PREEMPTIVE PROPHYLAXIS OF MIGRAINE

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Division of application No. 09/575,277, filed on May 22, 2000, which is a continuation-in-part of application No. 09/185,310, filed on Nov. 3, 1998, now Pat. No. 6,066,092, and which is a continuation-in-part of application No. PCT/US99/09414, filed on Apr. 29, 1999.

ABSTRACT
A method of preventing the headache phase of migraine in a human comprises administration of a 5HT1 receptor agonist or administration of an over-the-counter or nonprescription drug to said human exhibiting prodrome symptoms of migraine. Suitably, the method comprises administration of a migraine headache phase-preventing effective amount of the 5HT1 agonist or over-the-counter or nonprescription drug. There is disclosed a preemptive prophylaxis migraine method using the following cognitive tests: Simple Reaction Time; Running Memory Continuous Performance Task; Matching to Sample; Mathematical Processing Task; and interpreting the results as a percent of baseline indicator of need for prophylaxis.
Cognitive change as compared to baseline
Indicator of need for prophylaxis
Drop in one of stanine score
PREEMPTIVE PROPHYLAXIS OF MIGRAINE

CROSS REFERENCES

[0001] This application is a divisional of co-pending application Ser. No. 09/575,277, which application was, in turn, a continuation-in-part of co-pending application Ser. No. 09/185,310, filed 3 Nov. 1998, and a continuation-in-part of PCT Application No. PCT/US99/09414 filed 29 Apr. 1999 and which named the United States as a designated country.

TECHNICAL FIELD OF THE INVENTION

[0002] The present invention relates generally to the medical field and, more particularly, to a method for predicting the onset of a migraine headache and to a preemptive prophylaxis of the migraine headache. The preemptive prophylaxis is directed to prevent or reduce the headache phase and/or disability of migraine in humans by the administration of drugs during the prodrome phase of migraine.

BACKGROUND OF THE INVENTION

[0003] A headache may be one of several different varieties, each of which has its own unique pain characteristics which differ dramatically. The types of headache include tension, sinus, cluster, rebound and migraine. Migraine is a particularly painful headache that occurs from time to time. The pain is quite severe and often the person with migraine must stay in bed. Dietary, emotional and environmental factors may trigger an attack. In average, migraine sufferers experience an attack per month. Attacks last from four to seventy-two hours. Of interest is that the incidence of migraine appears to be on the rise. Because of the severity and incidence of migraine, prescription medicines have been invented to provide relief.

[0004] Migraine sufferers sometimes get a warning signal before the onset of the headache phase of a migraine. The warning signals apparent to the migraineur are classified as aura. The period of aura is preceded by a period classified as prodromal or premonitory period. The periods of aura, prodrome and premonitory are pre-headache. The International Headache Society (IHS) defines aura as neurological symptoms that usually develop over 5-20 minutes and last less than 60 minutes. Headache may occur directly or after an aura free interval of less than 60 minutes. Aura symptoms commonly include, but are not limited to, visual disturbances and numbness or tingling sensations. Less than 20% of patients have migraine with aura (IHS 1.2). See Headache Classification Committee of the International Society. Classification and diagnostic criteria for headache disorders, cranial neuralgia and facial pain. Cephalalgia (1988); 8: (Supp. 7): 1-96.

[0005] The IHS has defined prodromal symptoms as non-aura symptoms signaling the onset of a migraine attack. The symptoms typically occur a few hours to 48 hours before the onset of the headache phase of the migraine. Headache phase of migraine as used herein means the point in time when head pain is perceived by the sufferer. Prodrome or premonitory symptoms may occur in migraine with (IHS 1.1) and migraine without aura (IHS 1.2). The IHS prefers the term premonitory symptoms over prodrome due to historical use of prodrome to describe aura. Prodrome symptoms as used herein is synonymous to premonitory symptoms. See Headache Classification Committee of the International Society. Classification and diagnostic criteria for headache disorders, cranial neuralgia and facial pain. Cephalalgia (1988); 8: (Supp. 7): 1-96.

