Geographic Regional Differences in Rocuronium Bromide Dose–Response Relation and Time Course of Action

An Overlooked Factor in Determining Recommended Dosage


Background: Geographic location is not acknowledged as a stratifying factor that can directly affect drug potency, because drugs are still licensed with the same recommended dose for different geographic regions. The aim of the current study was to compare the potency and duration of action of rocuronium bromide in 54 patients in three countries with different life habits, diet, and ambient conditions, namely white Austrians, white North Americans, and Han Chinese in China.

Methods: Neuromuscular block of six consecutive 50-μg/kg rocuronium incremental doses followed by 300 μg/kg was evaluated using the Relaxometer mechanomyograph (Groningen University, Groningen, Holland). Dose–response curves were created using log-dose-probit transformation. The authors compared rocuronium bromide ED50, ED90, and ED95 (effective doses required for 50%, 90%, and 95% first twitch depression, respectively) as well as Dur25 and Dur50 (times from last incremental dose administration until 25% and 50% first twitch and 0.8 train-of-four ratio recovery, respectively) in patients of the three countries.

Results: Rocuronium ED50, ED90, and ED95 were significantly higher in Austrian patients (258 ± 68, 530 ± 159, and 598 ± 189 μg/kg) and Chinese patients (203 ± 59, 413 ± 107, and 475 ± 155 μg/kg) compared with American patients (148 ± 48, 316 ± 116, and 362 ± 149 μg/kg, respectively). Dur25 and Dur50 were significantly shorter in Austrian patients (22.3 ± 5.5 and 36.9 ± 12.8 min) and Chinese patients (30.4 ± 7.5 and 45.7 ± 15.9 min) compared with American patients (36.7 ± 8.5 and 56.2 ± 16.7 min, respectively).

Conclusions: The authors demonstrated a significant difference in rocuronium potency and duration of action among patients in the three countries. Larger studies are required for determining dosage recommendations for different geographic regions.

VARIABILITY of drug responsiveness could result from several factors. However, geographic location is not acknowledged as a stratifying factor that can directly affect drug potency. This has been a relatively neglected area of investigation, and drugs are still licensed with the same recommended dose for different geographic locations. In 1969, Ronald Katz1 described how, during the course of a 1-year sabbatical, he was surprised to observe clinically that the dose of tubocurarine that regularly abolished the twitch response to stimulation of the ulnar nerve in white patients in New York, New York, did not do so in white patients in London, England. He was fascinated by the fact that American anesthesiologists routinely used doses of tubocurarine that seem “homeopathic” to their English colleagues. What he found even more intriguing was the fact that the manufacturing company, in response to demand, prepared tubocurarine in two different concentrations, one for each country on both sides of the Atlantic Ocean. Recently, Fiset et al.2 demonstrated a similar transatlantic difference in vecuronium potency between Montreal, Quebec, Canada, and Paris, France.

The absence of apparent regional differences in various studies assessing potencies of other neuromuscular blocking drugs such as pancuronium, atracurium, mivacurium, and cisatracurium could be simply attributed to the fact that, to date, no structured study specifically aiming at examining regional difference was conducted for these neuromuscular blocking drugs. Investigations to elucidate the extent of such differences among different regions are confronted by many obstacles that were taken into consideration while designing our study, because we used mechanomyography that can precisely quantify the neuromuscular block; rocuronium was obtained from the same European production line (Organon, Oss, Holland), was transported under the strict transport instructions of the manufacturer, and was given within the shelf life; and one primary investigator (A.A.D.) supervised the strict implementation of the same study protocol in all three countries with use of the same equipment.

The aim of our study was to compare the potency and duration of action of rocuronium bromide in 54 patients in three countries with different life habits, diet, and ambient conditions, namely white Austrians, white North Americans, and Han Chinese of second-generation Han Chinese descent in China.

