Evaluation of Pharmacokinetics and Dose Proportionality of Diazepam After Intranasal Administration of NRL-1 to Healthy Volunteers

Sarina Tanimoto1, Luana Pesco Koplowitz2,∗, Richard E. Lowenthal1, Barry Koplowitz2, Adrian L. Rabinowicz3, and Enrique Carrazana3

Abstract
NRL-1 is a novel intranasal formulation of diazepam that is being evaluated as rescue medication in patients with epilepsy who experience bouts of increased seizure activity despite stable regimens of antiepileptic drugs. This phase 1, open-label, randomized, crossover study in healthy adult volunteers consisted of 3 single-dose periods (5, 10, and 20 mg) followed by a 2-dose period (2 × 10 mg) with a minimum 28-day washout between treatments. Blood samples were taken at prespecified time points after intranasal dosing, and bioanalytic analysis of diazepam and nordiazepam was conducted using a validated liquid chromatography–tandem mass spectrometry method. Plasma pharmacokinetic parameters were summarized using descriptive statistics, and dose proportionality (peak concentration [Cmax] and area under the plasma concentration–time curve [AUC0–∞]) was evaluated based on a power model within a 90% CI of 0.84 to 1.16. Comparisons were also conducted between single 10-mg dose and multidose (2 × 10 mg) treatments. NRL-1 administration resulted in rapid diazepam absorption (median time to peak concentration 1.4-1.5 hours). Plasma concentration-time profiles showed similar patterns of exposure that appeared to be dose dependent, with Cmax of 85.6, 133.6, and 235.3 ng/mL for the 5-, 10-, and 20-mg doses, respectively, although the lower 90% CI for Cmax and AUC0–∞ exceeded dose proportionality criteria. The coefficient of variation ranged from 59% to 67% for Cmax and 48% to 56% for AUC parameters. Dose-normalized AUC0–∞ values were comparable between the 2 × 10-mg and single 10-mg doses. Treatment-emergent adverse events were consistent with those expected for diazepam, with transient somnolence the most frequent adverse event (94.4%). These results support NRL-1 as a potential therapy for managing seizure emergencies.

Keywords
NRL-1, intranasal diazepam, pharmacokinetics, dose proportionality, seizure cluster, acute repetitive seizures

Epilepsy affects approximately 50 million people worldwide, and the prevalence in the United States is almost 3.5 million individuals including 3 million adults and 470,000 children.

Epilepsy is associated with a substantial disease burden that results from increased morbidity and mortality relative to the general population, its impact on patient and caregiver quality of life, and an increased need for healthcare resource use.

Although management of epilepsy relies on the use of antiseizure drugs (ASD) to prevent and control seizures, up to 40% of individuals being treated with ASD may have refractory epilepsy. Bouts of increased seizure, often referred to as seizure clusters, are frequently experienced by some patients with epilepsy. These bouts can be defined as intermittent increases of seizure activity while on stable regimens of ASDs, and they require immediate rescue medication that can readily and easily be administered to manage these

1 Pacific Link Consulting, San Diego, California, USA
2 DUCK FLATS Pharma, Elbridge, New York, USA
3 Neurelis, Inc, San Diego, California, USA

This is an open access article under the terms of the Creative Commons Attribution License, which permits use, distribution and reproduction in any medium, provided the original work is properly cited.

Submitted for publication 29 August 2019; accepted 26 November 2019.

Corresponding Author:
Sarina Tanimoto, MD, PhD, Pacific Link Consulting Services, 16091 San Dieguito Rd, PO Box 99333, Rancho Santa Fe, CA 92091-9800 (e-mail: sarina@pacificlinkconsulting.com)

∗Fellow of the American College of Clinical Pharmacology
seizure emergencies. Rescue therapy has generally relied on use of benzodiazepines, and although intravenous administration is frequently used in an inpatient setting, rectal diazepam (Diastat) has been the standard of care in an outpatient setting and is appropriate for administration by non–healthcare professionals. However, because rectal administration may be difficult under seizure conditions and may be embarrassing to patients and caregivers, additionally providing a large absorptive surface area with high vascularity and direct nose to brain drug delivery, thus bypassing presystemic surface area with high vascularity and direct nose to brain drug delivery, thereby reducing the need for a second rescue dose.

Intranasal administration can be considered an appropriate alternative with advantages that include being noninvasive and readily accessible to patients and caregivers, additionally providing a large absorptive surface area with high vascularity and direct nose to brain drug delivery, thus bypassing presystemic surface area with high vascularity and direct nose to brain drug delivery, thereby reducing the need for a second rescue dose.

