------------------------|------------|
| Italy              | Fluorescence-nested PCR, Immunohistochemistry  | Env         | 0                             | 0                                             | 0                                                        | 0                                      | 0                                               | 0                                                        | 105                                    | 34.5                                            | 3.45                                                   | Naccarato et al. (2019) |
|                    | Fluorescence-nested PCR                        | Env         | 0                             | 0                                             | 0                                                        | 0                                      | 0                                               | 0                                                        | 45                                     | 15                                              | 33                                                      | Zammarchi et al. (2006) |
|                    | Fluorescence-nested PCR, Real-time PCR, In situ hybridization | Env         | 20                            | 0                                             | 0                                                        | 26                                     | 5                                               | 19.2                                                     | 138                                    | 47                                              | 68.1                                                    | Mazzanti et al. (2011) |
|                    | PCR                                            | Env         | 0                             | 0                                             | 0                                                        | 0                                      | 0                                               | 0                                                        | 122                                    | 45                                              | 36.8                                                    | Pogo et al. (1999) |
| China              | Nested PCR                                     | Env         | 0                             | 0                                             | 0                                                        | 0                                      | 0                                               | 0                                                        | 241                                    | 44                                              | 18.25                                                   | Luo et al. (2006) |

PCR polymerase chain reaction, RT-PCR real-time PCR, Env envelope protein, LTR long terminal repeats
Results

In total, 40 relevant original articles (Glenn et al. 2012; Axel et al. 1972; Oskouee et al. 2014; Lawson et al. 2006, 2010, 2004; Bindra et al. 2007; Morales-Sanchez et al. 2013; Motamedifar et al. 2012; Park et al. 2011; Tabriz et al. 2013; Witt et al. 2003; Al Dossary et al. 2018; Cedro-Tanda et al. 2014; Etkind et al. 2000; Faedo et al. 2004; Ford et al. 2004, 2003; Hachana et al. 2008; Hafez et al. 2013; Levine et al. 1984; Luo et al. 2006; Mazzanti et al. 2011; Melana et al. 2002, 2001; Mok et al. 2008; Naccarato et al. 2019; Nartey et al. 2017; Naushad et al. 2014, 2017a, b; Pogo et al. 1999, 2010; Reza et al. 2015; San et al. 2017; Shariatpanahi et al. 2017; Slaoui et al. 2014; Wang et al. 1995; Zammarchi et al. 2006; Zapata-Benavides et al. 2007) (Table 1) were found on PubMed which investigated the association of MMTV-like virus with BC in 24 different populations. Table 1 enlists all these articles with summarized information extracted from these articles including information about the studied population, technique used for the detection of MMTV-like virus, target gene, number (No) of screened samples (normal, adjacent/benign, and BC) with their respective identified population-wise positivity ratios. Out of all the 40 studies, in total, 23 studies (Glenn et al. 2012; Oskouee et al. 2014; Lawson et al. 2006, 2010, 2004; Bindra et al. 2007; Morales-Sanchez et al. 2013; Motamedifar et al. 2012; Park et al. 2011; Tabriz et al. 2013; Witt et al. 2003; Al Dossary et al. 2018; Cedro-Tanda et al. 2014; Etkind et al. 2000; Faedo et al. 2004; Ford et al. 2004, 2003; Hachana et al. 2008; Hafez et al. 2013; Luo et al. 2006; Mazzanti et al. 2011; Melana et al. 2002, 2001; Mok et al. 2008; Naccarato et al. 2019; Nartey et al. 2017; Naushad et al. 2014, 2017a, b; Pogo et al. 1999, 2010; Reza et al. 2015; San et al. 2017; Shariatpanahi et al. 2017; Slaoui et al. 2014; Wang et al. 1995; Zammarchi et al. 2006; Zapata-Benavides et al. 2007) to detect the presence of MMTV-like virus in the normal, adjacent/benign and BC samples using Env (Glenn et al. 2012; Oskouee et al. 2014; Lawson et al. 2006, 2010, 2004; Bindra et al. 2007; Morales-Sanchez et al. 2013; Motamedifar et al. 2012; Park et al. 2011; Tabriz et al. 2013; Witt et al. 2003; Al Dossary et al. 2018; Cedro-Tanda et al. 2014; Etkind et al. 2000; Faedo et al. 2004; Ford et al. 2004, 2003; Hachana et al. 2008; Hafez et al. 2013; Luo et al. 2006; Mazzanti et al. 2011; Melana et al. 2002, 2001; Mok et al. 2008; Naccarato et al. 2019; Nartey et al. 2017; Naushad et al. 2014, 2017a, b; Pogo et al. 1999, 2010; Reza et al. 2015; San et al. 2017; Shariatpanahi et al. 2017; Slaoui et al. 2014; Wang et al. 1995; Zammarchi et al. 2006; Zapata-Benavides et al. 2007) Ltr (Naushad et al. 2014, 2017a, b), and gp52 (Lawson et al. 2010; Levine et al. 1984) gene-specific primers. Additionally, from them, few studies (Glenn et al. 2012; Bindra et al. 2007; Morales-Sanchez et al. 2013; Al Dossary et al. 2018; Cedro-Tanda et al. 2014; Etkind et al. 2000; Ford et al. 2004, 2003; Lawson et al. 2010; Mazzanti et al. 2011; Melana et al. 2001; Mok et al. 2008; Naccarato et al. 2019; Nartey et al. 2017; Naushad et al. 2014, 2017a, b; Pogo et al. 1999, 2010; Reza et al. 2015; San et al. 2017; Shariatpanahi et al. 2017; Slaoui et al. 2014; Wang et al. 1995; Zammarchi et al. 2006; Zapata-Benavides et al. 2007) also employed the second techniques for validating their PCR positive results using second techniques such as immunohistochemistry (Lawson et al. 2010; Naccarato et al. 2019), in situ hybridization (Mazzanti et al. 2011), real-time PCR (Mazzanti et al. 2011), and DNA sequencing analysis (Glenn et al. 2012; Bindra et al. 2007; Morales-Sanchez et al. 2013; Al Dossary et al. 2018; Cedro-Tanda et al. 2014; Etkind et al. 2000; Ford et al. 2004, 2003; Melana et al. 2001; Mok et al. 2008; Nartey et al. 2017; Naushad et al. 2017; San et al. 2017; Slaoui et al. 2014; Zammarchi et al. 2006; Zapata-Benavides et al. 2007). On the other hand, few studies also utilized antigen detection method (Levine et al. 2017).
and adjacent/benign controls, respectively. The detection positivity ratios ranging from 0% (Oskouee et al. 2014) to 32% (Shariatpanahi et al. 2017) in BC samples, while 0% (Ford et al. 2003) to 24% (Nartey et al. 2017) in adjacent/benign samples and 0% (Lawson et al. 2010) to 32% (Glenn et al. 2012) in normal controls.

