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(54) **Tapentadol készítmények**

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(54) **Tapentadol compositions**

Tapentadol-Zusammensetzungen

Compositions de tapentadol

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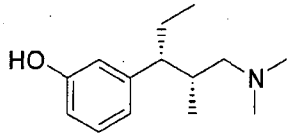
Description

Related Applications

[0001] This application claims priority from a U.S. provisional patent application serial no. 61/004,029 filed on November 23, 2007, which is incorporated herein by reference

Background of the Invention

[0002] Tapentadol, 3-(3-Dimethylamino-1-ethyl-2-methyl-propyl)-phenol (compound 1) is a centrally acting analgesic with a dual mode of action: μ-opioid receptor agonism and noradrenalinne reuptake inhibition. Its dual mode of action provides analgesia at similar levels of more potent narcotic analgesics such as hydrocodone, oxycodone, and morphine with a more tolerable side effect profile. Tapentadol was first disclosed in European patent no. EP 693,475 and is currently under FDA review.



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[0003] The traditional formulations of tapentadol for oral administration lead to a rapid release of the drug in the gastrointestinal tract and hence its analgesic action begins rapidly. However, a rapid reduction in the analgesic activity is observed. Therefore, treatment with tapentadol requires repeated administration of the pharmaceutical composition at relatively short intervals, often as high as four to ten times daily, to maintain the required concentration of active ingredient in the patient's blood plasma. The need for repeated dosing can lead to errors in administration and inability to maintain desirable concentration in the plasma, which are detrimental to patient compliance and the therapeutic objectives, particularly if the condition is chronic pain or a pain related condition. Accordingly, there is an unmet need to have slow or controlled release pharmaceutical compositions for oral administration of the active ingredient, tapentadol. Further, there is a need for compositions suitable for long term treatment of pain and pain related conditions particularly because such conditions persist among aged populations of the society.

[0004] Pregabalin (compound 2), a gamma-aminobutyric acid (GABA) analogue, is an anticonvulsant drug which is used as an adjunct therapy for partial seizures, for neuropathic pain, and in generalized anxiety disorder. Pregabalin was designed as a more potent successor to gabapentin and it is marketed by Pfizer under the trade name Lyrica®. Recent studies have shown that pregabalin is effective at treating chronic pain in disorders such as fibromyalgia and spinal cord injury.

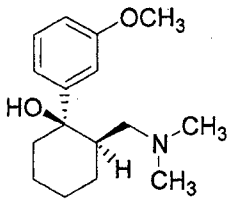
[0005] Gabapentin (compound 3) is another GABA analogue similar to Pregabalin and was initially synthesized to mimic the chemical structure of the neurotransmitter gamma-aminobutyric acid (GABA), but is not believed to act on the same brain receptors. Its exact mechanism of action is unknown, but its therapeutic action on neuropathic pain is thought to involve voltage-gated N-type calcium ion channels.

[0006] Most anti-inflammatory drugs such as non- steroidal anti inflammatory drugs (NSAIDs) have been associated with an increased risk of serious upper gastrointestinal complications. The risk is believed to be dose dependent and can be greater when more than one anti-inflammatory drug is administered. Hence, whenever possible, anti-inflammatory drugs should be administered in mono-therapy. This risk can be more pronounced in case of non-aspirin non- steroidal anti inflammatory (NA-NSAID) drugs. Intake of NA-NSAIDs as a group has been consistently associated with a four- to five-fold increase in upper gastrointestinal complications (UGIC). The evidence indicates that the risk is dose dependent. The estimated pooled cardiovascular Relative Risks (RR) in a recent meta-analysis were 3.0 (95% CI, 2.6-3.4) for low doses, 4.1 (95% CI, 3.6-4.5) for medium doses, and 6.9 (95% CI, 5.8-8.1) for high doses. Recent research indicates that NA-NSAID as a therapeutic class have a RR of 4.1 (95% CI, 3.6-4.8).

[0007] Meloxicam (compound 4), an oxicam derivative, is a member of the enolic acid group of non-steroidal anti-inflammatory drugs (NSAIDs). It is reported to be a selective COX-2 inhibitor. Meloxicam is chemically known as 4-hydroxy-2-methyl-N-(5-methyl-2-thiazolyl)-2H-1,2-benzothiazine-3-carboxamide-1,1-dioxide. It is commercially available under the trade name of MOBIC®. Meloxicam is indicated for relief of the signs and symptoms of osteoarthritis and rheumatoid arthritis, pauciarticular or polyarticular course Juvenile Rheumatoid Arthritis in patients 2 years of age and older.

[0008] Naproxen (compound 5) is another non-steroidal anti-inflammatory drug (NSAID) commonly used for the re-

duction of mild to moderate pain, fever, inflammation and stiffness caused by conditions such as osteoarthritis, rheumatoid arthritis, psoriatic arthritis, gout, ankylosing spondylitis, injury (like fractures), menstrual cramps, tendonitis, bursitis, and the treatment of primary dysmenorrhea. Naproxen, chemically known as (+)-(S)-2-(6-methoxynaphthalen-2-yl) propanoic acid, and naproxen sodium are marketed under various trade names including: Aleve, Anaprox, Naprogesic, Naprosyn, Naprelan, and Synflex. There have also been reports of naproxen producing disturbances in the gastrointestinal tract, like other NSAIDs. Additional NSAIDs include but are not limited to compound 6, Diclofenac; compound 7, Celecoxib; compound 8, Diflunisal; compound 9, Etodolac; compound 10, Fenoprofen; compound 11, Ibuprofen; compound 12, Indomethacin; compound 13, Ketoprofen, and compound 14, Ketorolac.



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[0009] Tramadol (compound 15) is a centrally acting synthetic opioid analgesic. It is chemically (±) cis-2-[(dimethylamino) methyl]-1-(3-methoxyphenyl) cyclo-hexanol hydrochloride. A commercially available form is the hydrochloride salt as Ultram tablets. Tramadol has been used for the management of moderate to moderately severe pain in adults. Tramadol is a non-NSAID analgesic that is not believed to cause the increased risk of stomach ulceration and internal bleeding associated with non-steroidal anti inflammatory drugs (NSAID). However, it still has some commonly reported side effects including nausea, constipation, dizziness, headache, drowsiness, and vomiting. Other reported side effects include itching, sweating, dry mouth, diarrhea, rash, visual disturbances, and vertigo. Thus, it would be desirable to prevent or reduce these side effects by prescribing lower doses of tramadol without compromising pain relief.

[0010] Literature reports indicate that NSAIDs such as naproxen can inhibit the excretion of sodium and lithium. Hence it is desirable to control their dosage to alleviate side effects in patients without comprising the extent of pain relief. Similarly, dizziness, somnolence, dry mouth, edema, blurred vision, weight gain, and "thinking abnormal" (primarily difficulty with concentration/attention) were more commonly reported by subjects treated with GABA analogues like pregabalin.

[0011] There is a strong unmet medical need for drugs that are free from side effects associated with, tramadol, pregabalin and NSAIDs. Considering that such drugs are often used over a long term by elderly patients to manage pain that is often chronic, compositions that can help reduce the dosage or frequency of either or both of drug types without comprising the therapeutic benefits would fill this medical unmet need.

[0012] Opioids have been combined with other drugs including non-opioid analgesic agents, to try to lower the amount of opioid needed to produce an equivalent degree of analgesia and reduce the side effects from opioids. It has been reported that some of these combination products also have a synergistic analgesic effect. See for example, A. Takemori, Annals New York Acad. Sci., 281,262 (1976) where compositions including combinations of opioid analgesics with non-analgesic drugs are reported to exhibit a variety of effects, e.g., sub additive (inhibitory), additive or super additive. Also, R. Taber et al., J. Pharm. Expt. Thera., 169(1), 29 (1969) describes a combination of morphine and methadone, another opioid analgesic. U.S. Patent No. 4,571,400 discloses a combination of dihydrocodeine, an opioid analgesic, and ibuprofen, a non-opioid analgesic. See also U.S. Patent Nos. 4,587,252 and 4,569,937, which disclose other ibuprofen opioid combinations. A. Pircio et al., Arch. Int. Pharmacodyn., 235, 116 (1978) report a 1:125 mixture of butorphanol, another opioid analgesic, and acetaminophen, a non-opioid analgesic has a greater effect than a 1:10 mixture.

[0013] Combinations of non-opioid analgesics have also been prepared to avoid the side effects associated with opioids, and the combinations are noted to have the benefit of requiring less of each ingredient and may provide additive effects. See, *e.g.* G. Stacher et al., Int. J. Clin. Pharmacol. Biooharmacy, 17, 250 (1979), U.S. Patent. No. 4,260,629, U.S. Patent No. 4,132,788. However, there have been warnings against the daily consumption of non-opioid analgesic mixtures and of the consumption of a single non-opioid analgesic in large amounts or over long periods (see, D. Woodbury and E. Fingl at page 349). In addition, ibuprofen, aspirin and some other NSAIDs may cause gastrointestinal side effects especially if used repeatedly. See, *e.g.,* M. J. S. Langman, Am. J. Med. 84 (Suppl. 2A): 15-19, 1988); P. A. Insel in "Goodman and Gilman's The Pharmacological Basis of Therapeutics." Gilman AG, Rall TW, Nies AS, et al. (eds). Pergamon Press, 8th Ed, 1990, Chapter 26, pp. 664-668.

[0014] Gilron et al., Journal of Pain, Saunders, Philadelphia, PA, US, Vol. 6, No. 3, page S41, XP004796276 relate to the results of a randomized controlled trial of a morphine-gabapentin combination in neuropathic pain. Tiippana et al., Anesthesia & Analgesia, Vol. 104, No. 6, pages 1545-1556, XP055059462 relate to a review of Efficacy and Safety

of perioperative Gabapentin and Pregabalin.

[0015] Neuropathic pain is believed to be caused by a primary lesion or dysfunction in the nervous system. Neuropathic pains have been categorized as peripheral neuropathic pain, due to lesion of the peripheral nervous system and central pain following lesions of the central nervous system. The prevalence of neuropathic pain is estimated to be about 1%. Neuropathic pain has been shown to be therapy resistant. However, a number of agents have been used to treat neuropathic pain including NSAIDs, opioids, antidepressants, anticonvulsants, excitatory amino acid antagonists, GABAergic agonists, Substance P antagonists, etc. Low doses of carbamazepine and amitriptyline have been recommended for neuropathic pain in general. The side effects of GABA agonists such as gabapentin, pregabalin, etc. have been documented in the literature. Hence, it is desirable to reduce their dosage to alleviate the patients of its side effects without comprising the extent of pain relief.