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Materials and Methods

A prospective controlled clinical consecutive study was conducted in three centers—Englewood Medical Center (Englewood, New Jersey), Dalian Medical University (Dalian, People’s Republic of China), and Graz Medical University (Graz, Austria)—in conformity with the guidelines of the “Consolidated Standards of Reporting Trials (CONSORT) Statement,”3 “Good Clinical Research Practice (GCRP) in Pharmacodynamic Studies of Neuromuscular Blocking Agents,”4 and editorial recommendations.5 After approval was obtained from Englewood Medical center institutional review board, Graz Medical University ethics committee, and Dalian Medical University ethics committee, all patients who agreed to participate in the study gave written informed consent. Potential participants with a history of neuromuscular disease, small joint arthritis, or body mass index < 20 or > 26 kg/m2,6 and patients receiving treatment with drugs thought to interfere with neuromuscular transmission were excluded from the study.

Eighteen white patients, nine males and nine females, with American Society of Anesthesiologists physical status I or II, aged 30–50 yr, and undergoing general anesthesia for scheduled elective surgery were first recruited at Englewood Medical Center. Shortly after that, each recruited patient was matched for sex, American Society of Anesthesiologists physical status, approximate age, and approximate body weight with 18 white patients in Graz University Hospital, followed by 18 matched Han Chinese patients in Dalian University Hospital. Anesthesia was induced with 1.5 μg/kg fentanyl and 2–3 mg/kg propofol until the eyelash reflex was obtunded. A laryngeal mask airway was inserted, and mechanical ventilation was adjusted to maintain 25–35 mmHg end-tidal carbon dioxide. Anesthesia was maintained with 100–150-mg/kg/h fentanyl and 2–4 μg·kg·min−1 remifentanil infusions. Patients were warmed using a forced-hot-air blanket to maintain core temperature above 36°C and skin temperature above 32°C. Estimated fluid and blood losses were replaced by Ringer’s solution.

The neuromuscular block at the adductor pollicis muscle was evaluated using the Relaxometer mechanomyograph (Groningen University, Groningen, Holland).7 After anesthesia induction, the force transducer was attached to the thumb, and the ulnar nerve was stimulated supramaximally at the wrist (pulse width 200 μs, square wave) via surface electrodes with train-of-four (TOF) stimuli (2 Hz for 2 s) at 12-s intervals. Data were continuously recorded using the AZG-Relaxometer 5.0 program (Groningen University, Groningen, Holland). T1 (first twitch of the TOF), expressed as percentage of T0 (first twitch of the TOF), was used for the evaluation of neuromuscular block.

After T1% baseline stabilization, six consecutive 50-μg/kg rocuronium incremental doses were administered into a rapidly running infusion. Each incremental dose was given after three consecutive T1 responses did not register a decline. T1% achieved after each rocuronium bolus was recorded. Rocuronium dose–response curves were created according to the cumulative dose–response method of Donlon et al.8 Individual patients’ dose–response relations were examined by least squares linear regression of the logarithm of each dose against a probit transformation of T1%, from which the doses required for 50%, 90%, and 95% T1% depression (ED50, ED90, and ED95, respectively) were calculated. Because probit values for 0% and 100% do not exist, 0% and 100% T1% depression were considered missing values.

After all six incremental doses, or if 100% twitch depression was achieved after any of the incremental doses, 300 μg/kg rocuronium was administered in addition to the remainder of the incremental doses. Patients were allowed to recover spontaneously from the neuromuscular block, and Dur25 (time from last incremental dose administration until 25% T1 recovery), Dur25–75 (time from 25% to 75% T1 recovery), Dur25–0.8 (time from 25% T1 to 0.8 TOF ratio recovery), and Dur0.8 (time from last incremental dose administration until 0.8 TOF ratio recovery) time-course-of-action variables4 were calculated.

Statistical Analysis

Because ED95 is the basis for determining the recommended dosage for neuromuscular blocking drugs, comparing rocuronium ED95 among patients of the three groups was considered the primary end point of our study. Based on data from a previous study2 of transatlantic difference in vecuronium dose response, where the mean ED95 was 44.2 ± 10.9 μg/kg in Montreal compared with 71.9 ± 30.5 μg/kg in Paris, our a priori power analysis for one-sided t test (α = 0.025 to accommodate for two comparisons according to Bonferroni correction) showed that a group size of 17 patients would be required to reveal a statistically significant difference between the groups with more than 80% power. One-way analysis of variance was used for statistical analysis. The Tukey-Kramer post hoc test was used to determine which intergroup differences were statistically significant. Data were expressed as mean ± SD. P < 0.05 was considered statistically significant.