In Pakistan, the association between MMTV-like virus and BC has been reported in only 03 studies (Naushad et al. 2014, 2017a, b) so far. They utilized PCR technique with primers specifically targeting the Env and LTR regions of the viral genome and documented varying MMTV-like virus detection positivity ratio ranging from 11.6% (Naushad et al. 2017) to 46.25% (Naushad et al. 2014) in BC samples.

A total of 05 studies (Oskouee et al. 2014; Motamedifar et al. 2012; Tabriz et al. 2013; Reza et al. 2015; Shariatpanahi et al. 2017) including 04 case–control studies (Oskouee et al. 2014; Motamedifar et al. 2012; Reza et al. 2015; Shariatpanahi et al. 2017) have been carried out in Iran so far, analyzing the association between MMTV-like virus and BC. These studies utilized PCR technique with primers specific for the Env region of the viral genome and documented varying MMTV-like virus detection positivity ratios ranging from 0% (Oskouee et al. 2014; Motamedifar et al. 2012; Tabriz et al. 2013) to 32.2% (Shariatpanahi et al. 2017) in BC samples, while 0% (Oskouee et al. 2014; Motamedifar et al. 2012; Reza et al. 2015) and 5% (Shariatpanahi et al. 2017) in the normal and adjacent/benign controls, respectively.