[0016] A number of treatments involving the administration of single drugs are currently recommended for pain relief. The single administration of narcotic and non-narcotic analgesics and NSAIDs has been shown to display pain alleviating properties. Some anti-epileptics, such as gabapentin and pregabalin, have also been reported to have pain alleviating properties in diabetic neuropathy.

[0017] Despite the benefits derived from current single drug pain relief regimens, these regimens have disadvantages. There is a strong unmet medical need for drugs that are free from side effects associated with tapentadol, tramadol, GABA analogue or NSAIDs. One area of concern relates to the incidence of unwanted side effects caused by many of the pain treatment regimens available today. Opioid analgesics, such as morphine, are sparingly prescribed for pain because of the well-known addictive effects and significant central nervous system (CNS) side effects and gastrointestinal side effects. Further tapentadol is known to elicit adverse effects, including nausea, vomiting, sleepiness, dizziness, itchiness, sedation, dry mouth, sweating and constipation.

[0018] However, a pharmaceutical composition comprising a slow release tapentadol and a second analgesic, wherein the second analgesic is tramadol, gamma-aminobutyric acid (GABA) analogue or an NSAID for treating a patient in need there of. Further, the prior art doesn't disclose a method of treating pain or pain related disorder comprising a method of administering to a mammal in need thereof, a pharmaceutical composition comprising a slow release tapentadol and a second analgesic, wherein the second analgesic is tramadol, gamma-aminobutyric acid (GABA) analogue or an NSAID is not disclosed. As there is a continuing need for analgesic medications that provide high efficacy pain relief with reduced undesirable effects.

Summary

[0019] The present invention provides a pharmaceutical combination comprising a slow release tapentadol and a second analgesic agent. The second analgesic agent can be a gamma-aminobutyric acid (GABA) analogue, whereby said GABA analogue is pregabalin or a pharmaceutically acceptable salt thereof. The invention further provides a method for treating pain and pain related disorders in a mammal, comprising administering to said mammal an effective amount of a composition comprising a slow release tapentadol and a second analgesic agent.

[0020] In another embodiment the invention provides a tapentadol / analgesic combination for treating moderate to severe painful conditions associated with diabetic neuropathy, rheumatoid arthritis, osteoarthritis and the like, by administering to a subject in need thereof, an analgesic pharmaceutical combination comprising from about 25 to about 400 mg of slow (controlled) release tapentadol and a second analgesic agent, wherein the second analgesic agent is about 5 to about 500 mg of a GABA agonist, with pharmaceutically acceptable carrier so as to provide better pain management. In the pharmaceutical composition, the tapentadol is in a controlled release form and a GABA analogue are present in an immediate release form, extended (controlled) release form or delayed release form along with pharmaceutically acceptable carrier.

[0021] An advantage of the disclosed compositions is a decreased dosing of the active ingredients, such as the GABA analogue to the patient which can promote better patient compliance. Further, the compositions comprising from about 25 to about 400 mg of slow release tapentadol and a second analgesic agent, wherein the second analgesic is a GABA agonist with pharmaceutically acceptable carrier so as to provide better pain management.

Brief Description of the Figures

[0022]

FIGURE 1 illustrates a comparison of the in vitro dissolution profiles of tapentadol HCl in slow release tapentadol HCl 100 mg and naproxen 250 mg tablets.

FIGURE 2 illustrates a comparison of the LS mean change from baseline in VAS score for the combination drug comprising slow release tapentadol 100 mg and naproxen 250 mg with those of tapentadol and naproxen mono

therapies based upon the average of Weeks 1-6.

FIGURE 3 illustrates a weekly LS mean changes from baseline for the four treatment groups.

FIGURE 4 illustrates a mean VAS pain score changes for four formulations; tapentadol 100 MG, pregabalin 250 MG, and slow release tapentadol 100 MG + pregabalin 250 MG fixed dose combination.

FIGURE 5 illustrates a mean VAS pain score changes for three formulations; Tramadol 50 mg, Tapentadol 100 MG, Placebo and Slow Release Tapentadol 100 MG + Tramadol 50 MG fixed dose combination.

FIGURE 6 illustrates a mean VAS pain score changes for the four formulations; tapentadol 100 MG, gabapentin 250 MG, and slow release tapentadol 100 plus gabapentin 250 MG fixed dose combination.

Detailed Description

[0023] One object of the present invention is to provide methods, which can be used in the treatment of pain and pain related diseases wherein the methods comprise administration of a therapeutically effective amount of a slow release tapentadol and a second analgesic agent, wherein the second analgesic agent is a gamma-aminobutyric acid (GABA) analogue to a patient in need thereof.

[0024] The two analgesic agents *e.g.* a slow release tapentadol and a second analgesic agent may be co-administered in a single medicament or they may be administered separately as two medicaments. Further, the first drug may (tapentadol) be administered in a regimen, which additionally comprises administration of the second drug separately or in a composition with the first drug.

[0025] In yet another embodiment, the invention provides a slow release tapentadol and a second analgesic agent are administered in suboptimal dosages.

[0026] In yet another embodiment, the invention provides a slow release tapentadol and a second analgesic agent are administered in amounts and for a sufficient time to produce a synergistic effect.

[0027] In yet another embodiment, the invention provides a composition where the second active agent is included in an immediate release coating.

[0028] In yet another embodiment, the invention provides a bilayer composition where one layer includes the tapentadol and one layer comprises the second active agent.

[0029] In yet another embodiment, there is provided a method of treating moderate to severe pain by administering to a subject in need thereof, a pharmaceutical composition comprising 5-500 mg tapentadol or a second analgesic, wherein the second analgesic is from about 5 to about 500 mg of a GABA agonist in admixture with pharmaceutically acceptable carrier.

[0030] Still further, a titration dosing regimen for the administration of slow release tapentadol to patients. The titration dosing regimen provides a significant reduction in the occurrence of adverse effects from the introduction of slow release tapentadol dosing, thus increasing patient compliance and medication tolerability.

[0031] In yet another embodiment, the invention provides a composition of tapentadol and a second analgesic agent, or a pharmaceutically acceptable salts thereof for use in medical treatment (for example, treatment of pain, e.g., neuropathic pain).

[0032] In yet another embodiment, the invention provides a method for the use of tapentadol and a second analgesic agent, or a pharmaceutically acceptable salts thereof to prepare a medicament for treatment of pain in a mammalian species (for example, a human).

DETAILED DESCRIPTION OF THE INVENTION

[0033] The term "Analgesic" as used in this invention means to include any drug used to relieve pain including paracetamol (acetaminophen), the non-steroidal anti-inflammatory drugs (NSAIDs) such as the salicylates, narcotic drugs such as morphine, synthetic drugs with narcotic properties such as tramadol, GABA analogues like pregabalin , gabapentin and various others other classes of drugs not normally considered analgesics are used to treat neuropathic pain syndromes; these include tricyclic antidepressants and anticonvulsants

[0034] The term "band range" for purposes of the present invention is defined as the difference in *in vitro* dissolution measurements of the controlled release formulations when comparing the dissolution profile (curve) obtained by the formulation upon completion of the manufacturing of the coated product (prior to storage) and the dissolution profile obtained after the coated product is exposed to accelerated storage conditions, expressed as the change in percent of the active agent released from the coated product at any dissolution time point along the dissolution curves.

[0035] The term "co-administration" as used herein means administration of the two drugs (agents) together (*e.g.*,

simultaneously as a mixture) or administration can be sequential. The sequential administration of the tapentadol can be prior to or after administration of the second analgesic agent, within minutes of each other or up to about 48 hours after the administration of the other agent. Preferably the administration of the tapentadol will be within about 24 hours of administration of the second analgesic agent and more preferably within about 12 hours of administration of the second analgesic agent.

[0036] The term "effective amount" as used herein means a dosage to produce a selected effect. For example, an effective amount of an analgesic is an amount which is sufficient in order for the pain of the patient to be reduced when compared with no treatment.

[0037] The term "GABA analogue" as used in this invention means pregabalin, its pharmaceutically equivalent salts, isomers, polymorphs, hydrates, complexes or clatharates and the like.

[0038] The term "NSAID" as used in this specification means any non-steroidal anti-inflammatory drug. Non-limiting examples include Celecoxib, Diclofenac, Diflunisal, Etodolac, Fenoprofen, Flurbirofen, Ibuprofen, Indomethacin, Ketoprofen, Ketorolac, Mefenamic acid, Meloxicam, Nabumetone, Naproxen, Oxaprozin, Piroxicam, Sulindac and Tolmetin and their pharmaceutically equivalent salts, isomers, polymorphs, hydrates, complexes, or clatharates and the like.

[0039] In yet another embodiment, the invention provides a composition where the second active agent is included in an immediate release coating. The pharmaceutical composition of any of claims 1-5, wherein the composition exhibits an in vitro dissolution profile such that after 2 hours, from about 0% to about 30% by weight of tapentadol is released, after 4 hours, from about 5% to about 22% by weight of tapentadol is released, after 6 hours, from about 15% to about 30% by weight of tapentadol is released, and after 8 hours, more than about 40% by weight of tapentadol is released; when measured using the USP Basket Method at 75 rpm in 900 ml 0.1 N HCl at 37° C.

[0040] The term "medicament" as used herein means a pharmaceutical composition suitable for administration of the pharmaceutically active compound to a patient.

[0041] The term "pharmaceutically-acceptable salt" refers to salts that retain the biological effectiveness and properties of the disclosed compounds and which are not biologically or otherwise undesirable. In many cases, the disclosed compounds are capable of forming acid or base salts by virtue of the presence of amino or carboxyl groups or groups similar thereto. The preparation of the salts and suitable acids or bases is known in the art.

[0042] The term "suboptimal dosage" us used herein means a dosage which is below the optimal dosage for that compound when used in single-compound therapy.

[0043] The term "additive effect" as used herein means the effect resulting from the sum of the effects obtained from the individual compounds.

[0044] The term "synergistic effect" as used herein means an effect which is greater than the additive effect which results from the sum of the effects of the two individual compounds.

[0045] The term "treatment of a disease" as used herein means the management and care of a patient having developed the disease, condition or disorder. The purpose of treatment is to combat the disease, condition or disorder. Treatment includes the administration of the active compounds to eliminate or control the disease, condition or disorder as well as to alleviate the symptoms or complications associated with the disease, condition or disorder.

[0046] The term "prevention of a disease" as used herein is defined as the management and care of an individual at risk of developing the disease prior to the clinical onset of the disease. The purpose of prevention is to combat the development of the disease, condition or disorder, and includes the administration of the active compounds to prevent or delay the onset of the symptoms or complications and to prevent or delay the development of related diseases, conditions or disorders.

