Migraines are highly prevalent primary headaches that can start in childhood or adolescence. It affects mostly women and manifests clinically as moderate to severe headache attacks, frequently of unilateral fronto-temporal locations. The pain quality is pulsating and/or pressure-type, incapacitating and worsens with physical activities, usually associated with nausea (may also be accompanied by vomiting), intolerance to light and/or noises and/or smells.

Headache may last from 4 to 72 hours when it is not treated or is ineffectively treated. Frequency of pain occurrence is variable and some patients can have it on a weekly basis, while others will have it less than once a month.

There are several drug options for the treatment of acute migraine attacks. The choice of one or another will depend on pain intensity, its frequency and associated symptoms, co-morbidity, progres-
sion profile and individual patients response. Several patients find pain relief using simple analgesics such as acetyl salicylic acid, acetaminophen, dipirone, mefenamic acid or some combination of these. In fact, a recent multicenter study found that the association of aspirin, acetaminophen and caffeine was efficient to treat most of the acute migraine attacks. Nonsteroidal anti-inflammatory drugs (NSAIDs) like diclofenac, naproxen and ibuprofen also proved to be efficacious in the acute treatment of migraine. Lysine clonixinate (LC) is a NSAID which belongs to the family of the non-salicylates and to the subgroup of anthranilic derivatives. It has a chemical structure resembling the flufenamic acid, although it is derived from nicotinic acid. Its efficacy in the treatment of acute migraine attacks was suggested by previous non-controlled studies, both in oral as well as in parenteral administration. Lysine clonixinate (LC) is a NSAID which belongs to the family of the non-salicylates and to the subgroup of anthranilic derivatives. It has a chemical structure resembling the flufenamic acid, although it is derived from nicotinic acid. Its efficacy in the treatment of acute migraine attacks was suggested by previous non-controlled studies, both in oral as well as in parenteral administration. Lysine clonixinate (LC) is a NSAID which belongs to the family of the non-salicylates and to the subgroup of anthranilic derivatives. It has a chemical structure resembling the flufenamic acid, although it is derived from nicotinic acid. Its efficacy in the treatment of acute migraine attacks was suggested by previous non-controlled studies, both in oral as well as in parenteral administration.

The aim of this study was to evaluate the efficacy of oral lysine clonixinate compared to placebo in the treatment of acute migraine.

**METHOD**

**Study group**

We studied 64 patients with migraine, prospectively diagnosed according to the criteria of the International Headache Society (IHS). Fifty five were women and 9 were men, aged from 18 to 53 years. Patients were randomized and divided into two groups (A and B), which received placebo or CL. Patients were instructed to take 2 tablets of medication (CL 125 mg/tablet or placebo) once the pain reached moderate or severe intensity in 6 consecutive attacks. Regardless of pain intensity, another tablet of LC or placebo had to be taken 40 minutes later by all patients. Patients were instructed to record in a diary their pain intensity just prior to taking the medication, and 1, 2 and 4 hours later. The patients also recorded the occurrence of vomiting after the drug ingestion and the consumption of other rescue medications after 4 hours. All patients in this study were permitted to take prophylactic drugs but not NSAIDs.

**RESULTS**

Group A included 31 patients, who had 152 moderate and 34 severe attacks, and were treated with placebo. Vomiting occurred in 11 of the moderate and 7 of the severe attacks. Thirty three patients in group B having 150 moderate and 48 severe attacks were treated with lysine clonixinate. Vomiting followed the ingestion of the medication in 8 moderate and in 16 severe attacks. With regard to the treatment of moderate attacks, 27.6% of group A and 65.4% of group B (p<0.0001) presented mild or no pain after 1h. After 2 hours, 29.8% of group A and 84.5% of group B (p<0.0001) had mild or no headache. After 4 hours, 34.7% of group A and 91.5% of group B (p<0.0001) had mild or no pain (Fig 1). Consumption of rescue medications after 4 hours was observed in 68% of group A and in 5.6% of group B (p<0.0001) (Fig 2). With regard to the severe attacks, 18.5% of group A and 18.8% of group B (p=0.9819) had mild or no headache; after 2h, 33.4% of group A and 31.3% (p=0.8645) of group B indicated mild or no pain; after 4h, 33.4%
of group A and 37.5% of group B (p=0.9521) reported mild or no pain) (Fig 3). Figure 4 shows that 77.8% patients of group A and 68.8% patients of group B consumed rescue medications after 4 hours.

**DISCUSSION**

Lysine clonixinate (2-(3-chloro-0-toluidine) piridin-3-lysine carboxilate) is a NSAID effective in the treatment of several pain syndromes such as renal colic, nerve entrapment pain, muscular pain and tooth ache10,12,13. The efficacy of other NSAIDs in the acute treatment of migraine, as well as that of lysine clonixinate, have been demonstrated in previous studies5-9. This study confirms the efficacy of lysine clonixinate compared to placebo in reducing both intensity and duration of moderate migraine attacks, as well as the consumption of rescue medications after 4 hours. The speed of action of lysine clonixinate is favorable and can't be compared with that of ibuprofen 7, diclofenac 5, tolfenamic acid 14, NSAID also demonstrated as being efficient in the acute treatment of migraine, since the methodology and design of this study, when the patients took another capsule 40 minutes later, was unique and not done with the above mentioned NSAID.

In another hand, the efficacy of the triptans, selective 5-HT1B/1D agonists and therefore specific drugs for the acute treatment of migraine, in reducing the intensity of migraine attacks is demonstrated at as early as 1 hour15,16,17,18. LC also has been proven effective after 1 hour in moderate attacks. However comparative postulations with regard to early efficacy of both drugs can't be accomplished due to different study designs, methods and patient selection.

Nonsteroidal anti-inflammatory drugs seem to act via inhibition of prostaglandin synthesis and platelet aggregation as well as the inhibition of serotonin release from platelets 19. Despite its efficacy and fast speed of action in many types of pain syndromes, it has been shown that lysine clonixinate also has a central pharmacological action, specifically in the thalamus and spinal cord10,13. Sierralta12 have demonstrated an interaction between lysine clonixinate and the central serotoninergic system.

Lysine clonixinate was not superior to placebo in reducing the intensity and duration of severe headaches, nor the consumption of rescue medications after 4 hours. Which sites of action were important in obtaining the results achieved in moderate attacks remains uncertain. Remission of the pain probably resulted from association of all those mechanisms of action and the reason why it failed to control severe attacks is unknown. Based upon pharmacokinetics of triptans, efficient drugs employed in the treatment of the acute migraine attack, it is suggested that it is important to penetrate the blood-brain barrier in order to obtain a rapid and prolonged effect 16. That is observed with lysine clonixinate10,12,13.

Headache recurrence, defined as the return of the headache within 24 hours after it has become mild or absent, may occur with triptans 17,20 and other headache drugs and should be studied with this NSAID. It is also necessary to evaluate the possibility of using lysine clonixinate in patients with ischaemic coronary artery disease and uncontrolled hypertension in whom triptans are contraindicated, as well as in patients with history of peptic disease in whom NSAIDs are contraindicated.

The authors have not observed any significant side effects using LC, although nausea, vomiting, allergic reactions, postural hypotension, vertigo and insomnia have been described in the literature 8,9,13.

This study indicates that oral administration of 250 mg lysine clonixinate followed by another 125 mg tablet 40 minutes later, is efficient and well tol-
erated in the treatment of moderate attacks of migraine. The authors would not recommend its use in severe attacks of migraine.

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**REFERENCES**