Effectiveness of Menosan® *Salvia officinalis* in the treatment of a wide spectrum of menopausal complaints. A double-blind, randomized, placebo-controlled, clinical trial

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**ARTICLE INFO**

**Objective:** To evaluate efficacy and safety of fresh *Salvia officinalis* extract tablets in relieving typical symptoms in menopausal women and to gain insight in the mode of action by measuring altered cerebral wave intensities.

**Methods:** Randomized 80 menopausal women from 48 – 65 years of age received Menosan® tablets [3.400 mg ethanolic extract of freshly harvested *Salvia officinalis* L.] or placebo under double-blind conditions for 4 weeks. An efficacy analysis evaluated the developments of the menopausal rating scale [MRS], hot flush severity score [HFS] and quantitative electroencephalography [qEEG] intensities in the per protocol population. Results were further corroborated by data from the intention to treat population including late postmenopausal women.

**Results:** Salvia off. distinctly reduced MRS by 39.2% from 15.3/6.87 to 9.3/5.75 and significantly in comparison to placebo (p = 0.002). The HFS score decreased by 55.3% from 15.9/13.77 to 7.1/7.41, reaching significance on week 3 onwards (p = 0.028). Clinical effects of Salvia off. correlated with relevant reduction of frontal lobe beta2 wave qEEG intensities at electrodes F3/4/7/8 and are underpinned by secondary parameters and ITT analysis. Salvia off. within 4 weeks significantly reduced the somato-vegetative (e.g. hot flushes) and psychological MRS subscale (e.g. physical and mental exhaustion) subscale (p < 0.05) without a significant effect on the genito-urinary subscale. A positive impact of Salvia off. compared to placebo was furthermore seen on sleep quality, discontent and fatigue (p < 0.05) as evidenced by sleep and profile of mood state questionnaires. Tolerability was uniformly rated as very good for Salvia off. extract and placebo, with an overall incidence of three adverse events in total, none of which treatment-related.

**Conclusion:** The results support the use of Salvia off. for the specific treatment of a wide range of somato-vegetative and psychological symptoms as experienced by menopausal women and correlate this effect to a restoration of associated dysbalanced brain waves.

The study was registered as EudraCT-No 2016-005033-77.

1. **Introduction**

A majority of perimenopausal women complain about challenging vasomotor symptoms like hot flushes, as well as fatigue, cognitive and mood alterations. It is widely known that estrogens neuro-modulate central neural circuits and exert an intricate influence on thermoregulation where the loss of a reliable estrogen rhythm weakens temperature homeostasis via dysmodulation of serotonergic and noradrenergic receptors (Rossmannith and Ruebberdt, 2009). Whilst standard therapy is still considered to be hormone replacement [HRT], its time restriction of use and association with a wide range of side effects including tumor growth, lead many women to seek effective and safe alternatives. *Salvia officinalis* has been shown to be depending on mechanisms of action other than estrogenicity and to regulate neurotransmitters implicated in hot flushes as well as state of mood and cognition. *Salvia officinalis* in the form of a hydroalcoholic extract has demonstrated inhibitory action on serotonergic, adrenergic, muscarinic and μ-opioid receptors in-vitro (Tober and Schoop, 2019).

*Salvia officinalis* furthermore has traditionally been used for the treatment of hyperhidrosis and recent studies indicate wider benefits. Hence, the
present study aimed to provide placebo controlled efficacy confirmation for Salvia off. in the treatment of climacteric symptoms as observed in a prior uncontrolled study (Bommer et al., 2011), and to confirm in-vitro data concerning neurotransmitter activity targeted by Salvia off. by testing associated brain waves. Quantitative electroencephalograms already explored pharmacological effects through neuro-modulation to allow for a correlation of brain activity changes with menopausal clinical symptoms (Staikou et al., 2012; Campbell et al., 2011). Based on these findings and the in vitro data obtained for the specific Salvia off. extract under investigation, examination of brain waves under therapy have been undertaken to elucidate the mode of action of Salvia off.