[0047] The term "pain and pain related conditions" as used herein is defined as any pain due to a medical condition including neuropathic pain, osteoarthritis, rheumatoid arthritis, fibromyalgia, and back, musculoskeletal pain, Ankylosing spondylitis, juvenile rheumatoid arthritis, migraines, dental pain, abdominal pains, ischemic pain, postoperative pain or because of an anesthetic or surgical contrition

[0048] The term "extended release material" as present in the inner solid particulate phase or the outer solid continuous phase refers to one or more hydrophilic polymers and/or one or more hydrophobic polymers and/or one or more other type hydrophobic materials, such as, for example, one or more waxes, fatty alcohols and/or fatty acid esters. The "extended release material" present in the inner solid particulate phase may be the same as or different from the "extended release material" present in the outer solid continuous phase.

[0049] The term "slow-release" or "controlled release" as used herein applies to any release from a formulation that is other than an immediate release wherein the release of the active ingredient is slow in nature. This includes various terms used interchangeably in the pharmaceutical context like extended release, delayed release, sustained release, controlled release, timed release, specific release and targeted release etc.

[0050] The term "candidate for sustained release" encompasses all the characteristics of a drug which make it a candidate for formulating it into an extended release fashion like a short elimination half life and consequent dosing of more than once a day, a single dose product given in an extended fashion to achieve better clinical results and avoid side effects associated with an immediate release etc.

[0051] The term "binding agent" as used in this specification, refers to any conventionally known pharmaceutically acceptable binder such as polyvinyl pyrrolidone, hydroxypropyl cellulose, hydroxyethyl cellulose, hydroxypropyl methylcellulose, ethylcellulose, polymethacrylate, polyvinylalcohol, waxes and the like. Mixtures of the aforementioned binding agents may also be used. The preferred binding agents are water soluble materials such as polyvinyl pyrrolidone having a weight average molecular weight of 25,000 to 3,000,000. The binding agent may comprise approximately about 0 to about 40% of the total weight of the core and preferably about 3% to about 15% of the total weight of the core. In one embodiment, the use of a binding agent in the core is optional.

[0052] The term "pharmaceutically acceptable derivative" means various pharmaceutical equivalent isomers, enantiomers, complexes, salts, hydrates, polymorphs, esters etc of tapentadol or tramadol or a GABA analogue or an NSAID.

[0053] The term "therapeutically effective amount" means an amount that elicits a biological response in a mammal including the suboptimal amount.

[0054] The term "hydrophilic polymers" as used in this specification include, but are not limited, to hydroxypropylmethylcellulose, hydroxypropylcellulose, sodium, carboxymethyl- cellulose, carboxymethylcellulose calcium, ammonium alginate, sodium alginate, potassium alginate, calcium alginate, propylene glycol alginate, alginic acid, polyvinyl alcohol, povidone, carbomer, potassium pectate, potassium pectinate, etc.

[0055] The term " hydrophobic polymers" as used in this specification include, but are not limited, to ethyl cellulose, hydroxyethylcellulose, ammonio methacrylate copolymer (Eudragit™ RL or Eudragit™ RS), methacrylic acid copolymers (Eudragit™ L or Eudragit™ S), methacrylic acid-acrylic acid ethyl ester copolymer (Eudragit™ L 100-5), methacrylic acid esters neutral copolymer (Eudragit™ NE 30D), dimethylaminoethylmethacrylate-methacrylic acid esters copolymer (Eudragit™ E 100), vinyl methyl ether/malefic anhydride copolymers, their salts and esters (Gantrez™) etc.

[0056] Other hydrophobic materials which may be employed in the inner solid particulate phase and/or outer-solid continuous phase include, but are not limited, to waxes such as beeswax, carnauba wax, microcrystalline wax, and ozokerite; fatty alcohols such as cetostearyl alcohol, stearyl alcohol; cetyl alcohol myristyl alcohol etc; and fatty acid esters such as glyceryl monostearate, glycerol monooleate, acetylated monoglycerides, tristearin, tripalmitin, cetyl esters wax, glyceryl palmitostearate, glyceryl behenate, hydrogenated castor oil, etc.

[0057] Non-limiting examples of NSAIDs for the compositions include Celecoxib, Diclofenac, Diflunisal, Etodolac, Fenoprofen, Flurbirofen, Ibuprofen, Indomethacin, Ketoprofen, Ketorolac, Mefenamic acid, Meloxicam, Nabumetone, Naproxen, Oxaprozin, Piroxicam, Sulindac and Tolmetin and their pharmaceutically equivalent salts, isomers, polymorphs, hydrates, complexes, or clatharates and the like.

[0058] The disclosed pharmaceutical compositions can exhibit an in vitro dissolution profile such that after 2 hours, from about 0% to about 30% by weight of tapentadol is released, after 4 hours, from about 5% to about 22% by weight of tapentadol is released, after 6 hours, from about 15% to about 30% by weight of tapentadol is released, and after 8 hours, more than about 40% by weight of tapentadol is released; when measured using the USP Basket Method at 75 rpm in 900 ml 0.1 N HCl at 37° C.

[0059] The tramadol material is any one of (1R, 2R or 1S, 2S)-(dimethyl aminomethyl)-1-(3-methoxyphenyl)-cyclohexanol (tramadol), its N-oxide derivative ("tramadol N-oxide"), and its O-desmethyl derivative ("O-desmethyl tramadol") or mixtures thereof. It also includes the individual stereoisomer, mixtures of stereoisomer, including the racemates, pharmaceutically acceptable salts of the amines, such as the hydrochloride salt, solvates and polymorphs of the tramadol material. Tramadol is commercially available from Gnmenthal or may be made by the process described in U.S. Pat. No. 3,652,589 and it is herein incorporated by reference.

[0060] A combination comprising a slow release tapentadol and a second analgesic agent, wherein the second analgesic agent is a gamma-aminobutyric acid (GABA) analogue. The disclosed compositions preferably contain a therapeutically effective amount of tapentadol or a pharmaceutically acceptable salt thereof, wherein the tapentadol is in the range of from about 5 to about 800 mg, preferably from about 50, to about 600 mg, more preferably from about 100 to about 400 mg and more preferably from about 200 to about 300 mg (calculated as tapentadol hydrochloride) per dosage unit and a therapeutically effective amount of a second analgesic agent, wherein the second analgesic is from about 5 to about 500 mg of gamma-aminobutyric acid (GABA) analogue.

[0061] The disclosed composition can be, for example, as granules, spheroids, pellets, multiparticulates, capsules, patches tablets, sachets, controlled release suspensions, or in any other suitable dosage form incorporating such granules, spheroids, pellets or multiparticulates.

[0062] The compositions can be, *e.g.*, coated tablets wherein the coating includes at least one water-insoluble, water permeable film-forming polymer, at least one plasticizer and at least one water-soluble polymer, and the second active agent. In a preferred form the coating can have at least one water-insoluble, water-permeable film-forming polymer varies from about 20% to about 90% of the coating dry weight, the proportion of the at least one plasticizer varies from about 5% to about 30% of the coating dry weight, and the proportion of the at least one water-soluble polymer varies from about 10% to about 75% of the coat dry weight.

[0063] A preferred water-insoluble, water-permeable film-forming polymer is ethylcellulose. A preferred water-insoluble polymer is polyvinylpyrrolidone. A preferred plasticizer is dibutyl sebacate.

[0064] The one or more of active ingredient in the combination according to the present invention may suitably be incorporated in a matrix. This may be any matrix, known to a person skilled the art, that affords slow release tapentadol over at least about a twelve hour period and preferably that affords in-vitro dissolution rates and in vivo absorption rates of tapentadol within the therapeutically effective ranges. The combination according to the present invention may preferably use a slow release matrix. Alternatively, normal release matrices having a coating which provides for slow release of the tapentadol may be used.

[0065] The slow release matrix employed in the combination of this invention may also contain other pharmaceutically acceptable ingredients which are conventional in the pharmaceutical art such as diluents, lubricants, binders, granulating aids, colorants, flavorants, surfactants, pH adjusters, anti-adherents and glidants, *e.g.,* dibutyl sebacate, ammonium hydroxide, oleic acid and colloidal silica. Any known diluent *e.g.* microcrystalline cellulose, lactose and dicalcium phosphate may be used to prepare this combination. Suitable lubricants are e.g. magnesium stearate and sodium stearyl fumarate. Suitable binding agents are e.g. hydroxypropyl methyl cellulose, polyvidone and methyl cellulose. Suitable disintegrating agents are starch, sodium starch glycolate, crospovidone, and croscarmellose sodium.

[0066] The surface actives that are suitable for this invention are Poloxamer 188.RTM, polysorbate 80 and sodium lauryl sulfate. The suitable flow aids for this invention are talc colloidal anhydrous silica. Non-limiting water soluble polymers that may be used to prepare the matrix include PEG having weight average molecular weights in the range of from about 1000 to about 6000. The combination comprising the slow release tapentadol according to the invention may conveniently be film coated using any film coating material conventional in the pharmaceutical art but preferably an aqueous film coating is used.

[0067] Alternatively, the combination comprising a slow release tapentadol and a second analgesic, wherein the second analgesic is a gamma-aminobutyric acid (GABA) analogue as per this invention may comprise a normal release matrix having a slow release coating. Preferably the combination comprises film coated spheroids containing the active ingredient and a spheronising agent. The spheronising agent may be any suitable pharmaceutically acceptable material which may be spheronised together with the active ingredient to form spheroids. A preferred spheronising agent as per this invention is microcrystalline cellulose. The microcrystalline cellulose used may suitably be, for example, Avicel™ PH 101 or Avicel™ PH 102 (FMC Corporation). The spheroids may optionally contain other pharmaceutically acceptable ingredients conventional in the pharmaceutical art such as binders, bulking agents and colorants. Suitable binders may include water soluble polymers, water soluble hydroxyalkyl celluloses such as hydroxypropylcellulose or water insoluble polymers (which may also contribute controlled release properties) such as acrylic polymers or copolymers for example ethylcellulose. Suitable bulking agents include lactose.

[0068] The spheroids are coated with a material which permits release of the active ingredient at a slow rate in an aqueous medium. Suitable slow release coating materials that may be used in this invention include water insoluble waxes and polymers such as polymethylacrylates (for example Eudragit™ polymers) or water insoluble celluloses, particularly ethylcellulose. Optionally, water soluble polymers such as polyvinylpyrrolidone or water soluble celluloses such as hydroxypropylmethylcellulose or hydroxypropylcellulose may be included. Optionally other water soluble agents such as polysorbate 80 may be added.

