

Effects of omega-3 and vitamin D supplementation in patients with breast cancer: a systematic review

Subtitle: Supplementation and breast cancer patients

Lailton Oliveira da Silva^{*1} , Victor Da Silva² , Anderson Weiny Barbalho Silva³ , José Juvenal Linhares⁴ 

¹ Nutritionist. Master's degree in Health Sciences, in the Postgraduate Program in Health Sciences (PPGCS-UFC). Sobral, Ceará – Brazil. lailtonutri@hotmail.com

² Doctor. Master's degree in Health Sciences, in the Postgraduate Program in Health Sciences (PPGCS-UFC). Sobral, Ceará – Brazil. dasilva.vitor2022@gmail.com

³ Biologist. Federal University of Ceará (UFC), Faculty of Medicine, Postgraduate Program in Biotechnology, Sobral (CE), Brazil. andersonweiny@sobral.ufc.br

⁴ Doctor. Federal University of Ceará (UFC), Faculty of Medicine, Postgraduate Program in Health Sciences, Sobral (CE), Brazil. juvenallinhares@gmail.com

* Correspondence to: Lailton Oliveira da Silva. Federal University of Ceará, Campus Sobral, Master's Course in Health Sciences - Sobral. Av. Cmte. Maurocélvio Rocha Pontes, 100. (85) 99607-420. lailtonsilvaoliveira@hotmail.com

Abstract: Breast cancer (BC) is the leading cause of death among women with cancer worldwide. Nevertheless, key challenges must be addressed, as new therapies can be employed to support the main treatment and enable a better quality of life, survival and prognosis. In this paper, we described the role of omega-3 and vitamin D supplementation in patients during the main treatment. This is a systematic review of the literature from randomized clinical trials, following the PRISMA and PICOS guidelines for elaborating the guiding question and constructing the results. Our findings in this review reveal and support that omega-3 and vitamin D supplementation can help women with metastatic disease who are not receiving hormone therapy during BC treatment for HR+ (hormone receptor positive). They improve immunity and antioxidant capacity and decrease cardiometabolic effects. This intervention is safe and can be employed as an adjuvant to the other main treatments.

Keywords: Breast neoplasms, Omega-3, Vitamin D, Supplementation

Received: Jan. 31, 2024; Revised: Mar. 21, 2024; Accepted: May 8, 2024; Published: May 11, 2024

Copyright ©2024 Lailton Oliveira da Silva, et al.

DOI: <https://doi.org/10.55976/fnds.22024124666-75>

This is an open-access article distributed under a CC BY license (Creative Commons Attribution 4.0 International License)

<https://creativecommons.org/licenses/by/4.0/>

Introduction

Breast cancer (BC) is the leading cause of death among women worldwide. According to estimates by the International Agency for Research on Cancer (IARC), almost 3.1 million women worldwide diagnosed with BC in 2023, with a higher incidence in parts of Europe, North America and some countries in South America (Brazil) and the African continent [1].

However, mortality behaves oppositely, as countries in South America and Africa present a worsening of this scenario compared to the other continents mentioned. Some factors related to this are: developing countries still have lower rates of early diagnosis, a low Human Development Index (HDI) and associated risk factors such as obesity, physical inactivity, smoking and alcohol consumption [1, 2].

Low-cost alternative therapies can be employed to benefit the clinical treatment of these patients [2]. Therefore, some research is being conducted to understand the benefits of supplementation of nutritional compounds during the treatment of breast neoplasms. The most studied nutrients in recent years include omega-3 and vitamin D [3-5].

Polyunsaturated fatty acids such as omega-6 (n-6), represented by: arachidonic acid and linoleic; and omega-3 (n-3), represented by: alpha-linolenic acid, eicosapentaenoic acid and docosahexaenoic acid, have several beneficial properties for healthy humans as they are linked to cell membrane structures and can act on hormone binding and cellular transport activities [5, 6].

In cancer treatment, it has been found that n-3 modulates the inflammatory and immune response and thus improve prognosis. Additionally, it can play an important role in cell differentiation and growth, and inhibit tumor growth in some molecular subtypes [7-9].

Vitamin D is available in two forms, vitamin D2 (ergocalciferol), which is found primarily in plant compounds such as algae and mushrooms, and vitamin D3 (cholecalciferol), which can be obtained, through the diet by consuming animal sources such as fish (tuna and salmon), meat, eggs, milk and dairy products, or through sun exposure [10].

The metabolically active form of vitamin D in the body is 1-25(OH)₂ or calcitriol. For the effects of calcitriol's biological activity to occur, its vitamin D receptor (VDR) is required, which is expressed in various tissues, including breast tissue. Therefore, calcitriol can benefit patients by inhibiting vascular endothelial growth factor (VEGF) and thus interfering with the angiogenesis process [4,11,12].

Although there are several benefits of vitamin D and n-3 in BC treatment, these benefits remain uncertain in certain populations and molecular subtypes [13,14]. Thus, the aim of this study is to describe and analyze the benefits of supplementation in BC patients during treatment.

Methodology

This systematic review follows the guidelines of the Preferred Reporting Items for Systematic Reviews and Meta-Analysis (PRISMA) [15], was registered in PROSPERO (International Prospective Register of Systematic Reviews) CRD42023442158, and its guiding question, formed by the acronym PICOS, is What is the evidence of the benefits of omega 3 and vitamin D supplementation during the treatment of women with BC?

