



Effects of an online home-based exercise intervention on breast cancer survivors during COVID-19 lockdown: a feasibility study

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Abstract

Purpose The strict lockdown implemented due the COVID-19 pandemic is generating a great impact on wellbeing and health-related quality of life (HRQoL) in people with cancer. We aimed to evaluate the efficacy and feasibility of an online home-based exercise intervention performed during a lockdown period analysing its effects on body composition, physical fitness, and HRQoL in breast cancer survivors.

Methods Fifteen women with breast cancer receiving hormonal therapy (55.5 ± 6.7 years) were included in the study. The exercise intervention consisted of two weekly sessions of remotely supervised functional training (60 min per day) and two weekly sessions of unsupervised aerobic training (20–30 min/session; 60–85% of maximum heart rate) for a total of 16 weeks. DXA absorptiometry was used for the assessment of body composition. Functional assessment included cardiorespiratory fitness (CRF) by Rockport walking test, upper and lower body strength (grip strength, arm curl test, and chair stand test), walking speed (brisk walking test), and agility (8-foot up-and-go test). The HRQoL was evaluated with the QLQ-BR23 questionnaire. The adherence to the intervention was measured as the percentage of online classes attended.

Results Rate of adherence for the online exercise intervention was $90 \pm 17\%$. The exercise intervention induced significant ($p < 0.05$) improvements in physical fitness: CRF (+9%), right arm and lower limb strength (+10% and +18%, respectively) and lower limbs lean mass (+2% and +3.5% for left and right leg, respectively).

Conclusion This feasibility study suggests that an online home-based exercise intervention during COVID-19 lockdown could improve physical fitness and body composition in breast cancer survivors even in a context of heightened concern for future health.

Keywords Breast neoplasms · Coronavirus · Exercise · Lockdown · Physical function

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Introduction

The coronavirus disease 2019 (COVID-19) pandemic has caused an unprecedented international crisis. The World Health Organization (WHO) declared this situation a public health emergency of international concern [1]. Initial measures adopted by different countries to control of COVID-19 included important mobility restrictions with governments applying strict lockdowns that lasted several weeks or even months. Unfortunately, these measurements have had negative implications in population health and lifestyle. In fact, several studies have reported a reduction in physical activity (PA) levels [2, 3] with an increase in sedentary behavior produced by the afore-mentioned mobility restrictions.

In the area of oncology, the lockdown among cancer patients has entailed a great negative impact in their global wellbeing and consequently in their health-related quality of life (HRQoL). On this basis, a recent study reported that about 9 out of 10 of cancer patients showed an impairment in their security and daily functionality due to the COVID-19 pandemic [4, 5]. The benefits associated to physical exercise in cancer survivors as an adjuvant therapy have been widely described [6, 7]. In fact, over the last decades, supervised exercise interventions have shown multiple biopsychosocial benefits for people with cancer [8, 9]. However, due to this pandemic scenario, the global governments and the public health policies have framed “staying at home” as a preventive strategy. In this current situation associated with a global lockdown and restricted mobility, it is necessary to identify and explore new approaches and intervention strategies that enable adequate PA promotion among people with cancer. A possible solution tailored to this population could be the development of an online home-based exercise intervention [10, 11]. Specifically, for patients with cancer and survivors who are more vulnerable due to their global health status with an increased risk of exposure and virus infection.

In cancer survivors, home-based exercise has been considered as a safe and effective strategy to improve markers of health such as body composition, fatigue, functional capacity and HRQoL [12–17]. Focusing on breast cancer, home-based exercise programs have been developed during the last decades [18]. In these programs, patients are assessed and followed on a periodic basis, and perform exercise routines with the support of brochures, practical guides, or electronic materials [19]. Consequently, counseled exercise intervention could be an effective option for certain circumstances since a recent study suggested that a home-based exercise intervention developed during home confinement improved QoL in women who had suffered breast cancer [20]. Nonetheless, this was a pilot study and should be

interpreted with caution due to the low number of participants ($n = 2$) [20].

One of the main limitations of an online home-based exercise intervention is that most are performed without supervision and consequently basic training variables such as the technique of the performed exercises, intensity or volume are not controlled and adapted for each participant [21]. Therefore, the previously mentioned variables could increase the risk of injury and adverse effects, generating a low training adherence. All these aspects underline the importance of developing a supervised exercise intervention adapted to the circumstances of each oncologic participant, in order to produce quantifiable improvements in physical function and HRQoL benefits in cancer survivors during COVID-19 lockdown. Hence, the aim of this study was to evaluate the efficacy and feasibility of an online home-based exercise intervention performed with a face-to-face and synchronous supervision during a lockdown period and, specifically, to analyze its effects on body composition, physical fitness and HRQoL in breast cancer survivors receiving hormonal therapy.