[0069] Further in an alternative embodiment, a flux-enhancing agent can also be included in the membrane or slow release coating can include one of the above-described polymers. The flux enhancing agent can increase the volume of fluid imbibed into the core to enable the dosage form to dispense substantially all of the tapentadol through the passage and/or the porous membrane. The flux-enhancing agent can be a water-soluble material or an enteric material. Examples of the preferred materials that are useful as flux enhancers include but not limited to sodium chloride, potassium chloride, sucrose, sorbitol, mannitol, polyethylene glycols (PEG), propylene glycol, hydroxypropyl cellulose, hydroxypropyl methycellulose, hydroxypropyl methycellulose phthalate, cellulose acetate phthalate, polyvinyl alcohols, methacrylic acid copolymers, poloxamers (such as LUTROL™ F68, LUTROL F127, LUTROL F108 which are commercially available from BASF) and mixtures thereof. A preferred flux-enhancer used in this invention is PEG 400.

[0070] The flux enhancer may also be a water miscible/soluble drug such as Tapentadol or its pharmaceutically acceptable salts, or the flux enhancer may be a drug that is soluble under intestinal conditions. If the flux enhancer is a drug, the present pharmaceutical composition has an added advantage of providing an immediate release of the drug that has been selected as the flux enhancer. The flux enhancing agent dissolves or leaches from the membrane or sustained release coating to form channels in the membrane or sustained release coating which enables fluid to enter the core and dissolve the active ingredient. In the preferred embodiment, the flux enhancing agent comprises approximately 0 to about 40% of the total weight of the coating, most preferably from about 2% to about 20% of the total weight of the coating.

[0071] A commonly known excipient such as a plasticizer may also be used for preparing the membrane or slow release coating Some commonly known plasticizers include but not limited to adipate, azelate, enzoate, citrate, stearate, isoebucate, sebacate, triethyl citrate, tri-n-butyl citrate, acetyl tri-n-butyl citrate, citric acid esters, and all those described in the Encyclopedia of Polymer Science and Technology, Vol. 10 (1969), published by John Wiley & Sons. The preferred plasticizers are triacetin, acetylated monbglyceride, grape seed oil, olive oil, sesame oil, acetyltributylcitrate, acetyltri-

ethylcitrate, glycerin sorbitol, diethyloxalate, diethylmalate, diethylfumarate, dibutylsuccinate, diethylmalonate, dioctylphthalate, dibutylsebacate, triethylcitrate, tributylcitrate, glyceroltributyrate and the like. Though the exact amount used depends on the type of plasticizer used, typically amounts from 0 to about 25% are used, and preferably from about 2% to about 15% of the plasticizer can be used based upon the total weight of the membrane or sustained release coating.

[0072] Generally, the membrane or slow release coating around the core will comprise from about 1% to about 20% and preferably about 2% to about 10% based upon the total weight of the core and coating.

[0073] The membrane or sustained release coating surrounding the core can further comprise a passage that will allow for controlled release of the drug from the core in a preferred embodiment. As used herein the term passage includes an aperture, orifice, bore, hole, weakened area or a credible element such as a gelatin plug that erodes to form an osmotic passage for the release of the tapentadol from the dosage form. Passage used in accordance with the subject invention is well known and are described in U.S. Pat. Nos. 3,845,770; 3,916,899; 4,034,758; 4,077,407; 4,783,337 and 5,071,607.

[0074] The following examples 1-13 are shown for illustrating the invention related to combination comprising a slow release tapentadol and a second analgesic, wherein the second analgesic is a gamma-aminobutyric acid (GABA) analogue.

[0075] The following examples 1-8, 11 and 13 are outside of the scope of the present invention.

Example 1:

[0076]

TABLE 1

Combination of slow release tapentadol 100 mg and naproxen 250 mg tablets	
First Active Ingredient mg/tablet	Example 1
Tapentadol Hydrochloride	100.0
Microcrystalline Cellulose	10.0
Colloidal Silicon Dioxide	1.5
Polyvinylpyrrolidone	4.5
Hydrogenated Vegetable Oil	5.0
Water*	Q.S
Coat	
Ethylcellulose Aqueous Dispersion	15.00
Polyvinylpyrrolidone	5.0
Polyethylene Glycol	2.0
Water*	Q.S
Second Active Ingredient	
Naproxen	250.0
Povidone K 30 USP	12
Microcrystalline cellulose	25
Croscarmellose sodium	15
Magnesium Stearate	3
Water*	Q.S.
* Removed during processing	

Manufacturing Process

[0077] The combination comprising a slow release tapentadol hydrochloride tablets and naproxen were manufactured in two phases using standard coating processes. In phase I, the Tapentadol Hydrochloride was formulated into a core that was further coated with slow release coat to get a slow release tapentadol core. In Phase II, this slow release coated Tapentadol hydrochloride core was coated with an immediate release layer comprising Naproxen as per details are given below;

[0078] **Phase I, Core preparation:** Tapentadol HCl is mixed with microcrystalline cellulose and colloidal silicone

dioxide and one or mixture of filler and granulated using suitable method known in the art using a binder solution comprising Polyvinylpyrrolidone or polyvinyl alcohol. The granulated tapentadol hydrochloride was dried and screened. This is further lubricated using hydrogenated vegetable oil with or without glidant. The lubricated blend is compressed into tablets using a compression machine.

[0079] **Coating Solution and Coating:** The coating solution is prepared using aqueous dispersion of water insoluble water permeable polymer of Ethylcellulose with water soluble polymer of Polyvinylpyrrolidone or hydroxypropyl methyl cellulose. Polyethylene glycol mixture prepared using propeller stirrer and the same is homogenized using suitable homogenizer. The core tablets are coated using coating solution using standard coater like O' Hara pan coater tip set at 4" at a spray rate of 25 mL/gun/min, exhaust temperature of around 45'C, an atomization pressure from 10-35 psi at a pan speed of 5-8 rpm, using airflow 350 CFM.

[0080] **Phase II:** In phase II, Naproxen formulation prepared using granulation technique known in the art and then blended with disintegrant and lubricant. Tapentadol slow release tablets prepared in Phase I is coated with lubricated blend of Naproxen formulation using compression coating machine where Tapentadol slow release tablets is used as a core and an immediate layer of Naproxen formulation forms an outer layer.

[0081] The naproxen coating was applied to slow release coated 100 mg tapentadol hydrochloride tablets using the above mentioned coater. Over this naproxen coated seal coated 100 mg tapentadol hydrochloride tablets, color coating was done using similar coat. The spraying was done at a temperature of 46-47' C, atomization pressure of 40-60 psi at a spray rate of 180 grams per minute/three guns. The pan speed was at 4-8 rpm and air volume of 1000±100.

[0082] Finally, color coated tablets were dried and optionally polished using Cindrella wax and the finished final tablets were packaged in a HDPE bottle with a suitable desiccant and subjected appropriate stability and clinical studies. In Example 1, a pharmaceutical composition comprising 100 mg of slow release tapentadol and 250 mg naproxen was formulated as per Table 1.

[0083] The manufacturing processes exemplified here are only meant for illustration and the pharmaceutical composition can be prepared using a number of methods well established in the art. Other exemplified formulations are listed below in tables 2-5;

Example 2:

[0084]

TABLE 2

Combination of slow release tapentadol 100 mg and naproxen 250 mg tablets	
First Active Ingredient mg/tablet	**Quantity mg**
Tapentadol HCl	100.0
Polyvinyl Alcohol	2.0
Colloidal Silicon Dioxide (Abrosil™ 200)	1.0
Sodium Stearyl Fumarate	1.0
Water*	Q.S
Core Weight	104.0
Coat	
Ethylcellulose (Ethocel™ PR 100)	9.20
Polyvinylpyrrolidone (Kollidon™ 90F)	4.14
Dibutyl Sebacate	2.66
Denatured Alcohol*	Q.S
Second Active Ingredient mg/tablet	
Naproxen	250.0
Povidone K 30 USP	12
Microcrystalline cellulose	25
Croscarmellose sodium	15
Magnesium Stearate	3

(continued)

Second Active Ingredient mg/tablet	
Water*	Q.S.
* Removed during processing	

Example 3

[0085]

TABLE 3

Combination of slow release tapentadol 100 mg and naproxen 250 mg tablets	
First Active Ingredient mg/tablet	**Quantity mg**
Tapentadol HCl	100.0
Polyvinyl Alcohol	2.0
Colloidal Silicon Dioxide (Abrosil™ 200)	1.0
Sodium Stearyl Fumarate	1.0
Water*	Q.S
Core Weight	104.0
Coat	
Ethylcellulose (Ethocel™ PR 100)	9.87
Polyvinylpyrrolidone (Kollidon™ 90F)	3.47
Dibutyl Sebacate	2.67
Denatured Alcohol*	Q.S
Second Active Ingredient mg/tablet	
Naproxen	250.0
Povidone K 30 USP	12
Microcrystalline cellulose	25
Croscarmellose sodium	15
Magnesium Stearate	3
Water*	Q.S.
* Removed during processing	

Example 4

[0086]

TABLE 4

Combination of slow release tapentadol 100 mg and naproxen 250 mg tablets	
First Active Ingredient mg/tablet	**Quantity mg**
Tapentadol HC1	100.0
Polyvinyl Alcohol	2.0
Colloidal Silicon Dioxide (Abrosil® 200)	1.0
Sodium Stearyl Fumerate	1.0
Water*	Q.S
Core Weight	104.0

(continued)

Coat	
Ethylcellulose (Ethocel® PR 100)	9.87
Polyvinylpyrrolidone (Kollidon® 90F)	3.73
Dibutyl Sebacate	2.67
Denatured Alcohol*	Q.S
Second Active Ingredient mg/tablet	
Naproxen	250.0
Povidone K 30 USP .	12
Microcrystalline cellulose	25
Croscarmellose sodium	15
Magnesium Stearate	3
Water*	Q.S.
* Removed during processing	

Example 5

[0087]

TABLE 5

Combination of slow release tapentadol 200 mg and naproxen 500 mg tablets	
First Active Ingredient mg/tablet	**Quantity mg**
Tapentadol HCl	200.0
Polyvinyl Alcohol	4.0
Colloidal Silicon Dioxide (Abrosil™ 200)	2.0
Sodium Stearyl Fumarate	2.0
Water*	Q.S
Core Weight	2.8
Coat	
Ethylcellulose (Ethocel™ PR 100)	12.28
Polyvinylpyrrolidone (Kollidon™ 90F)	6.50
Dibutyl Sebacate	3,75
Denatured Alcohol*	Q.S
Second Active Ingredient mg/tablet	
Naproxen	500.0
Povidone K 30 USP	20
Microcrystalline cellulose	40
Croscarmellose sodium	25
Magnesium Stearate	6
Water*	Q.S.
* Removed during the process	

[0088] Core Preparation; Tapentadol HCl and colloidal silicon dioxide were mixed and passed through a 1.0 mm screen. Polyvinyl alcohol was dissolved in purified water. The mixed tapentadol HCl and colloidal silicon dioxide powder was granulated with the aqueous solution of polyvinyl alcohol in a fluidized bed granulator, Glatt GPCGI and then dried. After granulation, the granules were blended with sodium stearyl fumarate and then passed through a 1.0 mm screen. The blend was then compressed into tablet cores using a Manesty Betapress.

