Overview of virus and cancer relationships. Position paper

ABSTRACT

The role of certain viruses in the etiology of some tumors is today indisputable, but there is a lack, however, of an overview of the relationship between viruses and cancer with a multidisciplinary approach. For this reason, the Health Sciences Foundation has convened a group of professionals from different areas of knowledge to discuss the relationship between viruses and cancer, and the present document is the result of these deliberations. Although viruses cause only 10-15% of cancers, advances in oncology research are largely due to the work done during the last century on tumor viruses. The clearest cancer-inducing viruses are: HPV, HBV, HCV, EBV, and, depending on the geographical area, HHV-8, HTLV-1 and HIV. HPV’s, for example, are considered to be the causative agents of cervical carcinomas and, more recently, of a proportion of other cancers. Among the Herpes viruses, the association with the development of neoplasms is well established for EBV and HHV-8. Viruses can also be therapeutic agents in certain neoplasms and, thus, some oncolytic viruses with selective tropism for tumor cells have been approved for clinical use in humans. It is estimated that the prophylaxis or treatment of viral infections could prevent at least 1.5 million cancer deaths per year.

Keywords: Virus, Cancer, Tumours, Human Papillomavirus, HPV, Hepatitis B, Hepatitis C, HBV, HCV, Epstein–Barr Virus, EBV, Human Herpes Virus 8, HHV-8, Human T Lymphotropic Virus I, HTLV-1, Human Immunodeficiency Virus, HIV, Vaccines, Oncolytic virus, Oncogenes, Cancer mortality, Cancer prevention, Cancer–economy

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INTRODUCTION

The success in human longevity achieved in recent decades is impressive and could be even greater if the major challenges of neurodegenerative diseases, cardiovascular disease and cancer could be controlled.

Of the three challenges above, cancer treatment is perhaps achieving more success than the other two, not only because of better prevention and earlier diagnosis, but also because of increasingly effective and better-tolerated treatment.

Viruses offer a first approximation to the aetiological diagnosis in the case of cancer and the role some of them play in the pathogenesis of certain tumours is today indiscutiable. Their prevention, such as in the case of HPV and cervical cancer, has radically changed our expectations for the future and allows us to envision the eradication of this tumour. Unfortunately, the virus-cancer relationship is not so clear-cut in many other tumours and when we look for global data on that relationship, we find scattered and inaccurate, irregular literature of a very uneven nature and quality.

For this reason, the Health Sciences Foundation has proposed examining the virus-cancer relationship more generally, in a multidisciplinary forum, aiming to obtain answers to a series of simple questions from experts in different aspects of the problem. This meeting has included oncologists, gastroenterologists, internists, gynaecologists, specialists in infectious diseases and HIV, microbiologists, basic biologists, representatives of patient associations and the media, and specialists in preventive medicine and ethics.

A first block of questions was aimed at putting the dimension of the Virus-Cancer relationship, the cancers most clearly caused by viruses, the great historical milestones of this relationship and some aspects of oncogenesis into perspective.

In a second block, some potential solutions ranging from vaccines to the treatment of hepatitis C were addressed and the possible role of bacteria and parasites as alternative oncogenic agents was also discussed. We also wanted to understand the importance that cancer patient associations place on this relationship, the understanding that the media has of the problem and the ethical aspects that all this raises.

MATERIAL AND METHODS

The members of the Board of Trustees of the Health Sciences Foundation proposed a series of questions on the relationship between viruses and cancer that would answer the main uncertainties of the problem at present. The questions were accepted by a multidisciplinary panel of experts. Each question was reported by one of the experts and subsequently the text and its conclusions were discussed by the working group until conclusions were reached that were accepted by all.

We now turn to the questions and their development. The views expressed are not intended to set out any particular recommendation or therapeutic indication and represent only the opinion of the group's speakers and not necessarily that of the institutions in which they work.

QUESTION: WHAT ARE THE MAJOR HISTORICAL MILESTONES IN THE VIRUS AND CANCER RELATIONSHIP?

In the early 1970s, most tumours were assumed to be caused by viruses, so much of the research into cancer revolved around tumour viruses. This assumption was eventually discarded and today it is estimated that only 10-15% of cancers are caused by viruses. However, research into viruses and cancer, the milestones of which are summarised below, proved decisive in understanding the aetiology of the disease [1-4].

Discovery of the avian sarcoma virus

In the early 1900s, Peyton Rous, working at the Rockefeller Institute in New York, showed that a type of sarcoma affecting chickens was transmissible through homogenate filtrates that could only contain viruses. Years later, the causative virus was isolated and named Rous Sarcoma Virus (RSV). This discovery marks the beginning of research into tumour viruses.

Transformation in cell cultures

During the 1960s, Harry Rubin and Howard Temin managed to establish a new experimental model to study tumour viruses at the California Institute of Technology. It consisted of the infection of fibroblasts in culture with the Rous Sarcoma Virus. The infection caused radical changes in the phenotype of the cells, which acquired capacities characteristic of tumour cells, such as: reduced dependence on growth factors, immortality, anchorage-independent growth, motility, invasiveness,
loss of contact inhibition, metabolic reprogramming and the ability to form tumours in syngeneic or immunosuppressed mice. These changes were called cellular transformation. The new experimental system ushered in a new era in tumour virus research [5, 6].

**Discovery of DNA tumour viruses**

In 1960, at the Rockefeller Institute in New York, Richard Shope identified the virus that causes papillomas in rabbits. Papillomas are benign tumours that rarely progress to squamous cell carcinomas of the skin. Unlike Rous Sarcoma Virus, whose genetic material is RNA, papillomaviruses are made of DNA. There are currently more than 100 human papillomaviruses (HPV) described. The discovery of HPV was followed by the discovery of more tumour viruses whose genetic material is DNA, including: i) Polyomavirus, which when infecting mice caused tumours of various origins, hence its name, ii) *Simian Virus 40* (SV40), which is a lytic virus in some primate cell types and silent in rodent cells and is therefore called non-permissive. In non-permissive cells, SV40 produces cellular transformation, although at a very low frequency [7-9].

Papillomavirus, polyomavirus, and SV40 are known as papovaviruses. Their DNA is a small circular molecule.

Other viruses with oncogenic capabilities include: i) Adenovirus, which infects the airways and can produce tumours when it infects hamster cells, ii) Herpesvirus, which produces different tumours in human cells, iii) Epstein Barr Virus (EBV), which can produce Burkitt’s lymphomas or nasopharyngeal carcinomas. Compared to the papovaviruses, the latter three are very large.

**Integration of the virus into the host cell genome**

Several lines of evidence showed that the tumour viruses had to remain active in the tumour cells to maintain the transformed phenotype. The inactivation of the virus produced a return to the normal cellular state.

Different viruses use different mechanisms to keep tumour cells from dividing after division. Some, like EBV or HPV, become integrated, becoming a part of the cell genome. Almost all cervical tumours are produced by HPVs, which make up the part of their genome necessary for cell transformation (the E6 and E7 oncogenes) [10].

Herpesviruses use two strategies to perpetuate their genome in tumour cells. Some, such as human herpesvirus-6 (HHV-6) integrate into the telomeric regions of the chromosomes, which are transcriptionally silenced, making the virus able to remain latent for long periods of time. Others, such as the herpesvirus that produces Kaposi’s sarcoma (HHV-8), replicate episodically, i.e., extrachromosomally, but are able to bind to chromosomes temporarily to ensure replication.

While DNA virus integration solved how they could remain in the tumour cells, it created a seemingly unsolvable problem. How did RNA viruses like Rous Sarcoma Virus remain? The solution to this problem led to one of the most surprising discoveries in biology: retrotranscription. Howard Temin and David Baltimore showed that RNA viruses copy their genome into double-stranded DNA, which integrates into the host cell’s genome. Retrotranscription violated the dogma proposed by Francis Crick by which the flow of biological information follows the DNA-RNA-Protein direction. In fact, retrotranscription is a mandatory step in the life cycle of retroviruses; as the RNA viruses that undergo this process are called. In contrast, the integration of DNA viruses is a very rare event (< 1 in 1,000 infections).

**Identifying the first of a long list of oncogenes**

Rous Sarcoma Virus, like most retroviruses, has a reduced genome (~10 kb). In it are encoded the genes necessary for its retrotranscription and integration, as well as the proteins that form part of its capsid. This simplicity made them very useful tools in identifying the elements required for cell transformation. By comparing viral variants capable of replication but not transformation, with variants capable of transformation but not replication, it was concluded that a single gene, called src, was responsible for cell transformation.

Characterising the src oncogene, in 1974 Harold Varmus and Michael Bishop, working at the University of California, San Francisco, obtained a totally unexpected result [11-13]. They found copies of src in cells that had not been transformed by Rous Sarcoma Virus. Actually, src is a gene present in totally normal cells that was hijacked by the tumour virus Rous Sarcoma Virus. This result was an authentic revolution and led to several conclusions that have revolutionised the way we think about cancer, some of which are detailed below:

- The cellular version of src, known as c-src had to play a role in the homeostasis of healthy cells.
- An ancestral form of the Rous Sarcoma Virus acquired the src after infecting a normal cell. In fact, there is ALV (avian leukaosis virus) which is infectious but not tumoral and has the same genes as Rous Sarcoma Virus, except src.
- The identification of c- and v-src opened the possibility that in the normal genome there were numerous proto-oncogenes that could have been co-opted by other tumour retroviruses.
- A single oncogene, v-src was sufficiently pleiotropic to modify the 20–30 characteristics that differentiate a normal cell from a tumour cell.

The discovery of src initiated a race to identify oncogenes carried by retroviruses, which led to the characterisation of oncogenes involved in the pathogenesis of numerous tumours. These are two examples:

- The avian myelomatisis virus, which produces transformation of marrow cells by means of the v-myc oncogene. In this case the original non-tumour virus was also the AVL.
- The Harvey and Kirsten sarcoma viruses, which led to the identification of the H-Ras and K-Ras oncogenes.

Currently the list of oncogenes associated with retroviruses exceeds two dozen [14].
QUESTION: AT PRESENT, WHAT PROPORTION OF MALIGNANT TUMOURS IN GENERAL HAVE A PROVEN OR PROBABLE CAUSALITY IN A VIRAL INFECTION?

It is accepted that one out of six tumours in the world is probably associated with an infection. It is important to highlight that in the series published on this subject, the sources of information and the criteria for detecting infectious agents are heterogeneous and this conditions the reliable approach to the subject.

The International Agency for Research on Cancer (IARC) in Lyon [15] through the Epidemiology and Infections Group has made significant contributions that underpin some of the current knowledge in this area. In a study by Plummer et al [16], published with data from 2012, it is established that of the 14 million new cases of cancer per year, 2.2 million (15.4%) were attributed to infections. Excluding one bacterium (Helicobacter pylori with 770,000 cases), which proved to be predominant, the next four most important infectious agents worldwide were HPV (640,000), Hepatitis B Virus (HBV, 420,000), Hepatitis C Virus (HCV, 170,000) and EBV (120,000). Furthermore, in sub-Saharan Africa, Kaposi's Sarcoma (caused by HHV-8) emerged as the second largest contributor to the cancer burden. In 2008, there were 7.5 million deaths from this type of tumour. Two-thirds of these deaths occur in poor countries. To these can be added the role of certain retroviruses linked to oncogenic capabilities such as HTLV-1 and Human Immundeficiency Viruses.

It should be noted that the role of the virus as an aetiological agent of tumours is conditioned by its prevalence, by its detection methods and by the estimation of the attributable risk in exposed populations. The accuracy of the measurement of exposure is critical as it ensures that the different methods for calculating attributable risk provide results that are comparable to reality. The experience of the "Global burden of cancers attributable to liver flukes" group shows that infections capable of becoming carcinogenic have demonstrated that biomarkers that are sensitive and capable of distinguishing high-risk from low-risk infections are essential in elucidating the true strength of their role in cancer [18, 19].

In the light of the contributions made in this regard, it seems appropriate to refer to an assessment of the cancer burden in France in 2015 attributable to infectious agents [20]. The authors conduct a systematic review of representative series on the prevalence of infectious agents in the main associated cancer types. They tracked original studies published up to September 2016 in PubMed and performed a random-effects meta-analysis. Cancer incidence data were obtained from the French Network of Cancer Registries, allowing the calculation of national incidence estimates in accordance with validated methods. Of the 352,000 new cancer cases in France in 2015, 14,336 (4.1% of all new cancer cases) were attributable to infectious agents. The largest contribution by infectious agents was represented by HPV and Helicobacter pylori, responsible for 6,333 and 4,406 new tumour cases (1.8 and 1.3% of all new cancer cases) respectively.