Material and methods

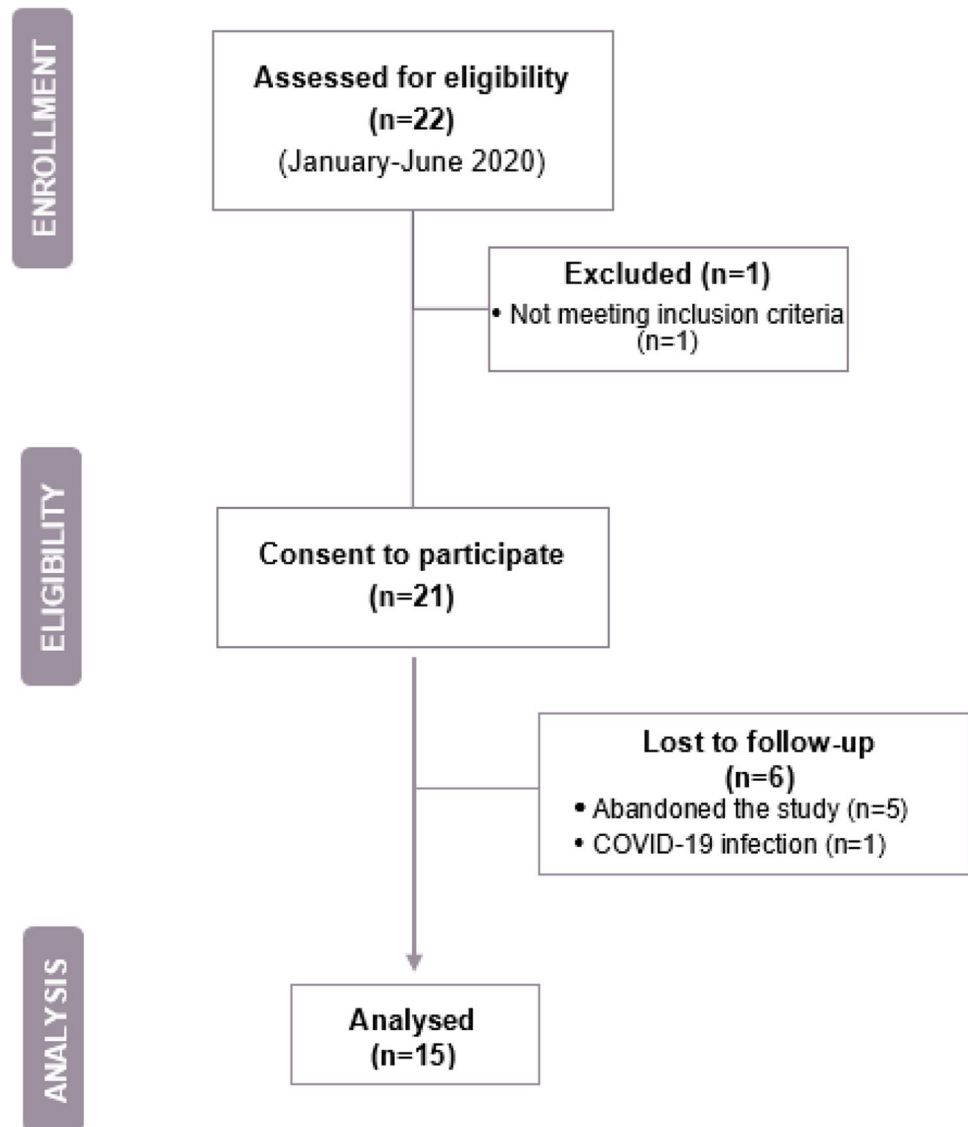
Study design

This is a quasi-experimental study with a pre-post design without a control group. The intervention had a duration of 16 weeks with the baseline assessment taking place in February 2020 while the follow-up assessment took place in June 2020. Initially, the intervention was designed to be performed with a face-to-face and supervised intervention. However, due to the unforeseen pandemic circumstance and the sudden lockdown, it was changed to an online home-based exercise intervention.