[0089] **Coating Preparation;** The ethyl alcohol and isopropanol were weighed and mixed. Dibutyl sebacate and ethylcellulose were added to and dissolved in the ethyl alcohol and isopropyl alcohol while stirring using a propeller stirrer, Coframo RZR1. The ethylcellulose and dibutyl sebacate were allowed to dissolve completely. The polyvinylpyrrolidone was added. The solution was stirred until all components were dissolved. The solution was passed through a high pressure homogenizer, Mini DeBee 2000 with a #7 nozzle, Bee International. The tablet cores were coated using the coating solution in a perforated coating pan, O'Hara Labcoat I11 36" Pan, Vector LCDS. The coating parameters are listed in Table 6;

Table 6

Coating Parameters	
Inlet Temperature:	48.5-49.5' C
Outlet Temperature:	38.5-39.5' C
Bed Temperature:	37.5-38.5' C
Spray Rate:	300g/minute
Atomizing Air/ Pattern:	25/25 psi
Distance gun/Bed:	6"
Distance between guns:	6"
Pan speed:	12rpm
Coating Amount Diameter:	
Thickness:	6 mm
Cup Height:	4.65 mm
Surface:	1.02 mm
Percentage:	112 mm^2
Amount:	16 mg

Example 6

[0090] In yet another example, the invention discloses a pharmaceutical composition which can effectively be used in the treatment of pain and pain related diseases wherein the compositions comprise a therapeutically effective amount of a slow release tapentadol and an NSAID to a patient in need can be formulated in other ways. For Example, the combination comprising a slow release tapentadol and an NSAID such as Naproxen was prepared as a bilayer tablet as exemplified below:

Layer 1:

Tapentadol HCl	200 mg
Microcrystalline cellulose	10-25%
Polyvinyl alcohol	3-5%
Ethylcellulose (5- 20 cp)	10-20%
Hydroxyethyl cellulose	5-15%
Colloidal silicon dioxide	2-5%
Sodium stearyl fumarate	1-2%

Layer 2:

Naproxen	250 mg
Microcrystalline cellulose	5-20%
Povidone	10-15%
Crosscarmellose sodium	5-10%
Magnesium stearate	0.5-2%

[0091] Preparation of Layer 1: Tapentadol Hydrochloride, microcrystalline cellulose and colloidal silicon dioxide were granulated with polyvinyl alcohol and dried. The dried granules are mixed with Ethylcellulose and Hydroxyethylcellulose

and lubricated with Sodium stearyl fumarate.

[0092] Preparation of Layer 2: Naproxen mixed with microcrystalline cellulose was granulated with povidone. Granules are dried and mixed with Crossearmellose sodium and finally lubricated with Magnesium stearate.

[0093] Compression: Layer 1 and Layer 2 are loaded into the hopper of Bilayer rotary compression machine and compressed with a desired hardness.

Example 7

[0094]

Table 7

Meloxicam Combination	
First Active Ingredient mg/tablet	
Tapentadol Hydrochloride	100.0
Lactose	65.0
Ethyl Cellulose	16.0
Cetostearyl Alcohol	43.0
Magnesium Stearate	2.0
Talc	4.0
Hydroxyethyl Cellulose	
Water *	qs
Coat	
Hydroxypropyl methyl cellulose	0.75
Hydroxy methyl cellulose	3.75
Opaspray	2.60
PEG 400	0.60
Talc	0.30
Water *	qs
Second Active Ingredient	
Meloxicam	7.5
Povidone K 30 USP	1.0
Lactose	25.0
Sodium starch Glycolate	7.5
Poloxamer 188	3.0
HPMC	1.5
PEG 8000	0.4
Titanium Dioxide	0.4
Wax	0.2
* Removed during processing	

Example 8

[0095]

Table 8

Meloxicam Combination	
First Active Ingredient mg/tablet	
Tapentadol Hydrochloride	100.0
Lactose	66.0
Ethyl Cellulose	0.0

(continued)

Meloxicam Combination	
First Active Ingredient mg/tablet	
Cetostearyl Alcohol	44.0
Magnesium Stearate	2.0
Talc	4.0
Hydroxyethyl Cellulose	14.0
Water *	qs
Coat	
Hydroxypropyl methyl cellulose	0.75
Hydroxy methyl cellulose	3.75
Opaspray	2.60
PEG 400	0.60
Talc	0.30
Water *	qs
Second Active Ingredient	
Meloxicam	7.5
Povidone K 30 USP	1.0
Lactose	25.0
Sodium starch Glycolate	7.5
Poloxamer 188	3.0
HPMC	1.5
PEG 8000	0.4
Titanium Dioxide	0.4
Wax * Removed during processing	0.2

Manufacturing Process, Slow release tapentadol hydrochloride and meloxicam

[0096] The combination comprising a slow release tapentadol hydrochloride tablets and meloxicam were manufactured using standard granulation and coating processes. Tapentadol hydrochloride and lactose were granulated together in a granulator and sprayed with ethylcellulose and water. The granulated tapentadol hydrochloride was dried and screened. The tapentadol hydrochloride granules were mixed with Cetostearyl alcohol. Talc and magnesium stearate were mixed with the tapentadol hydrochloride and the granules were compressed into tablets. The compressed tablets were coated using the coating constituents The above prepared coated slow release tapentadol tablets were further seal coated with Opadry Clear (YS-1-7006) solution using standard coater like O' Hara pan coater tip set at 4" at a spray rate of 25 mL/gun/min, exhaust temperature of around 45'C, an atomization pressure from 10-35 psi at a pan speed of 5-8 rpm, using airflow 350 CFM. The meloxicam coating was applied to coated 100 mg tapentadol hydrochloride tablets using the above mentioned coater. Over this 7.5 mg meloxicam coated seal coated 100 mg tapentadol, hydrochloride tablets, color coating was done using similar coat. The spraying was done at a temperature of 46-47' C, atomization pressure of 40-60 psi at a spray rate of 180 grams per minute/three guns. The pan speed was at 4-8 rpm and air volume of 1000 ± 100.

[0097] Finally, color coated tablets were dried and polished using Cindrella wax and the finished final tablets were packaged in a HDPE bottle with a suitable desiccant and subjected appropriate stability and clinical studies.

Example 9

[0098]

Table 9

Pregabalin Combination	
First Active Ingredient mg/tablet	
Tapentadol Hydrochloride	100.0

(continued)

Pregabalin Combination	
First Active Ingredient mg/tablet	
Lactose	65.0
Ethyl Cellulose	16.0
Cetostearyl Alcohol	43.0
Magnesium Stearate	2.0
Talc	4.0
Hydroxyethyl Cellulose	
Water *	qs
Coat	
Hydroxypropyl methyl cellulose	0.75
Hydroxy methyl cellulose	3.75
Opaspray	2.60
PEG 400	0.60
Talc	0.30
Water *	qs
Second Active Ingredient	
Pregabalin	250
Povidone K 30 USP	1.0
Lactose	25.0
Sodium starch Glycolate	7.5
Poloxamer 188	3.0
HPMC	1.5
PEG 8000	0.4
Titanium Dioxide	0.4
Wax	0.2
* Removed during processing	

Example 10

[0099]

Table 10

Pregabalin Combination	
First Active Ingredient mg/tablet	
Tapentadol HCl	200.0
Polyvinyl Alcohol	4.0
Colloidal Silicon Dioxide (Abrosil® 200)	2.0
Sodium Stearyl Fumarate	2.0
Water*	Q.S
Core Weight	2.8
Coat	
Ethylcellulose (Ethocel® PR 100)	12.28
Polyvinylpyrrolidone (Kollidon® 90F)	6.50
Dibutyl Sebacate	3,75
Denatured Alcohol*	Q.S

(continued)

Second Active Ingredient	
Pregabalin	250 mg
Povidone K 30 USP	12 mg
Microcrystalline cellulose	25 mg
Croscarmellose sodium	15 mg
Magnesium Stearate	3 mg
Water*	Q.S.
* Removed during processing	

Example 11

[0100]

Table 11

Gabapentin Combination	
First Active Ingredient mg/tablet	
Tapentadol Hydrochloride	100.0
Lactose	65.0
Ethyl Cellulose	16.0
Cetostearyl Alcohol	43.0
Magnesium Stearate	2.0
Talc	4.0
Hydroxyethyl Cellulose	
Water *	Qs
Coat	
Hydroxypropyl methyl cellulose	0.75
Hydroxy methyl cellulose	3.75
Opaspray	2.60
PEG 400	0.60
Talc	0.30
Water *	qs
Second Active Ingredient	
Gabapentin	250
Povidone K 30 USP	1.0
Lactose	25.0
Sodium starch Glycolate	7.5
Poloxamer 188	3.0
HPMC	1.5
PEG 8000	0.4
Titanium Dioxide	0.4
Wax	0.2
* Removed during processing	

Example 12

[0101]

Table 12

Pregabalin Combination	
First Active Ingredient mg/tablet	
Tapentadol HCl	200.0
Polyvinyl Alcohol	4.0
Colloidal Silicon Dioxide (Abrosil® 200)	2.0
Sodium Stearyl Fumarate	2.0
Water*	Q.S
Core Weight	2.8
Coat	
Ethylcellulose (Ethocel® PR 100)	12.28
Polyvinylpyrrolidone (Kollidon® 90F)	6.50
Dibutyl Sebacate	3,75
Denatured Alcohol*	Q.S
Second Active Ingredient	
Pregabalin	250 mg
Povidone K 30 USP	12 mg
Microcrystalline cellulose	25 mg
Croscarmellose sodium	15 mg
Magnesium Stearate	3 mg
Water*	Q.S.
* Removed during processing	

Example 13

[0102]

Table 13

Tramadol Combination	
First Active Ingredient mg/tablet	
Tapentadol Hydrochloride	100
Microcrystalline Cellulose	10
Colloidal Silicon Dioxide	1.5
Polyvinylpyrrolidone	4.5
Hydrogenated Vegetable Oil	5
Water *	Q.S
Coat	
Ethylcellulose Aqueous Dispersion	50
Opaspray YS -1 7006	5
Water*	Q.S
Second Active Ingredient	
Tramadol	100
Povidone K 30 USP	10
Microcrystalline cellulose	15
Croscarmellose sodium	9