An additional aspect that supports the robustness of the contributions is that three main groups of infectious agents and cancer modalities were used for the estimation of tumours attributable to the infection. Firstly, cancer sites where infectious agents are assumed to contribute to almost 100% of all cases (cervical cancer due to high-risk HPV). Secondly, cancer sites attributable to infection whenever the infectious agent is detected in the tumour tissue using sensitive and validated detection methods (oral, oropharyngeal, anal, laryngeal, vulvar, vaginal and penile cancers due to high-risk HPV; nasopharyngeal cancers, Hodgkin's and non-Hodgkin's lymphomas due to EBV and - although it is a bacterial infection - gastric MALT lymphoma due to H. pylori). Thirdly, cancer cases where infectious agents increase the risk of developing a tumour, but are not responsible for all new cancer cases, even when antibodies to the infectious agent are detected in the serum.

The proportion of new cancer cases in France associated with infectious agents is comparable to that of other high-income countries [21, 22]. The estimates of this study are overall slightly higher (4.1%) than those of the specific analyses for the United Kingdom [21] and Australia [22] in 2010, where gender categorisation showed that 2.5% and 2.4% of all new cancer cases among men were attributable to infectious agents in both countries, respectively, and 3.5% and 3.7% of all new cancer cases among women were also attributable to infectious agents. These differences with previous studies are due, in part, to differences in the risk of infection between areas, as well as differences in what is considered the gold standard for measuring the prevalence of infection in association with specific cancers [23].

There are important global differences, depending on health and sociocultural conditions, in how these infections are responsible for tumour genesis. Thus, in Australia and New Zealand the infectious origin of a cancer is in 3.3% of cases compared to 32.7% in Sub-Saharan Africa [24]. Improving the quality and quantity of knowledge seems essential so as to prioritise preventive programmes and reduce the disease burden.

Conclusion

It is estimated that one in six tumours in the world is probably or demonstrably associated with an infection. The role of viruses as an aetiological agent of tumours is conditioned by their prevalence, detection methods and the estimate of attributable risk in exposed populations. The most relevant agents are HPV, HBV, HCV, EBV and depending on the zone HHV-8, HTLV-1 and the Human Immundeficiency Viruses.
Infections by certain microorganisms (viruses, bacteria, and parasites) are risk factors for developing certain cancers. According to the latest update of the IARC monograph on biological agents associated with cancer, the following microorganisms were classified as carcinogenic (group 1) (Table 1).

In 2016, the estimate of the burden of cancers associated with infectious agents was published [17]. In this estimate, the number of new cases diagnosed in 2012 attributed to the infections previously described in the table by country was calculated by combining the incidence rates of the different cancers with estimates of the attributable fractions of the different infectious agents. The attributable fractions for each infection are calculated with the infection prevalence in cancer cases and the relative risk. These attributable fractions are highly variable depending on the infectious agent and country. Of 14 million new cancer cases diagnosed in 2012, according to GLOBOCAN 2012, 2.2 million (15%) were attributable to infections. This percentage varies greatly by region, from 5% in the USA, Canada, Australia and some European countries, to over 50% in some Sub-Saharan African countries.

The most important infectious agents were Helicobacter pylori (770,000 cases), HPV (640,000 cases), HBV (420,000 cases), HCV (170,000 cases) and EBV (120,000 cases). For many of these infections-cancers there are clearly effective preventive strategies such as vaccines and early detection programmes. Vaccination and early detection programmes with treatment should be established to reduce the incidence of these cancers associated with infections.

QUESTION: WHICH ARE THE TUMOURS WITH THE MOST CLEARLY DEMONSTRATED VIRAL AETIOLOGY?

Infections by certain microorganisms (viruses, bacteria, and parasites) are risk factors for developing certain cancers. According to the latest update of the IARC monograph on biological agents associated with cancer, the following microorganisms were classified as carcinogenic (group 1) (Table 1).

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**Conclusion**

Overall, 15% of cancers are associated with an infectious causal agent. The most important viral agents according to the number of new cancer cases diagnosed are Human Papilloma Virus, Hepatitis B Virus, Hepatitis C Virus and Epstein–Barr Virus.

There are major differences in the incidence of these tumours according to geographical region and socio-economic development.

Vaccines and early detection programmes could dramatically reduce the incidence of cancer associated with viruses, such as HPV and hepatitis B and C viruses.

**Table 1** Infectious agents associated with cancer. Adapted from Monograph 100. Review IARC; The Lancet Oncology. 2009

<table>
<thead>
<tr>
<th>Group 1. Agent</th>
<th>Cancers with sufficient evidence of association in humans</th>
<th>Cancers with limited evidence of association in humans</th>
<th>Mechanisms</th>
</tr>
</thead>
<tbody>
<tr>
<td>Epstein-Barr virus (EBV)</td>
<td>Nasopharyngeal cancer, Hodgkin’s lymphoma, Burkitt’s lymphoma</td>
<td>Gastric carcinoma, lymphoepithelioma-like carcinoma</td>
<td>Cell proliferation, apoptosis inhibition, genomic instability, cell migration</td>
</tr>
<tr>
<td>Hepatitis B virus (HBV)</td>
<td>Liver cancer</td>
<td>Non-Hodgkin’s lymphoma, cholangiocarcinoma</td>
<td>Inflammation, liver cirrhosis, chronic hepatitis</td>
</tr>
<tr>
<td>Hepatitis C virus (HCV)</td>
<td>Liver cancer, non-Hodgkin’s lymphoma</td>
<td>Cholangiocarcinoma</td>
<td>Inflammation, liver cirrhosis, liver fibrosis</td>
</tr>
<tr>
<td>Kaposi’s Sarcoma associated Herpesvirus (HHV-8)</td>
<td>Kaposi’s sarcoma, primary cavity lymphoma</td>
<td>Castleman’s disease</td>
<td>Cell proliferation, apoptosis inhibition, genomic instability, cell migration</td>
</tr>
<tr>
<td>Human Immunodeficiency Virus (HIV)</td>
<td>Kaposi’s sarcoma, Hodgkin’s lymphoma, non-Hodgkin’s lymphoma, cancer of the cervix, anus, conjunctiva</td>
<td>Cancer of the vulva, vagina, penis, liver, non-melanoma skin cancer</td>
<td>Immunosuppression (indirect mechanism)</td>
</tr>
<tr>
<td>High-risk Human Papillomavirus (HPV)</td>
<td>Carcinoma of the cervix, vulva, vagina, penis, anus, oropharynx, oral cavity and larynx</td>
<td></td>
<td>Immortalisation, genomic instability, inhibition of DNA repair response, anti-apoptotic activity</td>
</tr>
<tr>
<td>Human T-cell Lymphotropic Virus (HTLV)</td>
<td>Adult T-cell leukaemia and lymphoma</td>
<td></td>
<td>Immortalisation and transformation of T-cells</td>
</tr>
<tr>
<td>Helicobacter pylori</td>
<td>Gastric Carcinoma</td>
<td></td>
<td>Inflammation, oxidative stress, altered cell proliferation and gene expression, mutation, methylation</td>
</tr>
<tr>
<td>Clonorchis sinensis</td>
<td>Cholangiocarcinoma</td>
<td></td>
<td>Inflammation, oxidative stress, cell proliferation</td>
</tr>
<tr>
<td>Opisthorchis viverrini</td>
<td>Cholangiocarcinoma</td>
<td></td>
<td>Inflammation, oxidative stress, cell proliferation</td>
</tr>
<tr>
<td>Schistosoma haematobium</td>
<td>Urinary Bladder Cancer</td>
<td></td>
<td>Inflammation, oxidative stress, cell proliferation</td>
</tr>
</tbody>
</table>
QUESTION: IS IT POSSIBLE TO BROADLY UNDERSTAND THE PATHOGENESIS OF VIRUS-MEDIATED ONCOGENESIS? WHAT ARE BRADFORD HILL’S PRINCIPLES OF CAUSALITY?

As described above, viruses can transform normal cells into tumour cells by activating oncogenes. In some cases, such as Rous Sarcoma Virus, the virus itself contains the oncogene. In others, such as EBV, the virus causes the proliferation of lymphoid cells that eventually remodel their genome so that the expression of the c-Myc oncogene becomes abnormally high. HPVs integrate pieces of their genome randomly into cells. When these pieces include the oncogenes E6 and E7, they transform the cells, as happens in the vast majority of cervical and some oropharyngeal tumours [2-4, 7, 14, 25-31].

In 1965, Sir Austin Bradford Hill [1] listed nine principles that must be demonstrated epidemiologically in order to establish a causal relationship between an event and the observed effect. Bradford Hill’s principles of causality have been used frequently in public health research. None of the principles proves a causal relationship per se; the more principles are fulfilled, the more the notion of a causal relationship is strengthened. Of particular relevance was the application of Bradford Hill principles in the demonstration of the causal relationship between tobacco use and cancer.

The following is a list of principles and how they demonstrate the recently proposed cause-and-effect relationship between HPV and oropharyngeal tumours [10]:

1. The power of the association. While a small association does not mean that there is no causal effect, a larger association makes it more plausible. For example, the risk of oral squamous cell cancer is 2.8 times higher in patients infected with certain strains of HPV.

2. Consistency. If the findings of different observers in different populations are consistent, the probability of causality increases. Different independent studies, conducted in entirely different populations, came to the same conclusion about the relationship between HPV and oral cancers.

3. Specificity. HPV is associated with oropharyngeal cancers, where HPV infections have been identified.

4. Temporality. The effect has to occur after the cause. HPV infections precede the development of oropharyngeal tumours. In fact, the longer the infection has lasted, the greater the risk of cancer.

5. Biological gradient. Greater exposure should generally lead to greater incidence of the effect. However, in some cases, the mere presence of the factor can trigger the effect. In the case of HPV infections the gradation is difficult to establish.

6. Plausibility. A mechanism that explains how the cause can generate the possible effect is very useful, although it depends on the state of our knowledge. The fact that HPV contains oncogenes, whose action mechanism is very well characterised, increases the possibilities of a causal relationship.

7. Coherence. Coherence between epidemiological and laboratory findings increases the likelihood of causality. HPV oncogenes can transform cells in the laboratory.

8. Experimental evidence. It’s very much related to the previous point. There is ample experimental evidence of the ability of HPV to direct malignant transformation in various experimental models.

9. Analogy. The effect is seen in analogous situations. Certain oncogenic variants of HPV, such as HPV16, have a clear causal relationship with oral cancer, while very low tumourigenic variants are difficult to find in this type of tumour.

Conclusion

The Bradford Hill principles are a widely used tool for establishing a causal relationship between an event and a disease. Some tumour viruses such as the Human Papilloma Virus meet these criteria and are therefore considered to be the causative agent of cervical and, more recently, oropharyngeal carcinomas.

QUESTION: HOW COULD THE INVOLVEMENT OF HPV IN GENITAL TUMOURS BE SUMMED UP IN FIGURES?

HPV is one of the main causative agents of cancer as it is implicated in 4.5% (640,000 cases) of all new cancer cases worldwide each year (8.6% women; 0.9% men). It is estimated that in 2012 it caused more than half of all cancers attributable to infections in women (570,000 cases) [32]. The vast majority of HPV-related neoplasms develop in the anogenital tract (primarily cancer of the cervix and less frequently cancer of the vagina, vulva, and anus).

Cervical cancer is the fourth most common cancer among women and the fourth leading cause of cancer death worldwide (the second in women aged 15-44). In 2012 the number of new cases was estimated at 528,000 and the number of deaths at 266,000 [33]. More than 84% of the cases are diagnosed in developing countries. The age-standardised incidence rate is 14.0 cases and the mortality rate is 6.8 deaths per 100,000 women/year, with large variations between geographical regions and countries, especially depending on the extent of screening.

All other anogenital cancers that affect both women and men have a lower incidence. Worldwide, 115,000 cases are diagnosed, of which 68,500 are attributable to HPV. In men, 30,000 cases are diagnosed (17,000 in the anus and 13,000 in the penis) and in women 38,500 cases (18,000 in the anus, 8500 in the vulva and 12,000 in the vagina) [32]. In both the penis and the vulva there is a clear distinction between basaloid or “warty” tumours clearly causally related to HPV and keratinizing tumours that are seen at older ages and are not related to HPV. Moreover, in these locations it is also more difficult to establish in which cases HPV is the attributable factor. In this sense, most studies accept in addition to the detection of DNA-HPV in the tumour, other transforming activity markers such as the presence of E6 or E7 mRNA or p16. Based on these data, it is estimated that the
cases attributable to HPV are approximately 88% of anal cancer, 51% of penile cancer, 78% of vaginal cancer, and between 15% and 48% of vulvar cancer [17].