[0006] Prodrome or premonitory symptoms may have physical and mental components. The symptoms have been classified by clinical presentation as excitatory and inhibitory symptoms. Excitatory symptoms include, but are not limited to, irritability, euphoria (being ‘high’), physical hyperactivity, excessive yawning, excessive sleepiness, increased sensitivity to light and sound, and craving for foods. Inhibitory symptoms include, but are not limited to, depression, mental withdrawal, behaviour sluggishness, feeling tired, poor concentration, muscle weakness, anorexia and fluid retention. See Headache Classification Committee of the International Society. Classification and diagnostic criteria for headache disorders, cranial neuralgia and facial pain. Cephalalgia (1988); 8: (Supp. 7): 1-96 and Anthony M, Rasmussen B K. In: Olesen J, Tfelt-Hansen P, Welch K M A (eds). The Headaches. New York: Raven Press, Ltd, 1993: 256-257. Prodrome/premonitory symptoms have been estimated to occur in up to 88% of migraine patients. See supra, Rasmussen.

[0007] 5-HT₃ receptor agonists have been found useful in the treatment of migraine. 5-HT₃ receptors are located, for example, in the dog saphenous vein and the 5HT₃ receptor agonists with which the present invention is concerned contract the dog saphenous vein. Such compounds may therefore be identified by their contractile effect on the dog isolated saphenous vein strip as described, for example, by Apperley et al, Br. J. Pharmacol, 68, 215-224 (1980). Compounds which are selective 5HT₃ receptor agonists have also been found to selectively constrict the carotid arterial bed of the anaesthetized dog.

[0008] Much work has been done in attempts to identify the subclasses of 5HT₁ receptors which are implicated in migraine. It is currently thought that 5HT₁D (formerly 5HT₁DR), 5HT₁D (formerly 5HT₁DP) and 5HT₁F receptors are particularly important. Tests in isolated cerebral arteries can be used to determine which of these receptor sub-types mediate the action of 5HT₁ agonist compounds, for example as described in Bouchelet, L et al, Mol. Pharmacol 1996, 50, 219-223.

[0009] Some 5HT₃ agonists, including 5HT₁D and 5HT₁F agonists have also been found to inhibit the trigeminal nerve. This can be assessed by measuring plasma protein extravasation in the dura mater following trigeminal nerve stimulation and administration of labeled albumin; active compounds produce inhibition of dural plasma protein extravasation in this model, which is described in Buzzo, M. G and Moskowitz M. A, Br. J. Pharmacol, 1990 99, 202-206.

[0010] A variety of compounds have been identified in the art as 5HT₁ agonists, for example by selective constriction of the dog isolated saphenous vein or constriction of the carotid arterial bed of the anaesthetized dog. These include indole derivatives such as those disclosed inter alia in: published British Patent Specification Nos. 2082175, 2081717, 2083463, 2124210, 2150932, 2162522, 2168347, 2168973, 2185020, 2186874, 2191488, 2008645, 2289464, 2289465, 2286185, published U.S. Pat. Nos. 5,288,748, 5,317,103, 5,382,592, 5,385,928, 5,387,593, 5,418,236, 5,433,915, 5,451,584, 5,466,688, 5,468,768, 5,519,025, 5,545,044, 5,602,128, 5,618,948, 5,637,611, 5,837,715; published Ger-
The empirical formula is $C_{39}H_{60}N_{8}O_{8}$, representing a molecular weight of 413.5. IMITREX® (sumatriptan succinate) Tablets are, for example, disclosed and/or claimed in U.S. Pat. No. 4,816,470, U.S. Pat. Nos. 5,037,845 and 5,863,559, which are incorporated by reference herein.

IMITREX® (sumatriptan succinate) Injection is indicated for (1) the acute treatment of migraine attacks with or without aura and (2) the acute treatment of cluster headache episodes. IMITREX® (sumatriptan succinate) Injection is not approved for use in the management of hemiplegic or basilar migraine. The name, structural formula, empirical formula and molecular weight of sumatriptan succinate are as hereinbefore described. IMITREX® (sumatriptan succinate) Injection is, for example, disclosed and/or claimed in U.S. Pat. No. 4,816,470 and U.S. Pat. No. 5,037,845, which are incorporated by reference herein.

