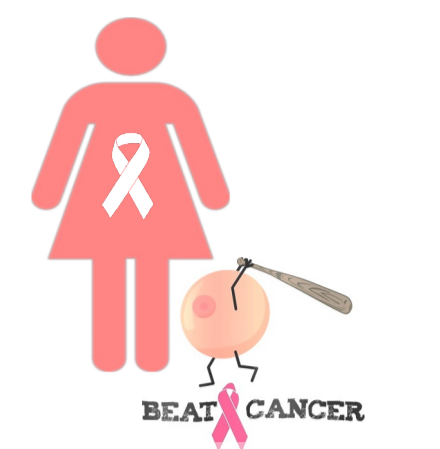




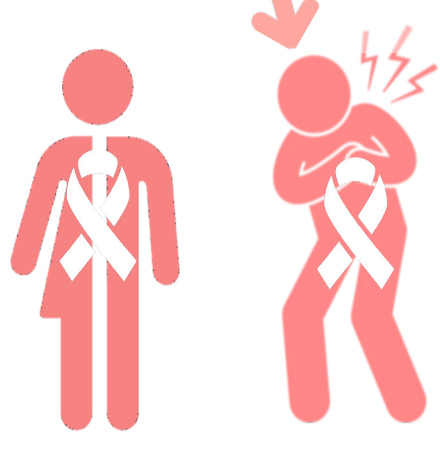
THE EFFECT OF PAIN NEUROSCIENCE EDUCATION AND BEHAVIOURAL GRADED ACTIVITY IN BREAST CANCER SURVIVORS: A RANDOMISED CONTROLLED MULTI-CENTER TRIAL

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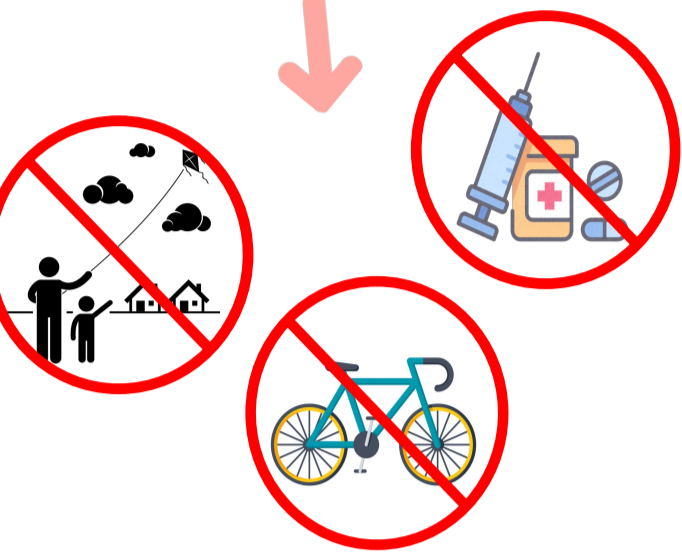
BACKGROUND



The survival rate of **Breast Cancer Survivors (BCS)** is rising up to **89.9%**



Debilitating side-effects can persist in BCS, of which Chronic pain (CP) is one of the most prevalent ones. Affecting 1 out of 2 BCS after treatment.



Pain reduces HRQoL and activity levels and pain medication has side-effects on the long term.



There is a need for non-pharmacological treatments, such as **Pain Neuroscience Education** (i.e., explaining the neurophysiology of pain), combined with **Behavioural Graded Activity** (i.e., increasing the patient's meaningful activities)

AIM

THE PRIMARY OBJECTIVE: Investigate whether PNE with BGA has an added value in **decreasing pain** compared to the usual care in BCS with chronic pain.

THE SECONDARY OBJECTIVES: Investigate whether PNE with BGA has the ability to **improve endogenous pain modulation and HRQoL** compared to the usual care in BCS with chronic pain.