The first step of the review was to determine the databases and keywords to be used. Therefore, electronic searches were carried out using the descriptors listed in Table 1 in three databases: PubMed, Direct Science and Capes. The keywords defined by the Health Sciences Descriptors (DeCS) in Portuguese were used as a search strategy in English according to the National Library of Medicine (NLM) with the corresponding MeSH to expand the retrieval of the largest possible number of studies.

Table 1. Complete search strategy in electronic databases, 2023

Terms	Descriptors
# 1 Cancer	Breast cancer
# 2 Intervention	Omega-3 fatty acids and/or vitamin D
# 3 Studies	Randomized Clinical Trials
Combination	# 1 AND # 2 OR #3 AND

The second step of this review was to define the eligibility criteria for the studies. These criteria were defined according to the PICOS acronym, which takes into account population, intervention, comparison and study design (Table 2). The studies eligible for this review were: (a) randomized clinical trials; (b) conducted in adults (>18 years old); (c) female; (d) during contingency management (CM) treatment and post-treatment; (e) published in English, Spanish and Portuguese in the last 10 years; (f) studies investigated the benefits of omega-3 and/or vitamin D supplementation in the treatment of patients with BC, with abstract and full text available from April 7, 2023 to June 21, 2023; (g) studies that described the dosage of omega-3 and vitamin D administered to patients.

Information about the number of articles retrieved in the search is described by the reviewers in Figure 1, which is shown in the flowchart, describing the number of articles retrieved at each stage of the search, selection, inclusion and exclusion process.

The search, selection, inclusion and exclusion stages were carried out by two trained reviewers. During the search process, the articles were initially classified and analyzed by title, manually by two blind and independent reviewers, and articles that did not meet the inclusion criteria were excluded.

Next, the abstracts were read and incongruent or duplicate articles were also removed. After screening, the articles were read in full to complete the process of study selection. Discrepancies were resolved by a third reviewer.

After conducting a search for studies for the systematic review and the inclusion criteria, studies that addressed the

effects of omega-3 and/or vitamin D supplementation on the health of patients with BC were included. Additionally, the reference lists of all relevant articles were examined to identify other eligible studies.

The investigation and discussion of the results, data regarding authors, year of publication, journal, number of databases and search period were extracted and assigned with the PRISMA Checklist with 27 items (yes or no). Information was then obtained on the objective, total sample size, age of participants, intervention groups, control group, study location, polymorphism studied, and characteristics of the intervention.

For the risk of bias analysis, the Cochrane tool (RoB 2) was used to evaluate all 7 articles included in this study. It was found that 1 (14%) study had a low risk of bias, 4 (57%) had some concerns about risk of bias and 3 (42%) had a high risk of bias (Figure 2).

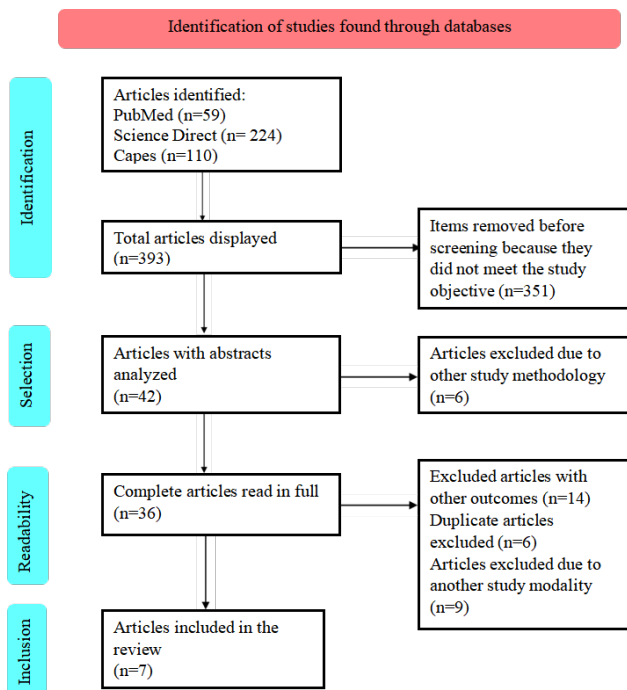


Figure 1. Flowchart of the study selection strategy, according to the PRISMA model, 2023

Study	D1	D2	D3	D4	D5	Overall
Da Silva Paixão <i>et al.</i> , 2017	⊖	⊕	⊕	⊕	⊕	⊖
De la Rosa Oliva <i>et al.</i> , 2019	⊕	⊖	⊕	⊕	⊕	⊖
Darwito <i>et al.</i> , 2019	⊕	⊕	⊕	⊖	⊖	⊖
Peppone <i>et al.</i> , 2019	⊕	⊖	⊕	⊖	⊕	⊖
Mohseni <i>et al.</i> , 2019	⊕	⊕	⊕	⊕	⊕	⊕
El-Bassiouny <i>et al.</i> , 2022	⊖	⊕	⊕	⊖	⊖	⊖
Going <i>et al.</i> , 2018	⊕	⊕	⊖	⊕	⊕	⊖

Domains:
D1: Bias arising from the randomization process.
D2: Bias due to deviations from intended intervention.
D3: Bias due to missing outcome data.
D4: Bias in measurement of the outcome.
D5: Bias in selection of the reported result.

