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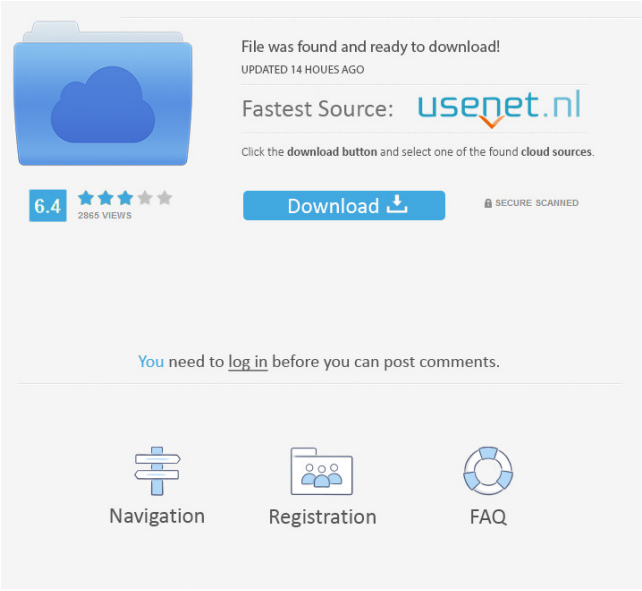

Gullfoss VST VST3 AAX v1.4.1-R2RBotulinum toxin A for neurogenic and nociceptive trigeminal pain in mice. The aim of this study was to assess the usefulness of botulinum toxin A (BTX-A) for the treatment of trigeminal pain in mice. Allodynia was induced by chemical injury of the trigeminal ganglion (TG) and subsequent infraorbital nerve (ION) injury. This allodynia was reversible with BTX-A. The dose of 300 units/kg of BTX-A injected in the TG had a more effective effect than

lower doses. The injection of BTX-A into the ION also induced reversal of the allodynia induced by chemical injury of the trigeminal ganglion and the infraorbital nerve. The administration of lower doses of BTX-A in the ION was also effective. There was no difference in the pain-relieving effect among the subnuclei of the TG. The pre- or posttreatment with BTX-A did not alter the effect of BTX-A. By the intrathecal administration of BTX-A, however, BTX-A had a dose-dependent effect on the mechanical

allodynia induced by ION injury. These results suggest that BTX-A is effective for the treatment of neurogenic and nociceptive trigeminal pain. Although we do not yet know the precise mechanism, an inhibition of pain transmission in the trigeminal nucleus caudalis may be involved in this analgesic effect.15 U.S. Code § 1382j - Rules of construction The term “State board” or “State board,” as used in sections 1382a and 1382j to 1382j–7 of this title, means the State board that has jurisdiction

over the transaction related to the product with respect to which the manufacture, use, or handling is proposed or in effect; except that it shall include a State board that has jurisdiction over the transaction related to the product and the manufacture, use, or handling of which is proposed by the manufacturer or a State board that is administered by the manufacturer. (Pub. L. 96–272, §3, Sept. 15, 1980, 94 Stat. 395, effective 180 days after Aug. 17, f678ea9f9e

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