Breast Cancer and Multiple Primary Malignant Tumors: Case Report and Review of the Literature

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Abstract. Multiple primary malignant neoplasms are multiple tumors with different pathogenetic origin. They may be synchronous or metachronous. The management of these conditions represents an interesting clinical scenario. A crucial aspect is the decision regarding which tumor to treat initially, and how to schedule further treatments according to individual tumor risk. This process involves a multidisciplinary physician team to ensure favorable outcomes. We describe a case report of a female patient affected by primary synchronous tumors of the breast and pectoral skin, which raised a series of diagnostic, etiological and therapeutic issues persuading us to carry out a critical review of the literature.

Multiple primary malignant neoplasms (MPMNs), or multiple primary cancers, are defined as multiple tumors with different pathogenetic origin. The phenomenon of MPMN was first described by Billroth at the end of the 19th century (1) and since then several cases of double or even triple primary malignant neoplasms have been reported (1-3). Neoplasms can be synchronous or metachronous and may appear in a single organ or in multiple organs concurrently. According to Moertel (2), synchronous neoplasms are defined as those that occur within 6 months from the diagnosis of a previous malignant tumor, and metachronous neoplasms are defined as neoplasms that appear 6 months after the first diagnosed tumor (2). Despite many changes in the definition have been proposed, the diagnosis of MPMNs that is widely accepted is based on the criteria described by Warren and Gates (3). These criteria require that: (i) each tumor must have a clear picture and histological confirmation of malignancy; (ii) each tumor must be topographically distinct and separated by healthy mucosa (at least 2 cm of normal mucosa between two tumors of the same region); (iii) the lesions must be not metastases of each other (3).

In this report, we describe our recent observation and treatment of a female patient affected by primary synchronous tumors of the breast and pectoral skin. This combination, to the best of our knowledge, has never been previously reported in the literature, and it raised a series of diagnostic, etiological and therapeutic issues persuading us to carry out a critical review of the literature.

Case Report

A 58-year-old postmenopausal woman, of Balkan origin which had settled in Rome for the past 5 years, was referred to the Department of Surgical Sciences, Sapienza University of Rome, Italy, for an erythematous, ulcerated lesion with necrotic areas of the chest wall involving the skin of the right breast (Figure 1). The patient gave written informed consent for use of her radiological imaging and photographic documentation. A detailed past medical history was collected, which resulted negative for primary comorbidities, genetic or environmental exposition to risk factors, as well as for family history of cancer.

Interestingly, the patient also presented crusts and ulceration on the right nipple, suffering by intense pain and bleeding from the nipple as well as from the necrotic areas of the chest wall. No weight loss, nor fatigue, fever or any other specific complaints were reported.

Clinical and Diagnostic Investigations

In the first instance, a mere cutaneous lesion was suspected and, therefore, a punch biopsy was performed, whose histological diagnosis documented a squamous cell carcinoma.
of the pectoral skin. Moreover, because of the ulcerations within the right nipple-areola complex, the patient underwent mammography and breast ultrasonographic imaging that documented the presence of multiple nodular opacities with irregular margins, variable diameter from 5 to 10 mm and calcifications in the external upper quadrant (Figure 2a). A highly suspicious axillary lymphadenopathy was also detected (Figure 2b). The imaging showed a multicentric breast disease with an associated suspicious in situ component, indicating the necessity of an ultrasound-guided breast core-needle biopsy (at two different sites). The histological examination documented a grade 3 invasive ductal carcinoma. Considering the coexistence of a cutaneous squamous cell carcinoma, immunohistochemical analyses were performed to verify the primitiveness of the breast lesions. The results were the following: human milk factor was non-probative; CKMNF116 resulted positive; p63, estrogen receptor (ER), progesterone receptor (PgR) and HER-2 neu receptor resulted negative. Therefore, according to the immunophenotype documented and the negativity of p63 expression, the primitiveness of breast tumor was confirmed. Staging investigations were performed and resulted negative for metastatic lesions.

Finally, to exclude a possible genetic predisposition to breast cancer, the patient was evaluated for the presence of BRCA1/2 gene mutations, but none was detected.

Treatment

After discussing the clinical picture of the patient in a multidisciplinary team, the patient underwent neoadjuvant chemotherapy. Due to the presence of two different tumors, a more aggressive approach was chosen compared to a normal chemotherapy protocol used in the case of clinical T4 breast cancer.