IMITREX® (sumatriptan succinate) Nasal Spray is indicated for the acute treatment of migraine attacks with or without aura in adults. IMITREX® (sumatriptan succinate) Nasal Spray is not intended for the prophylactic therapy of migraine or for use in the management of hemiplegic or basilar migraine. IMITREX® (sumatriptan) Nasal Spray contains sumatriptan, a selective 5-hydroxytryptamine, receptor subtype agonist. Sumatriptan is chemically designated as 5-[2-(dimethylamino)ethyl]-N-methyl-1H-indole-5-methanesulfonamide, and it has the structure shown in Formula (II):

The empirical formula is $C_{39}H_{60}N_{8}O_{8}$, representing a molecular weight of 413.5. Due to the presence of sulphuric acid in the nasal spray formulation, the hemisulphate salt of sumatriptan is formed in situ. IMITREX® (sumatriptan) Nasal Spray is, for example, disclosed and/or claimed in U.S. Pat. No. 4,816,470, U.S. Pat. No. 5,037,845, U.S. Pat. No. 5,705,520 and U.S. Pat. No. 5,554,639, which are incorporated by reference herein.

Naratriptan is marketed inter alia in the United States as AMERGE® (naratriptan hydrochloride) Tablets. In
many other countries, the trade mark NARAMIG® is used instead of AMERGE®. AMERGE® Tablets are indicated for the acute treatment of migraine attacks with or without aura in adults. AMERGE® Tablets are not approved for the prophylactic therapy of migraine or for use in the management of hemiplegic or basilar migraine. AMERGE® (naratriptan hydrochloride) Tablets contain naratriptan as its hydrochloride salt. Naratriptan hydrochloride is a selective 5HT1 agonist. It is chemically designated as N,N-dimethyl-3-(1-methyl-4-piperidinyl)-1H-indole-3-ethanesulfonamide monohydrochloride, and it has the structure shown in formula (III):

![Formula III]

The empirical formula is C19H21N3O2S.HCl, representing a molecular weight of 371.93. AMERGE® (naratriptan hydrochloride) Tablets are, for example, disclosed and/or claimed in U.S. Pat. No. 4,997,841, which is incorporated by reference herein.

Rizatriptan is marketed inter alia in the United States as MAXALT® (rizatryptan benzoate) Tablets and MAXALT-ML™ (rizatryptan benzoate) Orally Disintegrating Tablets. MAXALT® (rizatryptan benzoate) Tablets and MAXALT-ML™ (rizatryptan benzoate) Orally Disintegrating Tablets are indicated for the acute treatment of migraine attacks with or without aura in adults. MAXALT® (rizatryptan benzoate) Tablets and MAXALT-ML™ (rizatryptan benzoate) Orally Disintegrating Tablets are not approved for the prophylactic therapy of migraine or for use in the management of hemiplegic or basilar migraine. MAXALT® (rizatryptan benzoate) Tablets and MAXALT-ML™ (rizatryptan benzoate) Orally Disintegrating Tablets contain rizatryptan benzoate, a selective 5-HT1 receptor agonist. Rizatryptan benzoate is chemically described as: N,N-dimethyl-5-((1H-1,2,4-triazol-1-ylmethyl)-1H-indole-3-ethanamine monobenzoate. Its empirical formula is C14H16N3C6H5O2, representing a molecular weight of the free base of 269.4. MAXALT® (rizatryptan benzoate) Tablets are, for example, disclosed and/or claimed in U.S. Pat. No. 5,298,520 and U.S. Pat. No. 5,602,162, which are incorporated by reference herein. MAXALT-ML™ (rizatryptan benzoate) Orally Disintegrating Tablets are, for example, disclosed and/or claimed in U.S. Pat. No. 4,305,502, U.S. Pat. No. 4,371,516, U.S. Pat. No. 4,758,598, U.S. Pat. No. 5,298,520 and U.S. Pat. No. 5,602,162, which are incorporated by reference herein.

Zolmitriptan is marketed inter alia in the United States as ZOMIG™ (zolmitriptan) Tablets. ZOMIG™ (zolmitriptan) Tablets are indicated for the acute treatment of migraine with or without aura in adults. ZOMIG™ (zolmitriptan) Tablets are not approved for the prophylactic therapy of migraine or for use in the management of hemiplegic or basilar migraine. ZOMIG™ (zolmitriptan) Tablets contain zolmitriptan, which is a selective 5-HT1 receptor ago-nist. Zolmitriptan is chemically designated as (S)-4-[(3-[2-(Dimethylamino)ethyl]-1H-indol-5-yl)methyl]-2-oxazolidinone. The empirical formula is C19H21N3O2, representing a molecular weight of 287.36. ZOMIG™ (zolmitriptan) Tablets are disclosed and/or claimed in U.S. Pat. No. 5,466,699, which is incorporated by reference herein.