A single case–control study (Slaoui et al. 2014) has been conducted in Morocco so far to determine the association between MMTV-like virus and BC. They analyzed 42 BC samples and 18 adjacent/benign controls for the identification of MMTV-like virus in BC using PCR technique with primers specifically hybridizing in the Env sequence of the viral genome. They documented 57.1% and 33.3% MMTV-like virus detection positivity ratios in BC and adjacent/benign controls, respectively.

In Austria, only a single study (Witt et al. 2003) has been reported to date to relating MMTV-like virus presence with BC. They screened 50 BC samples using PCR technique with primers specifically targeting the Env sequence of the viral genome and they documented 0% MMTV-like virus detection positivity ratio in BC samples.

Until now, only a single case–control study (Hafez et al. 2013) has been carried out in Egypt to find out the etiological association between MMTV-like virus and BC. They analyzed 100 BC samples and 50 adjacent/benign controls using PCR technique with Env gene-specific primers for MMTV-like virus detection. They reported the higher MMTV-like virus detection positivity ratio in BC samples (36%) as compared to the adjacent/benign controls (4%).

In Vietnam, there has been a single case–control study (Ford et al. 2003) reported so far to find out if MMTV-like virus has any association with BC. They utilized PCR technique with primers specifically targeting the Env sequence of the viral genome and documented a 0.83% MMTV-like virus detection positivity ratio in BC samples and normal controls, respectively.

Until now, only a single case–control study (Melana et al. 2002) has been carried out in Argentina to elaborate the role of MMTV-like virus in the development of BC. They screened 74 BC samples paired with adjacent/benign controls using PCR technique with primers targeting the envelope sequence of the viral genome. They documented the higher MMTV-like virus detection positivity ratios in BC samples (31%) as compared to the adjacent/benign samples (0%) and normal controls (1.35%).

In Sweden, a single case–control (Bindra et al. 2007) has been carried out so far to identify the casual relationship between MMTV-like virus and BC using RT-PCR technique and they did not detect MMTV-like virus in any of the BC or control sample.

So far, a single case–control study (Al Dossary et al. 2018) has been carried out in Saudi Arabia to find out the association between MMTV-like virus and BC. They screened a total of 101 BC, 51 normal, and 93 adjacent/benign controls using PCR technique with primers specific for the Env region of the viral genome and documented MMTV-like virus positivity ratios was higher in the adjacent/benign controls (9.7%) as compared to the normal (0%) and BC samples (5.9%).

In Tunisia, a total of 03 studies (Hachana et al. 2008; Levine et al. 1984; Pogo et al. 2010) including 02 case–control studies (Levine et al. 1984; Pogo et al. 2010) have been reported so far to find out whether MMTV-like virus have any association with BC. They utilized antigen detection method and PCR technique with Env gene-specific primers for MMTV-like virus detection and documented MMTV-like virus detection positivity ratios with varying frequencies ranging from 3.9%
et al. (2007) including two case–control study (Morales-Sanchez et al. 2013; Cedro-Tanda et al. 2014; Zapata-Benavides et al. 2014) in adjacent/benign and normal samples, respectively.

To date, a total of 04 studies (Axel et al. 1972; Etkind et al. 2000; Pogo et al. 2010; Wang et al. 1995) including 3 case–control studies (Etkind et al. 2000; Pogo et al. 2010; Wang et al. 1995) have been carried out in the USA to elaborate the causal association between in adjacent/benign samples and BC. All these studies employed PCR technique with Env gene-specific primers and documented MMTV-like virus detection positivity ratios ranging from 36.9% (Etkind et al. 2000) to 45.5% (Pogo et al. 2010) in BC samples, while 6.9% (Wang et al. 1995) in the adjacent/benign samples and 0% (Etkind et al. 2000) to 1.8% (Wang et al. 1995) in the normal controls.

Until now, a total of 03 studies (Morales-Sanchez et al. 2013; Cedro-Tanda et al. 2014; Zapata-Benavides et al. 2007) including two case–control study (Morales-Sanchez et al. 2013; Cedro-Tanda et al. 2014) have been reported in Mexico to determine the causal relationship between MMTV-like virus and BC. All these studies utilized PCR with primers specific for the Env gene-specific primers and the documented MMTV-like virus detection positivity ratios varied from 0% (Morales-Sanchez et al. 2013) to 12.6% (Cedro-Tanda et al. 2014) in BC samples and 0% (Morales-Sanchez et al. 2013) to 16% (Cedro-Tanda et al. 2014) in adjacent/benign controls.

