

# Program to improve the quality of life due to unwanted loneliness in widowed women. Nature, an ally to accompany you

<b>Submission date</b> 10/07/2024	<b>Recruitment status</b> No longer recruiting	<input type="checkbox"/> Prospectively registered <input checked="" type="checkbox"/> Protocol
<b>Registration date</b> 12/02/2025	<b>Overall study status</b> Completed	<input type="checkbox"/> Statistical analysis plan <input type="checkbox"/> Results
<b>Last Edited</b> 03/03/2026	<b>Condition category</b> Mental and Behavioural Disorders	<input type="checkbox"/> Individual participant data <input checked="" type="checkbox"/> Record updated in last year

## Plain English summary of protocol

### Background and study aims

This project aims to combine tourism with the unique features of the Jerte Valley as a therapy to combat loneliness. It involves a series of nature-based activities that will position the Jerte Valley as a specialized tourist destination in therapeutic tourism, understood as tourism that aims to improve and maintain general health and specifically address loneliness, enhancing aspects that are impacted by this situation, such as quality of life, self-concept, resilience, etc. Particularly, this project addresses the issue of unwanted solitude among elderly widowed women, recognizing solitude in women as one of the social problems with the highest incidence and risk of social isolation, worsening the morbidity and mortality among women. Through a program of nursing interventions with other health agents and specialists in loneliness, we have developed a multi-component program based on therapeutic tourism. The main objectives of the presented project are to conduct a pilot experience of therapeutic tourism in the Jerte Valley and to analyse the efficacy of a multi-component program that combines nursing interventions, support from loneliness specialists, social anthropology, and psychology, along with nature activities, to foster socialization and improve the quality of life for these women. Lastly, the expected results of the program are the implementation of a pilot experience of therapeutic tourism in the Jerte Valley that could be generalizable to other vulnerable groups requiring special attention to the recovery of socialization. The study will involve the dissemination, in scientific, and cultural forums, and to the national loneliness observatory, of the program's efficacy.

### Who can participate?

Women between the ages of 60 and 85 years old who live independently in the community and have feelings of loneliness

### What does the study involve?

This study will:

SO1: Evaluate the variables of loneliness, despair, and quality of life before the intervention of the control group and the experimental group to establish the baseline.

SO2: Nursing interventions with emotional support in the experimental group over 6 weekends,

with the first four weekends being consecutive, taking a break on the fifth weekend, and then resuming for two more consecutive weekends, totaling 48 hours of interventions received by the end of the program.

SO3: Assess the variables of loneliness, despair, and quality of life after the completion of the intervention program.

SO4: Evaluate the intervention variable in both groups (control and experimental) one month after the intervention's completion to analyze the long-term maintenance of the results.

SO5: Disseminate the pilot experience of therapeutic tourism in the Jerte Valley through Conferences on therapeutic tourism in the Jerte Valley and the prevention of unwanted loneliness, reporting of scientific articles, and a documentary of the experience.

What are the possible benefits and risks of participating?

Benefits include being able to socialize as well as create support networks, and tools to combat loneliness and increase the quality of life. The study does not involve risks, the accompaniment of these women is fundamental, the group of professionals who will develop the interventions are trained to accompany and resolve any situation that may arise.

Where is the study run from?

University of Extremadura

When is the study starting and how long is it expected to run for?

January 2024 to September 2024

Who is funding the study?

The Diputación de Cáceres

Who is the main contact?

Belinda Basilio Fernández, [bbasfer@unex.es](mailto:bbasfer@unex.es)

## Contact information

### Type(s)

Public, Scientific, Principal investigator

### Contact name

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## Additional identifiers

## Clinical Trials Information System (CTIS)

Nil known

### Protocol serial number

Protocolo 1

## Study information

### Scientific Title

A multicomponent program to improve the quality of life of widowed women with unwanted loneliness: a study protocol for a pilot randomized controlled trial

### Study objectives

Multicomponent intervention to improve the quality of life in widowed women with unwanted loneliness

### Ethics approval required

Ethics approval required

### Ethics approval(s)

approved 21/03/2024, Bioethics and Biosecurity Commission of The University of Extremadura (Comisión de Bioética y Bioseguridad de la Universidad de Extremadura) (C/ Elvas s/n., Badajoz, 06006, Spain; +34 924289300; irocha@unex.es), ref: 46/2024

