Block of postjunctural muscle-type acetylcholine receptors in vivo causes train-of-four fade in mice

M. Nagashima¹, T. Sasakawa¹, S. J. Schaller¹,², and J. A. J. Martyn¹,*

¹Department of Anaesthesia, Critical Care and Pain Medicine, Massachusetts General Hospital, Shriners Hospitals for Children®–Boston, and Harvard Medical School, Boston, MA, USA, and ²Klinikum Rechts der Isar, Technische Universität Münche, Klinik für Anaesthesiologie, München, Germany

*Corresponding author. E-mail: jmartyn@mgh.harvard.edu

Abstract

Background: Train-of-four (TOF) fade during nerve-mediated muscle contraction is postulated to be attributable to inhibition of prejunctural nicotinic α3β2 acetylcholine receptors (nAChRs), while decrease of twitch tension is attributable to block of postjunctural muscle nAChRs. The validity of these presumptions was tested using specific prejunctural and postjunctural nAChR antagonists, testing the hypothesis that fade is not always a prejunctual phenomenon.

Methods: Pentobarbital anaesthetized mice had TOF fade measured after administration of: either 0.9% saline; the prejunctural α3β2 nAChR antagonist, dihydro-β-erythroidine (DHβE); the postjunctural nAChR antagonists, α-bungarotoxin (α-BTX) or α-conotoxin GI; and a combination of α-BTX and DHβE; or a combination of α-conotoxin GI and DHβE.

Results: Saline caused no neuromuscular changes. Administration of muscle nAChR antagonists, α-BTX or α-conotoxin GI caused significant decrease of twitch tension and TOF fade compared with baseline (P<0.01). DHβE alone caused no change of twitch tension or fade even after 90 min, but its coadministration with α-BTX or α-conotoxin GI significantly accelerated the onset of paralysis and degree of fade compared with α-BTX or α-conotoxin GI alone (P<0.01).

Conclusions: Occupation of postjunctural nAChRs alone by α-BTX or α-conotoxin GI causes fade. As the prejunctual effects of DHβE on fade became manifest only when co-administered with α-BTX or α-conotoxin GI, specific inhibition of prejunctural nAChR alone is not necessary and sufficient to cause fade. Fade observed during repetitive nerve stimulation can be because of block of either postjunctural nAChRs alone, or block of prejunctural and postjunctural nAChRs together.

Key words: acetylcholine; neuromuscular monitoring; neuromuscular block; neuromuscular transmission; receptors; nicotinic

Editor’s key points
- The fade of train-of-four (TOF) is thought to be a pre-synaptic effect.
- Specific pre- and post-junctional nicotinic acetylcholine receptor (nAChR) antagonists were used in mouse model.
- Blocking post-junctional nAChRs caused fade.
- Fade is not always a prejunctual phenomenon; block of postjunctural nAChRs can also cause fade.

Neuromuscular relaxants are used extensively in operating and emergency rooms and intensive care units.¹⁻³ After the use of relaxants, assessment of neuromuscular function is pivotal because relaxant-induced residual paralysis increases morbidity and mortality.⁴⁻⁶ Typically, the train-of-four (TOF) fade in muscle during 2 Hz nerve stimulation is used to assess neuromuscular function.⁵⁻⁷ Fade in muscle during repetitive nerve stimulation is postulated to be a prejunctural phenomenon, while depression of twitch tension can be either prejunctural in nature because of decreased release of acetylcholine (ACh) or a postjunctural consequence because of decreased availability of muscle nicotinic acetylcholine receptors (nAChR).²⁻⁶ Some investigators have posited that the prejunctional nAChRs enhance mobilization of
ACh by positive feedback to maintain tension at the same level during repetitive nerve stimulation; block of the prejunctional nAChRs by neuromuscular blocking agents attenuates ACh mobilization leading to fade.\textsuperscript{7,8} Based on ex vivo phrenic-diaaphragm and oocyte expression studies, the putative receptor mediating enhanced mobilization of ACh or fade is said to be the \(\alpha_3\beta_2\) nAChRs.\textsuperscript{7,9,10} Oocyte expression studies have confirmed that several clinically used relaxants do inhibit \(\alpha_3\beta_2\) nAChRs.\textsuperscript{10} The relationship of this inhibition of \(\alpha_3\beta_2\) to TOF fade, however, has not been demonstrated. Immunohistochemistry has demonstrated the presence of \(\alpha_3\) containing nAChR isoforms prejunctionally.\textsuperscript{12}

