

EFFECTS OF 4-THIAZOLIDINONE DERIVATIVES LES-2658 AND LES-1205 ON SLEEP – WAKEFULNESS CYCLE IN KINDLED RATS

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Abstract

The research is dedicated to in-depth study of neurotrophic and antiepileptic properties of original potential anticonvulsant agents from 4-thiazolidinones – LES-2658 (5-(3-nitro-benzylidene)-2-(thiazol-2-ylimino)-thiazolidin-4-one) and LES-1205 ([2,4-dioxo-5-(thiazol-2-ylcarbamoylmethyl)-thiazolidin-3-yl]-acetic acid ethyl ester), synthesized at the Department of Pharmaceutical, Organic and Bioorganic Chemistry of Danylo Halytsky Lviv National Medical University, Ukraine. Studying of sleep – wakefulness cycle characteristics in animals with chronic epileptic syndrome in conditions of 4-thiazolidinones derivatives LES-2658 and LES-1205 use was performed. The kindling syndrome was induced in Wistar rats via daily pentylenetetrazol (PTZ) (30 mg/kg, i. p.) administrations during three weeks and sleep – wakefulness cycle was studied under conditions of LES-2658 and LES-1205 administrations at doses 25.0 and 100.0 mg/kg i. p.. Total wakefulness, non - rapid eye movement sleep, rapid eye movement sleep, falling asleep latency, REM – onset latency and also number of REM sleep episodes have been determined by behavioral characteristics of experimental animals. It was established that 4-thiazolidinone derivatives Les-1205 and Les-2658 reduce REM sleep fragmentation and increase its duration in PTZ-kindled rats. Les-1205 compound at dose 100.0 mg/kg show a clear correcting influence on kindling – induced sleep disturbances.

Keywords: pentylenetetrazol kindling, sleep – wakefulness cycle, seizure syndrome, 4-thiazolidinones, diazepam.

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1. Introduction

Epilepsy is the most common neurological disorder and its prevalence worldwide is estimated at 0.5–1 % [1]. In most cases, pharmacotherapy is the only one method for satisfactory correction and control of convulsive states [2]. Despite the substantial progress in research and development of efficient anticonvulsants the search of new original anticonvulsant agents remains still open problem, especially in the context of pharmacoresistance and significant side effects under long use [3, 4].

The model of chronic epileptization of brain formed by repeated application of a subthreshold dose of pentylenetetrazol (PTZ) adequately reflects the signs of clinical forms of disease, including the disorders of sleep – wakefulness cycle [5, 6]. The study of the cycle allows estimate the brain mechanisms that control anxiety, to solve which mechanisms ensure its individual phases and to assess the influence of pharmacological agents on brain structure [5, 6]. Despite the well – es-

established relation between sleep and epilepsy [7], very few studies are available on the effect of an antiepileptic drug on the sleep – cycle.

4-Thiazolidinone derivatives are promising compounds in modern pharmacology for search of potential neurotropic agents with low acute and neurotoxicity properties [8–12]. Some 4-thiazolidinones demonstrate multi-level anticonvulsant activity with satisfactory toxicity parameters in conditions of such screening models as pentylenetetrazol (PTZ, metrazol) seizures, maximal electroshock test, rotorod test [8–12]. Earlier anticonvulsant activity of original 4-thiazolidinones Les-1205 – [2,4-dioxo-5-(thiazol-2-ylcarbamoylmethyl)-thiazolidin-3-yl]-acetic acid ethyl ester and Les-2658 – 5-(3-nitrobenziliden)-2-(thiazol-2-imino)-4-thiazolidinone (synthesized at the Department of Pharmaceutical, Organic and Bioorganic Chemistry, Danylo Halytsky Lviv National Medical University) have been identified in our studies [13–16]. Until recently, the studying of influence on the sleep – wakefulness cycle characteristics in rats with chronic epileptic syndrome formed by PTZ kindling were not used for determination of 4-thiazolidinones anticonvulsant activity mechanisms.

2. Aim of research

The study of 4-thiazolidinone derivatives LES-2658 and LES-1205 influence on the sleep – wakefulness cycle characteristics in rats with chronic epileptic syndrome formed by pentylenetetrazol (PTZ) kindling.

