Effects of Remacemide and its Metabolite FPL 12495 on Spike-Wave Discharges, Electroencephalogram and Behaviour in Rats with Absence Epilepsy

E. L. J. M. VAN LUIJTELAAR* and A. M. L. COENEN

NICI, Department of Psychology, University of Nijmegen, P.O. Box 9104, 6500 HE Nijmegen, The Netherlands

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Summary—The effects of the anti-convulsant drug remacemide and one of its active metabolites FPL 12495 were examined in a genetic model for generalized absence epilepsy, the WAG/Rij strain of rats. Number, mean and total duration of spike-wave discharges were measured following oral administration of remacemide and FPL 12495, together with parameters of background electroencephalographic activity (EEG) and spontaneous behaviour in the recording cage. A decrease in the number of the spike-wave discharges was found after remacemide administration. At the highest dose there was near total suppression of the spike-wave discharges. There were no important effects on behaviour and on spectral content of the background EEG, suggesting that remacemide has little side effects. A decrease in the number of spike wave discharges was also found after FPL 12495 gavage and there was a prolongation of the mean duration. Behavioural changes were only noticed after the highest dose. These were accompanied by changes in the spectral content and particularly by an increase in the amplitude of the delta and the high beta frequencies, together with a decrease in the spindle frequency range. FPL 12495 appeared to be more potent that remacemide in all its effects. These mainly in FPL 12495 are uncommon in the sense that so far no other investigated drug shows a decrease in the number together with an increase in the mean duration of the discharges. It seems that in contrast to other anti-epileptic drugs, FPL 12495 exerts a differential action on the two commonly distinguished mechanisms controlling number and duration.

Keywords—Remacemide, FPL 12495, absence epilepsy, WAG/Rij strain, genetic model, spike-wave discharges, EEG, spectral analyses.

Remacemide is a new anti-convulsant drug. In mice and rats it shows a specific and potent protection in the maximal electroshock seizure (MES) test (Garske et al., 1991; Palmer et al., 1992). It is also effective in suppressing audiogenic seizures and the drug inhibits NMDA and kainic acid induced convulsions and tonic seizures elicited by 4-amino pyridine (Cramer et al., 1993; Palmer et al., 1992). The anticonvulsant effects of remacemide appear to be somewhat specific for these tests since pentylenetetrazol, bicuculline, picrotoxin or strychnine induced convulsions are unaffected. The mechanism of remacemide is thought to be similar to phenytoin and other compounds which inhibit sustained repetitive firing of the fast Na+ channel (McLean et al., 1983). It has been proposed that remacemide may offer a suitable treatment for patients with generalized tonic-clonic and complex partial seizures. The desglycine metabolite of remacemide, FPL 12495, is an even more potent anti-convulsant than remacemide itself (Palmer et al., 1992) and is a considerably more potent antagonist of the fast Na+ channel and at the NMDA receptor site than remacemide (Harris et al., 1992).

The molecular structures of remacemide and FPL 12495 and their action in MES and in other models for convulsive epilepsy does not predict unambiguously the actions of these drugs against another type of generalized epilepsy, absence epilepsy, since various types of anti-convulsants react differentially in absence epilepsy. Drugs effective against tonic clonic epilepsy such as carbamazepine and phenytoin, do not decrease but aggravate absence seizures. Other anti-convulsants such as diazepam and loreclezole reduce spike-wave discharges while typical anti-absence drugs such as ethosuximide and trimethadione decrease spike-wave activity (Ates et al., 1992; Micheletti et al., 1987; Peeters et al., 1988; Peeters, 1991). If remacemide would mimic the effects of carbamazepine and phenytoin, one may predict that spike-wave activity would be enhanced after remacemide.
Genetic models are the pertinent choice among the models available for human absence epilepsy (Lösch, 1984) and such a model is the WAG/Rij strain of rats (Coenen et al., 1992a; van Luijtelaar and Coenen, 1986, 1989; van Luijtelaar et al., 1991). At an age of six months all members of this inbred strain show the electrophysiological and behavioural manifestations of generalized absence epilepsy. They exhibit spontaneously occurring spike–wave discharges mostly during periods with a low level of vigilance (Drinckenburg et al., 1991; van Luijtelaar et al., 1991). The pharmacological profile of these spike–wave discharges closely resembles the profile of human absence epilepsy (Peeters et al., 1988).