Judgement:
⊖ High
⊖ Some concerns
⊕ Low

Figure 2. Table representing the “traffic light” risk of bias of articles

Table 2. Criteria for inclusion and exclusion of studies according to PICOS, 2023

		Criteria for inclusion	Criteria for exclusion
P	Participants	Adult women undergoing treatment and post-treatment for BC, over the age of 18.	Women with comorbidities, and metastases, men with BC, and animal studies.
I	Intervention	Omega-3 and/or vitamin D supplementation in BC patients.	Use of medication, hormone therapy, complementary supplements, and multivitamins, among others.
C	Comparison	Control group. Case group.	No group for comparisons.
O	Results	Effects of omega-3 and/or vitamin D supplementation on the health of BC patients.	-
S	Studies	Randomized Clinical Trials	Literature reviews, case studies, dissertations, theses, book chapters, abstracts presented at conferences, expert opinions, editor's comments, and duplicate references.

Results

In the databases with the keywords and filters applied, 393 studies were found, with 224 articles in Science Direct, 110 in Periodic of Capes and 59 in PubMed. After reviewing the titles, 274 were excluded because they did not meet the objective of the study. Four articles were excluded in the second stage, because they did not meet the inclusion criteria, with other methodological approaches, animal studies, and studies that investigated individuals with other comorbidities, under 18 years of age, and without the intervention of physical activity and/or physical exercise in patients with BC. In the final analysis, 36 studies were read in full, and 29 were excluded because they presented different outcomes from the main investigation, reduplicated, or used different methodologies. As a result, 7 articles were included in this systematic review.

All study participants were diagnosed with BC in the

ductal and/or lobular regions of the breast. The age of the patients in the studies varied widely, mainly because BC most commonly affects women over the age of 50 [2]. Thus, the largest age range was 50 to 60 years, with four studies showing an average age of 51 years, while two studies were conducted with women under 50 years, whose average age was 47 years. Only one study did not include information on the average age of participants.

A total of 449 women were included in the studies, with the small study involving 29 participants and the largest including 150 participants. In terms of stage of BC, two studies recruited patients in stages II to III, four studies were conducted with patients in stages I to III and one study was conducted with women in stage I only.

During the intervention, studies were conducted at different stages of treatment, one after surgery, one before surgery, one carried out during chemotherapy (the article does not specify which chemotherapy was used), one study conducted during adjuvant chemotherapy, two during neoadjuvant chemotherapy, and one after surgery, chemotherapy and radiotherapy.

The year with the most publications was 2019 with four studies. This was followed by 2018 and 2017 with one study each, and the most recent study was published in 2022.

The studies were conducted on several continents: three in North America (USA, Mexico), one in South America (Brazil), two in Asia (Iran, Indonesia), and only one in Africa (Egypt).

All studies had a control group. In the omega-3 studies, the control groups were given capsules containing corn, sunflower and soybean oil. In the studies with vitamin D, the control group was administered with edible paraffin in one article, while in another article the control group was administered with a dose of vitamin D to act as a placebo. In the last study, no intervention was made in the control group.

With respect to omega-3 interventions, all selected studies used fish oil as a source. The period that the intervention group remained receiving supplementation ranged from 4 weeks (only one study) to 24 weeks (one study), with two studies performing the intervention for 6 and 7 weeks.

Regarding the frequency of intake, it was at least once daily in all studies, twice daily in three studies and once daily in only one study. In all studies with omega-3 supplementation, EPA and DHA were included in the composition.

Regarding vitamin D interventions, all selected studies used vitamin D3 as a source. The period that the intervention group remained receiving supplementation ranged from 6 weeks (only one study) to 12 weeks (one study), with one study providing the intervention for 8 weeks.

In terms of frequency, two studies conducted the approach at least once a day, and only one study performed the intervention once a week.

For a better explanation and visualization of the results described above, four Tables were created, Table 3 and Table 4 refer to the characteristics of the studies with omega-3, and Table 5 and Table 6 refer to the studies with vitamin D supplementation and describe the specific characteristics of each selected randomized clinical trial.

There is an important point to be clarified regarding the heterogeneity of the studies included in this review. The first fact to be mentioned concerns the different time points at which the treatment with supplementation was initiated for BC. Furthermore, the different mechanisms that each medication implemented in chemotherapy may affect the results of vitamin D and omega-3 supplementation are still not well understood.

It is also worth highlighting that the majority of studies presented a moderate risk of bias, and only a single selected article had a low risk of bias, which could affect the results described here.