Data on the burden of HPV-related neoplastic disease in 32 countries in Europe have recently been published for 2015. The estimated number of new cases per year for cervical cancer is 34,939 cases, for 31 countries as a whole (assuming that HPV is causal for practically 100% of cases). In vulvar cancer, there are an estimated 9,776 cases (1,554 if we take into account a fraction attributable to HPV of 15.8%). For vaginal cancer, there are an estimated 2,224 cases (1,562 if we have the 70.2% fraction attributable to HPV). In anal cancer, 4,663 cases (2,801 with a fraction attributable to HPV of 87.1%). Finally, for penile cancer, 4,231 cases (1,227 cases with a fraction attributable to HPV of 20.9%) [34].

Although anogenital premalignant lesions are much more frequent than cancer, their lack of recording makes it very difficult to estimate their prevalence. A recent analysis provides data on Europe for 2015. It is estimated that between 263,227 and 503,010 cases of grade 2+ cervical intraepithelial neoplasia (CIN) are diagnosed annually; 1,549 cases in women and 1,097 cases in men of grade 2/3 anal intraepithelial neoplasia (AIN), between 13,997 and 27,773 cases of grade 2 and 3 vulvar intraepithelial neoplasia (VIN), and between 2,596 and 4,751 cases of grade 2 and 3 vaginal intraepithelial neoplasia (VaIN).

High quality cancer incidence data are essential. Such information is currently only available to a small fraction of the world’s population. The absence of this information may lead to a misinterpretation of the actual disease burden, as well as to a possible delay or neglect in the implementation of preventive measures or an impossibility to evaluate such measures.

Conclusion

The neoplastic and pre-neoplastic disease burden related to HPV is very high, affecting women in particular.

In Europe, although most countries have cervical cancer screening programmes, HPV-related neoplasms account for more than 53,000 cases annually and precancerous lesions for approximately 680,000 - 844,000 cases.

HPV-related cancers, despite the availability of multiple preventive strategies, remain a major cause of morbidity and mortality in many parts of the world, particularly in less developed countries. A very high fraction of these neoplasms are preventable by screening and detection of precursor lesions and with the currently available vaccines.

QUESTION: WHAT WOULD BE THE IMPACT OF HPV AS A CAUSAL AGENT OF EXTRAGENITAL TUMOURS?

Beyond anogenital locations, HPV is associated with head and neck cancer. The term "head and neck cancer" has been widely adopted in recent literature to include lesions at several anatomical sites: the lip, oral cavity, nose and sinuses, nasopharynx, oropharynx, hypopharynx, and larynx [35]. HPV is predominantly associated with oropharyngeal cancer and within that with tonsil cancer.

The classic risk factors for head and neck cancer are tobacco and alcohol use, but recent decades have seen an increase in the proportion of HPV-associated oropharyngeal cancer cases. This increase may be due to changes in sexual behaviour as well as a decrease in the prevalence of smoking.

Annually, 24,000 new cases of HPV-associated oropharyngeal cancer are diagnosed in men and 5,500 in women [32]. Globally, the proportion of HPV-associated cases in oropharyngeal cancer is 30% (and of the tonsil about 50%) with a large difference depending on countries-regions, being more than 80% in countries such as the USA or Northern Europe, and less than 10% in countries in Africa or other regions [16, 19]. HPV 16 is the most prevalent type found in more than 80% of HPV-positive tumours. It should be noted that HPV-related oropharyngeal cancers differ from unrelated cancers in both epidemiological and clinical and molecular characteristics. Patients with positive HPV16 tumours appear to have better overall and disease-specific survival compared to the negative group. Laryngeal squamous cell papilloma and recurrent respiratory papillomatosis are well established HPV-induced tumours, primarily 6 and 11.

Conclusion

HPV causes a high proportion of head and neck cancers, particularly those of the oropharynx and within the oropharynx especially those of the tonsil.

QUESTION: IS IT POSSIBLE TO MEASURE THE IMPACT OF THE HEPATITIS B AND C VIRUSES ON THE INCIDENCE OF LIVER TUMOURS?

Within the natural history of chronic hepatitis of viral origin, the chronic hepatitis-cirrhosis-cancer sequence has been well known for decades. This incontrovertible data allows us to pose different questions of great relevance, such as whether it is possible to know the epidemiological impact of this association.

The question is not simple; unlike the unidirectional and almost linear relationship between other types of viral infection and the development of cancer, the oncogenic capacity of hepatotropic viruses is modulated by different factors, as listed below.

i) Cirrhosis is an oncogenic element "per se"

A differential and characteristic data of hepatocellular carcinoma is that in more than 85% of cases it occurs in a liver affected by cirrhosis. Strikingly, it is common to find hepatocellular carcinoma in patients with cirrhosis of non-viral cause (such as alcoholism and particularly metabolic fatty liver disease). This fact, besides having a marked influence on
the treatment strategy, indicates that the mechanisms associated with the development of cirrhosis (inflammation, angiogenesis, fibrogenesis, etc.) generate a micro-environment with its own oncogenic potential independent of the viral infection itself.

ii) Each virus has its own oncogenic potential

Regardless of the relatively similar natural history of chronic hepatitis associated with virus B and virus C, both viruses are very different in terms of their own character (DNA or RNA virus), their replicative cycle (with or without integration into the host cell genome) and the different degrees of production of necrosis, oxidative stress, expression of viral proteins etc. Furthermore, the relationship between the intensity of viral replication and the risk of cancer is clearly different; while it is clear in virus B, it does not exist in the case of virus C [36, 37]. Finally, the possibility of liver cancer in patients without cirrhosis is markedly more frequent in patients with hepatitis B, especially in Asia.

iii) There are significant geographical differences

The contribution of the different risk factors linked to the development of hepatocellular carcinoma has clear geographical differences that are related to the most frequent cause(s) of cirrhosis in each area. Thus, the incidence of hepatocellular carcinoma in Europe is mostly associated with patients with cirrhosis due to virus C, while in Asia this association is with virus B (Table 2).

iv) Chronic viral infection is associated with other risk conditions

It is well known that the progression of chronic liver disease is affected by different factors that are often present in patients with viral origin chronic liver disease, such as alcohol consumption, the presence of metabolic syndrome, or tobacco use. It should be noted that they are all well-known co-factors involved in the development of cancer. Therefore, the precise contribution of each of the factors involved (viral and non-viral) in the development of liver neoplasms is difficult to define [38].

In any case, there is no doubt that both by direct and indirect mechanisms (through the presence of cirrhosis, as mentioned above) the association between hepatotropic viruses and liver cancer is of unquestionable relevance, being possibly responsible for more than 60% of primary liver tumours.

QUESTION: WHAT HAVE HTLV-I AND HIV MEANT AS DIRECT OR INDIRECT CANCER-CAUSING AGENTS?

HTLV-1 and HIV are two viruses first described in the 1980s and directly or indirectly involved in human oncogenesis. Both share the same transmission routes (breast milk, sexual relations, blood products), the integration of their genome into the T cell and a long latency period, more pronounced in the HTLV-1, between infection and disease development.

Human T lymphotropic virus type 1 (HTLV-1)

HTLV-1 infects 10-20 million people worldwide and its main geographical distribution is Japan, the Caribbean, South America and sub-Saharan Africa. Only 5% of infected people develop disease and the 2 conditions that it causes are adult T-cell leukaemia-lymphoma (ATL) and HTLV-1-associated myelopathy or tropical spastic paraparesis [39].

People with HTLV-1 positive serology have a 2-5% risk of
Human Immunodeficiency Virus (HIV)

The origin of tumour diseases in patients with HIV infection is multifactorial, the most important factors being HIV itself, immunosuppression, co-infection with oncogenic viruses and increased survival due to the use of antiretroviral treatment (ART) (Figure 1). HIV is a necessary but not sufficient condition, as it is not strictly considered an oncogenic virus. However, fragments of its genome have been found in some patients with NHL, and the tat gene appears to be involved in the growth of Kaposi’s sarcoma cells. For all these reasons, it is considered an essential agent, but not the only one for the development of the tumours that appear in these patients [43].

Immunosuppression and uncontrolled HIV viremia increase the risk of developing neoplasms, such that an undetectable viral load and CD4+ lymphocytes >500 per mm³ are factors that protect against the development of some tumours, which explains why a decrease in the number of AIDS-defining malignancies (ADM) has been observed with the use of ART. On the other hand, paradoxically, there is an increase in non-AIDS-defining malignancies (NADM) due,
in part, to the increased survival of patients as a result of the use of ART, which prolongs life without completely restoring immune function (Figure 2). All of this conditions a greater exposure to environmental carcinogens (tobacco, ultraviolet light), a greater possibility of superinfection by oncogenic viruses and an increase in the possibilities of genetic mutations appearing in patients with a family history of cancer, giving rise to the appearance of neoplasms that
in other times did not develop due to the early mortality of the patients (Figure 1) [43, 44].

Another important point is that the presence of HIV causes a persistent stimulation of B lymphocytes, which alters anti-tumour immunity, facilitates the expression of oncogenic genes and allows superinfection by viruses with oncogenic capacity, such as EBV, HVH-8, HBV, HCV or HPV. All ADM are associated with viral diseases, as well as the most frequent ones within NADM except for pulmonary carcinoma directly related to smoking (Figure 2) [44].

Table 4 shows the differences and similarities between HTLV-1 and HIV.

**Conclusion**

Both viruses, HTLV-1 and HIV share the transmission routes, infect CD4+ T-lymphocytes and are involved in human oncogenesis. The HTLV-1 in a direct way, originating the T cell Leukaemia-Lymphoma and HIV, favouring the appearance of Tumours, both AIDS defining and non-defining.

**QUESTION: WHAT PROPORTION OF HIV PATIENTS WHO DIE DO SO BECAUSE OF AN HIV-ASSOCIATED NEOPLASM?**

Since the introduction of ART in 1996, mortality among people living with HIV (PLHIV) has decreased significantly (Figure 3) so that in 2016, 498 people died while in the initial years of the epidemic the numbers were much higher, reaching 5,857 deaths in 1995. However, immune reconstitution is generally not complete, so patients are exposed to the effects of chronic inflammation and persistent immune activation. Both processes favour the appearance of the so-called non-AIDS diseases, whose importance has progressively increased in recent years and which are one of the main causes of morbidity and mortality among PLHIV. This broad group of diseases includes most comorbidities, which in some cases become more important than the HIV infection itself. Among other processes, cardiovascular pathology, renal disease, osteoporosis and bone fractures, neurocognitive impairment and tumour pathology stand out.

Malignant neoplasms are a major cause of hospitalisation and death in PLHIV as a result of ART use and a decrease in opportunistic infections. Today, cancer can be diagnosed in 25-40% of patients, and while in the pre-ART era only 10% of deaths were of tumour origin, now up to 30% are. In our environment, according to data from the CORIS cohort, neoplasms were the most frequent cause of death by non-AIDS diseases between 2004 and 2015. In this same study it is important to note that 38.2% of the tumours diagnosed were AIDS-defining malignancies (ADM) and 61.8% were non-AIDS-defining malignancies (NADM) (Table 5) [45].

At the EACS Congress, the results obtained from the review of neoplasms diagnosed from 1986 to 2018 in a Spanish centre were communicated [46]. 643 out of 5,411 patients (12%) suffered at least one neoplasm. NADM increased over time with a decrease in the ADM/NADM ratio (Figure 4). 101 patients (15.9%) died due to tumour progression, with higher mortality among NADM 25.3% vs. ADM 10.6% (p<0.001). The highest mortality was seen in tumours of the pancreas (90%), lung (68%) and colon (69%).