It will be investigated in the present experiment whether remacemide and FPL 12495 have effects on generalized absence epilepsy as shown by rats of the WAG/Rij strain. In order to determine whether these drugs have also other effects, spontaneous behaviour and background EEG were additionally measured.

METHODS

Fifty-six male WAG/Rij rats, six months of age and weighing between 260–382 g were used as experimental subjects. Animals were individually housed and had always access to food and water. A 12–12 hr light–dark cycle with bright white lights on at 20.00 hr, was maintained throughout the experiment. The experiment took place in the dark phase of the day, which is the time of day with the highest incidence of spike–wave discharges (van Luijtelaar and Coenen, 1988). The experimental room was then illuminated by a weak red light for which albino rats are unsensitive but which allowed us to observe the rats’ behaviour.

EEG electrodes (Plastics One, MS 333/2-A) were implanted in the cortex under anaesthesia (Nembutal, 60 mg/kg i.p.). They were placed in the frontal and in the parietal/occipital region of the cortex with coordinates A 2.0, L 3.5 and A 6.0, L 4.0 respectively, with skull surface flat and bregma zero–zero. An earth electrode was placed in the cerebellum. The animals were allowed to recover for at least 10 days. Experiments were performed in freely moving animals singly housed in transparent EEG recording cages (25 × 25 × 30 cm).

Remacemide and FPL 12495 were supplied as the hydrochloride salt by Fisons Pharmaceuticals (Loughborough, U.K.). The compounds were dissolved in saline and remacemide was administered in doses of 20, 40 and 80 mg/kg and FPL 12495 in doses of 10, 20 and 40 mg/kg. Additionally, a control group was used and rats of this group received saline only. Each group consisted of 8 subjects. The rats were adapted to the recording conditions for 18 hr and a base-line EEG was recorded for 1 hr, starting at 10.00 hr. Then, animals were given remacemide, FPL 12495 or solvent in a dose volume of 1 ml orally and subsequently the EEG was recorded for 2 consecutive hours. Drugs were always administered at 11.00 hr. All animals were used only once.

The EEG signal was amplified and filtered by an Elema–Schönander polygraph and frequencies between 1 and 70 Hz were allowed to pass. The EEG was stored in digitized form on a magneto-optical disk (DATA Instruments, AT-CODAS). Spike–wave discharges were visually scored according to criteria elaborated elsewhere (van Luijtelaar and Coenen, 1986). Number, mean and total duration of spike–wave discharges were determined.

Additionally, the spectral content of the background EEG without spike–wave discharges, was determined during representative periods of passive wakefulness after the first hour following remacemide, FPL 12495 or saline administration. The minimum time period that was analysed for each subject was 35 sec. These epochs were digitized with a rate of 512 samples per second. The power spectrum was calculated by means of a Fast Fourier Transformation (FFT) with a bin density of 1 Hz, whereby the power spectrum was normalized. Z-scores are then obtained with a mean of zero and a variance of 1 in order to compensate for interindividual differences in amplification (Coenen and van Luijtelaar, 1989). The content of the following EEG frequency bands were determined: the delta-band (1–4 Hz), the theta-band (6–10 Hz), the sigma-band (11–14 Hz), the beta 1-band (15–30 Hz) and the beta 2-band (> 30 Hz).

Spontaneous on-going behaviour of the rats in their cages was observed through a window from an adjacent room. The duration of the following behaviour categories was quantified for 30 min starting 5 min after injection: locomotor behaviour interpreted as exploratory behaviour (walking, rearing, sniffing, digging), automatic behaviour (grooming, eating, drinking) and immobile behaviour (sitting, lying and standing still) (Coenen and van Luijtelaar, 1989; Vanderwolf, 1969). Data were recorded and analysed with the Observer (Noldus, 1991).

The overall effects induced by the compounds with respect to number and duration of spike–wave discharges, behaviour and EEG spectral analysis, were statistically analyzed by means of a one-factor (doses) ANOVA, while post-hoc comparisons according to Duncan were used to test differences between groups. The number of degrees of freedom were always 3 and 28. The effects on spike–wave activity were evaluated for 60 min periods.

