

# Analysis of the influence of electrostimulation and exercises on pelvic floor muscles tension in young, healthy women

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<b>Registration date</b> 25/07/2022	<b>Overall study status</b> Completed	<input type="checkbox"/> Statistical analysis plan <input type="checkbox"/> Results
<b>Last Edited</b> 25/07/2022	<b>Condition category</b> Other	<input type="checkbox"/> Individual participant data <input type="checkbox"/> Record updated in last year

## Plain English summary of protocol

### Background and study aims

The pelvic floor is an extremely important place in a woman's body. Like any other area of the organism, this one can also undergo various types of related dysfunctions with disturbed muscle tone present there. There are many factors that can contribute to the development of pelvic floor dysfunction, such as: physiological childbirth, menstrual cycle, previous lower urinary tract infections, comorbidities, medications taken, profession and many others. Therefore, it is necessary to look for therapeutic methods that will prove effective in strengthening the pelvic floor muscles in young, healthy women, and thus in the prevention of the occurrence of dysfunctions in the genitourinary system in the next stages of their lives.

The aim of the study is the electromyographic analysis of the effect of endovaginal electrostimulation and exercises on the pelvic floor muscle tone in young women, showing no dysfunction in the genitourinary system. The study will assess whether the applied physioprophyllactic procedures will be effective in increasing PFM tension, and thus in the prevention of dysfunction of this muscle group.

### Who can participate?

Women who have not previously given birth, aged 19-29 years, without pelvic floor dysfunction.

### What does the study involve?

Women will be randomly assigned to one of three groups: two test groups with pelvic floor electrostimulation and exercise, and a control group with exercise only. Bioelectrical activity of pelvic floor muscles are measured before and after treatment and during follow-up visits 1 and 3 months after the end of the study.

### What are the possible benefits and risks of participating?

Participants will receive a complete physioprophyllaxis program that can improve pelvic floor muscle tone. There is no risk involved in participating.

Where is the study run from?  
University of Opole (Poland)

When is the study starting and how long is it expected to run for?  
March 2022 to December 2023

Who is the main contact?  
Julia Duda  
julia.konrad@uni.opole.pl  
Prof. Jakub Taradaj  
j.taradaj@awf.katowice.pl

## Contact information

### Type(s)

Public

### Contact name

Mrs Julia Duda

### ORCID ID

<https://orcid.org/0000-0001-8094-5685>

### Contact details

68 Katowicka Street  
Opole  
Poland  
45-060  
+48 (0)788257367  
julia.duda@uni.opole.pl

### Type(s)

Scientific

### Contact name

Prof Jakub Taradaj

### ORCID ID

<https://orcid.org/0000-0002-1796-5832>

### Contact details

Academy of Physical Education  
72 Mikołowska Street  
Katowice  
Poland  
40-065  
+48 (0)668613945  
j.taradaj@awf.katowice.pl

## Additional identifiers

## Clinical Trials Information System (CTIS)

Nil known

## ClinicalTrials.gov (NCT)

Nil known

## Protocol serial number

KB-254/2022

# Study information

## Scientific Title

Electromagnetic electrostimulation and exercises in the physioprophylaxis of pelvic floor dysfunction

## Study objectives

1. Pelvic floor muscles exercises improve the bioelectrical activity of the pelvic floor muscles.
2. Endovaginal electrostimulation improve the bioelectrical activity of the pelvic floor muscles.
3. The frequency used in endovaginal electrostimulation affects the bioelectric activity of the pelvic floor muscles

## Ethics approval required

Old ethics approval format

## Ethics approval(s)

Approved 31/03/2022, Research Ethics Committee from Wroclaw Medical University (1 Pasteur Street, 50-367, Wroclaw, Poland; +48 (0)717841014; bioetyka@umed.wroc.pl), ref: KB-254/2022

## Study design

Prospective randomized clinical study with follow-up analysis

## Primary study design

Interventional

## Study type(s)

Prevention

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

Bioelectric activity of pelvic floor muscles

## Interventions

After baseline assessments, the participants will be randomly assigned (using an online tool) to:

1. Endovaginal electrostimulation using the 50 Hz frequency and pelvic floor muscles exercises
2. Endovaginal electrostimulation using the 10 Hz frequency and pelvic floor muscles exercises
3. Pelvic floor muscles exercises

The individuals receiving the treatment will be blinded. A computer-generated list of random numbers will be used and concealed from the researchers enrolling and assessing the participants. The outcome assessors and data analysts will be kept blinded to the allocation.