3. Material and methods

All investigative procedures and the animal facilities were conformed to the Guide of Care and Use of Laboratory Animals within European Convention for the Protection of Vertebrate Animals used for Experimental and Other Scientific Purposes (Strasbourg: Council of Europe 18.03.1986) and Law of Ukraine “On protection of animals from cruelty” (21.02.2006). For the study were used 54 male rats in age 5-6 months and weight ~180–220 gr (*Rattus Norvegicus* Var. *Alba*, *Wistar*). The animals were divided into 7 groups: one control (intact animals) group (n=6) and six experimental groups. For animals of all experimental groups the kindling syndrome was induced via daily pentylenetetrazol (PTZ) (30 mg/kg, i. p.) administrations during three weeks. Rats that the last three injections caused generalized tonic – clonic seizures were used for the experimental monitoring. The animals of experimental group I (n=9) were administered 0.9 % solution of NaCl; experimental group II (n=9) – animals were administered diazepam (“Gedeon Richter”, Hungary), at dose 1.5 mg/kg, i.p., 30 minutes before PTZ; experimental group III (n=7) and IV (n=8) – animals were administered LES-2658 at doses 100.0 and 25.0 mg/kg respectively, i. p., 30 minutes before PTZ; experimental group V (n=7) and VI (n=8) – animals were administered LES-1205 at doses 100.0 and 25.0 mg/kg respectively, i. p., 30 minutes before PTZ. The severity of seizures was evaluated according to five-point scale [5]. The tested rats were individually continuously monitored during 4 hours after 24 hours of the last application of PTZ under free behavior conditions and with identical degree of noise and lighting. All recordings were taken between 11.00 a. m. and 3 p. m. for a total recording time of 240 min. Total wakefulness (TW); total non-rapid eye movement sleep (NREM); total rapid eye movement sleep (REM); falling asleep latency (FAL) and REM – onset latency (RL) and also number of REM sleep episodes (RSE) have been determined by behavioral characteristics of experimental animals [6]. The durations of total wakefulness, NREM and REM sleep were given in % compared to the overall experiment duration (100 %); FAL and RL in minutes. Data were analyzed using the statistical criteria – ANOVA + Newmann-Keuls, and Mann-Whitney U-test. All values are expressed as means ± SEM. Differences were considered significant when: * p<0.05.

4. Results

The changes were observed in sleep pattern of experimental animals under kindling conditions.

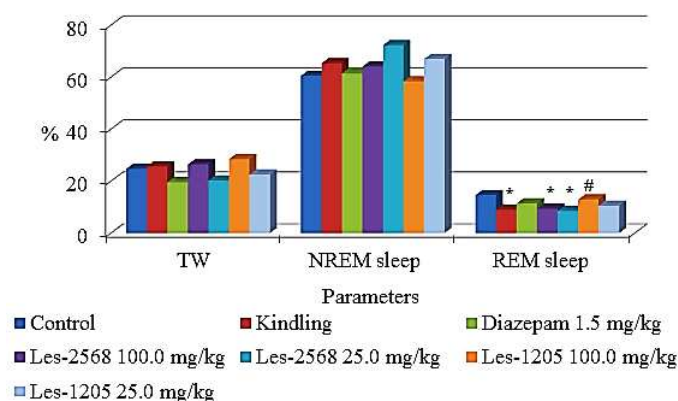


Fig. 1. The durations of total wakefulness, NREM and REM sleep under Les-2568, Les-1205 and diazepam administration in kindled rats. All parameters are given in % compared to the overall experiment duration (100 %). # – P<0.05 compared to the control group and * – P<0.05 compared to the kindled animals