Table 3. Specificities of studies testing omega-3 supplementation, eligible on sample, control/intervention group, and time of treatment, 2023

	Author/Year/ Country	Total sample size	The average age of groups	The average age of groups	Time of treatment of patients during the intervention
1	Da Silva Paixão et al., 2017 (BRASIL)	37 participants	Average age 51 years	Intervention Group: Received fish oil gel capsules. Control Group: Received mineral oil concentrate.	After surgery
2	Da Silva Paixão et al., 2017 (BRASIL)	52 participants	Average age 50,1 years	Intervention Group: Received fish oil gel capsules. Control Group: Received capsules containing sunflower oil.	During neoadjuvant chemotherapy
3	Darwito et al., 2019 (INDONÉSIA)	48 participants	Average age 47,5 years	Intervention Group: Received fish oil gel capsules. Control Group: Received an unspecified placebo.	During neoadjuvant chemotherapy
4	Peppone et al., 2019 (EUA)	81 participants	Average age 59,7 years	Intervention Group 1: Received a high dose of omega-6 (soybean oil). Intervention Group 2: Received omega-3 (fish oil). Control group: Low dose of omega-3.	After surgery, radiotherapy and chemotherapy

Table 4. Details of the selected articles investigated omega-3 supplementation, on the aspects of interventions, physical and psychological health, time of intervention, frequency, amount and source of omega-3, and main outcomes

Author/Year	Intervention Group	Total duration (number of weeks)	Frequency	Quantity (mg/g)	Source of omega-3	Main results
1 Da Silva Paixão et al., 2017 (BRASIL)	18 patients received n-3 PUFA	4 weeks	2 × daily (1 capsule)	1 capsule contained 470mg EPA and 390mg DHA	Fish oil	No difference was found between the two groups on Interleukin6 (IL-6), TNF- α , and IL-1 β . However, GI patients did not show oscillation in hs-CRP and CD4+ mononuclear lymphocyte levels. In the CG, an increase in hs-CRP was observed compared to the baseline (p = 0.024). Indicating a higher inflammatory profile in response to the tumor.
2 De la Rosa Oliva et al., 2019 (MÉXICO)	26 patients received n-3 PUFA	24 weeks	2 × daily (2 capsules)	1 capsule contained 2.4g EPA/DHA with a 1:2 ratio.	Fish oil	The IG showed a significant decrease in xerostomia episodes compared to the CG. (p = 0,032).
3 Darwito et al., 2019 (INDONÉSIA)	24 patients received n-3 PUFA	7 weeks	1 × a day (1 capsule)	1 capsule contained 2.4g EPA/DHA with a 1:2 ratio.	Fish oil	GI obtained lower expression of ki-67 and VEGF compared to CG (p = 0.032 and p = 0.041) indicating lower cell proliferation, angiogenesis, and lymph angiogenesis.
4 Darwito et al., 2019 (INDONÉSIA)	30 patients received n-3 PUFA	6 weeks	2 × daily (3 capsules)	1 capsule contained 325mg of EPA and 225 of DHA	Fish oil	GI receiving omega-6 had significant improvement regarding FRC and BIF total score compared to GI receiving omega-3 (p < 0.01 and p = 0.04) respectively. In addition, the omega-3 GI showed a statistically significant decrease regarding IL-6, PTGES2, and IFN γ , compared to omega-6 GI.

Subtitles: hs-CRP, High Sensitivity C-Reactive Protein; IG, Intervention Group; CG: Control Group.

Table 5. Specificities of studies testing vitamin D supplementation, eligible on sample, control/intervention group, and time of treatment, 2023

Author/Year/Country	Total sample size	The average age of groups	Intervention/Control Group	Time of treatment of patients during the intervention
1 Mohseni et al., 2019 (IRÃ)	52 participants	Average age 47 years	Intervention Group: Received vitamin D capsules. A placebo Group: Received edible paraffin.	During chemotherapy
2 El-Bassiouny et al., 2022 (EGITO)	150 participants	Average age 52 years	Intervention Group: Received vitamin D capsules. A placebo Group: No intervention was performed.	During adjuvant chemotherapy
3 Going et al., 2018 (EUA)	29 participants		Group 1: Received high doses of vitamin D. Group 2: received low-doses of vitamin D (behaving as placebo).	Before surgery

Table 6. Details of the selected articles investigated vitamin D supplementation, on aspects of interventions, physical and psychological health, timing of intervention, frequency, amount and source of vitamin D, and main outcomes

	Author/ Year	Intervention Group	Total duration (number of weeks)	Frequency	Quantity (mg/g)	Source of omega-3	Main results
1	Mohseni et al., 2019 (IRÁ)	26 patients received Vitamin D	8 weeks	Once per week	50.000 UI	Vitamin D3	No difference was found between the two groups on inflammatory markers. However, GI patients had a significant result regarding TAC compared to GP (p=0.017).
2	El- Bassiouny et al., 2022 (EGITO)	75 patients received Vitamin D	12 weeks	Once a day	5.000 UI	-	GI patients had a significant difference compared to CG regarding serum LDH, IL-6, and cTnT levels (p < 0.001).
3	Going et al., 2018 (EUA)	15 patients received Vitamin D	6 weeks	Once a day	10.000 UI	Vitamin D3	Patients who received high doses of vitamin D showed decreased serum levels of 27HC. However, there was no significant difference compared to the low-dose group.

Subtitles:hs-CRP, High Sensitivity C-Reactive Protein; IG, Intervention Group; CG: Control Group; TAC, Total Antioxidant Capacity; LDH, Lactate Dehydrogenase; cTnT, Cardiac Troponin T; IL-6, Interleukin 6; 27HC, 27- Hydroxycholesterol.