Altered cerebral beta2 wave intensities at F3/4/7/8 positions had been proposed as physiological markers for distinction of pre-from menopausal and early postmenopausal women, while excluding late postmenopausal women. Testing a putative adaptogenic effect of Salvia off. by modulating brain wave dysregulation associated with the menopausal transition as seen most prominently in perimenopause and especially early postmenopause, we aimed to investigate efficacy of a proprietary Salvia off. extract in a confirmatory efficacy analysis in this latter group (per protocol, PP) to more specifically address benefits of Salvia off. for typical menopausal ailments. Results were finally backed up with data including women in the late postmenopause with additional impact of age-related symptomatology to test for generalizability of treatment benefits (intention to treat, ITT). The definition of the per protocol group essentially correlates with the age definitions provided by the Staging of Reproductive Aging Workshop/STRAW (Frisancho et al., 2017), establishing the marking line between early and late postmenopause at 5 years after the final menstrual period (Harlow et al., 2012).

2. Materials and methods

This double-blind, randomized and controlled clinical study (EudraCT-No 2016-005033-77) was authorized by the Federal Institute for Drugs and Medical Devices (BfArM), Bonn/Germany, on April 18, 2017, approved by the ethics committee of the Medical Association of Hessen (Landesärztekammer) on April 27, 2017, and conducted at NeuroCode, Wetzlar/Germany in accordance with the Declaration of Helsinki.

Verum tablets Menosan® (Swiss authorization no. 61664, held by A.Vogel AG, Roggwil/Switzerland) contained 280 mg thujone-free spiss extract (Salvia off. Lamiaceae) equivalent to 3400 mg tincture. Inert placebo tablets were identically sized and colored to match verum tablets.

The randomization list was prepared by Hans Carlos Hoffmann (Asslar-Berghausen/Germany), using Kernel’s random number generator as entropy source with a block size of 8. Sealed emergency envelopes containing the randomization codes were retained by the statistician and by the investigator.

Informed consent was obtained from patients screened for eligibility by the following inclusion criteria: women between 48 and 65 years old, climacteric since ≥6 months, climacteric symptoms confirmed by the Menopause Rating Scale (MRS) total score ≥10, at least 5 hot flushes/24 h or intense hot flushes daily, disturbing sweating (Visual Analogue Scale [VAS] 0–10) and inconspicuous examination (i.e. no study relevant pathological findings, acute or chronic diseases or allergies). Exclusion criteria were ovarianectomy, chronic resistance to therapy in the indication area, epilepsy or severe depression (Hamilton Depression rating scale [HAMD] ≥30), anamnestic or current alcohol, drug, nicotine, coffee or tea abuse, intake of clinically relevant medications during the last two weeks before screening as well as during the active study period, particularly of Hormone Replacement Therapy [HRT], menopause- or cognitive performance-related herbal preparations and medication with central nervous action (e.g. psychotropics or antiepileptics), participation in another clinical study within the last 60 days, Body Mass Index/BMI ≥17.9 or ≤35.1, pregnancy, lactation, negative or invalid menopause quick test, noncompliance and cancellation of informed consent.

On screening day -7, demographic data were obtained and eligible patients were allocated to Salvia off. or placebo groups according to randomization list and in ascending order of patient’s appearance. Vital signs were checked during clinical examination, a 12 channel qEEG was performed and samples were collected for laboratory tests (hematology, urinalysis). Quick tests for drugs, alcohol (repeated on day 28), menopause and absence of pregnancy were performed. During the following 7 day baseline and throughout the four treatment weeks, participants recorded hot flushes’ number and intensity (ranging from 1 to 4 according to the definition of Sloan et al., 2001) and yielding by multiplication the Hot flush severity [HFS] score. First exposure to study medication was on day 0 under supervision when also different questionnaires were completed and participants underwent extensive qEEG examination before and after intake of study medication. Completed was the Menopause Rating Scale [MRS] calculated as the sum of the scores of the somato-vegetative subscale with the 4 symptoms hot flushes, heart discomfort, sleep problems, and joint and muscular discomfort, the Menopause- or cognitive performance-related herbal preparations and medication with central nervous action, in sexual and mental exhaustion, and the genitourinary subscale with the 3 symptoms sexual problems (change in sexual desire, in sexual activity and satisfaction), bladder problems, and vaginal dryness, furthermore the Hamilton Anxiety Rating Scale [HAMA], Profile of Mood State [POMS], and sleep quality questionnaire [SF-B/R]. The same qEEG and filling of questionnaires were repeated on day 28 after self-recorded daily intake of study medication. Returned tablets were counted to measure compliance. A closure visit with assessment of physical and safety parameters was conducted within 2 days after day 28 (Figure 1).