Already during the second cycle of chemotherapy, the staging computed tomography showed that the neoplastic lesions of the right breast were no longer appreciable.

Breast re-evaluation by magnetic resonance imaging, after three chemotherapy cycles, showed an important dimensional reduction of the cutaneous and breast lesions with re-epithelialization of the ulcerated areas.

The patient completed the neoadjuvant chemotherapy with three more treatment cycles and then underwent a combined surgery of right mastectomy, ipsilateral axillary lymph nodes dissection, excision of the remaining cutaneous squamous cell carcinoma, and reconstruction with a split-thickness skin graft taken from the contralateral leg with an electric dermatome, to correct for the loss of substance (Figure 3a and 3b). The surgery lasted 105 min and the estimated blood loss was 100 ml. The patient was discharged in the third post-operative day in a good clinical condition.

The histopathological examination of the surgical specimen revealed a complete pathological response of the right breast carcinoma, documenting only a small residue of ductal carcinoma in situ, and a complete regression of the cutaneous squamocellular lesion.

No early neither late post-operative complications were documented, and the graft was well tolerated. Due to the absence of both cutaneous and breast infiltrating tumors, it was decided to keep the patient in follow-up without further oncological treatments.

The patient maintained a good performance status and she is continuing on a program of 6-month clinical and radiological follow-ups as recommended by the multidisciplinary team.

Review of the Literature

MPMNs incidence is estimated to be between 0.73% and 11.7% (4). Patients with a primary tumor have a 10% higher risk of developing a second tumor with respect to the general population (5, 6). Causes of MPMNs are not yet clear, but they may be related to four principal factors: intrinsic factors, extrinsic factors, genetic factors, and therapeutic factors.

Intrinsic factors include susceptibility, immune status, and endocrine and embryonic development. The loss of immune surveillance and immune defenses both congenital and acquired increases the possibility to develop a MPMNs (7).

Extrinsic factors are represented by environmental factors and personal lifestyle, including occupational diseases, long exposure to ultraviolet rays and industrial pollution (7). In addition, smoking and alcohol, and low physical activity are important carcinogenic factors.

Genetic factors may favor the development of MPMNs. Members of families with BRCA gene mutations have an increased risk of early onset breast and ovarian cancer (8).

Therapeutic factors, including carcinogenic radiation treatment and chemotherapy, may significantly increase the
risk of developing MPMNs (9). It is surprising how chemotherapeutic agents might paradoxically have an “oncogenic” role, since on the one hand they increase survival, and on the other they may cause an acquired immunosuppression.

Spratt et al. (9) reported that new therapy protocols for cancer (i.e. alkylating agents, topoisomerase II inhibitor and/or radiotherapy) have caused increased rates of survival, determining a higher chance for patients to develop a second neoplasm and apparently increasing the incidence of cancer. Above all, in the case of metachronous tumors, the main cause of the development of a second tumor is the acquired immunosuppression. Several immunosuppressive and/or cytotoxic agents are potential carcinogens and may either have a direct oncogenic effect or play a role as cofactor in oncogenesis (9).

In our review we considered all the published case reports of MPMNs where breast cancer is present and have evaluated the correlation of the different single tumors with breast cancer.
Table I shows a summary of the previous studies describing MPNMs.

**Breast cancer and skin cancer.** Twelve cases of tumors involving the skin and the breast in the same person have been described (10-21); in particular, only two cases (11, 12) presented the coexistence of squamous cell carcinoma and breast cancer, similarly to our case, which turns out to be the third case report documented since 1980. As discussed by Pastore *et al.* (12), our patient presented a synchronism of a wide squamous cell carcinoma of the right breast synchronous to a right breast cancer.

The most frequent described MPMN is breast cancer and melanoma (13-20).

This association is not surprising, because several genetic and hormonal factors are responsible for the association between breast cancer and melanoma, in particular the germ-cell mutations or somatic mutations in genes such as *BRCA2* (22).

**Breast cancer and gastro-intestinal cancer.** The association between breast cancer and gastro-intestinal cancer may be considered as a sporadic or germinal association (23-46). While for tumors of the high gastro-intestinal tract, such as esophagus, stomach and liver cancer, the association with breast cancer is probably sporadic (8 cases have been described) (14, 23-29), in case of colorectal cancer several studies emphasized the coexistence of common predisposing extrinsic and genetic factors.