Data were collected at the beginning and the end of the study through face-to-face interviews. This was possible because pre-assessment was carried out 2-week before lockdown in Spain, and post-assessment took place in June, when social distancing and isolation measures were much more flexible. The intervention, however, did take place during the state of emergency and period of home confinement in Spain (from February to May 2020).

Participants and recruitment strategy

Twenty-one breast cancer women survivors were selected to participate in the study. Six abandoned the study (5 due to personal reasons and 1 was excluded because she had to stop the training due to COVID-19). Therefore, the final sample included 15 women (Fig. 1). All the participants belonged

Fig. 1 Flowchart of participants through the study

to the *Asociación de mujeres aragonesas de cáncer genital y de mama* (AMACGEMA) of Zaragoza (Spain). All participants were recruited for the study via email. A non-probabilistic convenience sampling was developed through an informative meeting that took place in January 2020 in the AMACGEMA venue.

The aims of the project and evaluations were explained in the mentioned meeting. Those that meet the following inclusion criteria were selected to participate in the present study: (i) age between 18 and 75 years old; (ii) diagnosed of breast cancer (stages I–III); (iii) > 6 months post-treatment (surgery, chemotherapy or radiotherapy); (iv) to be in active hormonal therapy; (v) no exercise medical contraindications (e.g., cardiovascular disease or neuromuscular disorders); and (vi) signing the written informed consent.

The exclusion criteria were (i) participants with cancer cachexia (body mass index < 20 kg/m²) [22]; (ii) being

diagnosed of another primary and/or secondary tumor or being in metastatic stage; (iii) cardiac and or lung diagnosed problems (e.g., myocardial infarction, arrhythmia, angina, heart failure or chronic obstructive pulmonary disease); (iv) to have participated in an exercise program during the previous 6 months; or (v) to have severe immunosuppression or fever.

Sociodemographic variables

A self-constructed questionnaire contained 10 items was designed to describe the participants' sociodemographic characteristics, education level, marital status, and other disease-related questions (received treatment, symptoms, and secondary effects).

Ethical committee

Eligible participants were informed about the project aims and purpose, and participation in the study was voluntary. All participants gave written informed consent prior their inclusion to the study. The present project was carried out in accordance with the principles of the Declaration of Helsinki and Good Clinical Practice guidelines. The project was approved by the Ethical Committee of the University of San Jorge (Zaragoza, Spain [n°006-19/20]). All participants were informed of the protocol of this study and agreed to participate voluntarily signing an informed consent.

Measures

Anthropometry and body composition

Height was measured with a stadiometer with 0.1 cm precision (SECA 225®, SECA, Hamburg, Germany), while weight was measured with a portable scale with a precision of 0.1 kg (SECA 861®, SECA, Hamburg, Germany). Waist and hip circumference were measured at the midpoint between the lowest rib and iliac crest at the end of a normal expiration and at the maximum diameter over the buttocks, respectively. Both measurements were performed with a non-elastic measuring tape with 0.1 cm precision (SECA 212®, SECA, Hamburg, Germany) with participants standing in anatomic position.

Dual-energy X-ray absorptiometry QDR-Explorer (QDR-Explorer, Hologic Corp., Software versión 12.4, Waltham, MA, USA) was used for the assessment of body composition. Body fat (kg and %) and lean body mass (kg) were obtained from whole-body scan. Lean body mass was obtained by subtracting bone mineral content from nonfat mass. The following sub-regions were also analyzed from the whole-body scan: lower and upper extremities and trunk.

Physical fitness assessment

The physical fitness related to functional capacity performance of participants was assessed with different tests.

A digital handgrip dynamometer (0.1 kg precision by Takey Smedley III T-19®, Scientific Instruments Co. Ltd., Niigata, Japan) was used to assess maximal handgrip strength. The “Arm Curl Test” was developed to assess upper limb strength [23]. Participants were asked to perform as many arm curls as possible in 30 s with a dumbbell of 2.5 kg weight.

The “chair stand test” was used to assess lower limb strength [23]. From a sitting position, participants were

asked to repeatedly stand up and sit back down as fast as possible for 30 s. The number of stands was recorded.

The “8-foot up-and-go test” was used to assess agility [23]. From a sitting position, participants were asked to stand up and walk to a cone placed at 2.45 m, turn and return to a seated position. The time required was reported.