![Figure 3](image_url)  
**Figure 3** | Evolution of mortality in people living with HIV in Spain from 1991 to 2016 (data taken from the National Institute of Statistics)
In other series outside our scope, the results are similar. Thus, Engels et al. [48] found that approximately 10% of deaths in PLHIV receiving ART during the period 1995 to 2009 were due to a neoplasm with a progressive increase over time. A large proportion of the deaths were due to non-Hodgkin’s lymphomas, lung cancer and hepatocellular carcinoma. As in our environment, it is observed that deaths due to NADM are another Spanish group [47] analysed the mortality in 157 PLHIV and found that 56 deaths were directly related to AIDS and 13 of them (23.2%) were due to a ADM. In the same series another 86 deaths occurred due to non-AIDS related processes and 14 (16.4%) were as a result of a NADM. Therefore, the overall mortality due to a neoplasm in this series was 17%. Another Spanish group [47] analysed the mortality in 157 PLHIV and found that 56 deaths were directly related to AIDS and 13 of them (23.2%) were due to a ADM. In the same series another 86 deaths occurred due to non-AIDS related processes and 14 (16.4%) were as a result of a NADM. Therefore, the overall mortality due to a neoplasm in this series was 17%.

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**Table 5** Mortality of tumour origin in different series of people living with HIV

<table>
<thead>
<tr>
<th>Author</th>
<th>Study</th>
<th>Years</th>
<th>Results</th>
</tr>
</thead>
<tbody>
<tr>
<td>Gotti et al [49]</td>
<td>Analysis of survival</td>
<td>1998-2012</td>
<td>866 tumours were diagnosed in 13,388 patients:</td>
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<tr>
<td></td>
<td></td>
<td></td>
<td>- 435 ADM (51%)</td>
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<tr>
<td></td>
<td></td>
<td></td>
<td>- 431 NADM (49%)</td>
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<td>40% deaths due to tumour at the end of the analysis</td>
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<tr>
<td></td>
<td></td>
<td></td>
<td>Died due to cancer</td>
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<tr>
<td></td>
<td></td>
<td></td>
<td>- 419 females (8.2%)</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>- 1,062 males (9.1%)</td>
</tr>
<tr>
<td>Cevallos et al [51]</td>
<td>Mortality</td>
<td>2007-2014</td>
<td>1,999 patients died (11.7%)</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>- 336 due to NADM (16.8%)</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>- 115 due to ADM (5.8%)</td>
</tr>
<tr>
<td>Croxford et al [52]</td>
<td>Mortality</td>
<td>2016</td>
<td>There were 206 deaths among 38,700 patients</td>
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<tr>
<td></td>
<td></td>
<td></td>
<td>NADM resulted in 40 deaths (19.4%)</td>
</tr>
</tbody>
</table>
increasing in importance while there is a decrease in the number of deaths due to AIDS itself.

All these studies show a decrease in mortality from AIDS-indicating diseases and a predominance of deaths from non-AIDS diseases in recent years, with neoplasms being one of the most frequent causes in all the series, even though the patients are on ART and virologically suppressed.

In Table 1 we see reflected the importance of death of tumour origin in PLHIV in other series different from those described in the text.

**Conclusion**

Non-AIDS defining malignancies are one of the most frequent causes of death in HIV-infected patients despite the fact that the sick receive effective antiretroviral treatment.

**QUESTION: WHAT IS THE INVOLVEMENT OF HERPESVIRUS IN THE CAUSALITY OF CANCER?**

The human herpes virus (HHV) group includes herpes simplex virus (HSV) 1 and 2, varicella zoster virus (VZV), cytomegalovirus (CMV), Epstein Barr Virus (EBV), HHV-6, 7 and 8. In general, they are characterised by producing primary infection at an early age and subsequently establishing latency in different tissues, from which they are able to reactivate, producing clinical manifestations again. In Table 6 we have summarised their most frequent clinical manifestations, including the tumour processes with which they have been associated.

Involvement in tumour processes is well established for EBV and HHV-8, and we will devote most of our review to them [53-55]. The rest of the herpes viruses are not considered to directly cause any type of cancer, although their role as co-factors is discussed. For example, there is some evidence to suggest that HSV-2 may increase the risk of cervical cancer after HPV infection and that it may play a role in the tumour genesis of serous ovarian carcinoma and in some prostate tumours [56]. Associations between HHV6 and different tumours have also been sought, but these have not yet been confirmed.

We will focus on the involvement of EBV in the development of tumour processes. EBV is the primary agent of infectious mononucleosis. It establishes latency asymptomatically in virtually all adults, in B cells, T cells, epithelial cells and myocytes and usually does not cause a cytopathic effect. However, it shows a clear ability to transform cells and is associated with the development of B-cell lymphomas, T-cell lymphomas, Hodgkin’s lymphomas and nasopharyngeal and other head and neck carcinomas [57]. In transplant recipients, it produces a Post-transplant lymphoproliferative disease (PTLD) that can be very aggressive [58].

- **Burkitt’s lymphoma** (BL), is the most common childhood malignancy in equatorial Africa (8-10 cases/10^5 annually) [57] and characteristically affects the jaw. The relationship between BL and EBV is practically 100% in endemic cases, 10-80% in sporadic cases and 30-40% in HIV-related cases. The cells usually express EBNA and CD20. It is considered that malaria may also be a co-factor in the production of this tumour, although the mechanism is not clear (immunosuppression of specific T cells, increased EBV viral load, or favouring reactivation of the virus in B lymphocytes, among other theories).
- **Nasopharyngeal carcinoma** is a rare tumour in most parts of the world, except in southern China, where it is one of the most common cancers [59]. Unlike Burkitt’s lymphoma, EBV is present in all anaplastic nasopharyngeal carcinoma cells, in both high- and low-incidence areas [60]. However, other co-factors are necessary (exposure to environmental carcinogens in food, genetic alterations, etc.), since the seroprevalence of EBV infection is much higher than the incidence of the tumour. Nasopharyngeal carcinoma cells express dormant proteins of EBV, including EBNA-1 and two membrane proteins, LMP-1 and LMP-2, along with the BamHI-A fragment of the EBV genome.
- **HIV+ patients** have a lymphoma incidence 60-100 times higher than expected and 70% of them are associated to EBV infection, usually of monoclonal origin. The profound immunosuppression of these patients facilitates the failure of EBV control by decreasing specific cytotoxic lymphocytes. **Non-Hodgkin’s lymphomas** in HIV+ patients often affect the CNS. In B-cell lymphomas, EBV-coded proteins promote cell immortalisation and proliferation by stimulating the NF-κB pathway and increasing the expression of anti-apoptotic genes. In HIV+ children, EBV infection has been linked to the development of smooth muscle tumours (leiomyoma and leiomyosarcoma). The non-Hodgkin’s lymphoma most frequently associated with EBV is diffuse large cell.
- **As early as the 1980s** it was shown that the EBV genome is found in more than 50% of **Hodgkin’s lymphomas**, including Reed-Sternberg cells [61]. It is also demonstrated in patients with HL-induced hemophagocytic syndrome. The proportion of HL with EBV is even higher in children under 10. The two most frequent markers in HL are EBER-1 and the latent membrane protein (LMP) [53].
- **EBV infection** is also associated with T-cell lymphomas, including fulminant forms of T-cell lymphoma after acute EBV infection. In the context of mature T/NK lymphomas, with the possible exception of extranodal NK/T cell lymphoma, the role played by EBV infection is even more complex than in B lymphomas and exceeds the intention of this review. It has been proposed that the type of infected cell, the EBV viral load in plasma, and the degree of viral lymphatic replication have prognostic value in T/NK cell lymphomas.
- **Another lymphoma associated with EBV infection** is **nasal angiocentric lymphoma**, which is endemic in Asia, Central and South America. This tumour affects the nasal septum, palate, gastrointestinal tract and, less frequently, the skin, testicles and peripheral nerves. EBV is detected in virtually all neoplastic cells that are probably derived from NK cells [62].
- **PTLD** is associated in most cases with EBV. The term includes a continuum of abnormalities ranging from benign polyclonal B-cell proliferation to malignant B-cell lymphoma. In these disorders the virus directly induces the transformation of naive or mature mem-
Overview of virus and cancer relationships. Position paper

E. Bouza, et al.

and one of the clearest examples of a tumour that requires infection by a virus and that is also modulated by the patient's immune status. The pathogenesis of the tumour requires infection with human herpesvirus 8 (HHV-8), also called Kaposi's sarcoma associated herpesvirus [63]. It is named after Moritz Kaposi, a Hungarian dermatologist who first described the entity in 1872.

Before briefly describing some of the characteristics of the KS, it should be noted that the HHV8 also causes two other tumours: cavity lymphoma or primary effusion lymphoma (PEL) and Castleman's disease.

KS is characterised by vessel proliferation (angiogenesis), accompanied by inflammation and proliferation of spindle cells. HHV-8 infects the cells and the gene products of the virus influence both the regulation of the cell cycle and the control of apoptosis. The infection induces a disordered proliferation of cells that mostly do not express a lytic cycle, but a latent infection. In addition, the function of the tumour suppressor genes is interrupted and recognition by the host immune system is prevented, thus promoting tumour growth. By performing a PCR in the biopsy, it is possible to amplify viral DNA sequences and also to identify, using immunohistochemistry, latent viral antigen in spindle cells (membrane proteins).

Other entities related to this virus are hemophagocytic lymphohistiocytosis, lymphomatoid granulomatosis or X-linked lymphoproliferative disease.

The treatment of all these tumours does not include the administration of antivirals since the cells are not in the lytic cycle, which is necessary for Acyclovir to be effective. The usefulness of cell therapy with specific cytotoxic T-lymphocytes against EBV and Rituximab is being studied.

Conclusion

Among herpes viruses the association with the development of neoplasms is well established for EBV and HHV-8. EBV is capable of promoting tumour transformation in multiple cells and is associated, among others, with B-cell lymphomas, including Hodgkin’s and non-Hodgkin’s lymphomas, T-cell lymphomas, nasopharyngeal carcinomas, other head and neck carcinomas, and post-transplantation lymphoproliferative disease.

QUESTION: WHAT IS THE RELATIONSHIP BETWEEN KAPOSI’S SARCOMA AND HERPES VIRUS?

Kaposi’s sarcoma (KS) is an angioproliferative tumour and one of the clearest examples of a tumour that requires infection by a virus and that is also modulated by the patient’s immune status. The pathogenesis of the tumour requires infection with human herpesvirus 8 (HHV-8), also called Kaposi’s sarcoma associated herpesvirus (KSHV)[63]. It is named after Moritz Kaposi, a Hungarian dermatologist who first described the entity in 1872.

Before briefly describing some of the characteristics of the KS, it should be noted that the HHV8 also causes two other tumours: cavity lymphoma or primary effusion lymphoma (PEL) and Castleman’s disease. Cavitory lymphoma is a B-cell non-Hodgkin’s lymphoma that has a preference for the peritoneal, pleural, and pericardial spaces. Castleman’s disease is also a very rare lymphoproliferative disorder that causes fever, splenomegaly, hepatomegaly, and lymphadenopathy, and can affect both HIV-negative and HIV-positive patients.

KS is characterised by vessel proliferation (angiogenesis), accompanied by inflammation and proliferation of spindle cells. HHV-8 infects the cells and the gene products of the virus influence both the regulation of the cell cycle and the control of apoptosis. The infection induces a disordered proliferation of cells that mostly do not express a lytic cycle, but a latent infection. In addition, the function of the tumour suppressor genes is interrupted and recognition by the host immune system is prevented, thus promoting tumour growth. By performing a PCR in the biopsy, it is possible to amplify viral DNA sequences and also to identify, using immunohistochemistry, latent viral antigen in spindle cells (membrane proteins).

Four types of KS are recognised depending on the type of patients affected: 1) Classic or sporadic KS, which corre-
sponds to the form originally described by Kaposi and which is extraordinarily rare (0.03% per year among HIV-positive males over 50 in the Mediterranean area). It usually occurs in males over 50-60 years of age and mainly affects people from the Mediterranean area (Sardinia, Sicily, etc.) or with Jewish ancestry; 2) Endemic KS prevalent in sub-Saharan Africa before the AIDS epidemic. It affects children and adults and it is believed that continuous exposure to certain types of soil facilitates the appearance of the disease; 3) iatrogenic KS, associated with the use of immunosuppressants and described mainly in solid organ transplant recipients; 4) finally, AIDS-associated KS (epidemic KS) [64]. With better control of the AIDS epidemic, incidence in the US has dropped from rates that reached 47 cases per year per million people. Since then, Kaposi's sarcoma has become less common, producing about 6 cases per million people each year. The different epidemiological and clinical aspects of these four types are summarised in Table 7.