Breast cancer and colon cancer represent primary tumors with high incidence, while metastatic breast cancer to the colon is a rare entity with the exception of the lobular histotype breast carcinoma which has resulted to be the most common histological type of breast cancer that metastasizes to the colon (47).

**Breast carcinoma and nervous system tumors.** The association between breast cancer and primitive brain tumor lesions is weak and less frequent (16, 48, 49).

The presence of a breast cancer and a brain lesion always needs further investigation aimed at diagnosing the primitiveness of the single lesions, considering that breast cancer appears to be the second leading cause of brain metastasis (50).

Less frequent but often metachronous lesions are, instead, the associations between breast cancer and peripheric nervous system tumors (51-53).

**Breast cancer and hematologic tumors.** Breast cancer patients may develop, with a higher risk with respect to the normal population, secondary malignancies, including leukemia and non-Hodgkin’s lymphoma (54). The second tumor is usually metachronous (22, 55-64) and appears 2–5 years following the eventual chemotherapy and radiotherapy (54), while synchronous breast cancer and lymphoproliferative diseases are much rarer (18, 37, 55, 58, 60, 63, 65). Several case reports describe the coexistence between breast cancer and lymphoma more frequently than expected, with no evidence that the second tumor was treatment-induced (60-65).

**Breast cancer and sarcoma.** The first case report of MPNMs including breast cancer and sarcoma was reported in 1936 (66). Another five case reports of these MPNMs have been presented in the literature (34, 67-70).

It was Warren who first highlighted how radiotherapy treatment for breast cancer was a risk factor for the development of sarcoma (66).

At the same time, it is important to remember that radiosensitivity is an intrinsic characteristic of the cells of the mammary gland, so that the radiotherapy treatment of a rhabdomyosarcoma of the upper arm can itself be a risk factor for breast cancer development (68).

Approximately 60% of cancer patients receive radiation therapy during the course of the disease (71) and even after 10 years from radiotherapy a second tumor may develop (72). The definition of radio-induced sarcoma is characterized by the following criteria proposed by Cahan (73) and then modified by Arlen (74): (i) radiant treatment at least three years before developing sarcoma, (ii) onset in the context of previously irradiated tissues, (iii) different histology compared to previously irradiated primary tumor.

Since most patients with breast cancer receive radiation therapy, it is essential to early diagnose this form of sarcoma.

**Breast cancer and lung cancer.** If a lung lesion is documented in a patient diagnosed with breast cancer, it is always necessary to exclude a lung metastasis, which represents the second most frequent site of metastasis of breast cancer after the bone (75).

However, it is not uncommon to find coexistence of a lung and breast cancer as primitive tumors in the same patient (11, 20, 28, 32, 34, 40, 48, 61, 64, 76-79). This association can be explained mainly considering that: i) breast cancer is the most frequent malignancy in women worldwide (80); ii) in the last years, the mortality of women with an early diagnosis of breast cancer has been significantly reduced and their longer overall survival increases the risk of developing a second tumor (81); iii) lung cancer is the second most common primary malignancy which may develop in patients who have a history of a previous breast malignancy (82).