The purpose of the current study was to further assess the pharmacokinetics and dose proportionality of diazepam after intranasal administration of NRL-1 to healthy volunteers.

Methods

Study Design

This phase 1, open-label, randomized, crossover study consisted of 3 single-dose periods encompassing 6 sequences followed by a 2-dose period. The study was conducted on an inpatient basis at the study site, and subjects remained at the site for ~36 hours in each treatment period, with a minimum 28-day washout between treatments. NRL-1 was administered as a 100-µL intranasal spray of either 50 mg/mL or 100 mg/mL formulation under fasted conditions after an overnight fast. In the single-dose periods, the 5- and 10-mg doses were administered into a single nostril, and the 20-mg dose required an administration in each nostril of the 100 mg/mL formulation. In the 2-dose period, 2 10-mg doses were given 4 hours apart in alternate nostrils.

Subjects

Eligible subjects were healthy male and female adult volunteers 18 to 55 years old, inclusive, with a body weight 51 kg to 111 kg, inclusive. Subjects were also required to have no clinically significant abnormal findings in their medical history or on physical examination, electrocardiogram (QTcF < 450 msec for men and QTcF < 470 msec for women), or clinical laboratory results during screening. Female subjects of childbearing potential had to agree to use a medically
acceptable method of contraception during the study duration, and those who were trying to conceive, were pregnant, or were lactating were excluded. Other key exclusion criteria were a history of major depression or a past suicide attempt or suicide ideation; a history of allergy or adverse response to diazepam; treatment with enzyme-altering drugs in the past 30 days, over-the-counter oral and/or nasal decongestants in the past 14 days, or prescription medications, including benzodiazepines, in the past 14 days, excluding hormonal contraceptives.

**Bioanalytical Evaluation**

Serial blood samples were collected into labeled K$_2$-EDTA tubes by direct venipuncture at prespecified time points. For the single-dose treatment periods, collection times were at predose and at 10, 20, 30, and 45 minutes, and 1, 1.25, 1.5, 1.75, 2, 4, 8, 12, 24, 36, 48, 72, 96, 144, 192, and 240 hours after dosing. For the 2-dose treatment period, blood was collected predose and at 10, 20, 30, and 45 minutes, and 1, 1.25, 1.5, 1.75, 2, 4, 4.167, 4.33, 4.5, 4.75, 5, 5.25, 5.5, 5.75, 6, 8, 12, 24, 36, 48, 72, 96, 144, 192, and 240 hours after the first dose. After collection, the plasma was separated by centrifugation (3000 rpm × 10 minutes at 0-4°C at a relative centrifugal force of 2000g), and equal aliquots were transferred to 2 labeled tubes and stored at approximately –20°C until analysis.

Sample analysis was performed at a central laboratory (Quest Diagnostics, Houston, Texas) using a validated liquid chromatography–tandem mass spectrometry method to determine sample concentrations of diazepam and its metabolite nordiazepam (desmethyldiazepam) over the ranges of 1 to 1000 ng/mL and 100 to 100,000 pg/mL, respectively; 1.00 ng/mL and 100 pg/mL represent the lower limits of quantitation for diazepam and nordiazepam, respectively.

Analyte analysis consisted of automated liquid-liquid extraction and separation using mobile phases of methanol/water with ammonium formate (mobile phase A) and methanol/water with acetic acid (mobile phase B). Separation was performed on an ACE 3 C18-PFP column (30 × 4.6 mm, 3 μm; Advanced Chromatography Technologies Ltd, Aberdeen, Scotland) with detection using the API-4000 system (Sciex, Framingham, Massachusetts).

Standard-curve and quality control samples were generated by spiking blank human K$_2$-EDTA plasma (Bioreclamation; Westbury, New York) with internal standards (diazepam, nordiazepam, diazepam-d$_5$, and nordiazepam-d$_5$) obtained from Cerilliant (Round Rock, Texas) to monitor assay performance.

For precision, the intra-assay results showed that the coefficient of variation (CV%) at the lower limit of quantitation was 4.0% for diazepam and 5.9% for nordiazepam; the corresponding intra-assay accuracies, expressed as percentage bias, were 3.9% and –0.5% for diazepam and nordiazepam, respectively.

**Safety and Tolerability**

Safety and tolerability were evaluated by the reporting of treatment-emergent adverse events (TEAEs) regardless of causality, with additional determination as to whether the events were related to treatment. Evaluations also included clinical laboratory tests, physical examination, vital-sign measurement, electrocardiography, clinical laboratory tests, and the Columbia Suicide Severity Rating Scale.