The “brisk walking test” was used to evaluate walking speed. Participants were asked to walk 30 m as fast as possible (running was not permitted). The time required was reported. Finally, cardiorespiratory fitness was (CRF) assessed by “Rockport walking test” [24]; the participants walked as fast as possible for one mile (1609 m). Post walk, heart rate, and time required were registered immediately after completion of the test.

All tests were repeated twice with a minute rest between attempts except for the “chair stand test”, “arm curl test”, and the “Rockport test” that were only performed once. The best attempt of each test was selected for further analyses.

Health-related quality of life

The European Organization for Research and Treatment of Cancer [25] Quality of Life Questionnaire and Breast Cancer specific module (EORTC-QLQ-BR23) [26] was used to evaluate HRQoL. This questionnaire contains 23 items that are divided into two dimensions. The first dimension includes four functional scales (body image [4 items], sexual functioning [2 items], sexual enjoyment [1 item], and future perspective [1 item]) while the second includes four symptom scales (systemic therapy side effects [7 items], breast symptoms [4 items], arm symptoms [3 items], upset by hair loss [1 item]). Each item is scored from 1 to 4. Scores were after linearly transformed to a 0 to 100 score. The highest scores are related to better functioning (except for sexual functioning and sexual enjoyment). On the other hand, higher scores in the symptom scale dimension are associated with a higher persistence of the symptoms.

Online home-based exercise program

The supervised exercise intervention had a duration of 16 weeks. The first 2 weeks were focused on exercise familiarization. All the sessions were developed through ZOOM (Zoom Video Communications, Inc., 2020) (Fig. 2).

The exercise intervention included four weekly training sessions divided into two functional supervised training sessions and two aerobic unsupervised training sessions. The supervised part consisted of the trainer leading the session by connecting online synchronously with the participants and directing the training. During the connection, the participants could watch the performance, were encouraged, and could interact and give feedback or ask questions. The trainer could