Infection with the HHV8 virus is a necessary condition for the development of KS, but not sufficient, since other cofactors are required for its appearance, such as genetic predisposition (sporadic form), and above all immunosuppression, either iatrogenic (organ transplant) or due to HIV infection. Thus, the prevalence of seropositivity in the population is much higher than that of Kaposi's sarcoma.

The most common clinical presentation of KS, especially in the early stages of the tumour, is skin and mucosal lesions. It is characterised by the appearance of raised violet lesions that mainly affect the lower extremities, the face (especially the nose), the oral mucosa and the genitals. They may be accompanied by significant oedema, reflecting vascular obstruction by lymphadenopathy or the action of cytokines involved in the pathogenesis of the tumour. Oral lesions appear in one-third of patients and mainly affect the gums and palate and may interfere with nutrition and speech. It is important to remember that KS can also have extracutaneous manifestations, mainly in advanced stages, affecting mainly the gastrointestinal and respiratory tracts. The gastrointestinal system is now less frequently affected. Its diagnosis is difficult in the absence of concomitant skin lesions. Clinical manifestations of gastrointestinal KS include weight loss, abdominal pain, nausea and vomiting, gastrointestinal bleeding, malabsorption, intestinal obstruction, and diarrhoea. As for lung disease, it can appear with highly variable radiological findings, including nodules, single or multiple, interstitial or alveolar infiltrates, pleural effusion, adenopathy or solitary nodules. The diagnosis is based on the characteristic appearance of the lesions in endoscopy or bronchoscopy and is confirmed by biopsy.

The incidence of KS in solid organ transplant recipients is 0-5-1% and represents approximately 6% of post-transplant tumours, excluding skin cancer. This incidence is much higher than that of the general population and is justified by the chronic immunosuppression required by transplant patients. Although KS is mostly described in HIV-positive patients, primary infection after transplantation has been shown to be the most important risk factor in our environment [65]. This primary infection can be acquired through the transplanted organ, or through the administration of blood products or other routes. About half of the KS diagnosed in transplant recipients respond only by reducing immunosuppression, or by switching to Sirolimus (Rapamycin) which has anti-tumour and anti-angiogenic effects [66].

In the patient with HIV infection, KS has been described in all risk groups, although with a higher incidence among homosexual or bisexual men, in relation to a higher seropositivity rate among these patients (close to 40%) [67]. KS rates in some studies reached 18%-50% 10 years after the diagnosis of HIV infection, if the CD4 figure was <200 cells/mm³ [68]. However, although the rate of HHV-8 infection has not changed, early

<table>
<thead>
<tr>
<th>Table 7</th>
<th>Classification of the different Kaposi’s Sarcoma</th>
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<tbody>
<tr>
<td>Type</td>
<td>Risk groups</td>
</tr>
<tr>
<td>CLASSIC (Sporadic)</td>
<td>Male/Female 3:1, Age&gt;60, Mediterranean, Central or Eastern European or Middle Eastern origin</td>
</tr>
<tr>
<td>ENDEMIC (Africa)</td>
<td>Adult males; children of both sexes, Equatorial Africa</td>
</tr>
<tr>
<td>IATROGENIC (Immunosuppression)</td>
<td>Exogenous IS, SOT, Age &gt;50, Cyclosporin A</td>
</tr>
<tr>
<td>AIDS-related</td>
<td>MSM in developed countries, Heterosexual men and women in Africa</td>
</tr>
<tr>
<td>Cutaneous involvement</td>
<td>Distal lower limbs, Similar to classic (sometimes more aggressive, with adult lymphoedema), Distal lower limbs or scattered, Localised or scattered</td>
</tr>
<tr>
<td>Profound involvement (visceral)</td>
<td>Rarely, Children: deep ganglia and organs, Common</td>
</tr>
<tr>
<td>Evolution</td>
<td>Normally indolent, Rare rapidly progressive, Indolent or locally invasive in adults, Aggressive in children, May disappear on modification of IS, May be aggressive, May disappear with HIV control</td>
</tr>
</tbody>
</table>

MSM: men who have sex with men; IS: immunosuppression; SOT: solid organ transplant
and effective antiretroviral treatment has significantly reduced the frequency of this tumour. The standardised incidence rate (SIR) for Kaposis’s sarcoma compared to the general population was 22,100, and declined to 3,640 with widespread use of antiretroviral treatment [69].

Some HIV+ patients with Kaposis’s sarcoma present with a systemic inflammatory syndrome due to cytokine release (KICS) that is characterised by fever, gastrointestinal, respiratory and neurological manifestations and is associated with an elevated KSHV tumour load.

Treatment of KS depends on the extent of the tumour and its location and symptoms. As with solid organ transplant recipients, HIV-positive patients may respond to immune recovery from the initiation of highly active antiretroviral therapy (HAART), although others may also require local or systemic chemotherapy. It has also been observed that KS is less frequent in patients who have received Ganciclovir or Foscarnet, demonstrating the relative sensitivity of the virus to these antivirals. The relative survival over five years is over 75%.

Conclusion

Kaposis’s Sarcoma is a tumour that requires prior infection with Human Herpes Virus type 8 as a necessary, but not sufficient, cause. Tumour development requires associated factors that may be genetic or due to immunosuppression. It exists in four clinical versions: Classic, Endemic, iatrogenic or HIV-associated Kaposis’s Sarcoma.

QUESTION: DO MICROORGANISMS OTHER THAN VIRUSES HAVE AN ONCOGENIC ROLE?

Viruses are certainly the most oncogenic infectious agents. However, there are other non-viral microorganisms that are clearly related. A bacterium: Helicobacter pylori; and three parasites (the so-called “Oriental liver flukes”): Schistosoma haematobium, Clonorchis sinensis and Opisthorchis viverrini,* are classified in Group I of carcinogens by the International Agency for Research on Cancer

Cancer-related bacteria

-Helicobacter pylori is a helical bacillus that lives exclusively in the human gastric epithelium. H. pylori infection can cause inflammation of the gastric mucosa that can progress, leading to gastritis, peptic ulcers and mucous-associated lymphoid tissue lymphoma (MALT). Because of its spiral shape, it can literally “screw itself in” to colonize the stomach epithelium, in addition to having the urease that allows it to neutralise its acidic environment. The bacteria generates free radicals, injects a cytotoxin known as CagA that alters the structure of the cells lining the stomach and generates locally high levels of molecules signalling inflammation such as TNF-alpha or interleukin 6 that could induce mutation of the epithelial cells of the stomach. Tropheryma whippelii, another bacteria in the digestive tract that causes Whipple’s disease, may be associated in a similar way with neoplasms.

Salmonella typhi has been linked with hepatobiliary and gall bladder tumours and -Chlamydia pneumoniae and Mycobacterium tuberculosis have been linked with lung cancer.

Chronic Bacterial Osteomyelitis and hidradenitis suppurativa are associated with squamous cell carcinoma of the skin also [70-74].

Cancer-related parasites

Millions of people in the world, especially those living in the tropics, suffer from some form of parasitic disease. The oncogenic role of certain parasites has been known for many decades. Three parasites, trematode helminths, cause urinary or hepatobiliary tumours [75, 76]:

-Schistosoma haematobium is very prevalent in sub-Saharan Africa and is a major cause of urinary bladder carcinoma. The disease is acquired after skin contact with fresh water infected with larvae. Adults invade the urinary venous plexuses and the eggs are expelled with the urine after passing through the bladder wall, causing haematuria, fibrosis and calcifications. Granulomas form on the wall of the bladder and evolve into nodules or masses that ulcerate, degenerating into both benign and malignant tumours. In addition, bladder lesions may increase the exposure of the epithelium to carcinogenic substances. Schistosoma japonicum may be oncogenic and produce colorectal cancer [77].

-Clonorchis sinensis is very prevalent in South East Asia (China, Thailand, Korea, Vietnam, Laos and Cambodia) and Opisthorchis viverrini (also found in Russia), to a lesser extent, causes cholangiocarcinoma [78, 79]. The disease is acquired by eating raw freshwater fish. Adults invade the bile ducts and occasionally the gallbladder and pancreatic duct, where they cause cholangitis and cholecystitis and can lead to cirrhosis. In massive and chronic infections, inflammation, scaling, metaplasia and ductal fibrosis can degenerate into cholangiocarcinoma and liver carcinoma, especially in patients exposed to carcinogenic agents such as nitrosamines. Opisthorchis felineus could perhaps be equally oncogenic.

- Other parasites may have a putative or indirect oncogenic role. The association between Plasmodium falciparum and the Epstein-Barr virus is not entirely clear in the oncogenesis of Burkitt’s lymphoma in Africa. Toxoplasma gondii has been detected in some patients with intraocular lymphoma. Strongyloides stercoralis may be associated with gastric lymphoma and adenocarcinoma of the colon.

A new form of neoplasm associated with a parasite, namely Hymenolepis nana, has been reported in 2015 in an immunosuppressed patient: the transformation of the parasite into a tumour that subsequently spreads [80]. It is the first and only known case of a person suffering from cancer originating from a parasite (the cancer cells were not human but parasitic).

But, how can a parasite cause cancer? Adult helminths are constantly moving, burrowing and laying eggs that secrete toxic proteins; the host tries to get rid of them, which induces chronic inflammation that can eventually degenerate into metaplasia.
However, some parasites may be protective in tumour genesis, inducing apoptosis phenomena and inhibiting signals of inflammatory regulation and angiogenesis. Certain proteins excreted by *Fasciola hepatica*, *Echinococcus granulosus*, *Trichinella spiralis*, *Toxoplasma gondii*, and *Trypanosoma cruzi* are capable of inhibiting tumour cell growth *in vitro*, and may be used as anti-cancer therapy in the future.

**Conclusion:**

A bacterium: *Helicobacter pylori*; and three parasites: *Schistosoma haematobium*, *Clonorchis sinensis* and *Opisthorchis viverrini*, are classified in Group I of carcinogens by the International Agency for Research on Cancer. *Helicobacter pylori* is clearly associated with gastric cancer and mucosa-associated lymphoid tissue lymphoma (MALT).

Some parasites such as *Schistosoma haematobium*, *Clonorchis sinensis* and *Opisthorchis viverrini* are classified by the WHO as group 1 oncogenic agents (those of proven aetiology). Other parasites have a less defined role such as *Plasmodium falciparum*.

A new and exceptional human tumour disease has been described after the transformation of *Hymenolepis nana* into a tumour.

**QUESTION: COULD THE CONCEPTS OF ONCOGENE AND ONCOPROTEINS BE SUMMARISED? AND THE CONCEPT OF GENOMIC INSTABILITY?**

Studies on chicken sarcoma in the first decade of the 20th century suggested that the origin of the tumours was viral. For example, pioneering studies by Peyton Rous showed how filtering pulverized chicken sarcoma through a sieve that did not allow cells to pass, only viruses, then injecting the extract, transmitted the sarcoma to the recipient chicken. Throughout almost two thirds of the century this viral theory of the origin of cancer was the main driving force behind cancer research and, in fact, a good number of viruses capable of triggering tumour processes were found. As oncologist Ludwik Gross said in 1970, “The viral origin of most malignant tumours has already been documented beyond any reasonable doubt. It would be quite difficult to assume a fundamentally different aetiology for human tumours” [81]. However, some observations did not fit with this model. For example, trying to explain the high incidence of tumours among miners, the Japanese pathologist Katsaburo Yamagiwa [82] observed that repeated exposure of the skin to charcoal induced sarcomas in rabbits. This suggested that the carbon must have some chemical agent that induced a change in the animal’s skin cells, and made them grow out of control, but how?

The answer finally came in the 1980s with the discovery of oncogenes. The concept of oncogenes derives from the presence of genes that are mutated in tumour cells, and which are responsible for transforming them into malignant ones. These genes perform normal functions in the cell such as stimulating growth, increasing survival, or defining identity. However, the presence of certain mutations in them means that, once translated into protein, they have an altered capacity capable of transforming a normal cell into a malignant one. The genes that, once mutated, give rise to oncogenes are called proto-oncogenes, and the proteins that are generated from these are called oncoproteins. The discovery of the first oncogene was published in 1982 by 3 independent laboratories, those directed by the doctors Robert Weinberg, Michael Wigler and the Spaniard Mariano Barbacid, [83] who managed to sequence transforming oncogenes isolated from bladder cancer cells [84].