To evaluate the potential for respiratory depression, oxygen saturation was measured using pulse oximetry at prespecified time points including within 1 hour predose, and postdose at 30 minutes and 1, 1.5, and 2 hours for the 20-mg and first 10-mg intranasal dose, and at 4.5, 5, 5.5, and 6 hours for the second 10-mg intranasal dose. Nasal irritation and sedation were both measured objectively using 6-point scales (0-5, with higher scores indicative of greater severity) as previously described, with the former reported by a trained observer and the latter reported by the subject (if awake) and a trained observer. Changes in olfaction were assessed using the NIH Toolbox Odor Identification Test, and acute pain following nasal administration was assessed using a 0-10 visual analogue scale (0 = no pain, 10 = extreme pain).

**Statistical and Pharmacokinetic Analyses**

A target of 36 enrollees was planned with the goal of 24 completers. This sample size was determined to provide 97% power to conclude dose proportionality for $C_{\text{max}}$ and AUC$_{0-\infty}$ over the range of 5 mg to 20 mg based on 90%CI for the slope ($\beta$) from the power model within 0.84 and 1.16 with $\alpha = 0.05$ and true slopes of 1. The analysis included a safety population, defined as subjects who received at least 1 dose of NRL-1, and a pharmacokinetic population, defined as subjects who completed at least 2 of the 3 single doses or the 2 10-mg treatments.

Pharmacokinetic parameters for diazepam and its metabolite, nordiazepam, were calculated from the individual plasma concentrations by noncompartmental methods using Phoenix WinNonlin version 7.0 (Certara Company, Princeton, New Jersey). The plasma pharmacokinetic parameters calculated were $C_{\text{max}}$, $t_{\text{max}}$, AUC$_{0-\infty}$, AUC$_{0-\text{tmax}}$, $t_{1/2}$, and apparent total body clearance (CL/F). For the 2-dose period, $C_{\text{max}(0-4h)}$ and $C_{\text{max}(4-8h)}$ represent the maximum concentration within the 4 hours after the first dose and the maximum concentration in the 4–8-hour time range, respectively, with similar representation of AUC$_{0-4h}$ and AUC$_{4-8h}$. 


Table 1. Summary Statistics of Pharmacokinetic Parameters After Single Dose of NRL-1

| Pharmacokinetic Parameter | 5 mg | 10 mg | 20 mg | 5 mg | 10 mg | 20 mg |
|---------------------------|------|-------|-------|------|-------|-------|
| **Diazepam**              |      |       |       |      |       |       |
| C_max, ng/mL              | 85.6 (67%) | 133.6 (64%) | 235.3 (59%) | 64.2 (106%) | 107 (82%) | 199.6 (65%) |
| t_max, h                  | 1.5 (0.3, 8) | 1.5 (0.8, 36) | 1.4 (0.5, 8) | ... | ... | ... |
| AUC_0-t, h                 | 2232 (51%) | 3803 (52%) | 7946 (52%) | 1910 (68%) | 3303 (62%) | 6864 (63%) |
| AUC_0-%/2, h               | 2411 (48%) | 4505 (56%) | 9168 (55%) | 2110 (62%) | 3854 (65%) | 7711 (72%) |
| t_1/2                      | 70 (42%) | 71 (44%) | 74 (49%) | 64 (47%) | 65 (48%) | 66 (49%) |
| CL/F, L/h                 | 2.8 (72%) | 3.1 (69%) | 3.2 (75%) | 2.4 (62%) | 2.6 (65%) | 2.6 (72%) |
| **Nordiazepam**           |      |       |       |      |       |       |
| C_max, ng/mL              | 11.7 (48%) | 19.3 (61%) | 36.6 (57%) | 10.5 (55%) | 16.1 (71%) | 31.6 (59%) |
| t_max, h                  | 96 (8.240) | 96 (12.240) | 96 (36.240) | ... | ... | ... |
| AUC_0-t, h                 | 1832 (46%) | 3115 (55%) | 6067 (53%) | 1638 (54%) | 2653 (66%) | 5285 (59%) |
| AUC_0-%/2, h               | 3229 (50%) | 4503 (37%) | 8838 (59%) | 2972 (46%) | 4258 (39%) | 7462 (73%) |
| t_1/2                      | 91 (51%) | 68 (29%) | 85 (32%) | 82 (52%) | 66 (30%) | 82 (28%) |

AUC indicates area under the plasma concentration-time curve; AUC_0-t, AUC from time of dose to last measurement; AUC_0-%/2, AUC from time of dose extrapolated to infinity; C_max, peak plasma concentration; CV%, coefficient of variation; t_max, time from dose to C_max; t_1/2, half-time of elimination.

Values are medians (minimum, maximum). Ranges in subscripts indicate time intervals of measurements.