Discussion

To our knowledge, this is the first detailed review conducted on RTC that evaluated the efficacy of omega-3 and vitamin D in patients with BC. Our findings in this review reveal and support that omega-3 and vitamin D supplementation can help women who are metastasis-free and not receiving hormone therapy during treatment for HR+ (Hormone Receptor Positive) BC. Thus, this intervention is safe and can be used adjunctively to other main treatments.

The main benefits of omega-3 supplementation include: attenuation of the inflammatory profile in response to the tumor due to the decrease in IL-6, IFN γ and PTGES2, decreased cell proliferation, angiogenesis, and lymphangiogenesis, through low expression of ki-67 and VEGF (Vascular Endothelial Growth Factor), and decreased xerostomia. In addition, immunity is improved by maintaining normal serum levels of CD4+ T lymphocytes (cytotoxic T lymphocytes) and hs-CRP (high-sensitivity C-reactive protein) [16-19].

Vitamin D can assist in the suppression of RH+ tumors, as well as mitigate cardiometabolic effects by reducing IL-6, LHD, and cTnT, improving total antioxidant capacity (TAC), and preventing tumor growth through 27HC (27-Hydroxycholesterol) and CYP27A1 [20-22].

Effects of omega-3 supplementation

Omega-3 polyunsaturated fatty acids (PUFAs) have several benefits in different types of cancer in adults and children [23-26]. The beneficial mechanisms include modulation of the inflammatory response, mitigation of

cell proliferation (metastasis and angiogenesis), and a role in gene expression that drives changes in cell metabolism. In addition, omega-3 PUFAs suppress the production of COX2, through the main omega-3 fatty acids (alpha-linolenic, EPA and DHA) [27, 28].

Tumors with positive hormone receptors, especially BC omega-3, can modulate estrogen metabolism, leading to a lower growth stimulus of hormone-dependent neoplastic cells, and decreasing arachidonic acid, which in turn leads to a decrease in derivatives of thromboxane, leukotriene and prostaglandin E2, which may reduce tumor cell survival [29, 30].

When n-3 PUFAs are used in patients receiving hormone therapy (aromatase inhibitors - AI), an improvement in pain and arthralgia can be observed, but the results are still contradictory [31-33].

In addition, no significant results were found in reducing the inflammatory profile of BC patients receiving AI. This result may be justified due to the duration of supplementation, as well as the side effects of AIs, which might inhibit the anti-inflammatory effect of omega-3 [33, 34].

At the same time, it is necessary to shed light on the dietary profile of these patients and how this food intake may interfere with treatment. In a recent study, it was possible to verify that BC patients are women who have a poor intake of essential nutrients and good fats, such as omega-6 and 3. This is reflected in their nutritional status and characterizes these women as overweight and obese, with a more inflammatory profile and a worse prognosis [35].

In addition, the Mediterranean diet (MD), which is rich in omega-3, is associated with a protective factor against

CM in Italian women. The study showed that women who had strong adherence to the Mediterranean diet had an approximately 20% reduced risk compared with women who had low adherence to the MD, with an OR of 0.82 (95% CI 0.71-0.95, $p = 0.008$) [36].

In a meta-analysis, an inverse association was observed between CM and ER-. Patients with this subtype of CM had a low adherence to DM, while patients with high adherence had a lower risk of 40% (OR 0.60, 95% CI 0.39 - 0.93, p -trend = 0.03) [37].

However, these results may be contradictory as no positive result was obtained in other populations [38]. It also corroborates that some molecular subtypes of CM seem to vary according to ethnicity and race. In addition to the genetic factor, it is a parameter that interacts with different environmental stimuli, such as lifestyle, food consumption, physical inactivity and smoking, among others [1].

The importance of vitamin D in the treatment of BC patients

Regarding vitamin D, it is worth noting that most patients with BC have low serum levels of 25-(OH) D (25-hydroxyvitamin D) and an obesogenic profile [39-41]. However, supplementation is an effective method to restore serum levels to baseline. It is important to consider whether vitamin D has a protective effect on patients before they develop CM and what the benefits of vitamin D has in the treatment of patients with CM, but the results are still uncertain [42].

Several studies have revealed that vitamin D can improve overall disease-free survival, and clinical prognosis, as well as mitigate inflammation and regulate antiproliferative pathways and cell differentiation [43-45].

The most widely accepted mechanism for the benefits of calcitriol in BC is that calcitriol exerts its role best when it binds to its receptor VDR. This VDR receptor is widely distributed in breast tissue and can modulate and control cellular mechanisms, such as cell proliferation, invasion, metastasis, angiogenesis and apoptosis [46,47].

A study conducted by Swedish researchers on women with an average age of 54 years found that a higher presence of this receptor in breast tissue indicated a better prognosis and a lower risk of death. In contrast, women with low levels of RVR expression had a high rate of cell proliferation as measured by ki-67 ($p < 0.001$), a larger tumor size ($p = 0.002$) and a high Nottingham grade ($p < 0.001$) [45].