Specific antagonists used to study \(\alpha_3\beta_2\) nAChRs ex vivo include dihydro-\(\beta\)-erythrodine (DH\(\beta\)E) or \(\alpha\)-conotoxin MII.\textsuperscript{11,13,14} When DH\(\beta\)E or \(\alpha\)-conotoxin-MII was used to block \(\alpha_3\beta_2\) nAChRs in phrenic-diaaphragm preparation of rat, fade was demonstrated, but only when the safety margin of neuromuscular transmission was concomitantly decreased with high magnesium concentrations. On the other hand, specific block of postjunctional muscle nAChRs with \(\alpha\)-bungarotoxin (\(\alpha\)-BTX) alone also induced TOF fade in rats.\textsuperscript{15} Thus, the exact role of prejunctional vs postjunctional nAChRs in the mediation of fade is unclear. The response to nAChR antagonists can differ between species. For example, waglerin-1, a potent antagonist of muscle nAChRs in mice has minimal effects in rats.\textsuperscript{16} Therefore, fade observed during postsynaptic nAChRs block\textsuperscript{15} may be restricted to rats, and not observed in another rodent species.

This study in mice, using specific postjunctional nAChR antagonists, \(\alpha\)-conotoxin GI (from conus snail Conus geographus) and \(\alpha\)-BTX (from cobra Bungarus multicinctus), tested the hypothesis that fade is not always a prejunctional phenomenon and that decreased availability of muscle nAChRs, is necessary and sufficient to cause fade. In separate experiments, the role of prejunctional, \(\alpha_3\beta_2\) nAChRs was tested using the specific antagonist, DH\(\beta\)E, alone and in combination with the specific postjunctional antagonists, \(\alpha\)-BTX or \(\alpha\)-conotoxin GI.

Methods

Animals

The study protocol was approved by the institutional animal care committee (approval number 2011N000013). Relevant aspects of the ARRIVE guidelines were followed. Male C57/Black 6 mice (Jackson laboratory, MA), weighing 23–30 g, were used. After at least one week of acclimatization at our animal facility, 7 groups of mice (n=5–6 per group) were randomly allocated to receive one of the following drug regimens during nerve-evoked gastrocnemius–soleus muscle contractions: (1) 0.9% normal saline (NS), (2) the specific postsynaptic muscle nAChR blocker (\(\alpha\)-BTX 0.175 µg g\(^{-1}\)) EMD Chemicals, Inc. Gibbstown, NJ), or (3) the specific postsynaptic muscle nAChR blocker (\(\alpha\)-conotoxin GI 30 ng g\(^{-1}\)) American Peptide Company, Sunnyvale, CA), (4) the specific presynaptic \(\alpha_3\beta_2\) nAChR blocker (DH\(\beta\)E 1.0 µg g\(^{-1}\)) Tocris Bioscience, Ellisville, MO), or a combination of presynaptic and postsynaptic nAChR antagonists: (5) \(\alpha\)-BTX 0.175 µg g\(^{-1}\) and DH\(\beta\)E 0.5 µg g\(^{-1}\), (6) \(\alpha\)-BTX 0.175 µg g\(^{-1}\) and DH\(\beta\)E 1.0 µg g\(^{-1}\), or (7) \(\alpha\)-conotoxin GI 30 ng g\(^{-1}\) and DH\(\beta\)E 1.0 µg g\(^{-1}\).

Neuromuscular function study

For the measurement of twitch tension and fade, the mice were anaesthetized with pentobarbital (50–70 mg kg\(^{-1}\) intraperitoneally (i.p.)). Supplemental intermittent doses of pentobarbital 10–20 mg kg\(^{-1}\) i.p., were empirically administered every 20–30 min to maintain adequate anaesthesia, ascertained by the withdrawal response to a nociceptive stimulus. Tracheostomy was performed for mechanical ventilation with air at 110 breaths per minute with a tidal volume of 6–8 ml kg\(^{-1}\) (MiniVent Type 845, Harvard Apparatus, Holliston, MA). A venous catheter (MRE-025, Braintree scientific, Braintree, MA) was inserted into the right jugular vein and temperature was monitored with a rectal probe and maintained at 36–38°C using a heating pad.