METHODS

INCLUSION

- ✓ Breast Cancer Survivor
- ✓ Pain Visual Analogue Scale $\geq 3/10$
- ✓ Cancer Free
- ✓ Treatment completed $\geq 3m$
- ✓ Dutch reading and speaking

OUTCOME MEASURES

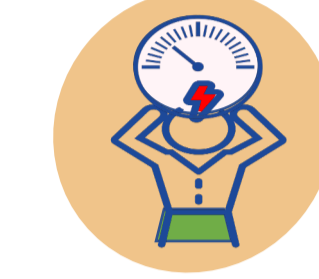
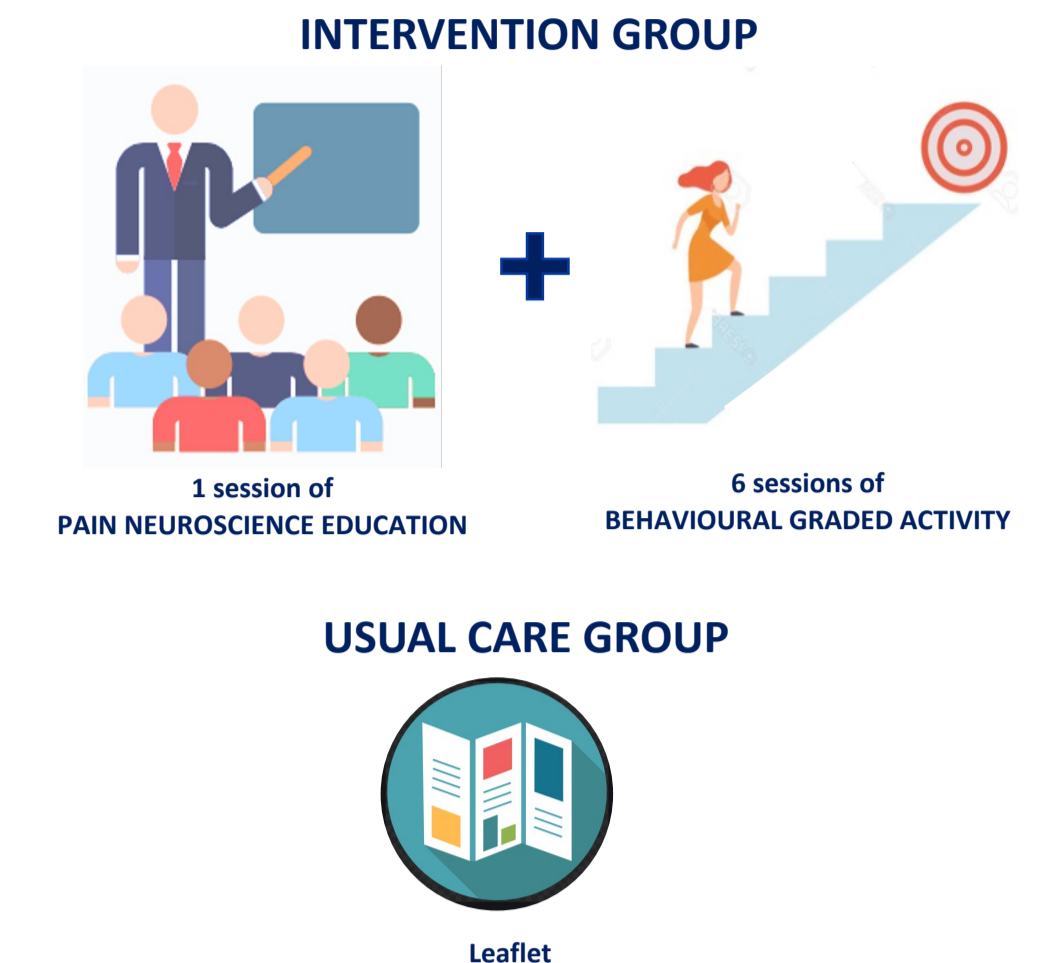