Other 5HT1 agonists in development include eletriptan, frovatriptan, almotriptan, ALX-0646, LY334370, U109291, IS159 and PNU-142633.

The above mentioned anti-migraine agents have not been approved for the prevention of migraine. In fact, the current product label for each of these agents states that they are not intended for prophylactic therapy of migraine, e.g., AMERGE® (naratriptan hydrochloride) Tablets. Clinically, these agents are used at the onset of the headache phase of the migraine, (CMAJ 1997;156:1273-1287.), and are commonly reserved for the treatment of severe migraine pain. (Silverstein S D, Lipton R B, Goadsby P J (eds). Headache in Clinical Practice. Oxford UK: Isis Medical Media Ltd, 1998, 74-77.) Thus, it would be desirable to provide an anti-migraine agent useful for the prevention of the headache phase of the migraine. It would be further desirable to be able to predict the onset of migraine before the head pain actually occurs and thereby permit the prophylactic administration of medicine.

The Automated Neuropsychological Assessment Metrics (ANAM) is a set of standardized batteries of cognitive tests, modified by neuropsychologists in the U.S. Armed Forces for precise measurement of cognitive processing efficiency of military personnel. The tests assess sustained concentration and attention, mental flexibility, spatial processing, cognitive processing efficiency, mood, arousal/fatigue level, and short-term, long-term and working memory. The ANAM is now in the public domain. The most recent version is ANAM V3.11a/96 which includes the following battery of tests:

- Subject Demographics Form
- Stanford Sleepiness or Sleep/Fatigue Scale
- Mood Scale 2
- Simple and Two-Choice Reaction Time
- Sternberg Memory Search Tasks
- Running Memory Continuous Performance Task
- Mathematical Processing Task
- Digit Set Comparison Task
- Logical Reasoning-Symbolic
- Tower of Hanoi (Tower Puzzle)
- Stroop Color/Word Interference
- Code Substitution (Letter/Symbol Comparison)
- Code Substitution (Immediate and Delayed Recall)
- Spatial Processing Task (Simultaneous)
- Matching to Sample
- Tapping (Left and Right Index Finger)
- Modified Orientation and Amnesia Test
It would be desirable to be able to use a subset of these tests to predict the onset of migraine before the head pain actually occurs and thereby permit the prophylactic administration of medicine.

The present invention is directed to meeting one or more of the above-stated desirable objectives.

SUMMARY OF THE INVENTION

The present invention provides a method of preventing the headache phase of migraine in a human comprising administration of a 5HT\textsubscript{1} agonist to said human exhibiting prodrome symptoms of migraine. Suitably, the method comprises administration of a headache phase preventing effective amount of the 5HT\textsubscript{1} agonist.

In a second aspect, the invention provides a 5HT\textsubscript{1} agonist for use in the manufacture of a medicament for administration to a human exhibiting prodrome symptoms of migraine in order to prevent the headache phase of migraine.

According to a third aspect, the invention provides a pharmaceutical composition comprising a 5HT\textsubscript{1} agonist as active ingredient for administration to a human exhibiting prodrome symptoms of migraine in order to prevent the headache phase of migraine.

In the second and third aspects, suitably a headache phase-preventing effective amount of a 5HT\textsubscript{1} agonist is administered. In preferred embodiments of the first, second and third aspects, the effective amount is from about 0.25 to 100 mg.

The 5HT\textsubscript{1} agonist employed in accordance with the first, second and third embodiment is typically administered orally, parenterally, by inhalation, intranasally or rectally. Parenteral, and oral administration are preferred, with oral administration being particularly preferred.

Suitably, the 5HT\textsubscript{1} agonist administered to said human exhibiting prodrome symptoms of migraine is selected from the group consisting of sumatriptan, naratriptan, rizatriptan, eletriptan, zolmitriptan, frovatriptan, almotriptan, ALX-0646, LY334370, U109291, IS159 and PNU-142633. It is preferably selected from the group consisting of sumatriptan, naratriptan, rizatriptan, zolmitriptan, eletriptan, frovatriptan and almotriptan. Sumatriptan and naratriptan are particularly preferred, with sumatriptan being especially preferred.