Estimation of the required sample size was established using archived qEEG data of pre-menopausal women free of complaints. The spectral power of frontal beta2 waves recorded ‘eyes open’ at electrodes F3/4/7/8 (Figure 2) of menopausal women was found to be related to hot flushes (Tegeler et al., 2015) and significantly higher than for fertile women aged 38 to 47. A 50% reduction of the spectral power by verum...
In total 80 Caucasian women suffering from menopause distress were enrolled (safety population) and randomly allocated to the *Salvia off.* or placebo treatment group. Two patients dropped out from ITT analysis with lacking efficacy data. 38 and 40 patients in *Salvia off.* and placebo groups, respectively, completed the study. Threof 48 menopausal patients aged 48 to 56 (mean 53 ± 2.19 years) met the PP analysis criteria and were applicable for PP analysis. The safety collective encompassed all patients with administration of at least one tablet of the study medication.

**3. Results**

In total 80 Caucasian women suffering from menopause distress were enrolled (safety population) and randomly allocated to the *Salvia off.* or placebo treatment group. Two patients dropped out from ITT analysis with lacking efficacy data. 38 and 40 patients in *Salvia off.* and placebo groups, respectively, completed the study. Threof 48 menopausal patients aged 48 to 56 (mean 53 ± 2.19 years) met the PP analysis criteria and were applicable for PP analysis. The safety collective encompassed all patients with administration of at least one tablet of the study medication.

Baseline demographics (age, BMI, body weight and height) were comparable for *Salvia off.* and placebo groups. Table 1a, b displays PP and ITT population. Overall compliance was 97.2 % for *Salvia off.* and 96.1% for placebo, respectively.

The primary efficacy analysis focused on the course of the clinical MRS and HFS scores under treatment. Table 2 shows a significant decrease of the MRS score for *Salvia off.* compared to placebo with a 39.2% decrease from 15.3 ± 6.87 to 9.3 ± 5.75 for *Salvia off.* and from 14.5 ± 6.04 to 13.0 ± 6.35 for placebo (p = 0.002) from day 0 to day 28. A very similar and significant reduction from 14.76 ± 6.18 to 11.13 ± 6.09 was seen for *Salvia off.* in the ITT analysis (p = 0.012) in contrast to placebo with a reduction from 14.63 ± 6.33 to 12.13 ± 5.86.

Figure 4 shows the courses of the HFS score for the two treatment groups. The obvious reduction of baseline HFS in favor of *Salvia off.* reaches statistical significance from week 3 onwards (p = 0.028) with a highly distinct reduction by 55% for *Salvia off.* from 15.9 ± 13.77 to 7.1 ± 7.41 and from 14.1 ± 9.33 to 10.3 ± 9.97 for placebo within four weeks, respectively. The percentage of patients, which experienced a clinically relevant, at least 50 % HSF score reduction from screening to week 4 was significantly higher for *Salvia off.* compared to placebo with 59.1% versus 26.9% respectively (p = 0.024). Results were again confirmed for the ITT collective which showed a clinically relevant ≥50 % reduction of the HSF score for 55% of *Salvia off.* patients compared to 32.5% placebo patients (p = 0.043).

Regarding the primary pharmacodynamic endpoint reduction of frontal beta2 power at electrode positions F3/4/7/8 from days 0–28, a...
relevant reduction was found for "eyes open" with significant differences between Salvia off. and placebo after 4 treatment weeks (Figure 5).

The analysis of secondary endpoints revealed a highly significant (p = 0.006) difference between Salvia off. and placebo for a clinically relevant ≥50% reduction of the MRS score and p values ranging from 0.008 to 0.049 for a 0%, or min. 20% or 60% reduction, respectively (Figure 6).

There was also a significant difference in reduction of the somato-vegetative and psychological MRS subscales (p = 0.0004 and p = 0.038, respectively) favoring Salvia off. (Figure 7), whilst the urogenital subscale showed only a trend regarding % of participants experiencing amelioration (data not shown). The single scores showed over the treatment period a significant difference (p < 0.05) in the reduction of the symptoms hot flushes, physical and mental exhaustion, sexual problems and joint and muscular discomfort whilst a trend was seen for sleep problems.