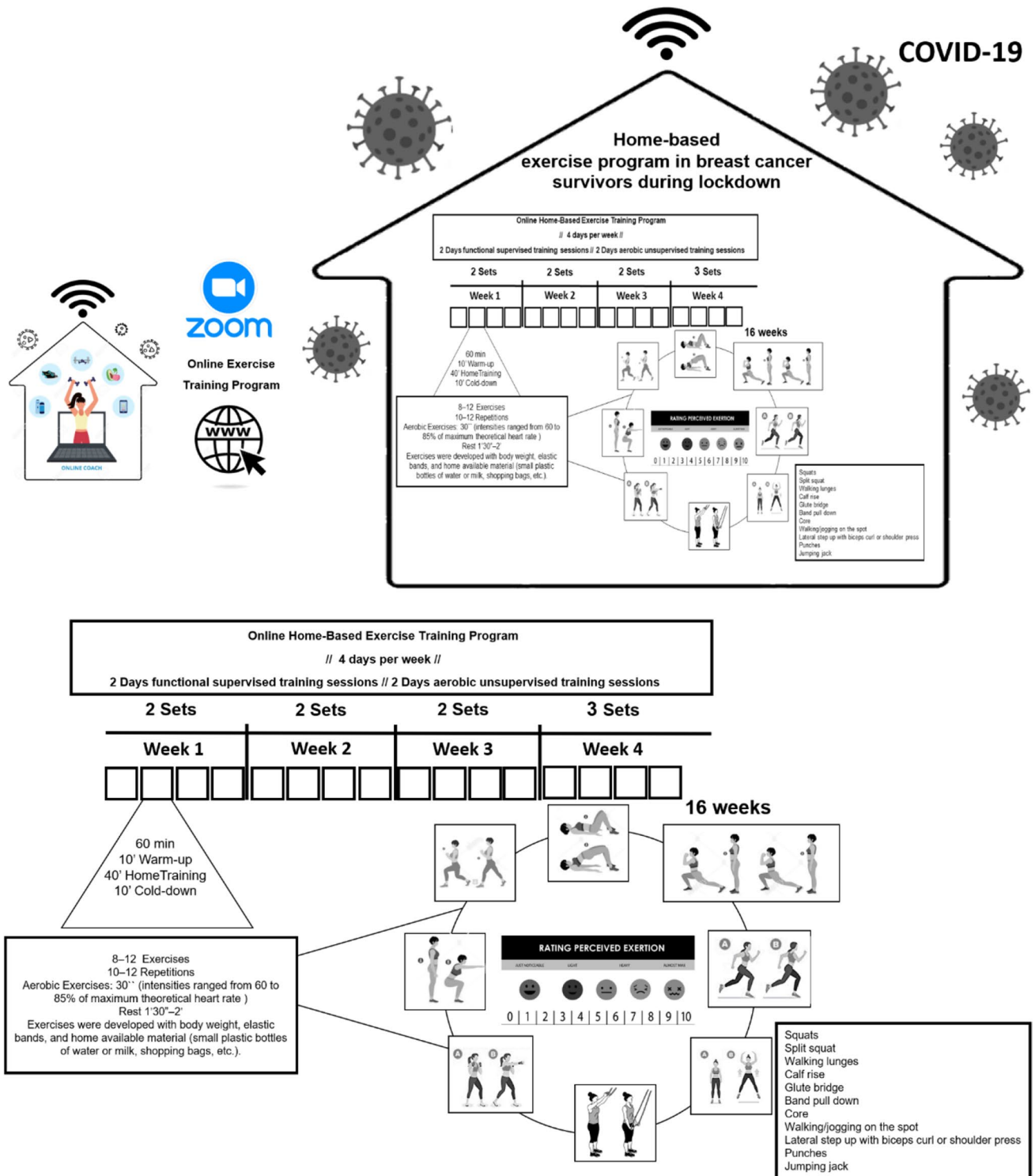


Fig. 2 Online home-based exercise program in breast cancer survivors during lockdown

also assess the participants performance and give individual or group feedback or corrections when necessary.

The supervised sessions had a duration of 60 min. Each session was divided into warm-up (10 min), a main part

of the training (40 min), and cool down (10 min). All the sessions included a combined circuit of 8–12 functional exercises to improve strength and CRF (squats, split squat, walking lunges, calf rise, glute bridge, band pull down, core,

walking/jogging on the spot, lateral step up with biceps curl or shoulder press, punches, jumping jack). Exercises were developed with body weight, elastic bands, and home available material (such as small plastic bottles of water or milk, or shopping bags). For the functional strength exercises, a 10–12 repetition protocol was developed, while the aerobic exercises lasted around 30 s. Two series were performed until week 12 and three series from week 13 to 16. A minimum rest of 90 s was established between exercises.

Participants were asked to use a heart rate monitor during all the sessions. The work intensities ranged from 60 to 85% of maximum heart rate (HR_{max}) obtained from the Rockport test and calculated through the Karvonen formula [27]. Rating of perceived exertion was also registered at the end of the warm-up and main training using the Borg scale [28].

Finally, to improve the specificity of the program and follow the WHO physical activity recommendations, we recommended participants to add each week two home aerobic unsupervised sessions (20–30 min/session; 60–85% HR_{max}). These aerobic sessions could include self-selected exercises such as dancing choreographies, indoor cycling, elliptical bike, or the use of other home available tools that would elicit a similar stimulus.

Safety and adherence

Daily adherence and safety to the intervention were monitored by means of a register of attendance. Moreover, to ensure the safety and welfare of participants involved in the research, the protect rights of human subjects and occurrence of adverse events were monitored. Finally, to determine adherence to the training, we considered a session to be completed when 80–90% of the planned exercises were successfully performed.

Statistical analyses

The Shapiro–Wilk test was used to test the normality of the outcomes. Mean, standard deviation, and confidence intervals 95% are presented for continuous outcomes. Categorical outcomes are expressed as absolute values and percentages. To test the differences between different moments of the intervention (pre-post) for the same group, the paired samples *t* test was used. The relative change (change percentage) was calculated to test the effect of the intervention. Additionally, the standardized effect size (ES) was calculated by Cohen's *d*, with values from 0.2–0.5 representing small effect; 0.5–0.8 moderate effect and > 0.8 large effect.

All the analyses were performed with SPSS v.21 (SPSS Institute Inc., IL, EUA). The level of significance was set at $p < 0.05$.

Results

Participants main characteristics

Sociodemographic characteristics (age, education level, and marital status) and variables associated with the disease (received treatment, disease recurrence, symptoms, and negative effects) are presented in Table 1.

Safety and adherence

No adverse events or health issues during the exercise intervention were noted. Adherence to the intervention averaged $90 \pm 17\%$.

Physical performance

Results for the baseline and post-intervention are presented in Table 2. Significant improvements were found for handgrip strength of the right arm, lower limbs strength, and CRF (all $p < 0.05$; ES: 0.84, 0.76, and 1.17, respectively).