Results

Population

A total of 36 subjects were enrolled and received at least 1 dose of drug. Among the subjects who comprised the safety population, 69.4% were male, 75.0% were black (13.9% white and 11.1% other), the mean ± SD age was 36.0 ± 9.9 years, and mean ± SD body weight was 85.7 ± 13.4 kg. Twenty-nine subjects completed the study, and of the 7 discontinuations, 6 (16.7%) were due to withdrawal by the subject and 1 (2.8%) was discontinued due to a protocol deviation (positive drug screen); none of the discontinuations was related to TEAEs. The pharmacokinetic population consisted of 33 patients.

Pharmacokinetics

Under single-dose administration, there was rapid absorption of NRL-1 across all doses (Table 1), with plasma concentration-time profiles that showed similar patterns of exposure that appeared to be dose-dependent (Figure 1A; Table 2). The arithmetic mean C_max values were 85.6, 133.6, and 235.3 ng/mL for the 5, 10, and 20 mg doses, respectively (Table 1). The arithmetic mean t_1/2 and CL/F were comparable among the 3 doses (Table 1).

The plasma concentration-time profiles for nordiazepam generally showed similar patterns to those of diazepam across the 3 NRL-1 doses (Figure 1B), with identical median t_max values of 96 hours. Nordiazepam C_max appeared to be dose-dependent (Table 1), although there was substantial interpatient variability, which ranged from 48% to 61% across the doses, and t_1/2 was lowest after 10-mg administration.

In the dose proportionality assessment for diazepam, the slope estimates were 0.81 (90%CI 0.62, 1.00) for C_max and 0.93 (90%CI 0.80, 1.06) for AUC_0-t. Both of the 90%CIs exceeded the formal dose proportionality criteria on the lower limit (Figure 2). The 90%CIs were also exceeded for the evaluated parameters of nordiazepam (C_max, AUC_0-t, and AUC_0-%/2; Figure 2).

The pharmacokinetic parameters observed during the multiple-dose phase are summarized in Table 2. Under conditions of multiple dosing, the diazepam geometric mean C_max value after the second dose (C_max,4-8h, 114.6 ng/mL; CV% 113%) was approximately 1.25 times higher than that after the first dose (C_max,0-4h, 91.5 ng/mL; CV% 100%). Similarly, the diazepam geometric mean for drug exposure was 1.54 times higher after the second dose (AUC_4-8h) relative to that of the first dose (AUC_0-4h, 327 h·ng/mL (CV% 97%) and 212 h·ng/mL (CV%, 98%), respectively. In the case of nordiazepam, the C_max and AUC geometric means for the second dose were, respectively, approximately 1.75 and 2.19 times higher than those for the first dose.

Comparison of the 2 × 10-mg dose with the single 10-mg dose using box plots showed that values for
Figure 1. Arithmetic mean plasma concentration-time profiles of diazepam (A) and nordiazepam (B) after single-dose administration of NRL-1.

$C_{\text{max},0-4\text{h}}$ were similar to that of the single-dose 10 mg $C_{\text{max}}$ (Figure 3A). The dose-normalized AUC$_{0-\infty}$ values were also comparable between the $2 \times 10$-mg and single 10-mg doses (Figure 3B).

Safety and Tolerability

Nearly all the subjects (97.2%) had at least 1 TEAE, and the TEAEs were considered treatment-related in 94.4% of the subjects (Table 3). There were no serious TEAEs and no discontinuations due to TEAEs. Somnolence was the most frequent TEAE (94.4%) and had the highest incidence with multiple dosing (61.1%). Most of the other common TEAEs occurred only with multiple dosing (Table 3).

There were no reports of pain on the pain visual analogue scale after administration of NRL-1, and nasal pain was not reported as a TEAE. No dose- or time-related trends were observed in nasal irritation, and the majority of subjects (65.5%-100%) had no nasal irritation (data not shown). Nearly all of the nasal irritation that was observed was rated as a score of 1 (“inflamed mucosa, no bleeding”), and the highest score was 2 (“minor bleeding which stops within 1 minute”), which was observed in the same subject at 1 and 2 hours after administration of the 5-mg dose. Changes in olfaction were generally small and did not appear to be dose dependent (data not shown).

There were no trends in sedation (data not shown), and all but 1 of the sedation scores were 2 (“napping or sleeping, but easily awakened”) or less; the 1 exception was a subject who had a score of 3 (“sleeping, awakened only with loud voice or shaking”) at 2 hours after administration of a 20-mg dose.