Brief Pain Inventory - BPI
European Organization for Research and Treatment of Cancer Quality of Life Questionnaire Core 30 - EORTC-QLQ-C30
 Douleur Neuropathique 4 - DN4
 Central Sensitisation Index - CSI
 International Physical Activity Questionnaire - IPAQ-SF
 Productivity Cost Questionnaire - PCQ
 Medical Cost Questionnaire - MCQ
 Pittsburgh Sleep Quality Index - PSQI
 Insomnia Severity Index - ISI
 Fatigue Severity Scale - FSS
 Pain Catastrophizing Scale - PCS
 Injustice Experience Questionnaire - IEQ
 Pain Vigilance and Awareness - PVAQ
 Depression, Anxiety and Stress Scale - DASS-21

INTERVENTION



n = 62
RANDOMISED
n = 60



Endogenous Pain Modulation
 Performed on three locations:
M. Pectoralis, *Most painful area* and *M. tibialis anterior*
Pain and Detection Thresholds:
 Pressure (algometer and cuff), Warmth and Cold
Temporal Summation:
 Pressure (algometer)
Conditioned Pain Modulation: Pressure (cuff)
 Stimuli - Pressure (algometer), Warmth and Cold

STATISTICAL ANALYSIS

Linear mixed models for repeated measures in SPSS

RESULTS



PRIMARY OUTCOME MEASURES - PAIN (BPI (0-10))

Measure	Post-Intervention	3 Months	12 Months
Pain severity	↑	↑	↑
Pain interference	↑	↑	↑
Worst pain	↑	=	=

SECONDARY OUTCOME MEASURES - HEALTH-RELATED QUALITY OF LIFE (EORTC-QLQ-C30 (0-100))

Measure	Post-Intervention	3 Months	12 Months
HRQoL	↑	↑	↑

SECONDARY OUTCOME MEASURES - ENDOGENOUS PAIN MODULATION

Measure	Post-Intervention	3 Months	12 Months
Cuff	=	=	=
Pressure pain thresholds	=	=	=
Warmth pain thresholds	=	=	=
Cold pain thresholds	=	=	=
Conditioned pain modulation	=	=	=
Pressure (Δkgf)	=	=	=
Warmth ($\Delta ^\circ C$)	=	=	=
Cold ($\Delta ^\circ C$)	=	=	=
Temporal Summation (ΔVAS)	=	=	=

EXPLANATORY OUTCOME MEASURES

Measure	Post-Intervention	3 Months	12 Months
Central sensitization (CSI (0-100))	↑	=	=
Neuropathic symptoms (DN4 (0-10))	=	=	=
Perceived injustice (IEQ (0-48))	↑	=	↑
Pain catastrophizing (PCS (0-52))	↑	↑	↑
Pain vigilance and awareness (PVAQ (0-80))	↑	↑	=
Stress (DASS21 Stress (0-21))	=	=	=
Anxiety (DASS21 Anxiety (0-21))	=	=	=
Depression (DASS21 Depression (0-21))	=	=	=
Physical activity (IPAQ (METs))	↑	=	=
Fatigue (FSS (1-7))	=	=	=
Sleep (PSQI (0-21))	=	=	=
Insomnia (ISI (0-28))	=	=	=

TABLE LEGEND

↑ improvement in favour of the PNE with BGA

Cohen's d effect sizes are interpreted as

↑↑↑ Large (0.80-1.29)

↑↑ Medium (0.50-0.79)

↑ Small (0.20-0.49)

= Negligible (<0.20)

Significant values ($p < 0.05$) are in GREEN

PROTOCOL REGISTRATION



CONCLUSION

PNE with BGA did result in a significant short-term reduction in **pain severity and interference** compared to usual care in BCS with chronic pain. Additionally, significant improvements in **maladaptive cognitions** were observed.

However, observed changes in pain **did not significantly improve patients' HRQoL and endogenous pain modulation.**