

# Clinical Trial Protocol

## Iranian Registry of Clinical Trials

09 Jun 2026

### The effect of Ex-PLISSIT model based sexual counseling on the intimacy and sexuality of married women with multiple sclerosis: a randomized controlled trial

#### Protocol summary

##### Summary

The aim of this study is to determine the effect of Ex-PLISSIT model based sexual counseling on the intimacy and sexuality of married women with multiple sclerosis. The participants are 42 married women aged 18 to 45 years with multiple sclerosis in Tabriz. They will randomly be allocated in two groups. In the intervention group, consultation will be held in one session for 90-60 minutes or more sessions if needed (i.e. intensive therapy) in accordance with the EX-PLISSIT model components. Control group will just receive routine care. The primary outcome are intimacy and sexuality before and two months after intervention measured by MSISQ-19.

#### General information

##### Acronym

##### IRCT registration information

IRCT registration number: **IRCT2017051833834N2**

Registration date: **2017-06-18, 1396/03/28**

Registration timing: **registered\_while\_recruiting**

Last update:

Update count: **0**

##### Registration date

2017-06-18, 1396/03/28

##### Registrant information

###### Name

Roghayeh Nourizadeh

###### Name of organization / entity

Tabriz University of Medical Sciences

###### Country

Iran (Islamic Republic of)

###### Phone

+98 41 3479 6770

###### Email address

nourizadehr@tbzmed.ac.ir

##### Recruitment status

**Recruitment complete**

##### Funding source

Tabriz University of Medical Sciences

##### Expected recruitment start date

2017-05-27, 1396/03/06

##### Expected recruitment end date

2017-10-22, 1396/07/30

##### Actual recruitment start date

empty

##### Actual recruitment end date

empty

##### Trial completion date

empty

##### Scientific title

The effect of Ex-PLISSIT model based sexual counseling on the intimacy and sexuality of married women with multiple sclerosis: a randomized controlled trial

##### Public title

The effect of Ex-PLISSIT model based sexual counseling on the intimacy and sexuality of married women with multiple sclerosis: a randomized controlled trial

##### Purpose

Treatment

##### Inclusion/Exclusion criteria

Inclusion criteria: The minimum literacy of secondary school; At least 6 months after diagnosis of MS; sexually active married women; women aged 18-45 years; Ability of listening and speaking; At least one year after marriage; Having a contact number to follow-up; Without major disability (EDSS <7); Exclusion criteria: Pregnant and lactating women; Hospitalization due to MS disease progression or relapse within the last month; Presence of other diseases affecting sexual function; Drug addiction and smoking; Using drugs affect sexual function; Being

treatment due to sexual dysfunction; Women with depression score  $\geq 16$  in short form of the Beck Depression Inventory

**Age**

From **18 years** old to **45 years** old

**Gender**

Female

**Phase**

N/A

**Groups that have been masked**

*No information*

**Sample size**

Target sample size: **42**

**Randomization (investigator's opinion)**

Randomized

**Randomization description****Blinding (investigator's opinion)**

Not blinded

**Blinding description****Placebo**

Not used

**Assignment**

Parallel

**Other design features****Secondary Ids**

empty

**Ethics committees****1****Ethics committee****Name of ethics committee**

Tabriz University of Medical Sciences (Nursing and Midwifery)

**Street address**

Azady Street / Golgasht (South Shariati)

**City**

Tabriz

**Postal code****Approval date**

2017-05-08, 1396/02/18

**Ethics committee reference number**

IR.TBZMED.REC.1396.115

**Health conditions studied****1****Description of health condition studied**

Multiple sclerosis

**ICD-10 code**

G35

**ICD-10 code description**

Multiple sclerosis

**Primary outcomes****1****Description**

Intimacy and sexuality

**Timepoint**

Before and two months after intervention

**Method of measurement**

MSISQ-19

**Secondary outcomes****1****Description**

Quality of sexual life

**Timepoint**

Before and two months after intervention

**Method of measurement**

SQOL-F

**Intervention groups****1****Description**

In the intervention group (21 women), consultation will be held in one session for 90-60 minutes or more sessions if needed (i.e. intensive therapy) in accordance with the model components.

**Category**

Treatment - Other

**2****Description**

In the usual care control group will be

**Category**

Treatment - Other

**Recruitment centers****1****Recruitment center****Name of recruitment center**

MS Society of East Azerbaijan

**Full name of responsible person**

Roya Azari Barzandig

**Street address****City**

Tabriz

**Sponsors / Funding sources****1****Sponsor****Name of organization / entity**

Tabriz University of Medical Sciences

**Full name of responsible person**

Dr. Mohammad Reza Rashidi

**Street address**

Vice President of Research, Tabriz University of  
Medical Sciences

**City**

Tabriz

**Grant name**

**Grant code / Reference number**

**Is the source of funding the same sponsor  
organization/entity?**

Yes

**Title of funding source**

Tabriz University of Medical Sciences

**Proportion provided by this source**

100

**Public or private sector**

*empty*

**Domestic or foreign origin**

*empty*

**Category of foreign source of funding**

*empty*

**Country of origin**

**Type of organization providing the funding**

*empty*

**Person responsible for general inquiries**

**Contact**

**Name of organization / entity**

Tabriz University of Medical Sciences

**Full name of responsible person**

Roya Azari Barzandig

**Position**

MSc of Counseling in Midwifery

**Other areas of specialty/work**

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Dr. Roghaiyeh Nourizadeh

**Position**

PhD in reproductive health

**Other areas of specialty/work**

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Roya Azari Barzandig

**Position**

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**Other areas of specialty/work**

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**Web page address**

**Sharing plan**

**Deidentified Individual Participant Data Set (IPD)**

*empty*

**Study Protocol**

*empty*

**Statistical Analysis Plan**

*empty*

**Informed Consent Form**

*empty*

**Clinical Study Report**

*empty*

**Analytic Code**

*empty*

**Data Dictionary**

*empty*