The first oncogene discovered was RAS, which we know today is a family of genes whose function is to regulate cell growth in response to growth factors. Many others were later discovered, notable cases such as MYC or cyclin E (CCNE) which, as in the case of SAR, control cell growth. We now know that many of the oncogenes are related to genes that control cell growth in such a way that oncogenic mutations promote the uncontrolled proliferation characteristic of tumours. While for several decades it was thought that most oncogenes were related to cell proliferation or their increased survival, tumour genome sequencing studies revealed, not without surprise, that in fact the class of genes most represented among oncogenes are those related to chromatin and epigenetics. Although in many cases we still do not fully understand how these alterations in epigenetic-related genes can trigger tumour processes, the most accepted idea is that these mutations change the identity of cells and generate undifferentiated states. This undifferentiated cell, which has lost its identity, is the one that subsequently -and thanks also to other mutations that promote its growth- gives rise to tumours. As expected, oncogenic mutations are often targets of tumour treatments, and the development of drugs that target all types of oncogenes, including those epigenetic-related genes, is currently a very active field within drug development and new anti-tumour therapies.

In addition to the presence of oncogenes, another characteristic present in practically all tumour cells is genomic instability. First described by Teodor Boveri at the beginning of the 20th century [85], genomic instability is defined as the presence of genomes with multiple chromosomal aberrations in tumour cells such as: the gain or loss of entire chromosomes, chromosomal translocations, or the loss or duplication of regions of one or more chromosomes. Genomic instability is caused by failures in DNA replication [86] or in the distribution of chromosomes during mitosis, processes that are often involved in tumour cells. Furthermore, it has recently been discovered that oncogenes are also a source of genomic instability, as they promote abnormal replication that leads to DNA damage. Fortunately, all cells have mechanisms that suppress genomic instability and normal cells keep their genome stable. The fact that genomic instability is a distinctive feature of tumour cells has opened the door to anti-tumour strategies based on selectively killing cells with genomic instability. One
known example is the poly(ADP)-ribose polymerase (PARP) inhibitors that have been shown to be effective in killing tumour cells with deficiencies in DNA repair in breast, ovarian and prostate tumours [87]. Another alternative that is now being clinically studied, and in which the laboratory of one of us played an important role, is that of ATR kinase inhibitors, which are mainly responsible for the type of genomic instability initiated by oncogenes [88].

**Conclusion**

Both the presence of oncogenes and genomic instability are features of tumour cells that distinguish them from normal cells, and thus they are being exploited for the development of therapies capable of selectively killing cancer cells.

**QUESTION: WHAT PROPORTION OF CANCER DEATHS CAUSED BY VIRUSES COULD BE PREVENTED BY EXISTING VACCINES?**

Cancer mortality series offer an uneven picture between regions of the world, due to the multiplicity of factors involved in their genesis. A global approach estimates that about 1.5 million cancer deaths could be avoided annually if the underlying infections that led to the development of the tumour were prevented or treated [17,22,89].

A pioneering contribution in this area is that of Catherine de Martel et al [89] who published their analysis of the incidence of 27 types of cancer in 184 countries in Lancet Oncology in 2012, describing the fact that two thirds of these deaths occur in less developed countries. These authors carried out an analysis to estimate the proportion of cancers globally that could be attributed to infections and have calculated, by categorising the planet into eight regions, the population that could be affected by new tumours and which could have been prevented by a specific intervention against exposure. They point out that of the 12.7 million new cancers worldwide in 2008, about 2 million were attributable to infections, and of these, 1.6 million (80%) were in developing countries. Of the 7.5 million deaths that occurred in 2008 attributable to an oncological process, an estimated 1.5 million were due to potentially preventable and treatable infections. An upwardly weighted estimate of 172 deaths worldwide could be prevented every hour from tumours caused by infectious agents potentially preventable by immunisation.

There are significant contrasts between different areas of the planet in how these infections are responsible for oncogenesis. Thus, in Australia and New Zealand the infectious origin of a cancer is identified in 3.3% of cases compared to 32.7% in Sub-Saharan Africa. In women, cervical cancer accounted for approximately half of the cancer burden related to infection; in men, liver and gastric cancers accounted for more than 80%. About 30% of cases attributable to infections occur in people under 50. In a smaller scope, a Spanish group has been able to describe, in a series of 190,203 cervical smears collected between 2012 and 2014, in women screened for Human Papillomavirus (HPV) in Castilla y León, that its prevalence in foreign women was 23.51%, significantly higher than in Spanish women (9.6%; p <0.001) and that the presence of morphological changes in the foreign women was also higher [90].

The application of preventive public health methods which already exist, such as vaccines (or anti-infective treatments in another conceptual framework), could have a significant effect on the global burden of cancer in the future. In the specific field of viral agents, it is worth noting that we already have vaccines with proven effectiveness against HPV and Hepatitis B Virus (HBV) [91,92]. “Increasing vaccine coverage should be a priority for health systems in high-burden countries,” as G Danaei [93] states in an editorial accompanying the aforementioned study [89].

A significant proportion of cancer cases attributable to infections can be prevented through improvements in vaccination and screening programmes. Expansion of vaccination (for HBV and HPV), screening and treatment programmes (e.g., for Hepatitis C Virus -HCV-) could greatly reduce mortality due to them.

An additional aspect of vaccination programmes is to describe their acceptance by the population, to carry out economic assessment studies that verify their suitability and availability from the perspective of the social funder, and to establish their accessibility.

In developed countries, in the context of patients co-infected with the Human Immunodeficiency Virus (HIV), very suggestive lines of work have been established by De Martel and Plummer’s own group [94]. These authors indicate that, although HPV and hepatotropic viruses such as HBV and HCV do not represent a differentially weighted in this category of patients in the USA, they do constitute the Human Herpesvirus-8 (HHV-8) at the expense of its aetiological involvement in Kaposi’s Sarcoma and Epstein–Barr Virus (EBV) due to its causal relationship with lymphoid pathology. There is no doubt that the research applied to vaccines that can be extended to the Herpesviridae family will represent a decisive impulse towards minimising their impact on patients co-infected with HIV.

**Conclusion**

Cancer mortality series offer an uneven picture between regions of the world, due to the multiplicity of factors involved in their genesis. From a global perspective, at a low estimate, around one and a half million deaths from cancer (of which two thirds occur in underdeveloped countries) could be avoided each year if the underlying infections that led to tumour development were prevented or treated.
QUESTION: WHAT IMPACT IN NUMBERS HAS HPV VACCINATION HAD ON REDUCING FEMALE GENITAL CANCER?

Vaccines against human papillomavirus (HPV) induce an elevated production of antibodies capable of neutralising the virus and preventing infection of the anogenital tract and, therefore, the possible development of precancerous lesions and cancer. Given the natural history of infection, which requires years or decades from initiation to development of cancer, protection against persistent infection and pre-malignant lesions is considered a surrogate marker of protection against cancer.

More than 10 years after the commercialisation of HPV vaccines, many studies provide data on their effectiveness (effectiveness in the general population and in real life) [95-99]. The fact that there is a reduction in the prevalence of vaccine genotypes or a reduction in the rate of pre-malignant lesions in the vaccinated population is certainly indicative of protection against cancer.

A recent systematic review and meta-analysis assesses the population-level impact of vaccine effectiveness in the real-world setting. Since 2007, a total of 99 countries and territories have initiated vaccination programmes. In 2016, WHO expanded the recommendation in favour of promoting the vaccination of multiple cohorts of girls over single-cohort vaccination. The analysis of 1,702 potential articles allowed the final inclusion of 65 articles carried out in 14 high-income countries (23 articles to assess HPV infection, 29 for anogenital warts and 13 for CIN2+ lesions). This meta-analysis of over 60 million individuals and 8 years of post-vaccination follow-up provides compelling evidence of the substantial impact of vaccination programmes on reducing HPV infection, and the diagnosis of anogenital warts and CIN2+ lesions [100].

Specifically, the prevalence of HPV 16 and 18 decreased significantly by 83% (RR 0-17, 95% CI 0-11–0-25) among girls aged 13-19, and 66% (RR 0-34, 95% CI 0-23–0-49) among women aged 20-24. Also, the prevalence of HPV 31, 33 and 45 decreased significantly by 54% (RR 0-46, 95% CI 0-33–0-66) among girls aged 13-19 years. Five to nine years after vaccination, diagnosis of CIN2+ lesions decreased significantly by 51% (RR 0-49, 95% CI 0-42–0-58) among girls aged 15-19 years and by 31% (RR 0-69, 95% CI 0-57–0-84) among women aged 20-24 years.

This meta-analysis demonstrates a stronger and faster direct impact, as well as a herd effect, in countries with high coverage and which included multiple vaccine cohorts. Therefore, these real-world data confirm that both the cause (high-risk HPV infection) and the immediate pre-cancer condition (CIN2+) are decreasing significantly among vaccinated populations and therefore the vaccine is effective in preventing cervical and other HPV-related cancers [100].

Another recent study in the U.S. assessed the type of HPV present in cytology samples from women aged 20-29 years assessed at Kaiser Permanente Northwest in 2007 and in two periods of the vaccine era: 2012-2013 and 2015-2016. The prevalence of vaccine HPV types went from 13.1% in 2007 to 2.9% in 2015-2016. After 9-10 years from the introduction of the vaccine, the prevalence of HPV vaccine types decreased by 78% among 20-24 years old and by 38% among 25-29 years old. This decrease was observed in both vaccinated and unvaccinated women, which is evidence of both direct and herd protection [101].

In the same vein, data from Scotland from nine cohorts of women aged 20 years have been published to evaluate the effectiveness of the bivalent vaccine in preventing precancerous lesions. The analysis confirms an 89% reduction in prevalent CIN3+; an 88% reduction in CIN2+; and a 79% reduction in CIN1 in vaccinated women born in 1995 and 1996 compared to unvaccinated women born in 1988. Unvaccinated women also showed a lower prevalence of CIN3+ lesions, supporting a herd protection effect. Because of this effect, a reduction in HPV 16/18 infections would not only impact women themselves but also the overall network of sexual relationships between men and women. The fact that the vaccination programme reduced all types of CIN3+ lesions, and not only those related to types 16 and 18 included in the bivalent vaccine, indicates strong cross-protection against other common oncogenic types such as HPV 31/33/45, in accordance with previous clinical trials [102].

Finally, data from registries showing a decrease in cervical cancer are still at a very early stage. Specifically, in Finland, monitoring of cancer registries in vaccinated women in clinical trials and their comparison with unvaccinated cohorts (period 2002/2007-2015) confirms that not a single case was diagnosed in the vaccinated cohort versus 10 cases of HPV-related cancer in the unvaccinated cohort (8 cases of cervical cancer, one vulvar cancer and one oropharyngeal cancer). However, in this analysis the vaccinated and unvaccinated cohorts were not comparable, which limits the interpretation of the data [97].

Another study of the annual incidence of invasive cervical cancer in the United States compares the four years prior to the introduction of the vaccine (2003–2006) with the vaccine era (2011–2014). There was a significant 29% reduction in average annual incidence rates during the study period (6.0 versus 4.1 per 1,000,000 people, rate ratio = 0.71, 95% CI: 0.64–0.78) among women aged 15 to 24 years, and 13.0% lower among women 25 to 34 years. During these years, vaccine coverage increased to 55.3% in the younger cohorts. Although incidence rates had already declined prior to the introduction of the vaccine, these strongly declining data suggest that it may be partly attributable to the introduction of the HPV vaccine [98].

In relation to the impact on non-genital tumours, in the case of both anal and oropharyngeal cancer, more than 80% of HPV-associated cases are caused by HPV16, and more than 90% by at least one of the types 16/18/31/33/45/52/58/6/11. The currently available vaccines would have a very high reduction impact on these two types of cancer.
QUESTION: IN AN IDEAL WORLD IN WHICH HEPATITIS C IS ERADICATED, COULD WE ESTIMATE THE DECREASE IN MORTALITY ATTRIBUTABLE TO LIVER TUMOURS? WHAT WOULD BE THE IMPACT ON TUMOUR MORTALITY IN A WORLD WHERE HEPATITIS B VACCINATION WAS UNIVERSAL?

The question posed is an interesting "epidemiology-fiction" exercise of great future importance and of enormous potential value in terms of public health.

The theoretical substrate of the question is essentially based on the fact that there is no reservoir other than human beings for viruses B and C, which encourages the logical hope that their elimination (by the implantation of universal vaccination or by means of antiviral treatment applied to all possible patients) would be capable of clearly attenuating, even until its disappearance, liver diseases associated with chronic viral infection, including primary liver cancer. This attractive concept would be particularly true in an ideal scenario of no new patients with hepatitis B due to vaccination, although the risk of cancer would persist in those patients already infected, in whom screening would still be essential.