When the 5HT\textsubscript{1} agonist is sumatriptan, a headache phase-preventing effective amount is typically from about 2 mg to 100 mg. If the sumatriptan is administered subcutaneously, a headache phase-preventing amount is suitably from about 2 mg to 6 mg. For oral administration, a suitable headache phase-preventing amount is 25 mg to 100 mg. When sumatriptan is administered intranasally, a suitable headache phase-preventing amount is in the range 5 mg to 20 mg. A suitable headache phase-preventing amount of sumatriptan for rectal administration is in the range 12.5 mg to 25 mg.

When the 5HT\textsubscript{1} agonist is naratriptan, a headache phase-preventing effective amount is typically from about 0.25 mg to 2.5 and suitably from 1 mg to 2.5 mg.

When the 5HT\textsubscript{1} agonist is rizatriptan, a typical headache phase-preventing effective amount is from about 1 mg to 10 mg, suitably 5 mg to 10 mg.

When the 5HT\textsubscript{1} agonist is zolmitriptan, and a typical headache phase-preventing effective amount is from about 0.5 mg to 5 mg, suitably 1 to 5 mg.

When the 5HT\textsubscript{1} agonist is eletriptan, a typical headache phase-preventing effective amount is from about 5 mg to 80 mg, suitably 20 mg to 80 mg.

When the 5HT\textsubscript{1} agonist is frovatriptan, a typical headache phase-preventing effective amount is from about 1 mg to 40 mg, suitably 2.5 mg to 40 mg.

When the 5HT\textsubscript{1} agonist is almotriptan, a headache phase-preventing effective amount is from about 2 mg to 150 mg, suitably 12.5 mg to 25 mg.

The present invention further provides a preemptive prophylaxis migraine method using the following cognitive tests: Simple Reaction Time; Running Memory Continuous Performance Task; Matching to Sample; Mathematical Processing Task; and interpreting the results as a percent of baseline indicator of need for prophylaxis. Preferably the tests are administered in the listed sequence. Advantageously the tests are preceded by the Stanford Sleepiness Scale and Mood Scale 2 tests.

In a preferred arrangement there is provided a preemptive prophylaxis migraine device including a microprocessor having a memory, a battery of tests loaded into the memory of the microprocessor and including a Simple Reaction Time, a Running Memory Continuous Performance Task, a Matching to Sample, and a Mathematical Processing Task; means for computing the score on a trial of these tests to establish a baseline and for storing the baseline in the memory; the means for computing being operative for computing the score of a subsequent trial of the tests and comparing the same to the stored baseline; and means for indicating a cognitive change.

In another aspect, the invention provides a method of preventing the headache phase of migraine in a human comprising administration of an over-the-counter or non-prescription medicine to said human exhibiting prodrome symptoms of migraine.

In another aspect of the invention, a method is provided for preventing the headache phase of migraine in a human comprising administration of a Cox-2 inhibitor to said human exhibiting prodrome symptoms of migraine.

These and other objects, aspects, features and advantages of the present invention will become apparent from the following detailed description when taken in conjunction with the referenced drawings.

BRIEF DESCRIPTION OF THE DRAWINGS

Reference is now made to the drawings which illustrate the best known mode of carrying out the invention and wherein the same reference numerals indicate the same or similar parts throughout the several views.

FIG. 1 is a plan view of a hand-held computer which is one apparatus for determining cognitive change in a human;

FIG. 1A is a plan view of a palm-top type computer which is another apparatus for determining cognitive change in a human;
FIG. 2 is a flow chart illustrating the steps and sequence of a method for performing preemptive prophylaxis of migraine; and

FIG. 3 is a chart illustrating the therapeutic phases of migraine and including the treatment options most suitable to each phase of migraine.

DETAILED DESCRIPTION

FIG. 1. A flow chart illustrating the steps and sequence of a method for performing preemptive prophylaxis of migraine device in the form of a hand-held computer, generally designated 10, and having a key pad 12 and a screen 14 which advantageously is at least four inches (10.16 cm.) square. A hinge 15 is provided so the screen 14 may be conveniently folded down upon the key pad 12 for storage or transporting. When open the computer 10 is conveniently about 5"x9" (12.7 cm. by 22.86 cm.) in size. The key pad 12 has a built-in set of two mouse buttons 16, 18, a start/stop or on/off button 22, an enter key 24, and Mood Scale 2 keys 1, 2 and 3. As used herein the terms “buttons” and “keys” are intended to mean the same thing. The computer 10 contains memory chips (not shown) which have a set of programmed cognitive tests 103-106 (hereafter described) and which record a person’s performance time in milliseconds on those tests. The computer program uses the score in milliseconds on the third trial of these cognitive tests as a baseline measurement, which is converted to a stanine score. Subsequent trials are similarly scored and converted to stanine.

