

BEISPIELSTUDIE

Randomized controlled trial in adults with headache recurrence after emergency department discharge, comparing the efficacy of oral sumatriptan to naproxen

KATEGORISIERUNG

Gattung

Forschungsprojekt mit lebenden Personen

Art

Unterart

Klinischer Versuch mit Arzneimitteln (einschliesslich Kombinationen nach Art. 2 Abs. 1 Bst. f und g MepV)

BACKGROUND

Migraine and other acute primary headaches are treated with a variety of parenteral medications in the emergency department. It is not clear which medication is best to prescribe to primary headache patients when they are discharged. This study compared the efficacy of oral sumatriptan to naproxen for treatment of post-ED recurrent primary headache.

METHODS

Adults aged 18 years or older who suffered from headache recurrence after discharge from an emergency department were randomized to either naproxen (Trade®) 500 mg or sumatriptan (Sumatriptan Spirig HC®) 100 mg. This was an open trial: patients and physicians knew who was given which medication. Both products were provided in the original package. The packages had trial-specific labels and were handed out by a hospital pharmacist. Patients were followed-up by telephone 48 hours after emergency department discharge. The primary outcome was change in pain intensity, measured during a two-hour period after ingestion. The change in the group that received 500 mg naproxen (Trade®) was compared to the change in the group that received 100 mg sumatriptan (Sumatriptan Spirig HC®). This difference was measured on a validated 11-point (0–10) verbal Numerical Rating Scale (NRS).

CATEGORISER-FRAGEN

Fällt das Forschungsprojekt in den Geltungsbereich des Humanforschungsgesetzes?

Ja

BECAUSE

This project was based on a study protocol which defined the exact procedures to be used. The study is designed to answer a disease related research question. It includes a relatively large number of persons and is not based on individual cases ("method-driven search for generalizable knowledge"); thus, it is classified as research according to HRA). Adults ("persons") who suffered from headache ("research concerning human diseases") after discharge from an emergency department were treated with either 500 mg naproxen (Trade®) or 100 mg sumatriptan (Sumatriptan Spirig HC®).

Handelt es sich bei dem Forschungsprojekt um ein Projekt mit lebenden Personen?

Ja

BECAUSE

Adults ("persons") who suffered from headache after discharge from an emergency department were included in this research project.

Handelt es sich bei dem Forschungsprojekt um einen klinischen Versuch im Sinne der KlinV oder der KlinV-Mep?

Ja

BECAUSE

In this research project, the investigator randomly assigned ("prospectively assigned") adults ("persons") who suffered from headache after discharge from an emergency department to receive either 500 mg naproxen (Trade®) or 100 mg sumatriptan (Sumatriptan Spirig HC®) ("health related intervention (therapeutic measure)"). The project assessed between-group difference in change in in pain intensity over the 2-hour period after they took the drugs ("to investigate its effect on health").

Wird in der Studie ein Arzneimittel (einschliesslich Kombinationen nach Art. 2 Abs. 1 Bst. f und g Medizinprodukteverordnung (MepV) vom 1. Juli 2020) untersucht?

Ja

BECAUSE

This clinical trial investigated the effects of naproxen (Trade®) 500 mg and sumatriptan (Sumatriptan Spirig HC®) 100 mg. (Both are "medicinal products").

Ist das Arzneimittel in der Schweiz zugelassen?

Ja

BECAUSE

This clinical trial investigated the effects of naproxen (Trade®) 500 mg and sumatriptan (Sumatriptan Spirig HC®) 100 mg. Both are "authorised" for the market in Switzerland, and are provided "as is", unchanged except for their labels. (Approval numbers for sumatriptan/[Sumatriptan Spirig HC®]: 58466, 58512, 58513 [Swissmedic]; and, for naproxen/[Trade®]: 51480 [Swissmedic]).

Wird in der Studie ein Placebo verwendet oder wird der Originalzustand des verwendeten Arzneimittels oder seine Verpackung verändert?

Nein

BECAUSE

This clinical trial investigated the effects of naproxen (Trade®) 500 mg and sumatriptan (Sumatriptan Spirig HC®) 100 mg. Both are "authorised" for the market in Switzerland, and are provided "as is". The medicinal product was provided in its original package but labelled with a trial-specific sticker (label). However, the label did not cover any pharmaceutically relevant information. This modification is not manufacturing according to the Therapeutics Product Act (TPA).

Wird das Arzneimittel in der Studie gemäss Fachinformation angewendet?

Ja

BECAUSE

This clinical trial investigated the effects of naproxen (Trade®) 500 mg and sumatriptan (Sumatriptan Spirig HC®) 100 mg on patients who suffered from headache after discharge from an emergency department. Naproxen and sumatriptan are approved in Switzerland for the treatment of headache. According to the summary of product characteristics , the maximum recommended dose for naproxen (Trade®) is 500 mg a day. For Sumatriptan (Sumatriptan Spirig HC®) it is 200 mg a day. Summary of product characteristics available here.