(continued)

Second Active Ingredient	
Magnesium Stearate	2
Water*	Q.S
* Removed during processing	

[0103] The disclosed compositions can also be prepared using methods that are well established in the art. It can be prepared using a general preparation outlined in tables 14and 15

Table 14

Formula		
First Active Ingredient	Range percent	Preferred Range %
Drug	50-98%	75-95%
Binder	0.1-40%	3-15%
Absorption Enhancer	0-20%	2-10%
Lubricant	0-5%	0.5-1%
Coat		
Polymer	50-99%	75-95%
Flux Enhancer	0-40%	2-20%
Plasticizer	0-25%	2-15%
Second Active Ingredient		
Drug	0.1-20%	1-10%
Binder	0.1-20%	1-15%
Surfactant	0-20%	0.1-15%
Pore Former	0-25%	0.1-15%
Polymer (Optional)	0-30%	0.1-20%

Table 15

First Active Ingredient	Percent of Core
Tapentadol HCl	90.54%
Povidone K 301 USP	4.38%
Sodium Tribasic Phosphate	4.58%
Magnesium Stearate	0.50%
Membrane	**Percent of membrane**
Cellulose Acetate (398-10)' 85% Triacetin 5% PEG 400 10%	85.00%
Triacetin	5.00%
PEG 400	10.00%
Second Active Ingredient	**Percent of second layer**
Second Analgesic	43.50%
Tween	2.00%
HPMC	54.50%

METHOD OF ADMINISTRATION

[0104] The present inventions further include a method of treating pain and pain related conditions. This was established using well controlled human clinical trials for each type of combination. A typical study determined the efficacy of combination comprising a slow release tapentadol and Pregabalin, a combination comprising a slow release tapentadol and

naproxen (NSAID) and a combination comprising a slow release tapentadol and tramadol. Each of these combinations was compared against monotherapy with the respective drugs for the treatment of pain and pain related conditions in patients.

Clinical Trials:

[0105] The following general methods were used in all of the studies;
Randomization: Randomization was performed with computer-generated random numbers in blocks of 10. Randomization codes of the monotherapy or combination therapy treatments were placed in sequentially numbered, opaque, sealed envelopes in the biopsy center. When a patient was recruited and consented, the next numbered envelope was opened by the operator, who had no knowledge of the randomization code before the treatment

[0106] Scale and Measurement: Pain was assessed using a visual analogue scale (VAS) graded from 0-100 mm. Pain scoring was performed every visit Week 1, Week 2, Week 3, Week 4, Week 5 and Week 6. The patients were taught with a standard method by the physician how to use a VAS scoring from 0 to 100 mm to grade the intensity of pain experienced during the treatment. On that scale, the left endpoint, 0, was defined as no pain and the right endpoint 100, as the worst pain the patient could imagine. There were no further marks on the line. The intensity of pain was indicated by the distance in millimeters from the left end. Patients were asked to grade the pain intensity after the administration of drugs and the pain was assessed by the patient in their weekly pain score diary. The doctors who prescribed the medicines obtained all the VAS forms for that specific patient.

[0107] Statistical Analysis: Data are expressed as means ± SD. Differences in VAS were analyzed with use of the unpaired Student t test and differences in pain score reductions between the two groups were analyzed with use of the Mann-Whitney U test for the median difference. Similar statistical tests were used for between group comparisons of other outcome variables as appropriate. A two tailed P value of less than. 0.5 was considered to demonstrate statistical significance. SAS software was used for the statistical analysis.

Study I, Tapentadol and NSAID Combination;

[0108] Patients: The efficacy was established using a 6-week, multi-center, double blind, randomized, parallel, three arm comparison of the efficacy of fixed dose slow release Tapentadol ER and Naproxen tablets, Tapentadol and Naproxen tablets in the treatment of knee osteoarthritis. The study enrolled 50 patients, from 18 to 75 years of age with moderate to severe pain associated with Functional Class 1-111 osteoarthritis. The enrollment was designed in such a way that a minimum of 30 patients completed the study in each arm. The patients who met the inclusion and exclusion criteria at screening, after signing the informed consent, entered a 7- day washout period where all analgesics were discontinued. During the visit 2 at the start of the first week of the study, the eligible patients who reported pain intensity 240 mm on a visual analog scale (VAS) in the index knee joint, had their VAS Baseline score measured and were randomly assigned to one of the three arms: The fixed dose tapentadol ER and Naproxen or to Tapentadol arm or to Naproxen arm. Table 16 shows the patient's profile of the study.

Table 16

Patient Characteristics				
Characteristic	**Treatment A**	**Treatment B**	**Treatment C**	**Placebo**
	N= 30	**N=30**	**N=34**	**N=30**
Mean age +/- SD (y)	42.2 ±13	44.4±11	43.2 ± 12	41.2 ± 12
Sex (M/F)	17/13	15/15	18/16	16/14

[0109] Drugs and Treatment Arms: The treatment arms and drug regimen is listed below.

Treatment Arm	**Drugs per day per patient**
1. Treatment A;	Placebo
2. Treatment B;	Tapentadol HCl 100 mg X 2
3. Treatment C:	Naproxen 250 mg X 2
4. Treatment D:	Combination of tapentadol HCl 200 mg + naproxen 500 MG.

[0110] Patients were assigned to either placebo or tapentadol HCl 100 mg or naproxen 250 mg or fixed dose combination of tapentadol 100 mg plus naproxen 250 mg. The patients with pain unresponsive to treatments or with unacceptable side effects were discontinued. The patients returned for efficacy measurement on Week 1, Week 2, Week 3, Week 4, Week 5 and Week 6.

[0111] The primary measure of efficacy was the Arthritis Pain Intensity VAS (visual analog scale) Score from patient visits. The arthritis VAS is the most commonly used, validated tool to assess pain intensity and one recommended by FDA to evaluate the analgesic potential of a drug product.

[0112] We considered that a clinically significant benefit of using fixed dose slow release tapentadol HCl and naproxen would be a reduction in the pain score (VAS) of at least 15% compared to the monotherapy with either tapentadol hydrochloride or naproxen. Alternatively, the dose reduction of at least 15% of either naproxen or tapentadol, when used as a co-administered combination, over the monotherapy is considered as a significant benefit.

[0113] The objectives of the inventions are met by the following results from the clinical trials. A total of 206 patients were randomized and evaluable for safety. Of these, 170 were evaluable for the intent- to-treat (ITT) population. The ITT population included all safety evaluable patients who had primary efficacy information recorded at the baseline visit (Visit 2) and at the Week 1 visit (Visit 3), the first primary efficacy variable collection point on treatment. The ITT population also included all patients who dropped out before the Week 1 visit due to lack of treatment efficacy. The median age of patients who enrolled was 61 years and the median duration of osteoarthritis was 10 years.

[0114] Fixed dose combination comprising a slow release tapentadol and naproxen produced statistically significant and clinically meaningful reductions, compared to the monotherapy using either tapentadol or naproxen, for the primary efficacy variable in pain intensity associated with knee osteoarthritis.

Tapentadol and GABA Analogue Combination, Study II

[0115] The study was of double-blind, randomized, placebo-controlled, and two-period cross-over design. After a 2-week run-in period, 32 diabetic patients (17 mean, 15 women with type 2 diabetes, age [mean ± SE] 61.7 ± 1.6 years, duration of diabetes 8,8 ± 1.5 years, duration of painful neuropathy 2.2 ± 0.4 years) were randomized to receive either placebo or tapentadol hydrochloride 100 mg or pregabalin 250 mg/gabapentin 250 mg or slow release tapentadol HCl 100 mg + pregabalin 250 mg or gabapentin 250 mg FDC for 4 weeks, exchanging their treatment for a further 4 weeks after a 2-week wash-out period. The patients administered the spray to both feet before bedtime. Biweekly pain and other sensory symptoms were assessed using a visual analog scale (VAS). The patient characteristics are shown in Table 17.

Table 17

Patient Characteristics	
Number of patients	22
Age (years)	63.7±1.8 (41-76)*
Sex	13 males, 9 females
BMI (kg/m2)	32.8±1.4
Type of diabetes	2 type 1, 20 type 2
Duration of diabetes (years)	9.1±1.5
Duration of neuropathy (years)	3.0±0.5
Duration of neuropathic pain (years)	2.5±0.4
Treatment order	10 ISDN, 12 placebo
HbA1c (%)†	
At study entry	7.8±0.3
At study completion	8.1±0.4
*Data are n or means, SE. Age Range; †HbA1c Reference Range 4.2-5.9%.	

[0116] Each patient had long history of difficult-to treat painful neuropathy and had tried various drugs such as acetaminophen, duloxitine, amitriptyline or gabapentin and had discontinued because the symptoms were unresponsive or due to unacceptable side effects. Eligible subjects included type 1 and type 2 diabetic patients not on any other medications for their neuropathic pain and with stable diabetic control. Exclusion criteria included erratic glycemic control, peripheral vascular disease (PVD) with absent foot pulses, presence of active foot ulceration, treatment with sublingual glyceryl trinitrate, patients on erectile dysfunction drugs, factors affecting the patient's evaluation of pain, and the presence of

other causes of peripheral neuropathies. No major changes made for diabetes treatment during the duration of the study.

[0117] Patients were assessed neurologically at the beginning of the run in period after which, the patients were randomly allocated to receive the placebo or tapentadol hydrochloride 100 mg or pregabalin 250 mg/gabapentin 250 mg or tapentadol HCl 100 mg + pregabalin 250 mg/gabapentin 250 mg FDC for 4 weeks. A 10-cm visual analog scale (VAS) was recorded biweekly by the patients for pain, where 0 means no pain at all and 10 means the most severe pain ever experienced. The treatment effect was defined to be the difference between the final score and the baseline score on the Lickert scale for each treatment phase.

[0118] The objectives of the inventions are met for the fixed dose combination comprising a slow release tapentadol and pregabalin/gabapentin produced statistically significant and clinically meaningful reductions, compared to the mono-therapy using either tapentadol or pregabalin, for the primary efficacy variable in pain intensity associated with diabetic neuropathy. We considered that a clinically significant benefit of using fixed dose slow release tapentadol HCl and pregabalin/gabapentin would be a reduction in the pain score (VAS) of at least 15 % compared to the other treatments.