FIG. 2. A flow chart illustrating the sequence of the method. From the seventeen tests of the original ANAM, four subtests were selected and sequenced for measuring cognitive processing efficiency of migraine sufferers, as follows:

1. Simple Reaction Time (SMRT), 103
2. Running Memory Continuous Performance Task (CPT), 104
3. Matching to Sample (M2SP), 105
4. Mathematical Processing Task (MATH), 106.

Also included are two preliminary measures of alertness and mood that are also part of the ANAM:

1. Stanford Sleepiness Scale, 101
2. Mood Scale 2, 102.

Description of Subtests:

1. The first step 101 is Stanford Sleepiness Scale which consists of seven statements that describe the present state of alertness or sleepiness and are numbered from one to seven, with one being highly alert and seven being close to sleep. Individuals rate their level of alertness prior to taking the first subtest of the battery. It provides a way to monitor fatigue over the course of repeated measures. Subjective ratings may be correlated with measured performance.

2. The second step 102 is Mood Scale 2 which consists of a list of thirty-six adjectives that are rated on a three-point scale. Using mouse button 16 participants respond to each adjective by indicating “yes,” “moderately,” or “no,” based on how they feel at the present time. The Mood Scale 2 categories include anger, happiness, fear (anxiety), depression, activity, and fatigue.

3. The third step 103 is Simple Reaction Time (SMRT) which presents a simple stimulus on the screen (*). In response, the individual presses the mouse button 16 each time the stimulus appears. The Reaction Time measures the speed of the motor response, the peripheral nerve conduction velocity. This represents the “hardware” of the nervous system in terms of input, followed by motor response. Actual cognitive processing time is not involved in this test.

4. The fourth step 104 is Running Memory Continuous Performance Test (CPT) which is a continuous letter comparison task. A randomized sequence of upper-case letters, A through Z, is presented one at a time in the center of the computer screen 14. The person presses button 16 if the letter on the screen matches the letter that immediately preceded it; and different button 18 if the letter on the screen is different than the immediately preceding letter. The task lasts approximately five minutes. The CPT was specifically designed to assess components of memory, attention, efficiency and consistency. This task is forced paced, with individuals having only a brief time in which to respond.

5. The fifth step 105 is Matching to Sample (M2SP) and consists of a number of trials that begins with a first design being presented in the center of the screen 14 for three seconds, followed by a showing that contains two designs. The person matches one of the two designs with the first design or sample by pressing the appropriate button 16 or 18. The design is a 4x4 checkerboard and varies by the number of cells that are shaded from one cell through twelve cells.

6. The sixth step 106 is Mathematical Processing (MATH) and involves arithmetic problems presented in the middle of the screen 14. Working from left to right, the person solves the addition and subtraction and decides if the answer is greater or less than the number 5.

As indicated, the scores are recorded by the computer 10 and the score on the third trial of these sequenced cognitive tests 103-106 are used as the baseline measurement. Subsequent trials measure cognitive change as compared to baseline. A drop of one in stanine score is an indicator of the onset of migraine and an indicator of need for prophylaxis. See FIG. 2, 107. This was empirically determined by the following research. The preemptive prophylaxis of migraine method was used to measure cognitive deficiency during a migraine in each of a group of ten migraineurs. The method was used to measure the return of cognitive efficiency after injection of sumatriptan, an anti-migraine medication, in each of the group of ten migraineurs. The method measured cognitive change, compared to the baseline stanine score, that predicted the onset of a migraine.

The above described preemptive prophylaxis of migraine device and method allows a migraine sufferer to
take medication to preempt the occurrence of head pain, associated symptoms and accompanying disability.