RESULTS

Remacemide

In Fig. 1(A), the number, mean and total duration of the spike–wave discharges are shown before and after the administration of saline, 20, 40 and 80 mg/kg remacemide. Each bar in the histogram represents the values in a block of 60 min of recording. There were no differences between the four groups before drug administration. There were no differences on the number of spike–wave discharges between the four groups in the
Fig. 1. (A) The effects of saline and three doses of remacemide on (a) number, (b) mean duration in sec, and (c) total duration of spike-wave discharges in sec, 1 hr before and 2 hr after injection. Means and standard errors of the means are depicted. (B) The effects of saline and three doses of FPL 12495 on (a) number, (b) mean duration in sec, and (c) total duration of spike-wave discharges in sec 1 hr before and 2 hr after injection. Means and standard errors are depicted.
first hour after the animals were given remacemide. However, there was a significant effect in the second hour after administration ($F = 4.85$, $P < 0.01$). The post-hoc test showed that the 80 mg/kg group had less spike-wave discharges than the saline and the 20 mg/kg group. Especially in the last 30 min remacemide became very effective: only one animal of the highest dose group showed just three spike-wave discharges, while the seven other rats had no spike-wave discharge left. Cochrans Q-test for dependent samples ($Q = 10.36$, $Fg = 3$, alpha = 0.02) confirmed that there were less animals with spike-wave discharges after 80 mg/kg remacemide than after solvent.

The mean duration of the spike-wave discharges was unaltered. The total duration of the spike-wave discharges showed also a significant effect in the second hour after administration ($F = 3.58$, $P < 0.05$). The post-hoc test showed that the total duration of spike-wave discharges of the 80 mg/kg group was shorter than that of the 0 and 40 mg/kg groups.

The behavioural data presented in Table 1 showed no significant effects. The data of the background EEG analyses are presented in Fig. 2(A). Although a decrease in the power of the spindle and beta-1 frequency band can be inferred from this figure, there were no significant changes in any of the frequency bands.

**FPL 12495**

In Fig. 1(B) the effects of FPL 12495 in doses of 10, 20 and 40 mg/kg are shown on the number, mean and total duration of the spike-wave discharges. There were no differences between the four groups prior to FPL 12495 administration. There was a significant effect on the number of spike-wave discharges in the first hour after drug administration ($F = 6.75$, $P < 0.001$): after 40 mg/kg the number was decreased compared to 10 mg/kg and solvent and also after 20 mg/kg the number was decreased compared to solvent.

The mean duration of the spike-wave discharges was affected by the drug in the first ($F = 3.90$, $P < 0.05$) and in the second ($F = 9.57$, $P < 0.001$) hour post injection. In the first hour the duration of 20 mg/kg group was longer than that of the 40 and 10 mg/kg groups. In the second hour the duration was prolonged after 40 mg/kg compared to 20, 10 and 0 mg/kg.

The total duration of the spike-wave discharges was only reduced in the first hour ($F = 4.45$, $P < 0.05$): the 40 mg/kg group differed from the 20, 10 and 0 mg/kg groups.

The behavioural data are presented in Table 1. The animals tended to show more exploratory behaviour, but the effects did not reach significance. The drug had an effect on the number of immobile periods ($F = 4.76$, $P < 0.01$): the post-hoc test showed that it was enhanced after 40 mg/kg only.

The data on the background EEG of FPL 12495 are presented in Fig. 2. A significant dose effect was found for the delta band ($F = 3.31$, $P < 0.05$): after 40 mg/kg group there was more delta activity than after 20 mg/kg or saline. There was a tendency for a dose effect in the theta band. In the spindle ($F = 4.49$, $P < 0.01$) and in the beta-2 band ($F = 5.38$, $P < 0.01$) significant dose effects could be demonstrated: the amplitude was smaller in the