Tapentadol and Tramadol Combination, Study III

[0119] The Fixed dose combination of slow release tapentadol hydrochloride and tramadol hydrochloride was tested in a human clinical trial. In this study III, 40 patients were used in a clinical trial were suffering from Chronic non-cancer pain (CNCP), defined as pain for longer than 6 months, including neuropathic pain, osteoarthritis, rheumatoid arthritis, fibromyalgia, and back and musculoskeletal pain were included in the clinical trial. No effort was made to segregate the patients based on the type of pain nor did we evaluate the extent of pain relief on the basis of pain type. Those patients with migraines, dental pain, abdominal pains (from chronic pancreatitis, kidney stones, etc.) and ischemic pain from vascular disease were excluded because they are usually not classified as CNCP. Patients with history of addiction (alcohol or drugs) were excluded from the trials. The patient characteristic and the VAS pain score are listed in Table 18 and Table 19, below;

Table 18

Patient Characteristics				
Characteristic	**Tramadol 100 mg**	**Placebo**	**Tapentadol 100 mg**	**Tapentadol 100 mg + Tramadol 50 mg Fixed Dose combination**
	N=10	**N=10**	**N=10**	**N=10**
Mean age +/- SD (y)	44.2±13	42.1±11	42.5±12	41.2±12
Sex (M/F)	23/10	20/17	22/11	19/18

Table 19

VAS Time hours		**VAS Pain Score**		
	Tramadol 100 mg	**Placebo**	**Tapentadol 100 mg**	**Tapentadol 100 mg + Tramadol 50 mg Fixed Dose combination**
	N= 10	**N=10**	**N=10**	**N=10**
0	32.4±21.2	33.9±19.1	33.7±18.2	31.9±19.1
3	23.1±20.1	32.9±19.1	24.8±19.1	25.1±17.8
6	18.8±20.6	31.9±21.1	21.1±22.9	22.2±18.6
9	14.9±17.2	30.8±20.3	14.2±23.1	15.3±20.6
12	13.1±21.2	31.6±18.6	11.2±20.9	11.3±20.2
15	13.9±19.2	32.1±21.5	11.0±16.2	9.5±18.9
18	15.4±21.3	32.9±19.7	9.5±17.8	9.1±22.1
21	14.5±19.2	31.9±18.3	10.4±16.2	9.3±17.3
24	13.9±19.2	32.8±20.9	11.6±16.2	10.8±17.5

[0120] The objectives of the inventions are met for the fixed dose combination comprising a slow release Tapentadol and Tramadol produced statistically significant and clinically meaningful reductions, compared to the mono-therapy using either tapentadol or tramadol, for the primary efficacy variable in pain intensity associated with diabetic neuropathy. We considered that a clinically significant benefit of using fixed dose slow release tapentadol HCl and tramadol (Example 13) would be a reduction in the pain score (VAS) of at least 15 % compared to the other treatments.

Results

[0121] The objectives of the inventions are met by the following results:

FIGURE 1 illustrates the in vitro dissolution profile of tapentadol HCl in slow release tapentadol HCl 100 mg and naproxen 250 mg tablets formulated according Example 1.

FIGURE 2 compares the LS mean change from baseline in VAS score for the combination drug comprising slow release tapentadol 100 mg and naproxen 250 mg (Example 1) with those of tapentadol and naproxen mono therapies based upon the average of Weeks 1-6. For the primary endpoint (LS Mean change from baseline over 6 weeks), there was a 49.0% change from baseline in the arthritis pain intensity VAS in the slow release tapentadol and naproxen 250 mg fixed dose combination compared 38% for tapentadol, 28% for naproxen and 21% for placebo groups

FIGURE 3 shows the weekly LS mean changes from baseline for the four treatment groups. The differences in drug response emerged when patients were receiving either tapentadol HCl 100 mg X 2 or naproxen 250 mg X 2 or the combination of 100 mg tapentadol & 250 mg naproxen (Example 1) X 2 at the very first week. The response to all drugs increased relative placebo from week 1 till week 6. The response to the combination drug was significantly higher than those due to mono therapy treatment with either naproxen or tapentadol.

FIGURE 4 shows the mean VAS pain score changes for the four treatments; tapentadol 100 MG, pregabalin 250 MG, and slow release tapentadol 100 MG + pregabalin 250 MG fixed dose combination (Example 10).

FIGURE 5 shows the mean VAS pain score changes for the three treatments; Tramadol 50 mg, Tapentadol 100 MG, Placebo and Slow Release Tapentadol 100 MG + Tramadol 50 MG (Example 13) fixed dose combination.

FIGURE 6 shows the mean VAS pain score changes for the four treatments; tapentadol 100 MG, gabapentin 250 MG, and slow release tapentadol 100 plus gabapentin 250 MG fixed dose combination (Example 11).

[0122] The abbreviations used herein have their conventional meaning within the chemical and biological arts.

[0123] Studies I and III are outside of the scope of the present invention.

Claims

1. A pharmaceutical composition comprising a slow release tapentadol and at least one pharmaceutically acceptable carrier, and a second active agent, wherein the second active agent is pregabalin, or pharmaceutically acceptable salts thereof.

2. The pharmaceutical composition of claim 1, wherein the tapentadol is tapentadol hydrochloride.

3. The pharmaceutical composition of claim 1 or 2, wherein the pregabalin is present in an immediate release form, extended (controlled) release form or delayed release form along with a pharmaceutically acceptable carrier.

4. The pharmaceutical composition of any of claims 1-3, comprising an immediate release coating wherein the immediate release coating comprises the pregabalin.

5. The pharmaceutical composition of any of claims 1-3, wherein the composition is a bilayer composition comprising a slow-release tapentadol layer, and a layer comprising the pregabalin.

6. The pharmaceutical composition of any of claims 1-5, wherein the composition exhibits an in vitro dissolution profile such that after 2 hours, from about 0% to about 30% by weight of tapentadol is released, after 4 hours, from about 5% to about 22% by weight of tapentadol is released, after 6 hours, from about 15% to about 30% by weight of tapentadol is released, and after 8 hours more than about 40% by weight of tapentadol is released; when measured using USP Basket Method at 75 rpm in 900 ml 0.1 N HCl at 37 °C.

7. The pharmaceutical composition of any of claims 1-6, wherein the composition is a coated tablet wherein the coating

comprises at least one water-insoluble, water permeable film-forming polymer, at least one plasticizer and at least one water-soluble polymer, and the second active agent.

8. The pharmaceutical composition of any of claims 1-7, wherein the proportion of the water-insoluble, water-permeable film-forming polymer is about 20% to about 90% of the coating dry weight, the proportion of the plasticizer is about 5% to about 30% of the coating dry weight, and the proportion of the water-soluble polymer is about 10% to about 75% of the coat dry weight.

9. The pharmaceutical composition of claim 8, wherein the water-insoluble, water-permeable film- forming polymer is ethylcellulose, the water-soluble polymer is polyvinylpyrrolidone and the plasticizer is dibutyl sebacate and/or wherein the core comprises a lubricant, a binder, a glidant, or any combination thereof and/or wherein the carrier comprises an adjuvant, a preservative, an antioxidant, a thickening agent, a chelating agent, an antifungal agent, an antibacterial agent, an isotonic agent, a flavoring agent, a sweetening agent, an anti-foaming agent, a colorant, a diluent, a moistening agent, a parietal cell activator, or a combination thereof.

10. The pharmaceutical composition of any of claims 1-9, wherein the tapentadol is present in an amount of from about 5 mg to about 400 mg.

11. The pharmaceutical composition of any of claims 1-10, wherein the pregabalin is present in an amount of from about 5 mg to about 500 mg.

12. Pharmaceutical composition of any of claims 1-11 for use in treating pain.

13. Pharmaceutical composition of claim 12 for use in treating neuropathic pain, osteoarthritis, rheumatoid arthritis, fibromyalgia, and back, musculoskeletal pain, ankylosing spondylitis, juvenile rheumatoid arthritis, diabetic neuropathy, migraines, dental pain, abdominal pains, ischemic pain, postoperative pain or because of an anesthetic or surgical contrition.

Patentansprüche

1. Eine pharmazeutische Zusammensetzung umfassend Tapentadol mit langsamer Freisetzung und wenigstens einen pharmazeutisch verträglichen Trägerstoff und einen zweiten Wirkstoff, wobei der zweite Wirkstoff Pregabalin oder eines seiner pharmazeutisch verträglichen Salze ist.

2. Die pharmazeutische Zusammensetzung gemäß Anspruch 1, wobei das Tapentadol Tapentadolhydrochlorid ist.

3. Die pharmazeutische Zusammensetzung gemäß Anspruch 1 oder 2, wobei das Pregabalin zusammen mit einem pharmazeutisch verträglichen Trägerstoff in einer Form mit sofortiger Freisetzung, einer Form mit verlängerter (kontrollierter) Freisetzung oder in einer Form mit verzögerter Freisetzung vorhanden ist.

4. Die pharmazeutische Zusammensetzung gemäß einem der Ansprüche 1-3, umfassend eine Beschichtung mit unmittelbarer Freisetzung, wobei die Beschichtung mit unmittelbarer Freisetzung das Pregabalin umfasst.

5. Die pharmazeutische Zusammensetzung gemäß einem der Ansprüche 1-3, wobei die Zusammensetzung eine Doppelschicht-Zusammensetzung ist, die eine langsam freisetzende Tapentadol-Schicht und eine Schicht umfassend das Pregabalin umfasst.

6. Die pharmazeutische Zusammensetzung gemäß einem der Ansprüche 1-5, wobei die Zusammensetzung ein *in vitro* Auflösungsprofil aufweist, das derart ist, dass nach 2 Stunden, von ungefähr 0% bis ungefähr 30 Gew.-% des Tapentadols freigesetzt werden, nach 4 Stunden von ungefähr 5% bis ungefähr 22 Gew.-% des Tapentadols freigesetzt werden, nach 6 Stunden von ungefähr 15% bis ungefähr 30 Gew.-% des Tapentadols freigesetzt werden und nach 8 Stunden mehr als ungefähr 40 Gew.-% des Tapentadols freigesetzt werden, wenn gemessen unter Verwendung des USP Basket-Verfahrens bei 75 UpM in 900 mL 0,1 N HCl bei 37 °C.

7. Die pharmazeutische Zusammensetzung gemäß einem der Ansprüche 1-6, wobei die Zusammensetzung eine beschichtete Tablette ist, wobei die Beschichtung wenigstens ein wasserunlösliches, wasserdurchlässiges filmbildendes Polymer, wenigstens einen Weichmacher und wenigstens ein wasserlösliches Polymer und den zweiten

Wirkstoff umfasst.