However, there are several elements that mitigate the undoubted potential impact on the incidence of hepatocellular carcinoma.

i) In patients with advanced hepatitis C, viral suppression does not negate the risk of liver disease progression

Different studies have clearly shown that the sustained viral response in patients treated with direct antivirals has a marked effect on the progression of liver disease in patients in the early stages of infection and that it reduces the risk of complications in patients with more advanced disease. However, in patients with significant portal hypertension (over 10 mmHg), i.e. those with more advanced disease and at risk of developing complications of cirrhosis, antiviral treatment fails to lower portal pressure below the risk threshold in a significant proportion of patients, the higher it is, the greater the degree of deterioration. Therefore, the risk of developing hepatocellular carcinoma (clearly influenced by the presence of cirrhosis with severe portal hypertension) is not eliminated despite the elimination of the virus in patients with advanced disease.

Therefore, as long as there are patients treated in whom there is basically advanced fibrosis, the risk of developing hepatocellular carcinoma is not zero. There are no data, however, to quantify the safe threshold of fibrosis below which screening would not be necessary.

ii) Is antiviral treatment of hepatitis C capable of triggering hepatocellular carcinoma?

This is a very controversial issue arising from the observation in a Spanish multicentre study of an increased frequency of hepatocellular carcinoma recurrence and in its severity in patients with hepatocellular carcinoma treated with radical intent and who immediately afterwards received antiviral treatment for hepatitis C. The researchers speculated about the possible impact of the abrupt suppression of the chronic immune response associated with the cure of the C virus infection with a loss of "anti-tumour surveillance" mediated by immune system cells. Information for and against this hypothesis has emerged since this study. In general, it can be considered that anti-viral treatment does not per se increase the risk of developing hepatocellular carcinoma in patients with cirrhosis with no known tumour. However, delaying the attempt to eradicate the virus in patients recently treated for hepatocellular carcinoma is recommended [103-106].

iii) There are other causes of cirrhosis and hepatocellular carcinoma

As discussed in previous questions, the presence of cirrhosis per se, regardless of its cause, is a pro-oncogenic factor. This is extremely important in patients who are carriers of fatty metabolic liver disease, which is a real epidemic in the Western world. Strikingly, this disease, whose pathogenic complexity and systemic inflammatory substrate are widely known, has its own pro-oncogenic mechanisms. From the epidemiological point of view, it is important to note that patients with hepatocellular carcinoma associated with fatty metabolic liver disease are more frequently observed than in other non-cirrhotic liver aetiologies, are diagnosed in more advanced stages, and are therefore less likely to receive treatment with curative intent. The future impact in terms of increased incidence of hepatocellular carcinoma associated with this disease is not known with certainty, but if the forecasts of increased prevalence worldwide are true, a dramatic increase can be expected in the medium term. In fact, recent studies [107] clearly reinforce this trend.

Ultimately, the impact of a universal elimination of hepatotropic viruses on the development of liver cancer would be of great importance, although it would not completely eliminate this dreaded disease.
QUESTION: IS THERE A PROSPECT FOR AN EPSTEIN-BARR VIRUS VACCINATION?

The relationship between viruses and cancer is clearly underestimated and little known, both at the professional level and at the population level, despite such strong data as the fact that the second carcinogen demonstrated worldwide at present is a virus (HPV) [108-115].

Successful and widespread vaccines have been achieved against HBV and HPV, but a widespread vaccine against EBV remains elusive. EBV, as already mentioned, is an oncogenic virus as proven by its association with Burkitt’s lymphoma and its ability to transform B cells in vitro [116], through the expression of Epstein-Barr nuclear antigens (EBNA) and latent membrane protein (LMP). EBV-infected cells are also capable of withholding the cellular lysis of acute infection and this also contributes to neoplastic development.

The lifelong persistence of EBV in humans is a complex multi-step process that begins with infection of the oropharynx, and culminates in the maintenance of EBV in circulating memory B cells. EBV probably uses the normal B-cell differentiation pathway to achieve persistence because it is able to implement various latency and lytic transcription programmes, assuming different antigenic states within infected individuals. However, despite the wide variety of antigens that predominate throughout the EBV life cycle, EBV candidate vaccines have traditionally focused on only a limited number of EBV antigens [117].

Both therapeutic and prophylactic vaccines have been explored.

Therapeutic vaccines aim to enhance and maintain the immune response in patients with EBV-associated disorders and have been studied mainly in nasopharyngeal carcinoma using dendritic cells with specific CD8 epitopes against LMP2 and using the MVA (Modified Vaccinia Ankara) platform including LMP2 and EBNA 1 antigens [118-120]. The data suggest that tumour growth can be controlled by the immune system [121,122]. These studies have led to trials with different therapeutic vaccines that have been well tolerated and the results of phase I and II trials are encouraging but suggest that they are not in themselves the solution for nasopharyngeal carcinoma (the main carcinoma on which these types of vaccines have been tested) as they have an effect on only a low proportion of patients and on a temporary basis [123-126].

The largest number of studies have focused on prophylactic vaccines.

In 1995, a first test was performed with the recombinant attenuated virus itself and did not induce antibodies in seropositive adults. The experts, faced with these results, decided that vaccination was not an appropriate approach to control this virus.

Over the past two decades, trials have been based on glycoprotein 350, which led to a phase II trial with a sample size of 181 people. With a three-dose protocol, antibodies remained for 18 months and the incidence of infectious mononucleosis was reduced by 78% but the vaccine did not reduce the risk of acquiring EBV infection [117].

Work has also been done with a vaccine based on glycoprotein 350 for its pre-kidney transplant application in children that was not very immunogenic [127,128].

Similar data were obtained to the Phase II trial with a vaccine using the toxoid conjugated EBNA3 nuclear antigen designed to elicit specific T-cell responses to EBV, but it also failed to prevent EBV infection. It should be noted that this vaccine is very simple to perform due to the simplicity of the antigen [129].

Antigen-antibody conjugates, or antigen-armed antibodies (AgAbs), which specifically deliver antigenic peptides to antigenic cells [130], have also been used as vaccine candidates without conclusive results.

EBV antigens have also been introduced in virus-like particles [107, 108] and in self-assembling nanoparticles. These platforms allow monomeric antigens to be similar to EBV, but as they lack DNA they are not infectious. Chimeric viruses are also achieved by the fusion of EBV antigens with the structural proteins of Newcastle disease virus. In these structures, for example, gp350 induces a greater response to vaccination [116, 131-137].

All these studies leave us a long way from achieving an effective vaccine against EBV and its primary diseases and, more importantly, against its ability to induce tumour processes.

Conclusion

Despite promising results, neither therapeutically oriented vaccines for nasopharyngeal carcinoma nor preventive vaccines to avoid primary infection by EBV have yet demonstrated sufficient levels of protection to be considered close to market entry.

QUESTION: WHAT ARE ONCOLYTIC VIRUSES? WHAT ARE THE PROSPECTS FOR TREATMENT WITH THEM?

An oncolytic virus is a virus that preferentially infects and kills cancer cells. As the infected cancer cells are destroyed, the oncolysis releases new viral particles that help destroy the remaining tumour.

The potential of viruses as anti-cancer agents was first discovered in the early 20th century, although coordinated research efforts did not begin until the 1960s. Several viruses, including adenovirus, reovirus, measles virus, HVS, Newcastle disease virus, and others, have been clinically tested as oncolytic agents.

Most current oncolytic viruses are designed to be selective against tumour cells, although there are naturally occurring examples, such as Reovirus and Senecavirus, that have been used in clinical trials.
The acquisition of virus selectivity by the tumour cell can be obtained by modifying the proteins of the viral capsid, giving it a high selectivity by the tumour cells. It can also be achieved by modifying the virus genome so that replication only takes place inside the tumour cell. This is achieved either by modifying the transcription in the target cell, or by attenuation, which involves introducing deletions in the viral genome to favour its entry into the tumour cells and eliminate its ability to penetrate healthy cells. It is also possible to introduce genes into oncolytic viruses that encode the synthesis of endostatin and angiostatin, natural proteins that inhibit angiogenesis, with the consequent necrosis of the tumour.

Oncolytic viruses not only cause direct destruction of tumour cells, but also stimulate the host's anti-tumour immune system responses. Most are genetically modified to increase tumour tropism and reduce virulence for non-neoplastic host cells [138]. Thus, they can stimulate a pro-inflammatory environment by improving antigen release/ recognition and subsequent immune activation to counteract immune evasion by malignant cells. In fact, oncological tics viruses also aim to take advantage of the tumour's tolerance mechanisms, which can facilitate viral infection and the death of cells that are not protected by the immune system. This allows for a theoretical domino effect involving chained viral transfer between the neoplastic cells and increased immune activation.

There are currently numerous viral species in different stages of research for immuno-oncological use. Possibly the best studied so far are herpes viruses, of which some strains have been found to have a tropism for natural tumour cells while others have been designed to improve selectivity [139-143]. Initial scans using herpes have shown promising results in murine glioblastoma. Additional efficacy has been seen in prostate cancer, using a recombinant smallpox-vaccine virus capable of regulating prostate-specific antigen and the expression of three co-stimulating factors involved in the presence of the antigen and activation of T cells [144, 145]. In addition, several strains of recombinant vaccine virus have shown promise as antineoplastic agents. One strain has anti-tumour angiogenesis, another has shown efficacy against hepatocellular carcinoma in animal models and the third improves the recognition of tumour cells [146-149]. Other viruses have been or are being explored as possible vehicles of immunomodulation in cancer, such as Newcastle disease virus, coxsackievirus, reovirus, and even the measles virus [150-153].

Among those approved for treatment by the FDA is Talimogene laherparepvec (OncoVEX GM-CSF), also known as T-vec, from Amgen which successfully completed phase III trials for advanced melanoma in March 2013. In October 2015, the U.S. Food and Drug Administration approved T-vec, under the brand name Imlygic®, for the treatment of melanoma in patients with inoperable tumours, becoming the first approved oncolytic agent in the Western world. It is based on the Herpes simplex virus (HSV-1). It has also been tested in a Phase I trial for pancreatic cancer and in a Phase III trial in head and neck cancer along with chemotherapy and radiotherapy with cisplatin.

Oncolytic viruses are generally associated therapeutically with other therapeutic strategies such as association with checkpoint inhibitors, Ipilimumab, Pembrolizumab, Atezolizumab, anti-tumour lymphocytes (CAR-T) or associations of oncolytic viruses with tumour epitopes: vaccines.

Clinical trials with oncolytic viruses in entities as different as colorectal cancer, glioblastoma, tumours with widespread metastases, pancreatic cancer, bladder cancer, ovarian cancer or primary tumours of the peritoneum are underway and represent a hope for the therapeutic future of these malignancies.

**Conclusion**

Oncolytic viruses are natural viruses, or artificially engineered in the laboratory, with selective tropism for tumour cells. They belong to different viral groups, acting selectively with different action mechanisms. Some have been approved for clinical use in humans and are already being used in some tumours, usually in association with chemotherapy or monoclonal antibodies.

**QUESTION: WHAT IS THE COST TO HEALTH CARE SYSTEMS OF TUMOURS DUE TO VIRUSES THAT COULD HAVE BEEN AVOIDED? WHAT ROLE SHOULD HEALTH AUTHORITIES PLAY IN APPROACHING THIS ISSUE?**

In addition to the human cost of cancer, the financial cost is considerable. Direct costs include treatment expenses as well as the cost of care and rehabilitation. Indirect costs include the loss of economic output due to work lost and premature death. There are also costs that are difficult to quantify such as non-medical costs. The exact global cost of cancer is unknown but is expected to increase due to the increase in the number of new cancer cases, as well as the rising cost of cancer therapies [17,154,155].

The health authorities must act on this issue, fundamentally modifying the prevention of risk factors and implementing preventive strategies.

The risk factors for the most immediate intervention are the decrease in genital HPV infections where, among other means, it is possible to intervene with vaccines and vaccination and the treatment of hepatitis virus infections [156,157].

**Conclusion**

The expenditure associated with cancers caused by infectious agents and particularly by viruses is difficult to quantify, but is high and clearly increasing.

The health authorities can intervene by means of prevention and vaccination campaigns, preferably with vaccination against HPV and Hepatitis B and through the treatment of Hepatitis C.
QUESTION: IS THE PRESS AWARE OF THIS ISSUE? WHAT ROLE SHOULD IT PLAY?