<table>
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<tr>
<th>Dose</th>
<th>Exploratory behaviour</th>
<th>Automatic behaviour</th>
<th>Immobile behaviour</th>
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<tr>
<td>Remacemide</td>
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<tr>
<td>0 mg/kg</td>
<td>26.1 ± 1.5%</td>
<td>32.5 ± 7.2%</td>
<td>41.4 ± 8.2%</td>
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<tr>
<td>20 mg/kg</td>
<td>28.0 ± 4.1%</td>
<td>27.1 ± 6.1%</td>
<td>44.8 ± 7.8%</td>
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<td>40 mg/kg</td>
<td>25.3 ± 6.0%</td>
<td>42.4 ± 9.0%</td>
<td>32.2 ± 10.8%</td>
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<tr>
<td>80 mg/kg</td>
<td>25.5 ± 5.3%</td>
<td>49.9 ± 2.9%</td>
<td>24.5 ± 6.8%</td>
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<tr>
<td>FPL 12495</td>
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<tr>
<td>0 mg/kg</td>
<td>26.1 ± 1.5%</td>
<td>32.5 ± 7.2%</td>
<td>41.4 ± 8.2%</td>
</tr>
<tr>
<td>10 mg/kg</td>
<td>29.3 ± 4.6%</td>
<td>40.0 ± 8.0%</td>
<td>30.7 ± 8.9%</td>
</tr>
<tr>
<td>20 mg/kg</td>
<td>23.1 ± 4.2%</td>
<td>40.3 ± 6.1%</td>
<td>36.6 ± 4.8%</td>
</tr>
<tr>
<td>40 mg/kg</td>
<td>40.5 ± 6.3%</td>
<td>26.5 ± 4.3%</td>
<td>31.2 ± 6.4%</td>
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spindle band after 40 mg/kg compared to saline or to 10 mg/kg, but larger in the beta-2 band after 40 mg/kg compared to saline.

**DISCUSSION**

Although there are some clear differences between the two compounds with respect to time of onset and potency, the present data show that both remacemide and FPL 12495 decrease the number of spike–wave discharges in this genetic model for generalized absence epilepsy. The decrease in the number of spike–wave discharges after specifically remacemide application was found however in the highest dose and in the second hour only. This late onset is probably due to the oral administration of remacemide and to the fact that this drug has to be metabolized in order to become active and to suppress spike–wave discharges. Remacemide is metabolized into FPL 12495 and at its turn FPL 12495 is further metabolized into FPL 14465 and FPL 14330. In fact remacemide has nine identified and five potential metabolites (Palmer et al., 1992). Both metabolites FPL 14465 and FPL 14330 can be found in cerebrospinal fluid and in the plasma (Palmer et al., 1992) in rats but it is unknown whether these and other metabolites suppress spike–wave discharges. The decrease in epileptic activity was also evident from the decrease in the total duration of spike wave discharges, again after the highest dose only and this was also particularly striking 90–120 min after administration. At that time only one single animal showed spike–wave discharges and only three in number: i.e. a mean of less than 0.5 spike-wave discharge compared to habitual twenty under base-line conditions. Since the plasma concentration after oral administration of remacemide peaks at 30 min and declines to approximately a quarter of that value in 2 hr, it seems likely that the firm suppression of the number of spike–wave discharges after 90–120 min can be more likely attributed to either a combined action of remacemide and active metabolites rather than to remacemide itself. The duration of the action of the active metabolites exceeded the 2 hr recording period and therefore it remains unclear how long remacemide or their metabolites suppress spike–wave discharges.