### Study design

Prospective cohort randomized controlled trial (RCT) with a mixed approach

### Primary study design

Interventional

### Study type(s)

Quality of life

### Health condition(s) or problem(s) studied

Quality of life related to unwanted loneliness in widows

### Interventions

This study will be carried out in the province of Cáceres, taking the population of widowed women. The sample size has been calculated using data from a previous study considering the mean and standard deviation of the UCLA Loneliness Scale-Revised standard deviation of the UCLA Loneliness Scale-Revised (UCLA LS-R) in women over 65 years of age. Because no studies defined the minimum clinical change it has been set at 4 points, it would require a minimum of 25 participants per group (50 in total) with a power of 80%, an  $\alpha$  value of 0.05 and an estimated loss rate of 20%. To perform this calculation, an online sample calculator an online sampling calculator was used to perform this calculation. A member of the research team who will not be directly involved in the trial will create a simple computer-generated randomization sequence using Research Randomizer software (Version 4.0, Urbaniak GC and Plous S., Middletown, CT, USA; <http://www.randomizer.org>). The assignment will be hidden with a password-protected file. Researchers involved in the data analysis processes will not know to which group each woman will be assigned (experimental or control).

## Intervention

Both the experimental group (regular users + multicomponent program) and the control group (regular users) will undergo pre-intervention, post-intervention and follow-up phases for 1 month. The difference between the two groups lies in the fact that the participants in the experimental group will be the only ones who will undergo the intervention based on a multicomponent psychosocial program.

The intervention program will be composed of 6 sessions that will be carried out on 4 continuous weekends beginning in April. They will consist of outings to the Jerte Valley to participate in therapeutic tourism. The sessions will be divided into a therapeutic component (first part of the session) with at least two members of the team and up to 4 maximum, who are professionals in the field of nursing, with experience in the field of bereavement and work with women. The second component of the session (second part of the part of the session) will be focused on the realization of sports in nature led by experts in the field and supported by the team and its accompaniment.

## Intervention Type

Behavioural

## Primary outcome(s)

The following primary outcome measures will be assessed before and after the intervention (in the control group, at the beginning and the end of the program):

1. Sociodemographic variables (age, educational level, socioeconomic status, whether living alone, etc.) measured using a sociodemographic data sheet during the study
2. Experiences regarding loneliness and the experience of the intervention measured using semi-structured interviews
3. Feeling of loneliness measured using a focus group with the experimental group
4. Loneliness measured using the University of California Loneliness Scale
5. Health status and well-being measured using the EQ5-5D-5L HRQoL (health-related quality of life) measurement instrument
6. HRQoL measured using the SF-12
7. HRQoL measured using the 5-D
8. HRQoL measured using the AQOL-8D
9. Hopelessness measured using the Beck Hopelessness Scale
10. Depression measured using the Yesavage Depression Scale
11. Physical activity during the last 7 days measured using the International Physical Activity Questionnaires (IPAQ)

## Key secondary outcome(s)

Interpretation of the participant's change in quality of life will be measured using the interview transcriptions during the study's qualitative research

## Completion date

30/09/2024

## Eligibility

### Key inclusion criteria

1. Be a woman between the ages of 60 and 85 years old
2. Living independently in the community
3. Feelings of loneliness

**Participant type(s)**

Patient

**Healthy volunteers allowed**

No

**Age group**

Mixed

**Lower age limit**

60 years

**Upper age limit**

85 years

**Sex**

Female

**Total final enrolment**

0

**Key exclusion criteria**

1. Comorbidity with other mental health problems such as severe mental disorders (schizophrenia or bipolar disorders), personality disorders, dementias, and major depressions
2. Comorbidity of physical problems that prevent them from performing adventure activities in nature, such as severe musculoskeletal injuries
3. Women who do not know the Spanish language well enough to hold a conversation

**Date of first enrolment**

26/02/2024

**Date of final enrolment**

21/03/2024

**Locations****Countries of recruitment**

Spain

**Study participating centre**

University for Seniors - Democratic Union of Pensioners- Jerte Valley Commonwealth- Ambroz Valley Commonwealth - Vera Commonwealth  
Avenida Virgen del Puerto,2

Plasencia  
Spain  
10600

## Sponsor information

### Organisation

University of Extremadura

### ROR

<https://ror.org/0174shg90>

## Funder(s)

### Funder type

Government

### Funder Name

Diputación Provincial de Cáceres

## Results and Publications

### Individual participant data (IPD) sharing plan

### IPD sharing plan summary

Data sharing statement to be made available at a later date

### Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
<a href="#">Protocol article</a>		24/01/2026	03/03/2026	Yes	No
<a href="#">Participant information sheet</a>			11/07/2024	No	Yes