Over the last ten years, the press and media have reduced the resources, spaces and personnel dedicated to generating, producing and writing health and medical content as a result of the economic crisis that affected them both through the closure of companies and newspaper titles, as well as through financial adjustments and staff cuts.

Training and specialisation levels have also been reduced in the health, sanitation and science sections for the same reasons.

Despite the variability of the economic crisis, the media have not recovered lost ground and continue to be immersed in situations of weakness and economic-financial fragility.

The impact of new technologies -no longer so new- and the possibilities they have offered for opening up the publication of content to society, as well as the appearance of new communication channels, have posed a risk for the media and threat of loss or dissolution of their preponderant role as the central and principal reference of information for society, having to compete with platforms such as the internet -in its broadest facet of distribution of all kinds of products-, social networks and so-called citizen journalism.

Despite these circumstances, the press and media have placed cancer and everything related to this disease at the forefront of relevance and interest with respect to health and healthcare issues and content, giving it a media prominence above other diseases, with a broad and compact informative approach that considers angles and approaches ranging from health policy, the variety of tumours, the role of patients, research and scientific advances, the work of medical-scientific societies and the oncology sector, the activity of health or pharmaceutical centres and companies, the commemoration of World Days, etc.

“Journalism loves cancer” would be the quick and easy headline.

Having thus explained and exposed the prominence of cancer in the press and media with a broad, diversified and comprehensive scope, it must also be said that viruses, infections, their prognosis, their effects and their medical and social impact as a public health issue are also a subject of strong media coverage, such as flu, AIDS and hepatitis.

However - and here we have the paradox of two related worlds that are disconnected both in the media and in society -, despite the many studies and evidence that link and concatenate the viruses with cancer, the information, dissemination, knowledge and content provided by the media about cancer on the one hand and viral infections on the other, information about the association between viruses and cancer in the media rarely connects these two aspects, unlike others such as cancer-smoking or cancer-ageing.

The role of vaccines, at a time when the anti-vaccine movements are rearming and advancing, first on social networks and then in society, is also decisive in this issue, and while the big question in the popular imagination continues to be whether a cancer vaccine will ever be discovered, medicine has some that serve to curb certain tumours.

Thus, the press and the media neither transmit nor have great knowledge about the binomial subject of study of this Symposium; they do not dedicate special time, space, resources or contents (informative or educational) to this specific angle of the many news stories they publish about cancer.

We suggest, therefore, some measures from the media to report on cancer and viruses

- Include in the news coverage on cancer - which is widespread and frequent - information related to its relationship with viruses, either as emergent or underlying information. It is an aspect that matters to society and so the media should not ignore it.
- Investigate and activate content on the virus / cancer binomial from the editorial planning boards, with reliable and specialised sources.
- To broaden the sense of disclosure on medicine, diseases, prevention and research, beyond the information resulting from announcements, reports or campaigns; and to emphasise this treatment of the contents.
- Combat health and cancer myths and ‘fake news’ with rigorous, clear and accurate verification practices to stop confusion, poor news and information gaps.
- The press and media are undoubtedly committed to the fight against cancer and include news coverage in their agendas and work strategies, but the relationship between oncological diseases and viruses must have greater incidence, reach, development and presence.

Conclusion

Despite frequent and relevant information in the media about cancer on the one hand and viral infections on the other, information about the association between viruses and cancer in the media rarely connects these two aspects, unlike others such as cancer-smoking or cancer-ageing.

QUESTION: HOW IS THE VIRUS CANCER RELATIONSHIP EXPERIENCED, FROM THE PERSPECTIVE OF THE SPANISH PATIENTS WITH CANCER GROUP, GEPAC?

The Virus and Cancer association is little known by the population and information from cancer patient associations is also partial and deficient in this respect. Therefore, a great deal of educational and informative work remains to be done in this area. This absence often forces one to turn to “Doctor Google” for information, which can lead to misinformation, since one does not know which sites are reliable, or does not have sufficient knowledge to fully understand and interpret the data that appear.
Probably what resonates most with patients, families and society at large are associations such as HPV, HBV or HIV and cancer. According to the U.S. National Cancer Institute, "Certain infectious germs, including viruses, bacteria and parasites, can cause cancer or increase the risk of tumour formation. Also, some infections weaken the immune system, making the body less able to fight other infections that cause cancer. Some viruses, bacteria and parasites also cause chronic inflammation that can lead to cancer [158-160]."

With no knowledge of these basic facts, it should not be surprising that the social demand and requirement by patients and associations for adequate vaccines, when they exist, is also not of an appropriate intensity. It is often ignored that protection with a condom is not always sufficient for the prevention of all types of sexually transmitted viruses, in addition to limiting sexual partners.

GEPAC wants to highlight the need for sex education for the prevention of STDs, to teach safe sexual practices, care for personal health and our sexual partners. If we are informed, we know all the ways of contagion and we are aware of the importance of medical check-ups if we live in dangerous situations; we can reduce the number of people affected by this type of illness.

It should be noted that most of the viruses that can pose a cancer risk can be transmitted through certain body fluids or blood. And not only that, this type of virus and thus disease, in addition to the health problems they entail, are linked to psychological repercussions that will depend on the severity of your situation, symptoms and treatment received.

In addition, it can lead to anxiety, depression, or feelings of guilt or fear of criticism due to the social stigma. Perhaps guilt is the most difficult emotion to handle in a situation where the contraction of a virus seems to be linked to risky sexual practices.

For all these reasons, we can conclude that only by educating young people and working with the groups most at risk will we be able to have informed adults in the future and thus reduce cases of infection.

**Conclusion**

*Patient associations should contribute to the dissemination of the relationship between viruses and cancer and of existing preventive measures, including vaccines, which have been shown to be effective in preventing avoidable cancers.*

**QUESTION: WHAT ARE THE REFLECTIONS ON THE VIRUS CANCER RELATIONSHIP FROM AN ETHICAL PERSPECTIVE?**

Classical medical training, the one we received in medical schools, has been and is the victim of a bias that vitiates many of our approaches to issues such as the one raised here, that of the relationship between viruses and cancer. It is a bias that did not exist in classical medicine but has become invasive since the early 19th century. The advances in experimental research made it possible from that moment on to describe with great precision the pathognomonic symptoms of some diseases, almost at the same time as their causal agents were found. Think, for example, of the microorganisms that were identified by the dozens in the late 19th century, and which led to the utopia of thinking that each morbid species would be produced by a specific agent: meningitis by meningococcus, pneumonia by pneumococcus, etc. On the other hand, the advances in experimental pharmacology allowed the identification of the first "specific" drugs that acted against the aetiological agents that produced diseases, leaving all other organic structures unharmed. It was the dream of Béchamp's "atoxyl" and Ehrlich's "magic bullets".

At the heart of all this was an unconscious presupposition, that of "specificity". Diseases were species like those in botany and zoology. A catalogue of diseases similar to Linnaeus's Systema Naturae, the catalogue of "morbid species", could therefore be drawn up. Each of them would have a "specific aetiology", which, for example, microbiology would discover, and "specific therapeutics" would also begin to be found, which by attacking the cause of the disease, would restore the state of health. It was therefore a linear process, starting with a specific cause, continuing with the clinical picture of a morbid species and ending with the application of a specific treatment. It was the chain of specificity. The role of medicine was to find the remedies, usually pharmacological, that would act on the specific agent and allow "restitutio ad integrum". A paradigmatic example of this is pulmonary tuberculosis. The clinical picture of this disease, which was very common at the time of the industrial revolution, was described with great precision by the great masters of the anatomical clinical method, including Laënnec. Koch identified the bacillus that caused it, and later, in the mid-20th century, the first specific treatment, streptomycin, was developed. From that moment on, every doctor knew what he had to do: use that antibiotic, and the tuberculosis and TB drugs that came soon after, to kill the Koch bacillus and thus cure the disease.

But the example of tuberculosis is very illustrative of the limitations of this way of understanding the disease. I still remember the debates about tuberculosis from my time as a medical student in the 1960s. Could it be said that TB was caused by Koch's bacillus? Wasn't it also true that pauperism, malnutrition, poor hygiene, etc., played a role? What was the cause of the tuberculosis? Was it just one or was it many? And how to articulate them?

I have brought this example to mind because it is very illustrative of the way in which school medicine has understood illness, and because it expresses very well the so-called "chain of specificity". TB is caused by Koch's bacillus. If we eradicated it, there would be no pulmonary tuberculosis, even with pauperism, malnutrition, etc. These, on the other hand, are not strictly medical issues, but social ones.

Let us now turn to our problem, that of viruses and can-
It is clear that cancer is the result of genetic dysregulation. This alteration is produced by various agents, including viruses. If we were able to eliminate these or correct the alterations, they produce in the genome by means of gene editing techniques, some, if not all, of the important types of cancer would disappear. That is the goal of medicine, and that is the aim of the research being done today. It is, again, the linear scheme, the chain of specificity.

Things are, however, more complex. There is no single causal factor. In fact, this term should disappear from our vocabulary, or at least become a discreet second term. Cause is what necessarily produces an effect, so that even if all the concomitant factors were to disappear, the effect would be produced by it alone. Something impossible to affirm in the empirical world. What we know are not causes, even if they seem to be, but functional correlations. In fact, all experimental research aims to identify them as precisely as possible. They are correlations, not causes. And among these correlations, there is no doubt that the Koch bacillus plays an important role, but so do immunity and environmental conditions, etc. In fact, not every person infected with Koch’s bacillus develops tuberculosis.

Rather than establishing a linear relationship between cause, clinical picture and therapy, it is necessary to make use of more complex schemes, of a circular or at least spiral nature. In the genesis of cancer, endogenous factors intervene, such as the genome we have, but also exogenous factors. The fact is that viruses are some of those exogenous factors that infect cells and alter their genetic information. There are many other exogenous factors that produce similar results. Think of radiation, for example, or tobacco. These are the factors that can be called direct. But there are many other indirect factors. Thus, tobacco is a direct agent in the case of certain lung cancers but acts as an indirect agent in hepatocellular carcinoma. And along with tobacco and alcohol, place sexual habits in the case of infection by the human papillomavirus, food habits in fatty liver and the increased likelihood of liver cancer, etc.

The conclusion of all this already long journey is that life habits are important factors in the genesis of cancer, in some cases indirectly, through viruses, and in others directly. This seems elementary, but it is far from clear, not only in the population, but even among doctors. We believe that what is important is the control of the gene alterations that trigger uncontrolled cell reproduction, and that the other is not a question of medicine. And we lose sight of the fact that rather than being curative, medicine should be preventative. The dizzying increase in its therapeutic capacity is making us forget this primary truth. The primary role of the physician is not cure but prevention; health education is preferable. Otherwise, we arrive at the absurdity that we live and suffer every day. This is the case of the diabetic who eats cakes because this can be fixed by increasing the dose of insulin, or the loss of respect for HIV because we already have effective therapies.

Prevention, or rather, health education, is not just about information. That is one of the great mistakes that have become almost universal. Educating is far more difficult and complicated than informing. Education is always about knowing how to manage one’s own life autonomously and prudently. It’s knowing how to make the right decisions. Or to put it another way, it’s making the choices we have to make. And here’s the ethics. The aim of this discipline is none other than to teach us how to make decisions, the right decisions, so that we do what we have to do. The big question of ethics is always the same: what should I do? And there can only be one answer: among all the possible courses of action in a given situation, only the decision that leads to the optimal course is good or correct, that is, the one which adheres to the values at stake most and injures them least. Only the optimum course is good. Everything else is bad. A good surgeon is the one who performs the best possible operation, and a good judge is the one who passes the best possible sentence, etc. To know how to do this, and do it well, is to be educated. This is not easy, because the human mind is full of dark areas, those that Freud called unconscious, that skew our decisions without us noticing. Good education is the art of managing all these factors properly so that our decisions are the best they can possibly be. This, and no other, is the goal of ethics.

**Conclusion**

Even though cancer does have a genetic basis, the triggers, including viruses, are always external to the genome itself. A very common mentality in medicine leads to linear thinking, reducing the aetiology to the genetic disorder and the therapy to the correction of that disorder. However, things are more complicated. The genome itself is the result of interactions with the environment, meaning that health depends not only on the genome but also on our lifestyle. And that is precisely the objective of ethics: to educate in the prudent and responsible management of the body, health, life and death.

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