A single study (San et al. 2017) has been conducted in Myanmar to date to relate the MMTV-like virus presence with BC. In total, they analyzed 58 BC samples were using PCR technique with primers specific for the Env region and documented a 1.7% 1.7% detection positivity ratio.

A total of 04 studies (Mazzanti et al. 2011; Naccarato et al. 2019; Pogo et al. 1999; Zammarchi et al. 2006) including a case–control study (Mazzanti et al. 2011) have been carried out in Italy so far, analyzing the etiological association between 1.7% and BC. These studies utilized the PCR technique with primers specific for the Env region of the viral genome and documented MMTV-like virus detection positivity ratios with varying frequencies ranging from 33% (Shariatpanahi et al. 2017) to 68.1% (Mazzanti et al. 2011) in BC samples, while 0% (Mazzanti et al. 2011) in normal and 19.2% (Mazzanti et al. 2011) in adjacent/benign controls.

In China, there has been a single study (Luo et al. 2006) reported so far to find out if MMTV-like virus is associated with BC. They utilized PCR technique with primers specifically targeting the Env sequence of the viral genome and documented 18.25% MMTV-like virus detection positivity ratio.

**Discussion**

BC is one of the most common types of cancer that infect millions of people worldwide each year. Although recent advancements in the diagnosis and treatment of the BC have helped to manage the disease but still the prevalence of BC is on a rise due to unknown underlying mechanisms (Akram et al. 2017).

To date, various individual studies have been carried out worldwide to find the state of association between MMTV-like virus and BC to further uncover the molecular pathways underlying BC but their results are conflicting. In addition, the statistical meta-analysis was also used by the researchers to analyze the previous individual studies for generating a more meaningful association between MMTV-like virus and BC but due to the shortcomings of statistical meta-analysis, researchers once again failed to establish a more reliable causal relationship between MMTV-like virus and BC.

In the present study, we evaluated the previous studies using a reliable, Bradford Hill criteria to find a causal relationship between MMTV-like virus and BC. In addition, we also evaluated the methodologies used by the previous studies to address the possibility of false-positive and false-negative results for better outcomes.

In total, 40 original articles (Glenn et al. 2012; Axel et al. 1972; Oskouee et al. 2014; Lawson et al. 2006, 2010, 2004; Bindra et al. 2007; Morales-Sanchez et al. 2013; Motamedifar et al. 2012; Park et al. 2011; Tabrizi et al. 2013; Witt et al. 2003; Al Dossary et al. 2018; Cedro-Tanda et al. 2014; Etkind et al. 2000; Faedo et al. 2004; Ford et al. 2004, 2003; Hachana et al. 2008; Hafez et al. 2013; Levine et al. 1984; Luo et al. 2006; Mazzanti et al. 2011; Melana et al. 2002, 2001; Mok et al. 2008; Naccarato et al. 2019; Narrey et al. 2017; Naushad et al. 2014, 2017a, b; Pogo et al. 1999, 2010; Reza et al. 2015; San et al. 2017; Shariatpanahi et al. 2017; Slouai et al. 2014; Wang et al. 1995; Zammarchi et al. 2006; Zapata-Benavides et al. 2007) were included in the present study. The MMTV-like virus positivity ratio reported in these studies was varied between 0% (Oskouee et al. 2014; Bindra et al. 2007; Morales-Sanchez et al. 2013; Motamedifar et al. 2012; Park et al. 2011; Tabrizi et al. 2013; Witt et al. 2003) and 78.9% (Axel et al. 1972) in BC samples. In most of the case–control studies (Glenn et al. 2012; Oskouee et al. 2014; Bindra et al. 2007; Morales-Sanchez et al. 2013; Motamedifar et al. 2012; Etkind et al. 2000; Ford et al. 2004, 2003; Hafez et al. 2013; Lawson et al. 2010; Levine et al. 1984; Mazzanti et al. 2011; Melana et al. 2002, 2001; Narrey et al. 2017; Naushad et al. 2014, 2017a, b; Reza et al. 2015; Shariatpanahi et al. 2017; Slouai et al. 2014; Wang et al. 1995), the positivity ratio for MMTV-like virus detection was higher in the BC samples as compared to the controls, while in some studies (Al Dossary...
et al. 2018; Cedro-Tanda et al. 2014), MMTV-like virus positivity ratio was higher in the controls as compared to the BC samples. Possible reasons for such population-specific inequalities in MMTV-like virus detection could be non-modifiable factors such as genetic makeup and socially controllable factors like health-seeking behavior and differential access to the health facilities.