[0085] It will be appreciated that the precise dose of a 5HT agonist administered to prevent the headache phase of migraine may depend on the particular compound used, the age and condition of the patient and the frequency and route of administration and will be at the ultimate discretion of the attendant physician. The compound may be administered in single or divided doses and may be administered one or more times, for example 1 to 3 times per day during exhibition of prodrome symptoms of migraine.

[0086] Also, a compound for use according to the invention may be formulated in a conventional manner using one or more pharmaceutically acceptable excipients. Thus, a compound for use according to the invention may for example be formulated for oral, sub-lingual, buccal, parenteral, rectal or intranasal administration or in a form suitable for administration by inhalation or insufflation (either through the mouth or nose).

[0087] For oral administration the pharmaceutical compositions may take the form of, for example, tablets or capsules prepared by conventional means with pharmaceutically acceptable excipients such as binding agents (e.g., pregelatinized maize starch, polyvinylpyrrolidone or hydroxypropyl methylcellulose); fillers (e.g., lactose, microcrystalline cellulose or calcium phosphate); lubricants (e.g., magnesium stearate, talc or silica); disintegrants (e.g., potato starch or sodium starch glycinate); or wetting agents (e.g., sodium laurel sulphate). The tablets may be coated by methods well known in the art. Preferred tablet formulations of sumatriptan are those described in European Patent No. 0503440.

[0088] Liquid preparations for oral administration may take the form of, for example, solutions, syrups or suspensions, or they may be presented as a dry product for constitution with water or other suitable vehicle before use. Such liquid preparations may be prepared by conventional means with pharmaceutically acceptable additives such as suspending agents (e.g., sorbitol syrup, methyl cellulose or hydrogelsated edible fats); emulsifying agents (e.g., lecithin or acacia); non-aqueous vehicles (e.g., almond oil, oily esters or ethyl alcohol); and preservatives (e.g., methyl or propyl P-hydroxybenzoates or sorbic acid).

[0089] For buccal administration, the compositions may take the form of tablets or lozenges formulated in conventional manner.

[0090] A compound for use according to the method may be formulated for parenteral administration by injection, conveniently intravenous, in intramuscular or subcutaneous injection. Formulations for injection may be presented in unit dosage form e.g., in ampoules or in multi-dose containers, optionally with an added preservative.

[0091] The compositions for parenteral administration may take such forms as suspensions, solutions or emulsions in oily or aqueous vehicles, and may contain formulatatory agents such as suspending, stabilizing and/or dispersing agents. Alternatively, the active ingredient may be in dry form such as a powder, crystalline or freeze-dried solid for constitution with a suitable vehicle, e.g., sterile pyrogen-free water or isotonic saline before use. They may be presented, for example, in sterile ampoules or vials.

[0092] A compound for use according to the method may also be formulated in rectal compositions such as suppositories or retention enemas, e.g., containing conventional suppository bases such as cocoa butter or other glyceride. Preferred compositions for rectal administration are those described in European Patent No. 0639072, which is incorporated by reference herein.

[0093] Tablets for sub-lingual administration may be formulated in any conventional manner.

[0094] For intranasal administration a compound for use according to the invention may be used, for example, as a liquid in the form of a spray or drops or as a powder. Suitably the preparation for intranasal administration is delivered in the form of a spray or aerosol from an insufflator or from a pressurized pack or nebulizer with the use of a suitable propellant. Preferably sumatriptan is administered intranasally in the form of its hemisulphate salt in a composition as described in European Patent No. 0490689, which is incorporated by reference herein.

[0095] For administration by inhalation the compound for use according to the invention is conveniently delivered in the form of an aerosol spray presentation from pressurized packs or a nebulizer, with the use of a suitable propellant, e.g., dichlorodifluoromethane, trichlorofluoromethane, dichlorotetrafluoroethane, carbon dioxide or other suitable gas. In the case of a pressurized aerosol the dosage unit may be determined by providing a valve to deliver a metered amount. Capsules and cartridges, e.g., of gelatin, for use in an inhaler or insufflator may be formulated containing a powder mix of a compound of use in the invention and a suitable powder base such as lactose or starch.

[0096] The method will now be further illustrated with reference to the following non-limiting examples.

**BIOLOGICAL EXAMPLES**

[0097] An open label pilot study was conducted to determine whether oral naratriptan 2.5 mg taken as a single dose during the prodrome could prevent the onset of the headache phase of migraine. Results of the study suggest that administration of 2.5 mg naratriptan during prodrome was effective in preventing the onset of the headache phase of migraine.