Blood oxygen saturation levels postdose were consistently within normal limits (94% to 100%). Suicide ideation and behavior were reported on the Columbia
Figure 2. Analysis of dose proportionality. Broken lines indicate the 90%CI criteria (0.84, 1.16) for meeting dose proportionality. AUC indicates area under the plasma concentration–time curve; AUC<sub>0-∞</sub>, AUC from time of dose to last measurement; AUC<sub>0-t</sub>, AUC from time of dose extrapolated to infinity; C<sub>max</sub>, peak plasma concentration.

Table 2. Summary Statistics of Pharmacokinetic Parameters After Multiple Dose (2 × 10 mg) of NRL-1

| Pharmacokinetic Parameter | Arithmetic Mean (CV%) | Geometric Mean (CV%) |
|---------------------------|-----------------------|----------------------|
| Diazepam                  |                       |                      |
| C<sub>max,0-4h</sub>, ng/mL | 123.0 (73%)          | 91.5 (100%)          |
| C<sub>max,4-8h</sub>, ng/mL | 161.4 (77%)          | 114.6 (113%)         |
| C<sub>max</sub>, ng/mL     | 180.2 (68%)           | 137.2 (97%)          |
| t<sub>max</sub>, h<sup>a</sup>   | 5.3 (0.5, 12)       |                      |
| AUC<sub>0-4h</sub>, h·ng/mL | 282 (71%)           | 212 (98%)            |
| AUC<sub>4-8h</sub>, h·ng/mL | 433 (70%)           | 327 (97%)            |
| AUC<sub>0-8h</sub>, h·ng/mL | 7297 (58%)          | 6067 (73%)           |
| AUC<sub>0-∞</sub>, h·ng/mL | 7936 (57%)          | 6831 (61%)           |
| t<sub>1/2</sub>, h<sup>a</sup> | 74 (45%)           | 67 (50%)             |
| CL/F, L/h                 | 3.4 (59%)            | 2.9 (61%)            |
| Nordiazepam               |                       |                      |
| C<sub>max,0-4h</sub>, ng/mL | 6.6 (71%)           | 5.2 (81%)            |
| C<sub>max,4-8h</sub>, ng/mL | 11.7 (78%)          | 9.1 (82%)            |
| C<sub>max</sub>, ng/mL     | 32.1 (55%)           | 26.7 (75%)           |
| t<sub>max</sub>, h<sup>a</sup> | 144 (24, 240)       |                      |
| AUC<sub>0-4h</sub>, h·ng/mL | 16 (64%)            | 13 (68%)             |
| AUC<sub>4-8h</sub>, h·ng/mL | 3.8 (77%)           | 29 (83%)             |
| AUC<sub>0-8h</sub>, h·ng/mL | 5625 (51%)          | 4796 (68%)           |
| AUC<sub>0-∞</sub>, h·ng/mL | 10 840 (23%)        | 10 595 (26%)         |
| t<sub>1/2</sub>, h<sup>a</sup> | 69 (21%)           | 67 (22%)             |

AUC indicates area under the plasma concentration–time curve; C<sub>max</sub>, peak plasma concentration; CV%, coefficient of variation; t<sub>max</sub>, time from dose to C<sub>max</sub>; t<sub>1/2</sub>, half-time of elimination.

<sup>a</sup>Values are medians (minimum, maximum). Ranges in subscripts indicate time intervals of measurements.

Figure 3. Box-and-whisker plots of diazepam C<sub>max,0-4h</sub> multiple-dose and C<sub>max</sub> of single-dose diazepam (A) and dose-normalized AUC<sub>0-∞</sub> multiple- and single-dose (B) after intranasal administration of NRL-1. The dashed line (- - -) is the median; the solid line (–) is the arithmetic mean. The ends of the “box” (⊥) are the first and third quartiles, and the whiskers show the lowest and highest data values still within 1.5 of the interquartile range of the lower and upper quartiles, respectively. Data values that do not fall between the whiskers are plotted as markers outside of the whiskers.

Suicide Severity Rating Scale by a single subject. This 1 subject, who reported suicidal ideation but no suicidal behavior at baseline and was subsequently enrolled in the study, reported suicidal ideation and behavior post-dose on day 0 after administration of 10 mg but on no other study days.