After careful evaluation of the results of identified studies through Bradford hill criteria showed that all the studies failed to fulfill the major postulates including strength, temporality, consistency, biological gradient, experiment, coherence, specificity, and analogy. Hence, we suggested that MMTV-like virus acts as a co-participant in the development of BC rather than having a causal relationship that might combine with the other viruses and such as human immunodeficiency virus (HIV) and hepatitis C virus (HCV), other factors including genetic abnormalities, smoking, alcohol consumption, and obesity may increase a person’s risk of developing BC by affecting the body’s immune system.

However, limitations and some of the major issues related with methodologies used in the included studies have been discussed below.

Possible causes of false-negative results

In few studies, the presence of MMTV-like virus was not detected in any of the analyzed samples. How we can be sure that their negative results of MMTV-like virus detection were not due to the poor quality of the extracted DNA? To overcome this issue, most of the studies (Glenn et al. 2012; Oskouee et al. 2014; Lawson et al. 2006, 2010, 2004; Bindra et al. 2007; Morales-Sanchez et al. 2013; Motamedifar et al. 2012; Park et al. 2011; Tabriz et al. 2013; Al Dossary et al. 2018; Cedro-Tanda et al. 2014; Etkind et al. 2004; Ford et al. 2004, 2003; Hachana et al. 2008; Hafez et al. 2013; Luo et al. 2006; Mazzanti et al. 2011; Melana et al. 2002, 2001; Mok et al. 2008; Naccarato et al. 2019; Narrey et al. 2017; Naushad et al. 2014, 2017a, b; Pogo et al. 1999, 2010; Reza et al. 2015; San et al. 2017; Shariatpanahi et al. 2017; Slouei et al. 2014; Wang et al. 1995; Zammarchi et al. 2006; Zapata-Benavides et al. 2007) used PCR technique for the initial detection of MMTV-like virus but only few studies (Glenn et al. 2012; Bindra et al. 2007; Morales-Sanchez et al. 2013; Al Dossary et al. 2018; Cedro-Tanda et al. 2014; Etkind et al. 2000; Ford et al. 2004, 2003; Lawson et al. 2010; Mazzanti et al. 2011; Melana et al. 2001; Mok et al. 2008; Naccarato et al. 2019; Narrey et al. 2017; Naushad et al. 2014, 2017a, b; Pogo et al. 1999, 2010; Reza et al. 2015; San et al. 2017; Shariatpanahi et al. 2017; Slouei et al. 2014; Wang et al. 1995; Zammarchi et al. 2006; Zapata-Benavides et al. 2007) validated their PCR positive results using second techniques such as immunohistochemistry (Lawson et al. 2010; Naccarato et al. 2019), In situ hybridization (Zammarchi et al. 2011), real-time PCR (Mazzanti et al. 2011), and DNA sequencing analysis (Glenn et al. 2012; Bindra et al. 2007; Morales-Sanchez et al. 2013; Al Dossary et al. 2018; Cedro-Tanda et al. 2014; Etkind et al. 2000; Ford et al. 2004, 2003; Lawson et al. 2010; Zammarchi et al. 2006; Zapata-Benavides et al. 2007) validated their PCR positive results using second techniques such as immunohistochemistry (Lawson et al. 2010; Naccarato et al. 2019), In situ hybridization (Zammarchi et al. 2011), real-time PCR (Mazzanti et al. 2011), and DNA sequencing analysis (Glenn et al. 2012; Bindra et al. 2007; Morales-Sanchez et al. 2013; Al Dossary et al. 2018; Cedro-Tanda et al. 2014; Etkind et al. 2000; Ford et al. 2004, 2003; Melana et al. 2001; Mok et al. 2008; Narrey et al. 2017a, b; San et al. 2017; Slouei et al. 2014; Zammarchi et al. 2006; Zapata-Benavides et al. 2007). Expression of PgR, HER2, Wnt-1, laminin receptor could be used as surrogate biomarker in MMTV-like virus-infected BC patients. Along with MMTV-like virus detection, these surrogate biomarkers were also analyzed by some studies (Faedo et al. 2004; Hachana et al. 2008; Lawson et al. 2010, 2004; Pogo
et al. 1999; Reza et al. 2015) to further validate their findings, out of which (Faedo et al. 2004; Lawson et al. 2010, 2004; Pogo et al. 1999; Reza et al. 2015) have validated their findings by analyzing these surrogate biomarkers, while the other studies (Hachana et al. 2008; Pogo et al. 1999) failed to validate their findings through surrogate biomarkers. Such deviations in the results of previous studies raise a big question mark on the selection of appropriate technique and their sensitivities.