Neuromuscular transmission and function were monitored by evoked mechanomyography using a Grass Stimulator 88, along with a Grass force transducer and software (Grass Technologies, West Warwick, RI). For this purpose, mice were placed in the dorsal recumbent position. The gastrocnemius–soleus muscle group tendons were tied with non-compliant silk and attached to separate Grass FT03 force displacement transducers. Both sciatic nerves were exposed at the thigh and tied with ligatures. Distal to the ligatures, stimulation electrodes (Subminiature Electrode, Harvard Apparatus, Holliston, MA) were attached for nerve-mediated stimulation of the muscle. The knees were stabilized rigidly with a clamp. A baseline tension of approximately 10 g was applied to the tendon of each muscle, which yielded optimal evoked tensions. The nerve-evoked muscle tension was recorded via a Grass P122 amplifier and displayed using the Grass Polyvibe Software.

During supramaximal stimuli, tetanic stimulation (50 Hz, 5 s) was applied and baseline mechanomyographic responses were stabilized over a period of 15 min using the TOF stimulation pattern (2 Hz per sec), every 20 s. At the end of this period the evoked muscle tension and fade developed during TOF stimulation were recorded as pre-drug administration (baseline) value. Subsequently, the test drugs alone or in combination were administered i.v. using a Hamilton syringe (Hamilton, Reno, NV). The drugs were reconstituted with 0.9% sterile saline and adjusted to a volume of 50 µl. After drug administration, 30 µl 0.9% sterile saline was administered to flush the drug through the venous catheter. The control group received 80 µl of saline. TOF fade was calculated as the T4/T1 ratio, where T4 and T1 are the fourth and first twitch tensions in the same train. Single twitch tension was defined as first twitch tension during TOF stimulation. All animals were killed at the end of the experiments with an overdose of pentobarbital (200 mg kg\(^{-1}\)).

Data and statistical analyses

Statistical analyses were performed using Mann–Whitney test when two groups were compared. The Kruskal–Wallis test or the Friedman test followed by Dunn’s multiple comparison test was used when more than two groups were compared. The results are expressed as median (range). P-values of <0.05 were considered significant.

Results

In the control group, intravenous injection saline did not change twitch tension or TOF fade compared with the pre-administration value for up to 90 min after injection (data not shown). Administration of DH\(\beta\)E alone also had no significant effect on twitch depression or TOF fade for up to 90 min, the end of the observation period (Fig. 1). This lack of neuromuscular effect after DH\(\beta\)E also confirms the stability of the preparation for up to 90 min. In the next group of mice, after stabilization of twitch tension, intravenous administration of \(\alpha\)-BTX caused twitch depression within a few minutes. At 75, 50, and 25% of baseline twitch
tension (25, 50 and 75% depression) of the first twitch of TOF (T1) after α-BTX, the fade associated with these twitch depressions were 0.85 (0.81–0.89), 0.76 (0.71–0.81), and 0.69 (0.64–0.71), respectively; all except the TOF at 75% of baseline twitch tension were significantly decreased compared to the pre-administration value (P = 0.425, P = 0.009, and P < 0.0001, respectively; Fig. 2). This dose of α-BTX caused 100% twitch depression over the time course, and the recovery of twitch tension was not observed during the observation period of 60 min after administration.

In another set of mice, intravenous DHE 0.5 or 1.0 µg g−1 was administrated together with α-BTX. With the administration of DHE 1.0 µg g−1 together with α-BTX, the fade was significantly accentuated at 75, 50, and 25% of baseline tension compared to α-BTX alone (P < 0.001, P = 0.003, and P = 0.003, respectively; Fig. 2). There was significant dose-dependent potentiation of the TOF fade by DHβE (linear trend; P < 0.001 at 75% T1, P < 0.001 at 50% T1, and P < 0.001 at 25% of baseline (T1) tension; Fig. 2).

In mice given α-conotoxin GI alone, significant twitch depression and TOF fade was seen compared with pre-administration values at 50% of baseline tension (TOF fade = 0.58 (0.54–0.72); P = 0.010) and at 25% of baseline tension (TOF fade = 0.55 (0.48–0.68); P = 0.002; Fig. 3). The administration of DHβE (1.0 µg g−1) together with α-conotoxin GI significantly potentiated the TOF fade caused by α-conotoxin GI alone (P = 0.008 at 75% of baseline tension, P = 0.008 at 50% of baseline tension, and P = 0.008 at 25% depression of baseline tension; Fig. 3).