[0098] Twenty patients recruited at 4 study sites were administered oral naratriptan 2.5 mg after onset of prodrome symptoms for a total of 63 prodromes. Sixty percent of the treated prodromes did not progress to headache. The pain severity of headaches that were not prevented were reduced as compared to baseline pain severity of migraine attacks not treated with naratriptan.

[0099] An additional 13 patients recruited at a fifth site were also treated. All 13 patients reported headache following each treatment. Upon review, all 13 patients reported onset of headache within 3 hours after administration of naratriptan. Eighty-seven percent of prodromes in this patient group were treated ≤2 hours before onset of headache. A similar finding was seen at the remaining 4 sites where 15 of the 25 headaches (60%) that were not prevented occurred ≤2 hours after treatment. This data suggests these patients waited too late to administer naratriptan and may be due to the fact that patients were instructed to record the onset of premonitory symptoms, but were asked to delay
treatment until they were sure a migraine headache was imminent. These data were considered independent of the data from the other 4 sites due to the dichotomy of response as compared to the 4 sites in which the response rates ranged from 58% to 66%.

[0100] FIG. 3 shows the phases of migraine. The entire migraine event is roughly divided into three time periods: pre-headache 31, headache 32 and postdrome 33. These time periods are further divided into phases. The pre-headache period is composed of Phase I, prodrome 34, and Phase II, aura 35. The headache period is divided into Phase III, early headache 36, and Phase IV, late headache 37. Finally, Phase V, postdrome 38, encompasses the entire postdrome period.

[0101] During the prodrome phase 4 of migraine, identified in FIG. 3, there are alternatives to the use of 5HT1 agonists to preclude the onset of the headache phase. Advantageously, these alternatives include over-the-counter (OTC) or nonprescription drugs such as acetylsalicylic acid (aspirin); the combination of 4'-hydroxyacetanilide, acetylsalicylic acid, and caffeine (e.g., Excedrin®); and other non-narcotic analgesics as the anti-migraine medication. Nonsteroidal anti-inflammatory drugs (NSAIDs) are also effective as an alternative to 5HT1 agonists during prodrome. Examples of suitable NSAIDs include ibuprofen and other NSAIDs (e.g. acetaminophen, naproxen, indomethacin) which inhibit both COX-1 and COX-2 enzymes, and wholly COX-2 inhibitors such as celecoxib, sold under the brand name Celebrex®; and rofecoxib, sold under the brand name Vioxx®. These so-called COX-2 inhibitors operate by inhibiting the COX-2 enzyme, which triggers symptoms of pain and inflammation, while leaving the COX-1 enzyme, which aids in maintaining the stomach lining, unaffected. As used herein, the term “COX-2 inhibitor” means an inhibitor which operates on the COX-2 enzyme and not the COX-1 enzyme. U.S. Pat. No. 6,048,850, the disclosure of which is herein incorporated by reference, describes one method of COX-2 inhibitor operation.

[0102] Other objects, features and advantages of the present invention will be apparent to those skilled in the art. While preferred medications, uses and steps of the method have been illustrated and described, this has been by way of illustration and the invention should not be limited except as required by the scope of the appended claims.

We claim:

1. A preemptive prophylaxis migraine method for preventing the headache phase of a primary migraine disorder, including the steps of:

   performing cognitive tests to determine prodromal symptoms of migraine; and
   
   administering a non-narcotic analgesic as an anti-migraine medication.

2. The preemptive prophylaxis migraine method as set forth in claim 1, wherein said non-narcotic analgesic is selected from the group consisting of acetylsalicylic acid (aspirin); nonsteroidal anti-inflammatory drugs (NSAIDs); and a combination of 4'-hydroxyacetanilide (acetaminophen), acetylsalicylic acid, and caffeine.

3. The preemptive prophylaxis migraine method as set forth in claim 2, wherein said nonsteroidal anti-inflammatory drugs include COX-2 inhibitors.

4. The preemptive prophylaxis migraine method as set forth in claim 3, wherein said COX-2 inhibitors include celecoxib and rofecoxib.

5. A preemptive prophylaxis migraine method as set forth in claim 2, wherein said nonsteroidal anti-inflammatory drugs include acetaminophen, ibuprofen, naproxen, and indomethacin.

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