Discussion

The use of DDM and vitamin E has resolved the issues related to the nonaqueous nature of diazepam by
enhancing both solubility and absorption of diazepam; NRL-1 is the only nonaqueous drug formulation that uses DDM. As shown by the pharmacokinetic results, administration of NRL-1 in healthy adult volunteers resulted in rapid absorption of diazepam, with a $C_{\text{max}}$ that was reached approximately 1.5 hours after a nominal lag time. Diazepam $C_{\text{max}}$ after intranasal administration of NRL-1 was lower than what may be expected based on the pharmacokinetics of intravenous (95-125 ng/mL) and oral (148-255 ng/mL) administration. However, $C_{\text{max}}$ and exposure (AUC) were generally consistent with the range that has been reported with rectal diazepam.19-23

Systemic exposure increased with higher doses. However, formal testing of dose proportionality was inconclusive due to the intersubject variability that is a well-recognized characteristic of the pharmacokinetics of diazepam.13,19,22,24,25 Despite this variability, the observed exposure over the dose range suggests that there is likely to be a predictable and consistent dose-response relationship. With further regard to the variability of NRL-1, it should be noted that the CV% of the geometric mean $C_{\text{max}}$ and AUC in the current study, which included subjects over a wide weight range, was consistent with what has been previously reported in a study that compared NRL-1 with rectal diazepam administered to healthy volunteers under “ideal” conditions (ie, fasting state with an enema the night prior to dosing and a second enema 1 hour prior to dosing the following morning).23 In that study, NRL-1 demonstrated less variability than rectal diazepam, which was characterized by pharmacokinetics that appeared to be weight dependent, with substantially higher CV% among heavier subjects. Taken together, these results suggest that, regardless of weight, NRL-1 provides more reliable pharmacokinetics than rectal administration.

In the double-dose period of the current study, the observed absorption and exposure indicated that it is easy to increase the dose as needed. Multiple doses of NRL-1 (2 × 10 mg separated by 4 hours) resulted in an initial diazepam $C_{\text{max,0-4h}}$ and AUC$_{0-4h}$ comparable to that observed for the single 10-mg dose. Dose-normalized AUCs were similar between the single and multiple 10-mg doses. The second dose (administered 4 hours following the first dose) resulted in accumulation in diazepam $C_{\text{max}}$ and AUC 4 hours postdose. These observations are in contrast to concentration effects that often drive exposure with dermal and transmembrane administration.26-28 It should also be noted that the approximately 2-fold higher $C_{\text{max}}$ and AUC values of nordiazepam after the second dose relative to the first dose were not reflective of accumulation, as the consistent rising of nordiazepam concentrations was delayed beyond the 8-hour postinitial dose. However, it should be noted that nordiazepam is an active metabolite, and despite the report that the absolute extent of conversion from diazepam is only about 55%,29 the long half-lives of both diazepam and its metabolite could potentially result in accumulation with frequent dosing, as suggested by a study of oral diazepam pharmacokinetics in obese individuals.14 Although that study showed greater accumulation of both diazepam and nordiazepam under conditions of chronic use among obese individuals, subjects were

### Table 3. Incidence of Treatment-Emergent Adverse Events in the Safety Population (N = 36)

| TEAE                              | Incidence, n (%) |
|-----------------------------------|------------------|
| Any TEAE                          |                  |
| Any TEAE                          | 17 (47.2)        |
| Serious TEAEs                     | 0                |
| Discontinuations due to TEAEs     | 0                |
| Treatment-related TEAEs           | 16 (44.4)        |
| Most common TEAEs (≥5% at any dose) |                  |
| Somnolence                        | 16 (44.4)        |
| Reticulocyte count increased      | 0                |
| Euphoric mood                     | 0                |
| Blood glucose increased           | 0                |
| Hemoglobin decreased              | 0                |
| Urine analysis abnormal           | 0                |
| Alanine aminotransferase increased| 0                |
| Blood pressure decreased          | 0                |

TEAE indicates treatment-emergent adverse event.
dosed daily for 1 month, a dosing frequency that would not be generally expected when used as a rescue medication; NRL-1 is an intermittent treatment for seizure clusters that is unlikely to create accumulation.

Although NRL-1 has not been evaluated in humans in the absence of DDM, it may be expected that the favorable pharmacokinetic characteristics that support this intranasal formulation are derived from the increased solubility and drug absorption associated with the proposed mechanism of DDM. This mechanism is based on enhanced absorption via relaxation of tight junctions combined with fluidization and greater penetration through cell membranes.

Overall, NRL-1 demonstrated a good safety profile in these healthy adult volunteers, with TEAEs that were consistent with what may be expected for diazepam. Although somnolence was the main TEAE, only low levels of transient sedation were reported, and no evidence for respiratory depression was observed. Furthermore, there were no clinically relevant changes in any of the clinical/laboratory tests. Administration of NRL-1 demonstrated good tolerability, with no reports of pain and only minimal nasal irritation and changes in olfaction that were not clinically relevant.