**Breast cancer and gynecological tumors.** A strong association between breast cancer and gynecological tumors, especially ovarian cancer, exists, as they are (62, 83-85) influenced by similar risk factors such as early menarchy, low number of pregnancies/nulliparity, obesity as well as high fat diet and fat intake >38% of daily calories.
| Year | Author                  | 1st Tumor                        | 2nd Tumor                        | 3rd Tumor                        | 4th Tumor                        | 5th Tumor | 6th Tumor | Synchronous or Metachronous primary cancers |
|------|------------------------|----------------------------------|----------------------------------|----------------------------------|----------------------------------|-----------|-----------|---------------------------------------------|
| 1980 | Tsuneyoshi M. Deppe G. | Breast carcinoma                 | Sarcoma                          | Colon adenocarcinoma            | Vulva squamous cell carcinoma    | -         | -         | Metachronous                                |
| 1984 | Sugiura K.             | Breast carcinoma                 | Acute monocytic leukemia         |                                  |                                  | -         | -         | Metachronous                                |
| 1985 | Bartal A.H.            | Breast carcinoma                 | Melanoma                         |                                  |                                  | -         | -         | Metachronous                                |
| 1985 | Patel S.               | Breast carcinoma                 | Lung tumor                       | Gastric cancer                  | Nervous system tumors (Brain glioblastoma) | -         | -         | Metachronous                                |
| 1986 | Ogasawara K.           | Breast carcinoma                 |                                  |                                  |                                  | -         | -         | Metachronous                                |
| 1988 | Shinjo K.              | Rhabdomyosarcoma                 | Breast carcinoma                 |                                  |                                  | -         | -         | Metachronous                                |
| 1989 | Kobayashi R.           | Lymphoma                         | Breast carcinoma                 |                                  |                                  | -         | -         | Metachronous                                |
| 1990 | Stierer M.             | Breast carcinoma                 | Nervous system tumors            |                                  |                                  | -         | -         | Metachronous                                |
| 1990 | Newbould MJ.           | Breast carcinoma                 |                                  |                                  |                                  | -         | -         | Metachronous                                |
| 1990 | Kohno A.               | B Cell malignant lymphoma         | Breast carcinoma                 |                                  |                                  | -         | -         | Metachronous                                |
| 1991 | Solan M.J.             | Breast carcinoma                 | Thyroid carcinoma                | Basal cell carcinoma (multiple)  | Breast carcinoma                 | -         | -         | Metachronous                                |
| 1992 | Lian D.L.              | Cervix tumor                     | Larynx squamous cell carcinoma   | Skin squamous cell carcinoma    | Breast carcinoma                 | -         | -         | Metachronous                                |
| 1992 | Marcos Sanchez F.      | Colon carcinoma                  | Renal carcinoma                  |                                  |                                  | -         | -         | Metachronous                                |
| 1994 | Frey B.M.              | Breast carcinoma                 | Lymphoma                         |                                  | Breast adenocarcinoma            | -         | -         | Metachronous                                |
| 1994 | Bumpers H.L.           | Larynx squamous cell carcinoma   | Lung squamous cell carcinoma     |                                  | Colon adenocarcinoma             | -         | -         | Metachronous                                |
| 1996 | Padmore R.F.           | Breast carcinoma                 | Melanoma                         |                                  | Renal cell carcinoma (Metachronous) | -         | -         | Metachronous                                |
| 1997 | Potsch C.              | Breast carcinoma (Synchronous)   | Lung carcinoma (Synchronous)     |                                  |                                  | -         | -         | Metachronous                                |
| 2001 | Shousha S.             | Breast carcinoma                 | Squamous cell carcinoma of the oesophagus | Breast carcinoma                 | Basocellular carcinoma (skin)    | -         | -         | Metachronous                                |
| 2003 | Eros N.                | Lymphoma NH                      |                                  |                                  |                                  | -         | -         | Metachronous                                |
| 2003 | Canda M.S.             | Breast carcinoma                 | Nervous system tumors (Brain glioblastoma) |                                  |                                  | -         | -         | Metachronous                                |
| 2003 | Roy S.D.               | Breast carcinoma                 | Axillary mantle cell lymphoma    |                                  |                                  | -         | -         | Metachronous                                |
| 2005 | Otrock Z.K.            | Lung adenocarcinoma              | Bladder carcinoma                | Prostate adenocarcinoma          | Breast carcinoma                 | -         | -         | Metachronous                                |
| 2006 | Fruscalzo A.           | Breast carcinoma                 | Leukemia lymphatic chronic       |                                  |                                  | -         | -         | Metachronous                                |
| 2006 | Filippakis G.M.        | Teratoma Ovaric                  | Breast carcinoma                 |                                  |                                  | -         | -         | Metachronous                                |
| 2006 | Kościuch J.            | Breast carcinoma                 | Gastric carcinoma                |                                  |                                  | -         | -         | Metachronous                                |
| 2007 | Agrawal R.             | Breast carcinoma                 | Cervix squamous cell carcinoma   |                                  |                                  | -         | -         | Metachronous                                |