Anthropometry and body composition

Anthropometric characteristics and body composition variables are presented in Table 3. Breast cancer survivors increased lean mass at both right and left legs from the beginning to the end of the exercise intervention ($p < 0.05$; ES 0.74 and 1.01 for left and right leg, respectively). No significant changes were found for the other variables evaluated (all $p > 0.05$).

Health-related quality of life

HRQoL results are presented in Table 4. Significant changes were found for the future perspectives domain which were lower at the end of the program (75.60 ± 29.45 vs 45.34 ± 36.10 ; $p < 0.05$; ES 0.90). No other significant differences were found for the rest of the assessed dimensions. Participants did not provide responses to the scales on sexual enjoyment and hair loss.

Discussion

The results of the present study show that an online home-based exercise intervention is an effective and safe strategy to improve physical fitness and lower limbs lean mass on breast cancer survivors.

Table 1 Main demographic and clinical characteristics at baseline ($n = 15$)

| | N(%) | |
|---|------------------|-----|
| Age (years) mean \pm SD | 55.53 \pm 6.73 | |
| Education level | | |
| Secondary studies | 2 (13.3) | |
| Professional degree | 8 (53.3) | |
| University or higher degree | 5 (33.3) | |
| Marital status | | |
| Single | 2 (13.3) | |
| Married | 10 (66.7) | |
| Divorced or separated | 2 (13.3) | |
| Widowed | 1 (6.7) | |
| Treatment | | |
| Surgery | 15 (100) | |
| Chemotherapy | 10 (66.7) | |
| Radiotherapy | 10 (66.7) | |
| Side effects at the beginning of the intervention | | |
| Fatigue | 8 (57.1) | |
| Depression/anxiety | 8 (57.1) | |
| Sleep disorders | 9 (64.3) | |
| Body weight changes | 9 (64.3) | |
| Pain | 8 (57.1) | |
| Lymphedema | 9 (64.3) | |
| Musculoskeletal | 12 (85.7) | |
| Neuropathy | 6 (42.9) | |
| Cronology of cancer before the exercise intervention | Mean | IQR |
| Time from diagnosis (months) | 28 | 56 |
| Time from end of treatment (months) | 25 | 107 |
| Time since the beginning of the hormonal therapy (months) | 27 | 111 |

Abbreviations: *IQR* interquartile range, *SD* standard deviation

Table 2 Effects of the exercise intervention on physical performance ($n = 15$)

| | Baseline Mean (SD) | Post Mean (SD) | Dif. Baseline-Post 95% CI | <i>p</i> -value | <i>ES</i> |
|---------------------------|-----------------------|-------------------|------------------------------|-----------------|-----------|
| Upper limbs strength | | | | | |
| Handgrip right arm (kg) | 19.7 (6.7) | 21.7 (5.9) | -2.03 (-3.50, -0.57) | 0.011 | 0.84 |
| Handgrip left arm (kg) | 17.1 (7.2) | 33.2 (44.6) | -16.09 (-43.64, 11.45) | 0.227 | 0.35 |
| Arm curl right arm (rep) | 16.3 (6.9) | 16.3 (7.9) | 0 (-3.97, 3.97) | 1.000 | 0.00 |
| Arm curl left arm (rep) | 15.4 (6.2) | 15.9 (7.7) | -0.46 (-6.60, 5.70) | 0.873 | 0.04 |
| Lower limbs strength | | | | | |
| Chair stand test (rep) | 16.2 (4.7) | 19.1 (3.5) | -2.92 (-5.26, -0.59) | 0.018 | 0.76 |
| Agility | | | | | |
| 8-foot up-and go test (s) | 5.0 (0.8) | 4.7 (0.5) | 0.31 (-0.15, 0.76) | 0.169 | 0.41 |
| Walking speed | | | | | |
| Brisk walking test (s) | 14.4 (0.8) | 14.6 (1.4) | -0.22 (-0.90, 0.47) | 0.508 | 0.18 |
| Cardiorespiratory fitness | | | | | |
| Rockport test (minutes) | 16.2 (1.2) | 14.8 (0.8) | 1.36 (0.69, 2.02) | 0.001 | 1.17 |

Data are mean \pm SD. Significant *p*-values are in bold. *CI* confidence interval, *ES* effect size, *SD* standard deviation