Electrical brain activity (days 0–28) in all brain areas within the 6 distinct frequency bands showed statistically significant spectral power

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**Table 1a. Patient characteristics in the per protocol population.**

| Characteristics | Placebo (n = 26) | Salvia off. (n = 22) | P value (Wilcoxon) |
|-----------------|------------------|----------------------|-------------------|
| Age (years)     | 53.1 (2.29)      | 53.1 (2.11)          | >0.05             |
| Body weight (kg) | 67.15 (9.93)     | 70.70 (11.35)        | >0.05             |
| Body height (cm) | 166.23 (5.62)    | 168.00 (5.46)        | >0.05             |
| BMI (kg/m²)     | 24.00 (4.18)     | 24.32 (3.71)         | >0.05             |

* Values for age, body weight, height and body mass index are presented as mean (SD).

**Table 1b. Patient characteristics in the ITT population.**

| Characteristics | Placebo (n = 40) | Salvia off. (n = 38) | P value (Wilcoxon) |
|-----------------|------------------|----------------------|-------------------|
| Age (years)     | 55.68 (4.30)     | 55.71 (3.75)         | >0.05             |
| Body weight (kg) | 68.98 (12.31)    | 71.12 (12.13)        | >0.05             |
| Body height (cm) | 165.68 (5.94)    | 167.45 (5.57)        | >0.05             |
| BMI (kg/m²)     | 24.78 (4.57)     | 24.71 (3.57)         | >0.05             |

* Values for age, body weight, height and body mass index are presented as mean (SD).

**Table 2. Menopause rating scale score day 0–28.**

|         | Salvia off. (A), n = 22 | Placebo (B), n = 26 | Difference |
|---------|-------------------------|---------------------|------------|
| Day 0   | Actual                  | %-Change            |            |
| Mean (SD)| 15.3 (6.87)            | 14.5 (6.04)         |            |
| Min-Max | 7.0–36.0                | 4.0–29.0            |            |
| Day 28  | Mean (SD)               | %-Change            |            |
|         | 9.3 (5.75)              | -37.1 (29.90)       | 13.0 (6.35) |
|         | [6.8; 11.9]             | [-50.4; -23.9]      | [10.4; 16.5]|
| 95% CI  | [6.8; 11.9]             | [-50.4; -23.9]      | [10.4; 15.6]|
| B - A   | [30.8 (32.4)]           | [30.8 (32.4)]       | [30.8 (32.4)]|

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Figure 4. Daily change of the Hot Flush Severity (HFS) Score over the 4 weeks treatment period compared to the arithmetic mean of the screening week for Salvia (green) vs Placebo (blue) *p ≤ 0.05.
during sleep (p < 0.05) whilst the other items, i.e. mental balance before sleeping, mental exhaustion before sleeping, sleep wake regulation, dream recall and feeling refreshed after sleep, didn’t change. A significant (p = 0.007) decrease in total HAMA score from 15.68 (day 0) to 9.82 (day 28) was observed for *Salvia off*. (p = 0.007), constituting a trend in favor of *Salvia off*. when compared with placebo where the total HAMA score was reduced from 15.27 to 11.19 (p = 0.02). According to protocol no analysis was performed on the level of HAMAS 14 single items, measuring psychic anxiety (mental agitation and psychological distress) as well as somatic anxiety (physical complaints related to anxiety), and encompassing anxious mood, tension, fears, insomnia, depressed mood, as well as intellectual, muscular, sensory, cardiovascular, respiratory, gastrointestinal, genitourinary, and autonomic symptoms and behaviour at interview, as the assay sensitivity for these particular parameters was judged not to be sufficiently precise and the intent was rather to gain an overall impression.

Tolerability was rated as very good by 100% of the patients of both treatment groups. Three adverse events in 3 patients (herpes zoster and road traffic accident in the *Salvia off*. and back pain in the placebo group) were reported and judged as unrelated and non-serious. No clinically relevant changes in laboratory parameters and vital signs were observed from day 0 to day 28.