Comparison of normal, benign, and malignant samples
Case–control studies are necessary to establish a causal relationship between the causative agent and the disease. Some of the studies we summarized analyzed only the BC samples (Axel et al. 1972; Lawson et al. 2006, 2004; Park et al. 2011; Tabriz et al. 2013; Witt et al. 2003; Faedo et al. 2004; Hachana et al. 2008; Luo et al. 2006; Mok et al. 2008; Naccarato et al. 2019; Naushad et al. 2014, 2017; Pogo et al. 1999; San et al. 2017; Zammarchi et al. 2006; Zapata-Benavides et al. 2007) and did not allow us to compare their results with normal or adjacent/benign controls. However, most of the studies (Glenn et al. 2012; Oskouee et al. 2014; Bindra et al. 2007; Morales-Sanchez et al. 2013; Motamedifar et al. 2012; Al Dossary et al. 2018; Cedro-Tanda et al. 2014; Etkind et al. 2000; Ford et al. 2004, 2003; Hafez et al. 2013; Lawson et al. 2010; Levine et al. 1984; Mazzanti et al. 2011; Melana et al. 2002, 2001; Naraty et al. 2017; Naushad et al. 2014; Pogo et al. 2010; Reza et al. 2015; Shariatpanahi et al. 2017; Slaoui et al. 2016; Wang et al. 1995) also analyzed the normal and adjacent/benign tissues along with BC samples and comparison of their results demonstrated that MMTV-like virus detection positivity ratios in BC samples were higher in Glenn et al. (2012), Oskouee et al. (2014), Bindra et al. (2007), Morales-Sanchez et al. (2013), Motamedifar et al. (2012), Etkind et al. (2000), Ford et al. (2004, 2003), Hafez et al. (2013), Lawson et al. (2010), Levine et al. (1984), Mazzanti et al. (2011), Melana et al. (2002, 2001), Naraty et al. (2017), Naushad et al. (2014), Pogo et al. (2010), Reza et al. (2015), Shariatpanahi et al. (2017), Slaoui et al. (2014), and Wang et al. (1995) studies, while lower in Al Dossary et al. (2018) and Cedro-Tanda et al. (2014) studies as compared to the normal and adjacent/benign controls. However, none of the studies has reported the association of MMTV-like virus with specific BC subtype and histologic grade.

Conclusion
The results of this comprehensive review are controversial. They failed to prove the causal relationship between MMTV-like virus and BC rather suggested it as co-participant in the pathogenesis of BC. However, due to limitations of the methodologies used by the previous studies to detect the presence of MMTV-like virus in BC, additional experiments are required to prove the MMTV-like virus etiology in BC.

Abbreviations
DNA: Deoxynucleobnucleic acid; Env: Envelope gene; EBV: Epstein–Barr virus; HPV: Human papilloma virus; LTR: Long terminal repeats; MeSH: Medical subject headings; MMTV: Mouse mammary tumor virus; PCR: Polymerase chain reaction; RT-PCR: Real-time polymerase chain reaction; RNA: Ribonucleic acid.

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YH formulated the idea; MU collected the data and performed the statistical analysis with the help of MA. MU wrote the first draft of the manuscript which is finalized by YH. All authors read and approved the final manuscript.

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