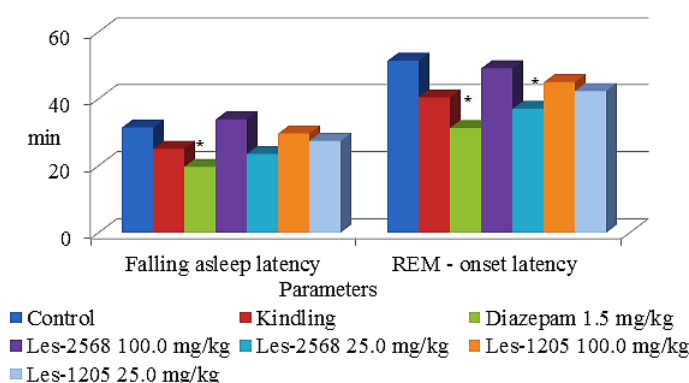


Fig. 2. The characteristics of falling asleep (FAL) and REM – onset (RL) latency under Les-2568, Les-1205 and diazepam administration in kindled rats. * – P<0.05 compared to the kindled animals

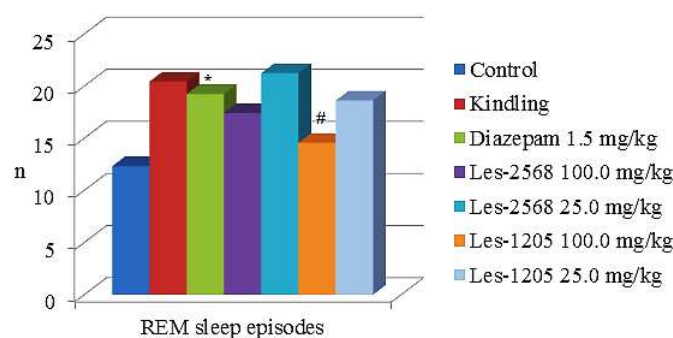


Fig. 3. The number of REM sleep episodes (RSE) under Les-2568, Les-1205 and diazepam administration in kindled rats. # – P<0.05 compared to the control group and * – P<0.05 compared to the kindled animals

5. Discussion

The tendency to elongation of NREM sleep was observed in kindled rats. Its duration was 65.29 ± 8.79 % and exceeded similar in the control group by 7.9 % ($P > 0.05$) (Fig. 1). At the same time, duration of REM sleep decreased by 38.4 % and was 9.04 ± 2.84 % with statistically significant ($P = 0.015$). Furthermore the tendency to decrease of FAL by 20.1 % (25.1 ± 9.48 min) as well as of RL by 20.3 % (40.41 ± 13.1 min) (for all $P > 0.05$) was observed in kindled rats compared to

group I (**Fig. 2**). Also, the number of RSE increased by 65.8 % (to 20.44 ± 5.46) and was statistically significant with $P < 0.05$ (**Fig. 3**).

It was established that administration of referenced drug and tested compounds LES-2658 and LES-1205 provoke changes in the in sleep pattern of kindled rats too.

The TW period was 19.6 ± 5.94 % and decreased by 23.6% ($P > 0.05$) as well as NREM sleep duration was 61.4 ± 7.91 % and increased by 6.05 % ($P > 0.05$) compared to kindled rats in rats when the diazepam was administrated at dose 1.5 mg/kg, i. p.. The duration of REM sleep was 11.45 ± 2.06 % and increased by 26.6 % ($P > 0.05$) but remained lower in comparison of intact rats group by 22.0 % ($P > 0.05$) (**Fig. 1**). Also, FAL and RL were 19.69 ± 6.48 min ($P = 0.021$) and 31.31 ± 8.36 min ($P < 0.05$) and increased by 37.3 % and by 39.1 % respectively compared to the control group (**Fig. 2**). The number of RSE was 1.56 times higher (19.25 ± 4.79) in comparison to the intact rats group ($P = 0.06$) (**Fig. 3**).