This study was performed in healthy adult volunteers, and the results cannot be directly extrapolated to patients with epilepsy and to pediatric patients in particular. However, preliminary evidence from a recent phase 1 study in patients with epilepsy suggests that the pharmacokinetics and safety of NRL-1 administered to children and adolescents under ictal/peri-ictal, and nonictal conditions are similar to those in adults.

Conclusions

We may conclude there to be good absorption and exposure of diazepam with intranasal administration of NRL-1 across the range of dosing; exposure with increasing dose was indicative of dose proportionality. NRL-1 was also associated with overall safety and tolerability consistent with what is known for diazepam, which has been well characterized. These results further support NRL-1 as a potential novel therapeutic approach to the management of seizure emergencies using a formulation that provides easy and socially acceptable administration during the intermittent occurrence of seizure clusters.

Conflicts of Interest

S.T. is an employee of Pacific Link Consulting Services, which provided consulting services to Neurelis, and has a leadership position with ARS Pharmaceuticals, Inc, which has an agreement with Neurelis for the use of Intravail. L.P.K. is an officer of DUCK FLATS Pharma, which provided consulting services to Neurelis. R.E.L. is an employee of Pacific Link Consulting Services, which provided consulting services to Neurelis, and has a leadership position with ARS Pharmaceuticals, Inc, which has an agreement with Neurelis for the use of Intravail. B.K. is an employee of DUCK FLATS Pharma, which provided consulting services to Neurelis. A.L.R. is an employee and has received stock options from Neurelis. E.C. has received personal compensation for consulting with Neurelis, Alexza, Marinus, and Zogenix and holds stock in Neurelis. E.C. has received compensation for serving on the Board of Directors of Marinus Pharmaceuticals, Epalex, and Hawaii-Biotech.

Acknowledgments

Medical writing support was provided at the direction of the authors by E. Jay Bienen, PhD, of The Curry Rockefeller Group. The Curry Rockefeller Group, LLC, provided additional editorial assistance including formatting and proofreading, funded by Neurelis, Inc.

Funding

This study was supported by Neurelis, Inc.

Data-Sharing Statement

Neurelis, Inc will not be sharing individual deidentified participant data or other relevant study documents.

References

1. G. B. D. Epilepsy Collaborators. Global, regional, and national burden of epilepsy, 1990-2016: a systematic analysis for the Global Burden of Disease Study 2016. Lancet Neurol. 2019;18(4):357-375.
2. Centers for Disease Control and Prevention. Epilepsy Data and Statistics. https://www.cdc.gov/epilepsy/data/index.html. Accessed April 22, 2019.
3. Laxer KD, Trinka E, Hirsch LJ, et al. The consequences of refractory epilepsy and its treatment. Epilepsy Behav. 2014;37:59-70.
4. Jafarpour S, Hirsch LJ, Gainza-Lein M, Kellinghaus C, Detyniecki K. Seizure cluster: definition, prevalence, consequences, and management. Seizure. 2019;68:9-15.
5. Kapoor M, Cloyd JC, Siegel RA. A review of intranasal formulations for the treatment of seizure emergencies. J Control Release. 2016;237:147-159.
6. Acorda Therapeutics. PLUMIAZ™ (diazepam) Nasal Spray [press release]. http://www.acorda.com/products/research-development/plumiaz. Accessed July 1, 2019.
7. Allan A, Cullen J. Best BETs from the Manchester Royal Infirmary. BET 1: intranasal lorazepam is an acceptable alternative to intravenous lorazepam in the control of acute seizures in children. Emerg Med J. 2013;30(9):768-769.
8. Maglalang PD, Rautiola D, Siegel RA, et al. Rescue therapies for seizure emergencies: new modes of administration. Epilepsia. 2018;59 Suppl 2:207-215.

9. Bancke LL, Dworak HA, Rodvold KA, Halvorsen MB, Gidal BE. Pharmacokinetics, pharmacodynamics, and safety of USL261, a midazolam formulation optimized for intranasal delivery, in a randomized study with healthy volunteers. Epilepsia. 2015;56(11):1723-1731.

10. Detyniecki K, Van Ess PJ, Sequeira DJ, Wheless JW, Meng TC, Pullman WE. Safety and efficacy of midazolam nasal spray in the outpatient treatment of patients with seizure clusters—a randomized, double-blind, placebo-controlled trial. Epilepsia. 2019;60(9):1797-1808.

11. Wheless JW, Meng TC, Van Ess PJ, Detryniecki K, Sequeira DJ, Pullman WE. Safety and efficacy of midazolam nasal spray in the outpatient treatment of patients with seizure clusters: An open-label extension trial. Epilepsia. 2019;60(9):1809-1819.