Some limitations in this review need to be better addressed, including in cohort studies and randomized trials, among others. Particularly in relation to the interventions of omega-3 and vitamin D, the optimal dose and the time of intervention, as there seems to be no minimum period for supplementation established in the literature. Furthermore, in some studies only a single serum dosage was administered, which may obscure the true value of 25-(OH) D and omega-3. Moreover,

the studies examined include interventions in different ethnic cities with different molecular subtypes of BC, as the different subtypes generate different pathologic and molecular responses, which consequently interferes with the prognosis and the natural course of the disease. It is also worth nothing that some metabolic factors may affect the bioavailability of these compounds, as well as nutritional status, dietary intake and sedentary lifestyle, among others.

From future perspectives, randomized clinical trials separated by a certain molecular subtype should be developed to investigate the efficacy of omega-3 and vitamin D supplementation during the treatment of women with BC. It is also important that new interventions are separated according to different stages of treatment, as chemotherapy drugs can mask the real positive effects of supplementation, or even decrease their molecular activities.

Conclusion

Omega-3 can assist in the adjuvant treatment of BC, as this nutrient has shown several positive effects on the health of these patients by improving immunity and inflammatory markers.

In respect of calcitriol, it can also be used during chemotherapy in RE+ patients. It can improve the cardiovascular metabolic profile and prognosis, as well as inhibit the progression of CM through 27HC.

Finally, longer longitudinal studies and randomized clinical trials in women with only one molecular subtype are needed to determine the optimal dose and timing of vitamin D and omega-3 intervention. This will further clarify the results found in this review and add to the literature. It is further emphasized that the more low-cost therapies investigated, the better the clinical treatment options for patients with breast neoplasms.

Authors' contribution

Lailton Oliveira da Silva and Victor da Silva collected the research data, in addition to filling in the results for tabulating this data, as well as wrote the article. Anderson Weiny Barbalho Silva and José Juvenal Linhares supervised the work, provided support services in data collection, as well as gave guidance on writing the research.

Conflict of interests

Nothing to declare.

Funding

The research did not receive funding.

References

- [1] IARC [Internet]. Estimated number of deaths in 2020, Word. Available from <https://gco.iarc.fr/today/home>. [Accessed 24th Jul 2023].
- [2] ACS [internet]. Dietary Supplements, 2021. Available from <https://www.cancer.org/cancer/managing-cancer/treatment-types/complementary-and-integrative-medicine/dietary-supplements.html> [Accessed 25th Jul 2023].
- [3] Carlberg C, Muñoz A. An update on vitamin D signaling and cancer. *Seminars in Cancer Biology*. 2022; 79: 217-230. doi: 10.1016/j.semcancer.2020.05.018.
- [4] Carlberg C, Velleuer E. Vitamin D and the risk for cancer: A molecular analysis. *Biochem Pharmacol*. 2022; 196: 114735. doi: 10.1016/j.bcp.2021.114735.
- [5] Saini RK, Prasad P, Sreedhar RV, et al. Omega-3 Polyunsaturated Fatty Acids (PUFAs): Emerging Plant and Microbial Sources, Oxidative Stability, Bioavailability, and Health Benefits-A Review. *Antioxidants* (Basel). 2021; 10: 1627-1636. doi: 10.3390/antiox10101627.