In view of the potentiation of fade by DHβE when co-administered with α-BTX or α-conotoxin GI the time course of the onset of the effects to decrease of tension to 75, 50 and 25% of baseline tension were examined. The administration of α-BTX alone caused a decrease of tension to 75, 50, and 25% of baseline tension were examined. The administration of α-BTX alone caused a decrease of tension to 75, 50, and 25% of baseline tension in approximately 10, 15 and 24 min, respectively. The combined administration of α-BTX and DHβE (1.0 µg g−1) resulted in a faster onset of effect of 7 min (P = 0.008), 10 min (P = 0.040), and 18 min (P = 0.716) for decrease of baseline tension to 75%, 50%, and 25% of baseline tension compared with a α-BTX alone. With α-conotoxin GI alone and in combination with DHβE, the onset of effect for decrease of tension to 75, 50, and 25% of baseline was approximately 4, 5, 6 min alone, vs 3 min (P = 0.004), 4 min (P = 0.009), 5 min (P = 0.030), respectively, for the combination (Table 1).

Discussion

These studies in mice demonstrate that block of the postsynaptic nAChRs by i.v. administration of α-BTX or α-conotoxin GI, per se, is capable of inducing fade. Injection of the presynaptic nAChR (α3β2) antagonist, DHβE, alone does not cause decrease of twitch tension or TOF fade but when administered with α-BTX or α-conotoxin GI, potentiates the TOF fade produced by the post synaptic antagonists. DHβE with α-BTX or α-conotoxin GI also accelerated the onset of neuromuscular paralysis.
α-BTX has postjunctional effects only,\textsuperscript{17} \textsuperscript{18} and does not bind to αβ2 nAChRs\textsuperscript{14} \textsuperscript{15} or to any prejunctional nAChRs, as demonstrated by the lack of modulation of [\textsuperscript{3}H]-ACh release during high frequency nerve stimulation in the rat phrenic nerve-hemidiaphragm.\textsuperscript{11} \textsuperscript{20} Thus, the fade observed after α-BTX is postjunctional phenomenon and is not because of any prejunctional effects. Some previous studies with α-BTX, however, have failed to show fade during ex vivo experiments.\textsuperscript{11} \textsuperscript{21} Concordant with our current observations, some other ex vivo experiments did show fade with prolonged incubation with α-BTX.\textsuperscript{22} This discrepancy in the lack of fade in some studies might be explained by the very slow onset time (150 min) when lower concentrations of α-BTX are used ex vivo.\textsuperscript{23} Regardless of the lack of fade in some of the ex vivo experiments, no study has documented any prejunctional effects of α-BTX.

In order to exclude the possibility that the fade observed with α-BTX was not limited to that drug only, another specific postjunctional antagonist, α-conotoxin GI which has no effects on αβ2 nAChRs, was tested.\textsuperscript{24} \textsuperscript{26} Previously, the administration of α-conotoxin GI had no effects on arterial blood pressure or heart rate, and did not modulate responses to vagus and preganglionic stimulation, all of which confirmed the absence of any autonomic side-effects.\textsuperscript{24} Therefore, the twitch depression and TOF fade observed after injection of α-conotoxin GI is a postjunctional effect and not related to autonomic effects. The fade observed with decreased availability of muscle nAChRs induced by α-BTX or α-conotoxin GI is consistent with the observations in the pathological state of myasthenia gravis, a postjunctional disease, where fade is observed because of autoantibody-mediated decrease of muscle nAChRs.\textsuperscript{24} \textsuperscript{28}

In view of the previous elegant documentations that DH\textsubscript{i}E (1 µM) effectively decreased prejunctional ACh release,\textsuperscript{11} or efficiently blocked ACh-induced currents in αβ2 nAChRs expressed in cells,\textsuperscript{29} our study did not quantify the effect of DH\textsubscript{i}E on ACh release. In the phrenic nerve, the DH\textsubscript{i}E (1 µM)-mediated decrease of ACh release was similar to that produced by α-conotoxin MII (0.1 µM) and d-tubocurarine (1 µM) (50–60%).\textsuperscript{11} Despite this decrease of ACh release by DH\textsubscript{i}E and α-conotoxin MII, no fade was observed until the margin of safety was decreased by very high concentrations of Mg\textsuperscript{2+} (6–7 mM).\textsuperscript{11} Normal Mg\textsuperscript{2+} concentration is 0.7–1.0 mM. DH\textsubscript{i}E 1 µM, which reduced [\textsuperscript{3}H]-ACh release 40% at 75% depression of baseline twitch tension (Figs 2 and 3), even before deep postjunctional block.