FPL 12495 is more potent than remacemide in decreasing the number of spike–wave discharges and acts also faster than remacemide. FPL 12495 reaches its maximum plasma value after 30 min and remains constant for several hours (Palmer et al., 1992). Surprisingly, FPL 12495 increases the mean duration of spike wave discharges. It is of interest that the effects on number and on mean duration of spike–wave discharges dissociate in two ways, firstly on time and secondly on direction. Firstly, the effects on the number are immediately present and stay limited to the first hour, while the effects on the mean duration tend to be present in the first hour and are larger in the second hour. This latter finding suggests that the effects on duration are due to a combined action of FPL 12495 and one or more of its active metabolites. Another possibility is that FPL 12495 is racemic and that its differential effects on time might be related to varying effects of the two isomers. Secondly, there is also a dissociation with respect to the direction of the induced changes in the anti-epileptic activity of FPL 12495. This compound suppresses the number of spike–wave discharges but it prolongs their mean duration; a decrease in the number together with an increase in the mean duration of the spike–wave discharges is striking and uncommon. Until now we and others have evaluated specific anti-absence drugs, anti-convulsants, broad spectrum anti-epileptics (Ates et al., 1992; Coenen and van Luijtenaar, 1989; Micheletti et al., 1987, Peeters et al., 1988, van Luijtenaar and Coenen, 1989), β-carbolines (Coenen et al., 1989, 1992b), quisqualate, kainate and NMDA receptor agonists and antagonists (Frey and Voits, 1991; Peeters et al., 1989; Peeters, 1991). GABA-ergic compounds (Micheletti et al., 1985; Peeters et al., 1989a), morphine-like analgesics (Frey and Voits, 1991) and opiates (Lason et al., 1994), dopaminergic agents (Buzsáki et al., 1990; Warter et al., 1988), central anticholinergics (Frey and Voits, 1991) and psychotropic and alpha-adrenergic agents (Kleinlogel, 1995; Vanderwolf, 1969). Although not all authors distinguished between the number and the mean duration of the spike–wave discharges, it was not remarked that any of these compounds decreased the number of spike–wave discharges associated with an increase of its mean duration. This means that the present data are rather uncommon in the sense that they indicate that a single compound and probably one of its metabolites may exert effects on number and mean duration in opposite ways.

It has to be remarked that the drug mechanisms involved in protection against spike–wave discharges might be quite different from those involved in protecting against maximal electroshock seizures. Phenytion and carbamazepine are very effective in the maximal electroshock test, but they increase the numbers of spike–wave discharges (Peeters et al., 1988; van Luijtenaar and Coenen, 1989). Remacemide and FPL 12945 reduce the number of spike–wave discharges and they are effective in the maximal electroshock test (Palmer et al., 1992; Stagnitto et al., 1990). It seems however, that remacemide is more potent in the electroshock test at which an ED50 as low as 21.4 mg/kg was noticed. Here only 80 mg/kg was effective.

The level of vigilance and the number of spike–wave discharges have an intimate relationship. Few spike–wave discharges can be expected during periods of high vigilance and active behaviour and vice versa (Coenen et al., 1992; Drinkenberg et al., 1991; van Luijtenaar et al., 1991). The present data show that the decrease in the number of spike–wave discharges after the highest dose of remacemide is not accompanied by clear changes in motor activity and by changes in the background EEG.
The highest dose of FPL 12495 induced a tendency for more locomotor activity while the number of immobile periods was enhanced with shorter durations, implying some behavioural activation. Enhanced activity prevents the occurrence of spike–wave discharges and it cannot be excluded that the decrease in the number of spike–wave discharges is secondary to the increase in locomotor activity. However, the behavioural changes occurred only after the highest dose of FPL 12495 while the effects on the number of spike–wave discharges were already present after the middle dose. This suggests that the effects of FPL 12495 on spike–wave discharges are more due to its anti-epileptic action and not to its mild behavioural changes.

The spectral analysis of the background EEG did not reveal significant changes after remacemide. This suggests that remacemide has no effects on the background EEG activity as so many psychoactive drugs including anti-epileptics (Duncan, 1987; Krijzer et al., 1993). The highest dose of FPL 12495 induced changes in the spectral content: an increase in the delta and in the high beta range and a decrease in the spindle band. Krijzer et al. (1993) observed a power increase in the 35–100 Hz and a power decrease from 15 to 25 Hz in the frequency spectrum of some psychostimulant drugs.

The effects on the spectral content might be related to a yet unknown common property of FPL 12495 and these psychostimulant drugs, perhaps a high affinity for the non-competitive NMDA receptor (Harris et al., 1992).

It can be concluded that remacemide has a similar anti-absence activity in this model and that its active metabolite, FPL 12495, also reduces absence activity in spite of a tendency to increase the mean duration of spike–wave discharges. FPL 12495 seems more potent than remacemide in reducing the number of spike–wave discharges. The combination of the decrease in number and increase in mean duration, in particular for FPL 12495, is rather uncommon and offers a way to separate the mechanisms responsible for the onset and the duration of spike–wave discharges. FPL 12495 has effects on behaviour and an EEG profile with some similarity to that of psychostimulant drugs. Remacemide has no large effects on behaviour and does not affect the background EEG and it should be tried in intractable types of absence epilepsy.

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REFERENCES