- [6] Redruello-Requejo M, Samaniego-Vaesken ML, Puga AM, et al. Intake, Determinants and Dietary Sources of Omega-3 and Omega-6 Polyunsaturated Fatty Acids in the Spanish Population: Study Results ANIBES. *Nutrients*. 2023; 15: 562-571. doi: 10.3390/nu15030562.
- [7] Vega OM, Abkenari S, Tong Z, et al. Omega-3 Polyunsaturated Fatty Acids and Lung Cancer: nutrition or Pharmacology? *Nutrition and Cancer*. 2021; 73: 541-561. doi: 10.1080/01635581.2020.1761408.
- [8] Fabian CJ, Kimler BF, Hursting SD. Omega-3 fatty acids for breast cancer prevention and survivorship. *Breast Cancer Research*. 2015; 17: 62-73. doi: 10.1186/s13058-015-0571-6.
- [9] Pizato N, Luzete BC, Kiffer LFMV, et al. Omega-3 docosahexaenoic acid induces pyroptosis cell death in triple-negative breast cancer cells. *Scientific Reports*. 2018; 8: 1952-1964. doi: 10.1038/s41598-018-20422-0.
- [10] Benedik E. Sources of vitamin D for humans. *International Journal for Vitamin and Nutrition Research*. 2022; 92: 118-125. doi: 10.1024/0300-9831/a000733.
- [11] Wang W, Hu W, Xue S, et al. Vitamin D and Lung Cancer; Association, Prevention, and Treatment. *Nutrition and Cancer*. 2021; 73: 2188-2200. doi: 10.1080/01635581.2020.1844245.
- [12] Martin-Gorgojo A, Gilaberte Y, Nagore E. Vitamin D and Skin Cancer: An Epidemiological, Patient-Centered Update and Review. *Nutrients*. 2021; 13: 4292-4306. doi: 10.3390/nu13124292.
- [13] Vanhevel J, Verlinden L, Doms S, et al. The role of vitamin D in breast cancer risk and progression. *Endocrine-Related Cancer*. 2022; 29: R33-R55. doi: 10.1530/ERC-21-0182.
- [14] Lee KH, Seong HJ, Kim G, et al. Consumption of Fish and ω -3 Fatty Acids and Cancer Risk: An Umbrella Review of Meta-Analyses of Observational Studies. *Advances in Nutrition*. 2020; 11: 1134-1149. doi: 10.1093/advances/nmaa055.
- [15] Page MJ, McKenzie JE, Bossuyt PM, et al. The PRISMA 2020 statement: an updated guideline for reporting systematic reviews. *International Journal of Surgery*. 2021; 88:105906. doi: 10.1016/j.ijssu.2021.105906.
- [16] Paixão EMDS, Oliveira ACM, Pizato N, et al. The effects of EPA and DHA enriched fish oil on nutritional and immunological markers of treatment naïve breast cancer patients: a randomized double-blind controlled trial. *Nutrition Journal*. 2017; 16: 71-82. doi: 10.1186/s12937-017-0295-9.
- [17] De La Rosa Oliva F, Meneses García A, Ruiz Calzada H, et al. Effects of omega-3 fatty acids supplementation on neoadjuvant chemotherapy-induced toxicity in patients with locally advanced breast cancer: a randomized, controlled, double-blinded clinical trial. *Nutricion Hospitalaria*. 2019; 36: 769-776. doi: 10.20960/nh.2338.
- [18] Darwito D, Dharmana E, Riwanto I, et al. Effects of Omega-3 Supplementation on Ki-67 and VEGF Expression Levels and Clinical Outcomes of Locally Advanced Breast Cancer Patients Treated with Neoadjuvant CAF Chemotherapy: A Randomized Controlled Trial Report. *Asian Pacific Journal of Cancer Prevention*. 2019; 20: 911-916. doi: 10.31557/2FAPJCP.2019.20.3.911.
- [19] Peppone LJ, Inglis JE, Mustian KM, et al. Multicenter randomized controlled trial of omega-3 fatty acids versus omega-6 fatty acids for the control of cancer-related fatigue among breast cancer survivors. *JNCI Cancer Spectrum*. 2019; 3: pkz005. doi: 10.1093/jncics/pkz005.
- [20] Mohseni H, Amani R, Hosseini SA, et al. Genetic Variations in VDR Could Modulate the Efficacy of Vitamin D3 Supplementation on Inflammatory Markers and Total Antioxidant Capacity among Breast Cancer Women: A Randomized Double-Blind Controlled Trial. *Asian Pacific Journal of Cancer Prevention*. 2019; 20: 2065-2072. doi: 10.31557/2FAPJCP.2019.20.7.2065.
- [21] El-Bassiouny NA, Helmy MW, Hassan MAE, et al. The cardioprotective effect of vitamin D in breast cancer patients receiving adjuvant doxorubicin-based chemotherapy. *Clinical Breast Cancer*. 2022; 22:

