



Commentary

Lasmiditan for the acute treatment of migraine: Subgroup analyses by prior response to triptans¹

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The use of Lasmiditan, a highly selective agonist for the 5-HT_{1F} receptor with penetration into the central nervous system, had its effectiveness proven for the acute treatment of migraine in two phase 3 studies, called SAMURAI and SPARTAN. In both studies, the percentage of patients who were pain-free after 2 h in lasmiditan groups was higher than in the placebo group.^{2,3}

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Bearing in mind that lasmiditan and triptans are different in structure and pharmacologically, the study in question aimed to investigate the response of lasmiditan for the acute treatment of migraine in relation to the previous response to triptans (good or insufficient) and in those who never use this class of drug, based on the SAMURAI and SPARTAN studies.

Regarding the response to the acute treatment of migraine with lasmiditan in those who have already used triptan, lasmiditan proved to be effective both in those with good response to triptan and in those with insufficient response in the following outcomes: pain-free in 2 h, absence of most uncomfortable symptoms in 2 hours and pain relief in 2 hours. The benefit was greater than placebo at all doses tested (50, 100 and 200 mg lasmiditan).

In those, who never used triptans, lasmiditan was effective in the acute treatment of migraine in doses of 100 and 200 mg when compared with placebo for the same outcomes reported above. Taking into account that lasmiditan has no vasoconstrictor effects in animal models and in human, the results of the current study suggest that lasmiditan, the first tested drug of the new therapeutic class called “ditan”, is a therapeutic option for patients with or without previous experience with triptans, as well as those with contraindications to that class of medication.

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Reference

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