### Table 1: DH\textsubscript{i}E accentuates the onset of neuromuscular block induced by postjunctional nAChR antagonists, α-BTX, or α-conotoxin GI.

<table>
<thead>
<tr>
<th>Doses of antagonists</th>
<th>Time to 25% inhibition of T1 (min)</th>
<th>Time to 50% inhibition of T1 (min)</th>
<th>Time to 75% inhibition of T1 (min)</th>
</tr>
</thead>
<tbody>
<tr>
<td>α-BTX (0.175 µg g\textsuperscript{-1}) (n=6)</td>
<td>10.4 (8.4–12.4)</td>
<td>14.8 (11.3–20.2)</td>
<td>24.5 (12.9–34.1)</td>
</tr>
<tr>
<td>α-BTX (0.175 µg g\textsuperscript{-1}) + DH\textsubscript{i}E (0.5 µg g\textsuperscript{-1}) (n=6)</td>
<td>9.8 (7.1–11.1)</td>
<td>16.0 (9.2–20.8)</td>
<td>29.8 (15.0–35.7)</td>
</tr>
<tr>
<td>α-BTX (0.175 µg g\textsuperscript{-1}) + DH\textsubscript{i}E (1.0 µg g\textsuperscript{-1}) (n=6)</td>
<td>6.4* (4.8–8.7)</td>
<td>9.7* (6.6–13.8)</td>
<td>18.2 (9.4–28.4)</td>
</tr>
<tr>
<td>α-conotoxin GI (30 ng g\textsuperscript{-1}) (n=5)</td>
<td>3.7 (3.0–4.4)</td>
<td>4.4 (3.8–5.3)</td>
<td>5.7 (4.5–6.7)</td>
</tr>
<tr>
<td>α-conotoxin GI (30 ng g\textsuperscript{-1}) + DH\textsubscript{i}E (1.0 µg g\textsuperscript{-1}) (n=5)</td>
<td>3.0** (2.2–3.0)</td>
<td>3.7** (2.7–3.9)</td>
<td>4.8* (3.3–5.0)</td>
</tr>
</tbody>
</table>

\*\textsuperscript{P}<0.05 and **\textsuperscript{P}<0.01 vs values of toxin postjunctional (α-BTX or α-GI) when given alone.
and α-BTX or α-conotoxin GI were combined together. The faster onset of effects seen when pre- and postjunctional effecting drugs are present is reminiscent of effects of neuromuscular blocking agents which have combined actions on the pre-and post-junctional nAChRs. The presynaptic and postsynaptic effects of muscle relaxants are essential in contributing to the total efficacy of neuromuscular depression. 4, 6 DHβE and α-conotoxin MII, in addition to binding to α3β2 nAChRs, also bind to α4β2 and α6 containing nAChRs, respectively. 11, 29 33 The presence of the latter nAChRs has not been demonstrated at the motor nerve terminal. Thus the accelerated onset of effect when DHβE is combined with postjunctional AChRs is unrelated to the block of α6 or α4AChRs by DHβE.

Our study examined if block of α3β2 nAChRs causes fade. Although decrease of ACh release by DHβE alone may not be necessary and sufficient to cause fade, there could be other receptor-mediated contributory factors that can induce fade. Mucarinc AChRs, in addition to purinergic and adrenergic receptors, are also present prejunctionally. 3 34 Muscle relaxants do have actions on non-nicotinic (mucarinc) AChRs and also other receptors. 4, 34 It is possible that muscle relaxants may act on other nicotinic, mucarinc and purinergic receptors, in addition to α3β2 nAChR, to cause fade. Nevertheless, our results with two specific postsynaptic ligands, α-BTX and α-conotoxin GI, strongly indicate that decrease of the postjunctional receptors available for neuromuscular transmission, in and of itself, is capable of producing fade. Block of prejunctional nAChRs by DHβE is not necessary and sufficient to cause fade in vivo. Therefore, fade is not necessarily a prejunctional phenomenon but possibly also reflects the decreased margin of safety of neuromuscular transmission, which can be postjunctional or combined pre- and post-junctional in origin.

Authors’ contributions
M.N.: designed, conducted this study, analysed the data, and wrote the manuscript. T.S. and S.J.S.: helped with the experiments and critically reviewed the manuscript. J.A.J.M.: helped design the study, supervised the work, analysed the data, and edited the manuscript. All authors approved the final manuscript.

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Declaration of interest

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