- 359-366. doi: 10.1016/j.clbc.2022.01.008.
- [22] Going CC, Alexandrova L, Lau K, et al. Vitamin D supplementation decreases serum 27-hydroxycholesterol and expression of CYP27A1 in tumors of breast cancer patients. *Cancer Research*. 2017; 77: 5635-5635. doi:10.1158/1538-7445.AM2017-5635.
- [23] D'Angelo S, Motti ML, Meccariello R. ω -3 and ω -6 Polyunsaturated Fatty Acids, Obesity and Cancer. *Nutrients*. 2020; 12: 2751-2767. doi: 10.3390/nu12092751.
- [24] Podpeskar A, Crazzolara R, Kropshofer G, et al. Omega-3 Fatty Acids and Their Role in Pediatric Cancer. *Nutrients*. 2021; 13: 1800-1813. doi: 10.3390/nu13061800.
- [25] Aredes MA, da Camara AO, de Paula NS, et al. Efficacy of ω -3 supplementation on nutritional status, skeletal muscle, and chemoradiotherapy toxicity in cervical cancer patients: A randomized, triple-blind, clinical trial conducted in a middle-income country. *Nutrition*. 2019; 67: 110528. doi: 10.1016/j.nut.2019.06.009.
- [26] Lee KH, Seong HJ, Kim G, et al. Consumption of Fish and ω -3 Fatty Acids and Cancer Risk: An Umbrella Review of Meta-Analyses of Observational Studies. *Advances in Nutrition*. 2020; 11: 1134-1149. doi: 10.1093/advances/nmaa055.
- [27] De Abreu Leite JT, Lobo LC, De Andrade LG. Ômega-3 no tratamento paliativo do câncer. *Revista Ibero-Americana de Humanidades, Ciências e Educação*. 2021; 7: 1547-1561. doi: 10.51891/rease.v7i10.2680.
- [28] Gomes ALM, Magalhães JA, Neves JP, et al. Efeitos da suplementação de arginina, glutamina e ômega-3 sobre a resposta inflamatória e estado nutricional de pacientes oncológicos. *Research, Society and Development*. 2020; 9: e193953285-e193953285. doi: 10.33448/rsd-v9i5.3285.
- [29] Hutchins-Wiese HL, Picho K, Watkins BA, et al. High-dose eicosapentaenoic acid and docosahexaenoic acid supplementation reduces bone resorption in postmenopausal breast cancer survivors on aromatase inhibitors: a pilot study. *Nutrition and Cancer*. 2014; 66: 68-76. doi: 10.1080/01635581.2014.847964.
- [30] Laudisio D, Castellucci B, Barrea L, et al. Mediterranean diet and breast cancer risk: a narrative review. *Minerva Endocrinol (Torino)*. 2021; 46: 441-452. doi: 10.23736/s2724-6507.20.03266-6.
- [31] Shen S, Unger JM, Crew KD, et al. Omega-3 fatty acid use for obese breast cancer patients with aromatase inhibitor-related arthralgia (SWOG S0927). *Breast Cancer Research and Treatment*. 2018; 172: 603-610. doi:10.1007%2Fs10549-018-4946-0.
- [32] Hershman DL, Unger JM, Crew KD, et al. Randomized Multicenter Placebo-Controlled Trial of Omega-3 Fatty Acids for the Control of Aromatase Inhibitor-Induced Musculoskeletal Pain: SWOG S0927. *Journal of Clinical Oncology*. 2015; 33: 1910-1917. doi: 10.1200/jco.2014.59.5595.
- [33] Lustberg MB, Orchard TS, Reinbolt R, et al. Randomized placebo-controlled pilot trial of omega 3 fatty acids for prevention of aromatase inhibitor-induced musculoskeletal pain. *Breast Cancer Research and Treatment*. 2018; 167: 709-718. doi: 10.1007/s10549-017-4559-z.
- [34] Hutchins-Wiese HL, Picho K, Watkins BA, et al. High-Dose Eicosapentaenoic Acid and Docosahexaenoic Acid Supplementation Reduce Bone Resorption in Postmenopausal Breast Cancer Survivors on Aromatase Inhibitors: A Pilot Study. *Nutrition and Cancer*. 2014; 66: 68-76. doi: 10.1080/01635581.2014.847964.
- [35] Silva LO, Sousa IML, Paim RTT, et al. Perfil nutricional e consumo alimentar de pacientes com câncer de mama: uma revisão integrativa. *RBONE*. 2023;17:185-91.
- [36] Turati F, Carioli G, Bravi F, et al. Mediterranean Diet and Breast Cancer Risk. *Nutrients*. 2018; 10: 326-339. doi: 10.3390/nu10030326.
- [37] Van Den Brandt PA, Schulpen M. Mediterranean diet adherence and risk of postmenopausal breast cancer: results of a cohort study and meta-analysis. *International Journal of Cancer*. 2017;140:2220-2231. doi: 10.1002/ijc.30654.
- [38] Butler LM, Wu AH, Wang R, et al. A vegetable-fruit-soy dietary pattern protects against breast cancer among postmenopausal Singapore Chinese women. *The American Journal of Clinical Nutrition*. 2010; 91: 1013-1019. doi: 10.3945/ajcn.2009.28572.
- [39] Rosso C, Fera N, Murugan NJ, et al. Vitamin D Levels in Newly Diagnosed Breast Cancer Patients According to Tumor Sub-Types. *Journal of Dietary Supplements*. 2022; 14: 1-13. doi: 10.1080/19390211.2022.2144582.
- [40] Bakhshaiesh TO, Nazeri E, Jafarbeik-Iravani N, et al. Vitamin D and breast cancer risk: A systematic review and meta-analysis in Iranian patients. *Annals of Medicine and Surgery (Lond)*. 2022; 80: 104162. doi: 10.1016/j.amsu.2022.104162.
- [41] Fernandes EMA, Linhares JJ. Papel dos níveis séricos de vitamina d e da síndrome metabólica e o risco de câncer de mama. *Arquivos Catarinenses de Medicina*. 2021; 50: 257-70.
- [42] Isbilen E, Kus T, Cinkir HY, et al. Better survival associated with successful vitamin D supplementation in non-metastatic breast cancer survivors. *Turkish Journal of Biochemistry*. 2021; 46: 509-516. doi: 10.1515/tjb-2021-0137.
- [43] Zhang X, Harbeck N, Jeschke U, et al. Influence of vitamin D signaling on hormone receptor status and HER2 expression in breast cancer. *Journal of Cancer Research and Clinical Oncology*. 2017; 143: 1107-

1122. doi: 10.1007/s00432-016-2325-y.

- [44] Li Z, Wu L, Zhang J, et al. Effect of vitamin D supplementation on risk of breast cancer: a systematic review and meta-analysis of randomized controlled trials. *Frontiers in Nutrition*. 2021; 8: 655727. doi: 10.3389/fnut.2021.655727.
- [45] Huss L, Butt ST, Borgquist S, et al. Vitamin D receptor expression in invasive breast tumors and breast cancer survival. *Breast Cancer Research*. 2019; 21: 84. doi: 10.1186/s13058-019-1169-1.
- [46] Cui X, Pertile R, Eyles DW. The vitamin D receptor (VDR) binds to the nuclear matrix via its hinge domain: a potential mechanism for the reduction in VDR-mediated transcription in mitotic cells. *Molecular and Cellular Endocrinology*. 2018; 472:18-25. doi: 10.1016/j.mce.2017.11.015.
- [47] Trivedi T, Zheng Y, Fournier PGJ, et al. The vitamin D receptor is involved in the regulation of human breast cancer cell growth via a ligand-independent function in the cytoplasm. *Oncotarget*. 2017; (8)16: 26687. doi: 10.18632/oncotarget.15803.