Table I. Continued
| Year | Author                  | 1st Tumor                        | 2nd Tumor                                | 3rd Tumor                                 | 4th Tumor                                | 5th Tumor                                 | 6th Tumor                                 | Synchronous or Metachronous |
|------|-------------------------|----------------------------------|------------------------------------------|-------------------------------------------|------------------------------------------|-------------------------------------------|-------------------------------------------|-----------------------------|
| 2008 | Cercato MC.             | Breast carcinoma (Synchronous)   | Squamous cell carcinoma of the larynx    | Kidney carcinoma (metachronous)           | Colon adenocarcinoma (metachronous)      | Synchronous                              | Metachronous                                                             |
| 2008 | Izawa N.                | Osteosarcoma                     | Colon adenocarcinoma                    | Histiocytoma                              | Lung cancer                              | Synchronous                              | Metachronous                                                             |
| 2012 | Ahmed H.                | Breast carcinoma                 | Osteosarcoma                             | Colon adenocarcinoma                      | Histiocytoma                              | Synchronous                              | Metachronous                                                             |
| 2012 | Ko K.                   | Breast carcinoma                 | Lung cancer                              | Osteosarcoma                              | Histiocytoma                              | Synchronous                              | Metachronous                                                             |
| 2012 | Anania G.               | Breast carcinoma                 | Caecum adenocarcinoma                   | Colon adenocarcinoma                      | Histiocytoma                              | Synchronous                              | Metachronous                                                             |
| 2013 | Băltătescu G.I.         | Breast carcinoma                 | Ovarian dysgerminoma                    | Sigma adenocarcinoma                      | Histiocytoma                              | Synchronous                              | Metachronous                                                             |
| 2013 | Sini G.                 | Breast carcinoma (bilateral)     | Nervous system tumors                   | Clear cell renal cell carcinoma          | Lung cancer                              | Synchronous                              | Metachronous                                                             |
| 2013 | Kurul S.                | Breast carcinoma                 | Melanoma                                 | Gallbladder carcinoma                     | Lung cancer                              | Synchronous                              | Metachronous                                                             |
| 2013 | Pogal J.                | Breast carcinoma                 | Colon adenocarcinoma                     | Bladder carcinoma                        | Synchronous                              | Metachronous                                                             |
| 2013 | Higgins L.              | Breast carcinoma                 | Skin squamous cell carcinoma            | Bladder carcinoma                        | Synchronous                              | Metachronous                                                             |
| 2014 | Ko K.                   | Breast carcinoma                 | Gastric cancer                           | Endometrial cancer                        | Metachronous                              | Metachronous                                                             |
| 2014 | Ahmed H.                | Breast carcinoma                 | Uterus carcinoma                         | Rectal adenocarcinoma                     | Synchronous                              | Metachronous                                                             |
| 2014 | Ko K.                   | Breast carcinoma                 | Breast carcinoma                         | Base tongue carcinoma                     | Synchronous                              | Metachronous                                                             |
| 2014 | Ahmed H.                | Breast carcinoma                 | Colon adenocarcinoma                     | GIST                                       | Synchronous                              | Metachronous                                                             |
| 2014 | Ahmed H.                | Breast carcinoma                 | Endometrium cancer                       | Breast carcinoma                          | Metachronous                              | Metachronous                                                             |
| 2015 | Zhao J.                 | Colon carcinoma                  | Nervous system tumors (Malignant peripheral nerve sheath tumor) | Bladder cancer                            | Synchronous                              | Metachronous                                                             |
| 2015 | McPherson J.R.          | Breast carcinoma                 | Lung cancer                              | Colon adenocarcinoma                      | Esophagus cancer                         | Synchronous                              | Metachronous                                                             |
| 2015 | Williamson C.W.         | Breast carcinoma                 | Giant cell tumor; Hepatocellular carcinoma | Lymphoma; Melanoma                        | Synchronous                              | Metachronous                              | Metachronous                                                             |
| 2016 | Lopez J.J.              | Breast carcinoma                 | Trichoolemmal tumor                     | Myeloma                                   | Synchronous                              | Metachronous                              | Metachronous                                                             |
| 2016 | Vennepureddy A.         | Breast cancer                     | Myeloma                                  | Monoclonal B-Cell lymphocytosis           | Synchronous                              | Metachronous                              | Metachronous                                                             |
| 2016 | Cohen P.R.              | Breast cancer                     | Hodgkin lymphoma                         | Lung cancer                               | Kidney cancer                            | Synchronous                              | Metachronous                                                             |
| 2016 | Ari A.                  | Breast carcinoma                 | Colon adenocarcinoma                     | Colon adenocarcinoma                      | Thyroid cancer                           | Synchronous                              | Metachronous                                                             |
| 2017 | Liu J.                  | Breast carcinoma                 | Mantle Cell Lymphoma (MCL)              | Synchronous                              | Metachronous                              | Metachronous                              | Metachronous                                                             |