Results

Patient demographics are presented in table 1. There were no significant differences among the three groups regarding the duration of T1% stabilization, skin temperature, core temperature, mean arterial pressure, estimated fluid and blood losses, and propofol and remifentanil doses during the neuromuscular monitoring period of the study.
Compared with the American patients, the dose–response curve was shifted to the right in the Chinese patients and further shifted to the right in the European patients (fig. 1). Our study showed that there was a statistically significant difference in ED50, ED90, and ED95 among the three groups. Rocuronium ED50, ED90, and ED95 were significantly higher in the European and Chinese patients compared with the American patients (table 2). There was a statistically significant difference in Dur25 and Dur0.8 among the three groups. Dur25 and Dur0.8 were significantly shorter in the European and Chinese patients compared with the American patients (table 3).

Discussion

Our results demonstrated that rocuronium potency was significantly higher in American patients than in similar European patients. A transatlantic difference in sensitivity of neuromuscular blocking drugs was previously reported. Katz et al.1 showed that tubocurarine resulted in significantly more intense neuromuscular block with duration of action that lasted almost twice as long in American patients in New York compared with a similar group of British patients in London. Fiset et al.2 demonstrated that vecuronium was more potent, with a significantly lower ED90, in Montreal patients compared with similar patients in Paris.

The difference in rocuronium sensitivity among the three countries of our current study raises the question of whether “native” sensitivity could be acquired. Collins et al.9 compared rocuronium potency in 18 white patients and 18 second-generation Han Chinese subjects living in Vancouver for a mean of 10 yr. They showed that the difference in ED50 did not reach statistical significance between the Chinese and white patients living in Vancouver. Interestingly, our results were quite different, because the ED50 in our Chinese subjects living in Dalian, People’s Republic of China, were significantly higher than our white subjects in New Jersey. This suggests that the ethnic Chinese subjects of the Collins et al.9 study, who lived in Vancouver for an average of 10 yr, could have acquired the same sensitivity as the native Vancouver white patients.9 Similarly, in the Fiset et al.2 study that demonstrated a significant difference in vecuronium potency between patients in Paris and Montreal, paradoxically half of the Montreal patients were born in Europe but were living in Montreal for more than 10 yr,

Table 1. Patient Demographics

<table>
<thead>
<tr>
<th></th>
<th>Austrian Group</th>
<th>Chinese Group</th>
<th>American Group</th>
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<tbody>
<tr>
<td>Age, yr</td>
<td>37.7 ± 13.2</td>
<td>44.8 ± 6.9</td>
<td>40.8 ± 12.9</td>
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<tr>
<td>Weight, kg</td>
<td>74.4 ± 8.0</td>
<td>68.1 ± 13.1</td>
<td>71.9 ± 12.7</td>
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<tr>
<td>Height, cm</td>
<td>176 ± 8</td>
<td>168 ± 10</td>
<td>177 ± 9</td>
</tr>
<tr>
<td>Body mass index, kg/m²</td>
<td>24 ± 1.1</td>
<td>24.1 ± 0.6</td>
<td>24.5 ± 1.2</td>
</tr>
</tbody>
</table>

No significant differences between the three groups. Data are mean ± SD; n = 18.

Table 2. Rocuronium Effective Doses

<table>
<thead>
<tr>
<th></th>
<th>Austrian Group</th>
<th>Chinese Group</th>
<th>American Group</th>
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</thead>
<tbody>
<tr>
<td>ED50, μg/kg</td>
<td>258 ± 68*</td>
<td>201 ± 59†</td>
<td>148 ± 48†</td>
</tr>
<tr>
<td>ED90, μg/kg</td>
<td>530 ± 159*</td>
<td>413 ± 107§</td>
<td>316 ± 116†</td>
</tr>
<tr>
<td>ED95, μg/kg</td>
<td>598 ± 189*</td>
<td>475 ± 155§</td>
<td>362 ± 149†</td>
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Tukey-Kramer post hoc test. Data are mean ± SD; n = 18.