The therapy programme lasts 8 weeks during which all the participants will perform pelvic floor exercises at home 3 times a week. Women will exercise according to the schedule (3 series, 9 contractions in series, contraction time 6s, rest time 12s, 3 quick contractions at the end of each series (1,2,3s), a 2-minute recovery between sets). Additionally, in two groups, women will undergo endovaginal electrostimulation 3 times a week. A single treatment will take 20 minutes. In group 1, it is planned to use a frequency of 50 Hz and in the second group, a frequency of 10 Hz. The remaining treatment parameters for both groups will be identical (pulse time 200  $\mu$ s, contraction time 5s, rest time 10s, intensity to the patient's feelings).

All participants will undergo assessment of the bioelectrical activity of pelvic floor muscles before the start of treatment, after intervention and 1 and 3 months after the therapy programme.

## **Intervention Type**

Behavioural

## **Primary outcome(s)**

Bioelectrical activity of pelvic floor muscle measured using an endovaginal electrode, an EMG apparatus and integrated computer software. An assessment of the bioelectrical activity of pelvic floor muscle using the Glazer protocol. The measurements will be performed before the therapy, after the therapy, i.e. 8 weeks later and 4 and 12 weeks after the end of intervention

## **Key secondary outcome(s)**

Assessment of the bioelectric activity of the pelvic floor muscles using the glazer protocol:

1. One 60-second rest (pre-baseline) - the women were instructed to feel the pelvic floor in a resting position.
2. Five 2-second phasic (flick) contractions with a 2-second rest in-between - the women were instructed to contract the PFM as quickly as possible, and then quickly and fully relax the PFM immediately after contraction.
3. Five 10-second tonic contractions with a 10-second rest in-between - the women were instructed to contract the PFM as strongly as possible, maintain the contraction for 10seconds, and then fully relax the PFM after contraction, remaining relaxed for 10seconds.
4. One 60-second endurance contraction - the women were instructed to contract the PFM at such a level as to hold it for 60 seconds.
5. One 60-second rest (post-baseline) - the women were instructed to feel the pelvic floor in a resting position.

The measurements will be performed before the therapy, after the therapy and 1 and 3 months after the end of the intervention.

## **Completion date**

30/12/2023

## **Eligibility**

### **Key inclusion criteria**

1. Women aged 19 - 29 years
2. Nulliparous
3. Women who have already had intercourse
4. Women without pelvic floor dysfunction

**Participant type(s)**

Healthy volunteer

**Healthy volunteers allowed**

No

**Age group**

Adult

**Lower age limit**

19 years

**Upper age limit**

29 years

**Sex**

Female

**Key exclusion criteria**

1. Women before 19 and after 29 years of age
2. Women who gave birth
3. Virgins
4. Pelvic floor dysfunction
5. Sensory deficits
6. Pacemakers and other electronic implants
7. Metal implants
8. Psychiatric disorders
9. Cancer
10. Infections
11. Fever
12. Acute inflammation
13. Hypersensitivity to electricity

**Date of first enrolment**

08/08/2022

**Date of final enrolment**

30/12/2023

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

Poland

**Study participating centre**

University of Opole Institute of Health Sciences  
68 Katowicka Street

Opole  
Poland  
45-060

## Sponsor information

### Organisation

Opole University

### ROR

<https://ror.org/04gbpnx96>

## Funder(s)

### Funder type

University/education

### Funder Name

Uniwersytet Opolski

### Alternative Name(s)

University of Opole, Opole University, UO

### Funding Body Type

Government organisation

### Funding Body Subtype

Universities (academic only)

### Location

Poland

## Results and Publications

### Individual participant data (IPD) sharing plan

The datasets generated during and/or analysed during the current study are available from the corresponding author on reasonable request ([j.taradaj@awf.katowice.pl](mailto:j.taradaj@awf.katowice.pl))

### IPD sharing plan summary

Available on request