Table 3 Effects of exercise intervention on anthropometry and body composition ($n = 15$)

| Variable | Baseline Mean (SD) | Post Mean (SD) | Dif. Baseline-Post 95% CI | <i>p</i> -value | <i>ES</i> |
|--------------------------|--------------------|----------------|---------------------------|-----------------|-----------|
| Weight (kg) | 65.6 (10.8) | 65.4 (11.2) | 0.25 (−0.65, 1.14) | 0.564 | 0.15 |
| BMI (kg/m ²) | 25.5 (3.8) | 25.4 (3.9) | 0.10 (−0.25, 0.45) | 0.551 | 0.16 |
| Fat mass (kg) | 25.0 (6.7) | 24.6 (6.6) | 0.37 (−0.19, 0.94) | 0.180 | 0.36 |
| Fat mass (%) | 38.2 (6.0) | 37.6 (5.9) | 0.52 (−0.12, 1.16) | 0.105 | 0.45 |
| Lean mass left leg (kg) | 5.6 (1.0) | 5.7 (1.1) | −0.18 (−0.31, −0.05) | 0.012 | 0.74 |
| Lean mass right leg (kg) | 5.8 (1.1) | 6.0 (1.2) | −0.25 (−0.91, −0.11) | 0.002 | 1.01 |
| Lean mass (kg) | 37.7 (5.6) | 37.9 (5.9) | −0.25 (−0.91, 0.39) | 0.410 | 0.22 |
| Waist circumference (cm) | 83.8 (9.5) | 82.4 (9.1) | 1.41 (−0.23, 3.07) | 0.086 | 0.52 |
| Hip circumference (cm) | 100.7 (7.3) | 101.3 (7.4) | −0.60 (−2.90, 1.70) | 0.579 | 0.16 |

Data are mean ± SD. Significant *p* values are in bold. *BMI* body mass index, *CI* confidence interval, *ES* effect size, *SD* standard deviation

Table 4 Effects of exercise intervention on QLQ-BR23 scores ($n = 15$)

| Item | Baseline Mean (SD) | Post-intervention Mean (SD) | <i>p</i> value | <i>ES</i> |
|-------------------------------|--------------------|-----------------------------|----------------|-----------|
| Body image functioning | 61.1 (32.1) | 65.6 (32.3) | 0.549 | 0.16 |
| Sexual functioning | 75.6 (22.6) | 83.2 (24.5) | 0.213 | 0.35 |
| Sexual enjoyment | NR | NR | NR | – |
| Future perspective | 75.6 (29.4) | 45.3 (36.1) | 0.005 | 0.90 |
| Systemic therapy side effects | 70.0 (22.2) | 73.4 (22.0) | 0.596 | 0.14 |
| Breast symptoms | 76.1 (20.4) | 81.5 (20.5) | 0.710 | 0.10 |
| Arm symptoms | 67.0 (27.5) | 67.5 (32.2) | 0.536 | 0.17 |
| Hair loss | NR | NR | NR | – |

Data are mean ± SD. Significant *p* value are in bold. *Abbreviations*: *ES* effect size, *NR* not reported, *SD* standard deviation

The COVID-19 pandemic has impacted every aspect of life producing increased levels of sedentary time and reduced physical activity levels. These changes will be deleterious for the health of the global population and will be critical for cancer survivors. Consequently, home-based exercise interventions could be an excellent choice for people affected by cancer.

The use of e-Health in oncology has started a new era that goes beyond the traditional health care model [29]. Home-based exercise interventions have been shown as a valid strategy that can provide benefits in CRF, strength, PA levels, HRQoL, and body composition on cancer survivors [15, 30]. Results obtained in the present study show that an online home-based exercise intervention performed in a context of lockdown enables the maintenance of physical function in female breast cancer survivors (our participants improved the score obtained in the chair stand test, that measures the ability of participants to stand and sit on a chair a task that everyone develops daily). Our results are in overall agreement and extend those of DeNysschen et al. [14], who with an 8-week home-based exercise intervention applied in female breast cancer survivors

($n = 26$), who were receiving prolonged hormonal treatment, showed improvements in handgrip strength, arm curl test and chair stand test, but not in CRF [14]. The differences in CRF between studies can be attributed to both the duration of the exercise training program (8 weeks vs 16 weeks) and the test used to assess the CRF (3 min step test vs Rockport test).