When the LES-2658 was administrated at dose 100.0 mg/kg, i. p., duration of TW and REM sleep were 26.46 ± 9.69 % ($P > 0.05$) and 9.43 ± 3.11 % ($P < 0.05$) and increased by 3.1 % and by 26.6 % respectively (**Fig. 1**). While NREM sleep duration was 64.11 ± 7.87 % and decreased by 1.8 % ($P > 0.05$) compared to kindled rats. Also, FAL and RL were 33.86 ± 13.36 min and 49.14 ± 9.82 min and increased by 34.9 % and by 21.6 % ($P > 0.05$ for all) respectively (**Fig. 2**). While the number of RSE was 17.43 ± 3.10 and decreased by 14.7 % ($P > 0.05$) compared to the kindled rats (**Fig. 3**). In case, when the LES-2658 was administrated at dose 25.0 mg/kg, i. p., duration of TW was 20.25 ± 7.43 % and decreased by 21.9 % ($P > 0.05$) in comparison of kindled rats (**Fig. 1**). While the NREM sleep duration was 72.31 ± 6.14 % and increased by 10.7 % ($P > 0.05$) compared to kindled rats and was higher than the same indicator in intact rats by 19.5 % ($P = 0.043$). Duration of REM sleep was 8.61 ± 2.27 % ($P < 0.05$) and remained lower by 41.4 % compared to the intact rats (**Fig. 1**).

Application of the Les-1205 at dose 100.0 mg/kg caused an increase TW duration (28.51 ± 8.48 %) compared to kindled rats by 11.1 % ($P > 0.05$) and compared to the II experimental group (diazepam used) by 45.5 % ($P < 0.05$) (**Fig. 1**). Herewith the duration of NREM sleep phase was 58.4 ± 9.0 % and slightly decreased by 10.6 % ($P > 0.05$). The duration of the REM sleep was 12.94 ± 2.58 % and increased by 43.1 % ($P < 0.05$) (**Fig. 1**). Under these conditions, FAL and RL were 29.64 ± 7.75 min and 45.0 ± 13.06 min ($P > 0.05$ for all) increased by 18.1 % and by 11.4 % respectively (**Fig. 2**). Also, the number of RSE was 14.57 ± 3.73 decreased significantly by 28.8 % with $P < 0.05$ (**Fig. 3**). When the Les-1205 was administrated at dose 25.0 mg/kg, i. p., duration of REM sleep phase was 10.57 ± 2.51 % and increased by 16.9 % ($P > 0.05$). Also, RL was 42.35 ± 11.29 min decreased by 8.9 %, ($P > 0.05$) in comparison of II experimental group (**Fig. 1, 3**).

For PTZ-induced kindling model such effects as decreasing of total sleep duration and increasing of its fragmentation were characteristic. These experimental results coincide with the research of other authors [4, 17, 18].

The use of 4-thiazolidinone derivative Les-1205 at dose 100.0 mg/kg in above-mentioned model of disorders of sleep – wakefulness cycle provoke opposite effects and is accompanied with increasing of the total sleep duration and decreasing of the number of its fragments. But for the derivative Les-2568 similar action was not observed and that can be explained by its less neurotrophic activity [8, 11].

Comparing of the impact of tested compounds with effects of diazepam showed that the for diazepam action was more typical reduction of FAL and RL in kindled rats and these data coincide with the results of other authors [19, 20]. Herewith, there was not observed an influence of diazepam on the REM sleep characteristics. The total duration of REM sleep was reduced and the number of fragments increased although was not statistically significant compared to the kindled rats group. Hence, this fact is in favor for opposite action of tested compounds and diazepam upon some components of sleep – wakefulness cycle.

So, the obtained and presented results on influence of 4-thiazolidinone derivatives Les-2658 and Les-1205 on sleep – wakefulness cycle in kindled rats cannot pretend to be the full solution of the search for new anticonvulsant agents. It is an attempt of studying and understanding anticon-

vulsant mechanisms and properties of 4-thiazolidinones as possible drug. And these results are promising for in depth-study using other different experimental seizure models.

6. Conclusion

1. Under developed kindling conditions the wakefulness and REM sleep phases were decreased while at one time NREM sleep phase was increased. Moreover there are reducing REM-onset latency and increasing fragmentation of REM sleep phase.

2. 4-Thiazolidinone derivatives Les-1205 and Les-2658 reduce REM sleep fragmentation and increase its duration in PTZ-kindled rats and compound Les-1205 at dose 100.0 mg/kg has been shown a clear correcting influence on kindling – induced sleep disturbances while diazepam at dose 1.5 100.0 mg/kg does not cause any corrective influence on the REM sleep of kindled rats.

3. Such ability of tested 4-thiazolidinones to recover of REM sleep parameters may indicate on importance in the implementation of anticonvulsant activity realization of this class of chemical compounds.

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