4. Discussion

*Salvia off*. proved to be a valid option to fulfill the increasing demand of women reluctant or contraindicated to HRT who seek a natural and safe alternative treatment of their vasomotor and other typical menopausal symptoms like alterations of mood and cognition. This is the first time that in a placebo controlled, confirmatory clinical trial efficacy for a *Salvia off*. preparation was shown not only in reduction of hot flashes but of other climacteric symptoms as well.

As previously shown (Bommer et al., 2011) and demonstrated by the present study *Salvia off*. improves not only vasomotor symptoms (MRS, HFS scores) but simultaneously exerts a positive impact on accompanying somato-vegetative and psychological symptoms thus addressing a broad range of menopausal complaints. Indeed similar effects have been shown in this prior uncontrolled study for 2 months with the preparation under investigation. It is of interest to note that although this study demonstrated in the course of a 4 weeks application significant effects of *Salvia off*. on MRS and HSF against placebo, a further increase of treatment benefit can be expected with prolonged treatment based on the results of a 3 month placebo-controlled study with a *Salvia off*. preparation which showed increase of treatment effects in flushing frequency and severity compared to placebo with increase of application period (Zeidabadi, 2020). The same is however not to be anticipated for vaginal dryness which in contrast to sexual problems didn’t yield significant effects compared to placebo in both studies, be it over a 4 or a 12 weeks period.

Results indicate a stronger effect of *Salvia off*. in the PP population compared to the ITT collective, which however still yielded significant results as evidenced by the data obtained for the clinical score of the MRS as well as for the HFS. Hence, *Salvia off*. clearly showed superior efficacy in women experiencing a typical menopause associated with symptoms lasting for up to 5 years after the final menstrual period (Bachmann, 2005).

With hot flushes and sleep problems constituting the most prevalent concerns in menopause, the impact of *Salvia off*. shown on improvement of sleep quality and psychosomatic problems during sleep constitutes a relevant asset, regardless of whether owed primarily to alleviation of night sweats and their detrimental impact on sleep or due to other, as yet to be elucidated intrinsic factors of a treatment with *Salvia off*. itself, even more so as life quality remains a core issue for postmenopausal women (Schneider, 2017; Conde et al., 2006; Ledesert et al., 1994). The relevant difference in reduction of discontent compared to placebo as well as the trend in favour of *Salvia off*. observed for reduction of downheartedness
might further indicate a positive impact on perception of life quality as conferred by the treatment with *Salvia off*. The highly significant reduction of the HAMA score by *Salvia off* over the course of 28 days showing a trend compared to placebo is backed up by former study results obtained for single dose applications of *Salvia off* in young healthy adults showing an anxiolytic effect of the verum treatment (Kennedy, 2006).

Memory improvement has been shown for *Salvia off* preclinically (Smach et al., 2015), and supports the clinical findings of enhanced cognition as seen in the present study. Such an effect, in contrast to vasomotor improvements (Imhof et al., 2018) was not confirmed in a recent clinical study with soy (Furlong et al., 2020). Clinical vasomotor improvements have furthermore been shown for a wide variety of phytopharmaceuticals, many relying on a phytoestrogenic way of action, including evening primrose oil (Farzaneh et al., 2013), red clover (Myers and Vigar, 2017; Ghazanfarpour et al., 2015, 2016; Shakeri et al., 2015), valerian (Ensiyeh et al., 2018), and black cohosh, the latter showing in a recent clinical study with soy (Furlong et al., 2020). Clinical vasomotor improvements (Imhof et al., 2018) was not confirmed in this study, indicative that less stress is produced and correlating with menopausal symptoms. Beta1 and 2 spectral power have been associated with glutamergic and GABA-ergic transmission, respectively, indicating that a normalization of the activity of these transmitters also might contribute to the action of *Salvia off*.

Mental rest is visualized by decreased delta power and mental endeavor by enhancement of theta power (Dimpfel et al., 2011) while in the current study, attenuation of both delta and theta power was measured at central, parietal and temporal electrode positions at “eyes open” and during mental challenges. Taking into consideration that delta waves are controlled by acetylcholine, such attenuation of central-parietal and temporal brain areas reflects an activation of cholinergic transmission, known to be implicated in the central thermoregulation while additionally changes of theta power have been related to noradrenergic transmission, two mechanisms matching in being involved in sweating and hot flushes both of which were shown to be controlled by *Salvia off* also under in vitro conditions (Tober et al., 2019).