Table I. Continued
Women with sporadic breast cancer have a high risk of developing sporadic ovarian cancer, and this risk increases if we consider genetic syndromes such as the inherited BRCA1 and BRCA2 gene mutations (8, 86, 87) and the Lynch syndrome type II, which also includes colon, endometrium and high gastrointestinal tract tumors (87). In hereditary tumor syndromes a single germline mutation in a single gene leads to a high risk of cancer (87). The association between primary breast and ovarian tumors is very strong and so in the oncological follow-up of breast cancer, the ovaries are always investigated to rule out an ovarian lesion. In fact, unlike breast cancer, ovarian cancer is less frequent but is characterized by poor prognosis, considering that the overall 5-year survival is 50% and in stages III and IV it drops to 15-20% (88).

Breast cancer and thyroid cancer. Women with a history of thyroid cancer have a greater risk of developing a second tumor when compared to the normal population, especially breast cancer (89, 90). In the case of multiple tumors, the second most frequent site is the breast gland (90) and the risk is higher if thyroid cancer appears in women aged less than 50 years (91).

These two tumors share numerous molecular mechanisms that might activate the carcinogenic process. Among these mechanisms, mutations in the thyroid receptor play an important role in breast hyperplasia (92).

Finally, there are genetic syndromes such as the Cowden syndrome or the Cowden-like syndrome, in which breast and thyroid cancers are associated (93). It is therefore not surprising that there are several case reports of breast and thyroid MPMNs, both synchronous (48, 94) and metachronous (10, 64).

Breast cancer and urological tumors. Bladder and breast cancers share extrinsic risk factors (including personal lifestyle, ultraviolet radiation, smoking and alcohol) and some genetic mutations in signal transduction pathways (95, 96). Damrauer et al. showed that basal-like and luminal-like subtypes of bladder cancer surprisingly reflect the hallmarks of breast biology (97).

If the genetic association of breast cancer with urological tumors can be considered sporadic (11, 12, 16, 31, 44, 61, 76, 86, 95, 96), the same cannot be said for bladder cancer (11, 12, 76, 95).

Discussion

Between 1980 and 2018 only 12 cases of MPMNs involving the breast gland or the skin have been described (10-21). Among these, only two (11, 12) cases report the coexistence of breast and cutaneous squamous cell cancers (Table I). The first case was described by Lian in 1992 (11), who focused on a patient suffering by six metachronous primary cancers (cervix, larynx, skin, left breast, bladder and right lung) (11).

The second study described by Pastore et al. in 2013 (12) reported the clinical case of a male patient who developed...
synchronous carcinoma of the bladder, squamous cell carcinoma of the forehead skin, and breast cancer (12).

We report the case of a patient who developed two histologically distinct malignancies, meeting the inclusion criteria of Warren and Gates to be defined as a MPMN (3). The immunohistochemical analyses, carried out on the two surgical specimens, showed the different primitiveness of the lesions, excluding a metastatic origin for one of them: a wide squamous cell carcinoma of the pectoral skin and an invasive ductal carcinoma of the breast. These can be considered as two synchronous and non-metachronous tumors since the diagnosis was performed within 6 months of each other (3).

When Billroth (1) in 1889 diagnosed for the first time a case of MPMN, the phenomenon of multiple tumors was still rare. Nowadays, the diagnosis of second primary neoplasms is not so infrequent. To our knowledge, the case that we describe is the second, after the one described by Pastore (12) reporting a synchronism between giant squamous cell carcinoma of the right breast skin and ipsilateral breast cancer. Synchronous tumors are much rarer and often attributed to known and unknown genetic predispositions (19).