* P < 0.001 compared with the American group. † P < 0.05 compared with the Chinese group. § P ≤ 0.01 compared with the Austrian group. § P ≤ 0.05 compared with the Austrian group. ED50, ED90, and ED95 = effective doses required for 50%, 90%, and 95% first twitch of train-of-four depression.
Table 3. Rocuronium Time-course-of-action Variables

<table>
<thead>
<tr>
<th></th>
<th>Austrian Group</th>
<th>Chinese Group</th>
<th>American Group</th>
</tr>
</thead>
<tbody>
<tr>
<td>Onset time, min</td>
<td>13.1 ± 3.1</td>
<td>12.5 ± 2.9</td>
<td>12.2 ± 1.3</td>
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<tr>
<td>DUR25–75, min</td>
<td>22.3 ± 5.5†</td>
<td>30.4 ± 7.5‡</td>
<td>36.7 ± 8.5†</td>
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<tr>
<td>DUR25–0.8, min</td>
<td>11.3 ± 3.1</td>
<td>12.7 ± 3.3</td>
<td>13.2 ± 4.7</td>
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<tr>
<td>DUR0.8, min</td>
<td>14.6 ± 9.3</td>
<td>15.3 ± 8.3</td>
<td>19.5 ± 9.4</td>
</tr>
<tr>
<td>DUR0.8–0.6, min</td>
<td>36.9 ± 12.8*</td>
<td>45.7 ± 15.9</td>
<td>56.2 ± 16.7†</td>
</tr>
</tbody>
</table>

Tukey-Kramer post hoc test. Data are mean ± SD; n = 18. *P < 0.001 compared with the American group. †P < 0.05 compared with the Chinese group. ‡P ≤ 0.01 compared with the Austrian group.

Onset time = time from beginning of incremental doses administration until first response of train-of-four (T1) maximum suppression; DUR25 = time from last incremental dose administration until 25% T1 recovery; DUR25–75 = time from 25% to 75% T1 recovery; DUR0.8 = time from last incremental dose administration until 0.8 train-of-four ratio recovery; DUR0.8–0.6 = time from 25% T1 to 0.8 train-of-four ratio recovery.

Hence, they manifested a different sensitivity than the European subjects.

However, Katz et al.1 in addition studied a third group of American citizens, operated on at the US Air Force Hospital in London, who lived in London and its environs for periods of 13–36 months. Interestingly, these American citizens demonstrated sensitivity that was similar to that seen in the New York patients and not the British subjects of the study.1 This probably implies that 3 yr was not a sufficient period to acquire the “native” sensitivity.

Pharmacogenetic studies have shown that ethnic or racial variations in hepatic metabolizing isoenzymes are important demographic variables that may account for differences in drug response among different populations.10 However, rocuronium is eliminated mostly unchanged via biliary excretion,11 whereas the difference in rocuronium potency between the two European and American white groups in our study was much wider than the difference between the ethnic Chinese and white groups, clearly indicating that ethnic or racial difference was not the most influential factor in our results; rather a multifactorial interaction of environmental factors such as pollution, life habits, diet, ambient conditions, and others could be the most likely explanation.

Because the aim of our study was to compare the dose–response relation between different regions and was not intended as an absolute potency estimate, we used the cumulative dose technique rather than the single-dose method, the accepted standard for creating dose–response curves4 that would have required a far larger sample size of 108 patients. The cumulative dose technique could be used for intermediate-acting neuromuscular blocking drugs such as rocuronium, provided that the incremental doses were given within a brief period of time,4,8 such as the 36 s of three consecutive T1 responses that we used in our study. This allowed no significant recovery between the incremental doses as indicated by the short incremental doses onset time among the three groups (table 3).

In conclusion, our results demonstrated a significant difference in rocuronium potency and duration of action among patients in the three countries. Larger studies are required for determining dosage recommendations for different geographic regions.

The authors thank Lisa Hofstatter (Anesthesia Head Nurse, Graz University Hospital, Graz, Austria), Fang Wei (Medical student, Dalian University First Affiliated Hospital, Dalian, People’s Republic of China), and Charles Zhou (Head of Beijing Talent Trade Co. Ltd., Beijing, People’s Republic of China) for their great efforts in collecting and preparing the data.

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