12. Maggio ET, Pillon DJ. High efficiency intranasal drug delivery using Intravail® alkylsaccharide absorption enhancers. Drug Deliv Transl Res. 2013;3(1):16-25.

13. Agarwal SK, Kriel RL, Brundage RC, Ivaturi VD, Cloyd JC. A pilot study assessing the bioavailability and pharmacokinetics of diazepam after intranasal and intravenous administration in healthy volunteers. Epilepsy Res. 2013;105(3):362-367.

14. Abernethy DR, Greenblatt DJ, Divoll M, Shader RI. Prolonged accumulation of diazepam in obesity. J Clin Pharmacol. 1983;23(8-9):369-376.

15. Posner K, Brown GK, Stanley B, et al. The Columbia-Suicide Severity Rating Scale: initial validity and internal consistency findings from three multisite studies with adolescents and adults. Am J Psychiatry. 2011;168(12):1266-1277.

16. Dalton P, Doty RL, Murphy C, et al. Olfactory assessment using the NIH Toolbox. Neurology. 2013;80(11 Suppl 3):S32-S36.

17. Sethuraman VS, Leonov S, Squassante L, Mitchell TR, Hale MD. Sample size calculation for the power model for dose proportionality studies. Pharm Stat. 2007(6):1.

18. Divoll M, Greenblatt DJ, Ochs HR, Shader RI. Absolute bioavailability of oral and intramuscular diazepam: effects of age and sex. Anesth Analg. 1983;62(1):1-8.

19. Cloyd JC, Lalonde RL, Beniak TE, Novack GD. A single-blind, crossover comparison of the pharmacokinetics and cognitive effects of a new diazepam rectal gel with intravenous diazepam. Epilepsia. 1998;39(5):520-526.

20. Henney HR 3rd, Sperling MR, Rabinowicz AL, Bream G, Carrazana EJ. Assessment of pharmacokinetics and tolerability of intranasal diazepam relative to rectal gel in healthy adults. Epilepsy Res. 2014;108(7):1204-1211.

21. Lamson MJ, Sitki-Green D, Wannarka GL, Mesa M, Andrews P, Pelloch J. Pharmacokinetics of diazepam administered intramuscularly by autoinjector versus rectal gel in healthy subjects: a phase I, randomized, open-label, single-dose, crossover, single-centre study. Clin Drug Invest. 2011;31(8):585-597.

22. Ivaturi V, Kriel R, Brundage R, Loewen G, Mansbach H, Cloyd J. Bioavailability of intranasal vs. rectal diazepam. Epilepsy Res. 2013;103(2-3):254-261.

23. Hogan RE, Gidal BE, Koplowitz B, Koplowitz LP, Lowenthal RE, Carrazana E. Bioavailability and safety of Valtoco™ (diazepam intranasal solution) compared to oral and rectal diazepam. Poster presented at the 72nd Annual Meeting of the American Epilepsy Society, November 30–December 4, 2018, New Orleans, LA.

24. Ivaturi VD, Riss JR, Kriel RL, Cloyd JC. Pharmacokinetics and tolerability of intranasal diazepam and midazolam in healthy adult volunteers. Acta Neurol Scand. 2009;120(5):353-357.

25. Ivaturi VD, Riss JR, Kriel RL, Siegel RA, Cloyd JC. Bioavailability and tolerability of intranasal diazepam in healthy adult volunteers. Epilepsy Res. 2009;84(2-3):120-126.

26. Schneider T, Vermeulen R, Brouwer DH, Cherrie JW, Kromhout H, Fogh CL. Conceptual model for assessment of dermal exposure. Occup Environ Med. 1999;56(11):765-773.

27. Tregear RT. Physical functions of skin. London, New York: Academic Press; 1966.

28. Wester RC, Maibach HI. Relationship of topical dose and percutaneous absorption in rhesus monkey and man. J Invest Dermatol. 1976;67(4):518-520.

29. Greenblatt DJ, Divoll MK, Soong MH, Boxenbaum HG, Hartzman JS, Shader RI. Desmethylzolpidem pharmacokinetics: studies following intravenous and oral desmethylzolpidem, oral clorazepate, and intravenous diazepam. J Clin Pharmacol. 1988;28(9):853-859.

30. Tarquinio D, Segal E, Wheless JW, Rabinowicz AL, Carrazana E, the DIAZ 001.04 Study Group. Pharmacokinetics and safety of Valtoco™ (NRL-1; diazepam nasal spray) in children with epilepsy during seizure (ictal/peri-ictal) and non-seizure (inter-ictal) conditions: results from a phase I, open-label study. Poster presented at the 48th Annual Meeting of the Child Neurology Society, October 23-26, 2019, Charlotte, NC.