In our case, we excluded any syndromic frameworks, such as Li Fraumeni (TP53) or Von Hippel Lindau syndrome, or genetic mutations (i.e. BRCA1/2), that might have predisposed to the development of multiple tumors. Moreover, despite the fact that germline TP53 mutations are rare in patients with primary tumors, it was not possible to exclude the possibility that a sporadic mutation in TP53 may have led to a predisposition to multiple tumors (98-100).

Our patient did not have a family history of malignancies, which led us to verify the presence of intrinsic and extrinsic risk factors that could justify the sporadic occurrence of MPMNs. According to the literature, principal risk factors known for MPMNs are tobacco, alcohol or exposure to radiation (101). We excluded the first two factors, but we could not exclude a previous prolonged exposure to sun light. Our patient presented a Fitzpatrick skin phototype II with a lead to the development of squamous cell carcinoma of the face, upper and lower limbs and may often evolve into a squamous cell carcinoma.

Our patient did not present any AK at the physical examination but we could not rule out that an AK may have been present before the squamous cell carcinoma.

In addition to sunlight, other non-sun related risk factors, such as exposition to insecticides, herbicides, and fungicides lead to the development of squamous cell carcinoma of the skin (101). Our patient was a housewife living in an urban context and hardly ever came into contact with non-sun risk factors.

Age is a risk factor for the development of MPMNs (9). Considering that longevity is increasing in the general population, a higher incidence of MPMNs is expected, particularly in patients with a positive history of malignancy (103). However, our patient did not have a personal history of previous tumors.

We cannot therefore attribute it with certainty to a known genetic syndrome or a sporadic occurrence justified by known risk factors. The difficulty in deciding the best therapeutic strategy has led us to review all the MPMNs case reports including breast cancer.

Whatever the cause and/or the established risk factors, some parameters need to be considered for the treatment of MPMNs: the stages of the different neoplasms, tumor behavior, patient’s age, life expectancy, and comorbidities.

While in metachronous tumors the therapeutic course always involves the sequential treatment of each single tumor, in synchronous tumors we found that a personalized and singular treatment is decided after an appropriate evaluation in multidisciplinary team meetings and a consensus on a therapeutic strategy (104), as it is done in the case of an inflammatory breast cancer (105). If in localized cancer it is possible to choose between surgical or chemo-/radio-therapy approach to treat coexisted tumors, in advanced cancer disease the antitumor therapy selection is difficult and often not supported by research evidence (104).

In our case, the multidisciplinary team decided to start with neoadjuvant chemotherapy to down-stage the two tumors, and to reach a better local control of the disease. Definitive histopathologic examination showed that chemotherapy had successfully down staged both diseases and, in addition to surgery, made possible a full recovery.

To our knowledge, our case is the first in the literature reporting simultaneous and complete pathological response of two different tumors using a single chemotherapy line protocol composed by carboplatin and taxane.

Conclusion

Nowadays, multiple primary cancers are not rare entities. Improved cancer care and increased survival have led to an increase in the number of patients developing multiple cancers.

While it is possible to eliminate risk factors such as smoking, alcohol consumption and unhealthy lifestyle, not-modifiable risk factors such as the genetic susceptibility, the state of immune deficiencies and the sometimes-irreversible effects of anticancer therapies cannot be managed.

Although no familial history or carcinogenetic factor could be evidenced, the occurrence of multiple tumors, especially the synchronous ones, is unlikely to be fortuitous.

Patients who develop multiple cancers may not fall into one of the hereditary syndromes, but it is possible that there are unknown and/or rare genetic predispositions that
determine the development of multiple tumors. For this reason, all multiple cancer patients should be followed-up for the development of subsequent cancers.

A complete pre-operative evaluation and interdisciplinary collaboration can always guarantee the best possible treatment for a patient.

Nowadays, the description of case reports is of fundamental importance since they strongly contribute to the progress of medical knowledge, generating new hypotheses and acting as stimulus for the scientific community to verify them. This study portrays a particularly rare clinical picture and provides first evidence of efficacy of a therapeutic protocol that deserves to be investigated in more detail.

Conflicts of Interest

The Authors have no conflicts of interest to declare regarding this study.

Authors’ Contributions

A.D.L. collected the data, conceived and wrote the paper; F.F. collected the data; M.V. collected the data; R.C. reviewed the paper; G.L.T. reviewed the paper; L.B. collected the data; M.M. reviewed the paper; M.I.A. collected the data, conceived and wrote the paper.

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