As Tober (Tober et al., 2019) has shown convincing effects for *Salvia off*, not on estrogen but on neurotransmitter level, qEEG measurements have been chosen to track *Salvia off*’s mode of action. This became consequently the first study to gain insight into the mode of action of *Salvia off* in menopause. In vitro data show a distinct influence of the extract on serotonergic, adrenergic, muscarinic, GABA-ergic, and μ-opioid receptors, exempt from an estrogenic action (Rahbe et al., 2013).

In this study we observed a reduction of the primary pharmacodynamic parameter frontal spectral beta2 power at electrodes F3/4/7/8 in the PP population after 4 treatment weeks with *Salvia off* compared to placebo, indicative that less stress is produced and correlating with menopausal symptoms. Beta1 and 2 spectral power have been associated with glutamergic and GABA-ergic transmission, respectively, indicating that a normalization of the activity of these transmitters also might contribute to the action of *Salvia off*.

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Alpha1 spectral power reflects changes in serotoninergic transmission, whereas alpha2 waves seem to be under the control of dopamine. Attenuation of alpha waves, as experimentally shown in the current study, both acutely as well as after the course of four study weeks, probably indicate activation of these two transmitter systems, which themselves are known to be modulated by estrogens and reflects mental activation.

Hence *Salvia off* induced both a state of mental activation as well as relaxation.

As for the safety analysis, the known excellent safety and tolerability profile of *Salvia off* once more was demonstrated. Since the proprietary *Salvia off* extract under investigation is devoid of thujone it is most suitable for long term therapy as usually requested by the natural course of the condition to be treated.

Overall, this double-blind, placebo controlled study proves the clinical efficacy of *Salvia off* with respect to reduction of most of the typical menopausal complaints, in particular hot flushes and elucidates their

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Figure 6. Treatment success for *Salvia off* and placebo regarding reduction of MRS in % of responders for a ≥0, ≥20, ≥50 and ≥60 reduction, respectively. **p ≤ 0.01.

Figure 7. Reduction of MRS somato-vegetative and psychological MRS subscales for *Salvia off* and placebo. ***p ≤ 0.001.
association with altered brain waves as evidenced by qEEG spectral power intensities.

A significant change of general electrical brain activity indicated higher mental capacity and improved coping with cognitive challenges accomplished by a reduction of endogenous stress production. qEEG data visualized the impact on central nervous transmitter systems which are involved in required neuroadaptive processes as a consequence of menopausal estrogen decline.

Results confirmed the data of the prior uncontrolled study with *Salvia officinalis* and suggest that *Salvia officinalis* and more specifically the thujone-free proprietary extract from freshly harvested *Salvia officinalis* leaves under investigation, is an effective and safe herbal approach for the long-term treatment of hot flushes and other climacteric symptoms in menopause.

Declarations

**Author contribution statement**

D. Wilfried: Conceived and designed the experiments; Analyzed and interpreted the data; Contributed reagents, materials, analysis tools or data; Wrote the paper.

C. D. G. Nina: Performed the experiments; Analyzed and interpreted the data; Contributed reagents, materials, analysis tools or data.

B. Silvia: Conceived and designed the experiments; Analyzed and interpreted the data; Wrote the paper.

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**Data availability statement**

Data included in article.

**Declaration of interests statement**

The authors declare the following conflict of interests: This study was sponsored by A.Vogel, Roggwil, Switzerland, manufacturer of the tested *Salvia officinalis* preparation Menosan® *Salvia*. S. Bommer is an A.Vogel employee.

Figure 8. Change at the combination of C4P4T4b6 of brainwave activity of distinct frequency ranges (delta, theta, alpha1 and alpha2, beta1 and beta2) on Day 28 (after 4 weeks’ therapy) compared to baseline Day A = 0 (set to 100%) under qEEG recording conditions “eyes open” [EO] and cognitive tests reading [RE], d2-concentration test [d2], memory test [ME], concentration performance test [CPT], reaction time test [RT], number identification test [NIT], and number connection test [NCT] under treatment with *Salvia officinalis* and placebo.
Additional information

No additional information is available for this paper.

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