It is important to understand that a reduction of muscle mass is currently considered a prognostic factor of the disease and is associated with a higher degree of functional limitations and dependency, and with lower cancer survival rates [31, 32]. Several studies have demonstrated a pronounced muscle strength decline (upper- and lower-body) in breast cancer survivors [32–34]. Nowadays, handgrip strength is known as one of the prognostic factors of sarcopenia [35, 36]. Focusing on breast cancer, handgrip strength has been directly associated with HRQoL and inversely with the adverse secondary effects of cancer treatment [37]. Our results, similarly to those of Denyssen et al. [14], have shown handgrip strength improvements, with an increase of the right arm (19.7 kg pre- vs. 21.7 kg post- intervention). Also, in a recent case study developed during the COVID-19 lockdown, one of the two

participants included in the study showed improvements in handgrip strength (27.4 kg pre- vs. 34.9 kg post-intervention) after a 16-week exercise intervention [38].

Functional training in cancer patients and survivors is an emerging topic. For years, most researchers have focused on determining the effectivity of exercise programs based on traditional strength exercises. Jones et al. [39], in a recent study developed in breast cancer survivors ($n = 51$), implemented a 12-week intervention based on a combined aerobic and resistance circuit, observing positive changes in physical fitness (CRF and upper and lower limbs strength), similar to those found in our study.

Regarding body composition, Lahart et al. [40] in a 6-month home-based PA intervention in breast cancer survivors found significant improvements in body weight and BMI. However, in our study only changes on legs lean mass from baseline to the end of the online program were detected. A possible reason for these differences between studies could be due to the duration of the studies (16 weeks vs 6 months). Moreover, the method to assess body composition was not the same (dual-energy X-ray absorptiometry vs bioelectrical impedance). It is also important to note that our study was performed in a context of COVID-19 lockdown in which people have been forced to stay at home and, consequently, probably had to modify their lifestyle [2, 41, 42].

Cancer patients express clinically increased psychological symptoms (depression and anxiety) during the disease [43] and an online intervention could be a viable strategy to improve HRQoL and motivation towards exercise and, consequently, reduce these symptoms [15]. The results of the present study obtained with the EORTC-QLQ-BR23 did not show a positive effect of exercise on body image or secondary treatment effects. Surprisingly, we found a significant decrease in future perspective scores. This could be partially explained by the confinement that all participants had to suffer while the intervention took place, a situation that could generate uncertainty, anxiety, or fear to the near future. In the same line, a recent study demonstrated that the COVID-19 pandemic negatively affected the emotional health of women diagnosed with primary breast cancer who showed higher general emotional vulnerability due to the disruption of treatments generated by COVID-19 [44].

There are some limitations in this study that should be noted. First, our sample size was limited. Furthermore, no sample size calculation was performed, as the cohort analyzed in the present study was a convenience sample (i.e., all participants belonged to the AMACGEMA meeting the eligibility criteria). However, it is important to note that this study has been undertaken thought special circumstances due to the COVID-19 pandemic. Therefore, it was extremely difficult to have a greater number of participants. Second, the absence of a control group did not allow to establish a solid relationship of these findings. However,

given that physical exercise in breast cancer survivors is highly recommended because of its health benefits, and the circumstances of pandemic and confinement made it highly likely that regular physical activity would decrease, we considered that it was unethical to leave a group without a training program. Another limitation is the lack of control on nutritional variables or dietary habits, which can largely influence body composition.

Nonetheless, the present study has several strengths such as the possibility of adapting the study methodology to the circumstances of the COVID-19 pandemic and the associated lockdown. Also, noteworthy is the high adherence of participants to the online home-based exercise intervention under these circumstances.

Conclusion

This feasibility study suggests that an online home-based exercise intervention performed during a confinement period is safe and beneficial for breast cancer survivors who showed improvements in physical fitness and lower limbs lean mass. Further larger studies (particularly, controlled trials) are needed to evaluate which is the best intervention to improve both physical and psychological health-related markers.

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Data availability All data that support the findings of this study are available from the corresponding author upon reasonable request. Source data are provided with this paper.

Code availability Not applicable.

Declarations

Ethics approval The present project was carried out in accordance with the principles of the Declaration of Helsinki and Good Clinical Practice guidelines. The project was approved by the Ethical Committee of the University of San Jorge, Spain (n°006-19/20).

Consent to participate Not applicable.

Consent for publication Not applicable.

Conflicts of interests The authors declare that they have no conflict of interest.

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