8. Die pharmazeutische Zusammensetzung gemäß einem der Ansprüche 1-7, wobei der Anteil des wasserunlöslichen, wasserdurchlässigen filmbildenden Polymers ungefähr 20% bis ungefähr 90% des Trockengewichts der Beschichtung beträgt, der Anteil des Weichmachers ungefähr 5% bis ungefähr 30% des Trockengewichts der Beschichtung beträgt und der Anteil des wasserlöslichen Polymers ungefähr 10% bis ungefähr 75% des Trockengewichts der Beschichtung beträgt.

9. Die pharmazeutische Zusammensetzung gemäß Anspruch 8, wobei das wasserunlösliche, wasserdurchlässige filmbildende Polymer Ethylcellulose ist, das wasserlösliche Polymer Polyvinylpyrrolidon ist und der Weichmacher Dibutylsebacat ist und/oder wobei der Kern ein Gleitmittel, ein Bindemittel, ein Fließregulierungsmittel oder jede derer Kombinationen umfasst und/oder wobei der Trägerstoff einen Hilfsstoff, ein Konservierungsmittel, ein Antioxidationsmittel, ein Verdickungsmittel, einen Komplexbildner, ein Antimykotikum, einen antibakteriellen Wirkstoff, ein isotonisches Mittel, einen Aromastoff, ein Süßungsmittel, ein Antischaummittel, einen Farbstoff, einen Verdünner, ein Feuchtmittel, einen Parietalzellenaktivator oder eine Kombination davon umfasst.

10. Die pharmazeutische Zusammensetzung gemäß einem der Ansprüche 1-9, wobei das Tapentadol in einer Menge von ab ungefähr 5 mg bis ungefähr 400 mg vorhanden ist.

11. Die pharmazeutische Zusammensetzung gemäß einem der Ansprüche 1-10, wobei das Pregabalin in einer Menge von ab ungefähr 5 mg bis ungefähr 500 mg vorhanden ist.

12. Pharmazeutische Zusammensetzung gemäß einem der Ansprüche 1-11 zur Verwendung in der Behandlung von Schmerz.

13. Pharmazeutische Zusammensetzung gemäß Anspruch 12 zur Verwendung in der Behandlung von neuropathischem Schmerz, Osteoarthritis, rheumatoider Arthritis, Fibromyalgie und Rückenschmerz, Schmerzen des Bewegungsapparates, Spondylarthritis ankylosans, juveniler rheumatoider Arthritis, diabetischer Neuropathie, Migräne, Zahnschmerzen, Abdominalschmerzen, ischämischem Schmerz, postoperativem Schmerz oder Schmerzen aufgrund einer anästhetischen oder chirurgischen Quetschung.

Revendications

1. Composition pharmaceutique comprenant un tapentadol à libération lente et au moins un vecteur pharmaceutiquement acceptable, ainsi qu'un deuxième principe actif, où le deuxième principe actif est la prégabaline, ou les sels pharmaceutiquement acceptables de celle-ci.

2. Composition pharmaceutique selon la revendication 1, où le tapentadol est le tapentadol chlorhydrate.

3. Composition pharmaceutique selon la revendication 1 ou 2, où la prégabaline est présente sous une forme à libération immédiate, une forme à libération prolongée (contrôlée) ou une forme à libération retardée avec un vecteur pharmaceutiquement acceptable.

4. Composition pharmaceutique selon l'une quelconque des revendications 1 à 3, comprenant un enrobage à libération immédiate, où l'enrobage à libération immédiate comprend la prégabaline.

5. Composition pharmaceutique selon l'une quelconque des revendications 1 à 3, où la composition est une composition bicouche, comprenant une couche de tapentadol à libération lente et une couche comprenant la prégabaline.

6. Composition pharmaceutique selon l'une quelconque des revendications 1 à 5, où la composition présente un profil de dissolution *in vitro* tel qu'après 2 heures, entre environ 0 % et environ 30 % en masse de tapentadol sont libérés, après 4 heures, entre environ 5 % et environ 22 % en masse de tapentadol sont libérés, après 6 heures, entre environ 15 % et environ 30 % en masse de tapentadol sont libérés, et après 8 heures plus d'environ 40 % en masse de tapentadol sont libérés ; lors d'une mesure en utilisant la procédure USP Basket Method à 75 rpm dans 900 ml de HCl 0,1N à 37 °C.

7. Composition pharmaceutique selon l'une quelconque des revendications 1 à 6, où la composition est un comprimé

enrobé, où l'enrobage comprend au moins un polymère filmogène insoluble dans l'eau et perméable à l'eau, au moins un plastifiant et au moins un polymère hydrosoluble, et le deuxième principe actif.

8. Composition pharmaceutique selon l'une quelconque des revendications 1 à 7, où la proportion du polymère filmogène insoluble dans l'eau et perméable à l'eau est comprise entre environ 20 % et environ 90 % de la masse sèche de l'enrobage, la proportion du plastifiant est comprise entre environ 5 % et environ 30 % de la masse sèche de l'enrobage, et la proportion du polymère hydrosoluble est comprise entre environ 10 % et environ 75 % de la masse sèche de l'enrobage.

9. Composition pharmaceutique selon la revendication 8, où le polymère filmogène insoluble dans l'eau et perméable à l'eau est l'éthylcellulose, le polymère hydrosoluble est la polyvinylpyrrolidone et le plastifiant est le sébacate de dibutyle et/ou où le noyau comprend un lubrifiant, un agent liant, un agent glissant ou n'importe quelle combinaison de ceux-ci et/ou où le vecteur comprend un adjuvant, un conservateur, un antioxydant, un agent épaississant, un agent chélatant, un agent antifongique, un agent antibactérien, un agent isotonique, un arôme, un agent sucrant, un agent antimousse, un colorant, un diluant, un agent humidifiant, un activateur de cellules pariétales, ou une combinaison de ceux-ci.

10. Composition pharmaceutique selon l'une quelconque des revendications 1 à 9, où le tapentadol est présent à une teneur comprise entre environ 5 mg et environ 400 mg.

11. Composition pharmaceutique selon l'une quelconque des revendications 1 à 10, où la prégabaline est présente à une teneur comprise entre environ 5 mg et environ 500 mg.

12. Composition pharmaceutique selon l'une quelconque des revendications 1 à 11 pour utilisation dans le traitement de la douleur.

13. Composition pharmaceutique selon la revendication 12 pour utilisation dans le traitement de la douleur névropathique, de l'arthrose, de la polyarthrite rhumatoïde, de la fibromyalgie, et de la douleur dorsale musculo-squelettique, de la spondylarthrite ankylosante, de la polyarthrite rhumatoïde juvénile, de la neuropathie diabétique, des céphalées, de la douleur dentaire, des douleurs abdominales, de la douleur ischémique, de la douleur postopératoire ou due à une contrainte anesthésique ou chirurgicale.

FIGURE 1
Tapentadol Dissolution Profile from Example 1

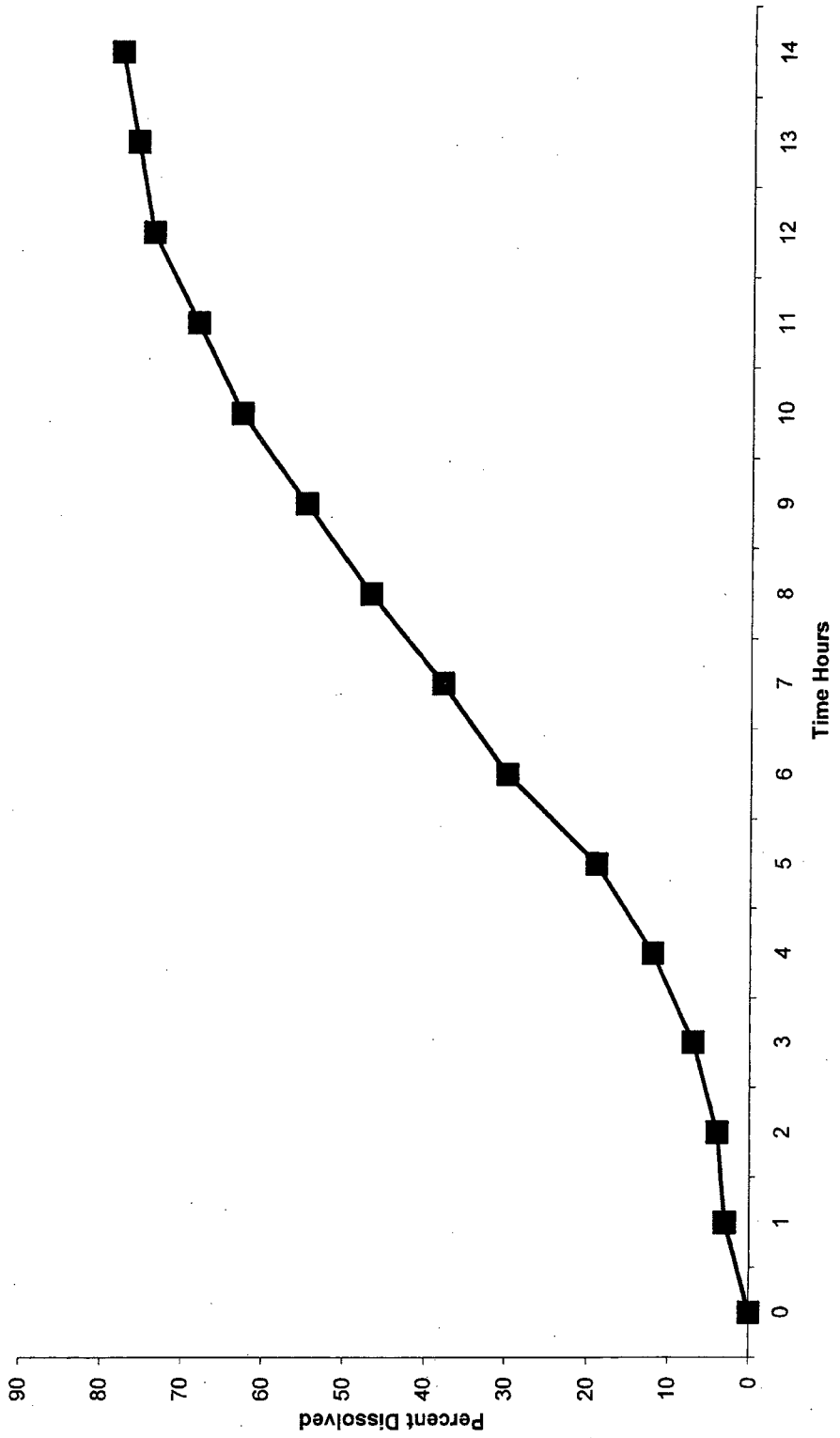


FIGURE 2,
LS Mean Change (%), in Arthritis Pain Intensity VAS score
Slow Rel Tapentadol 100 mg + Naproxen 250 mg

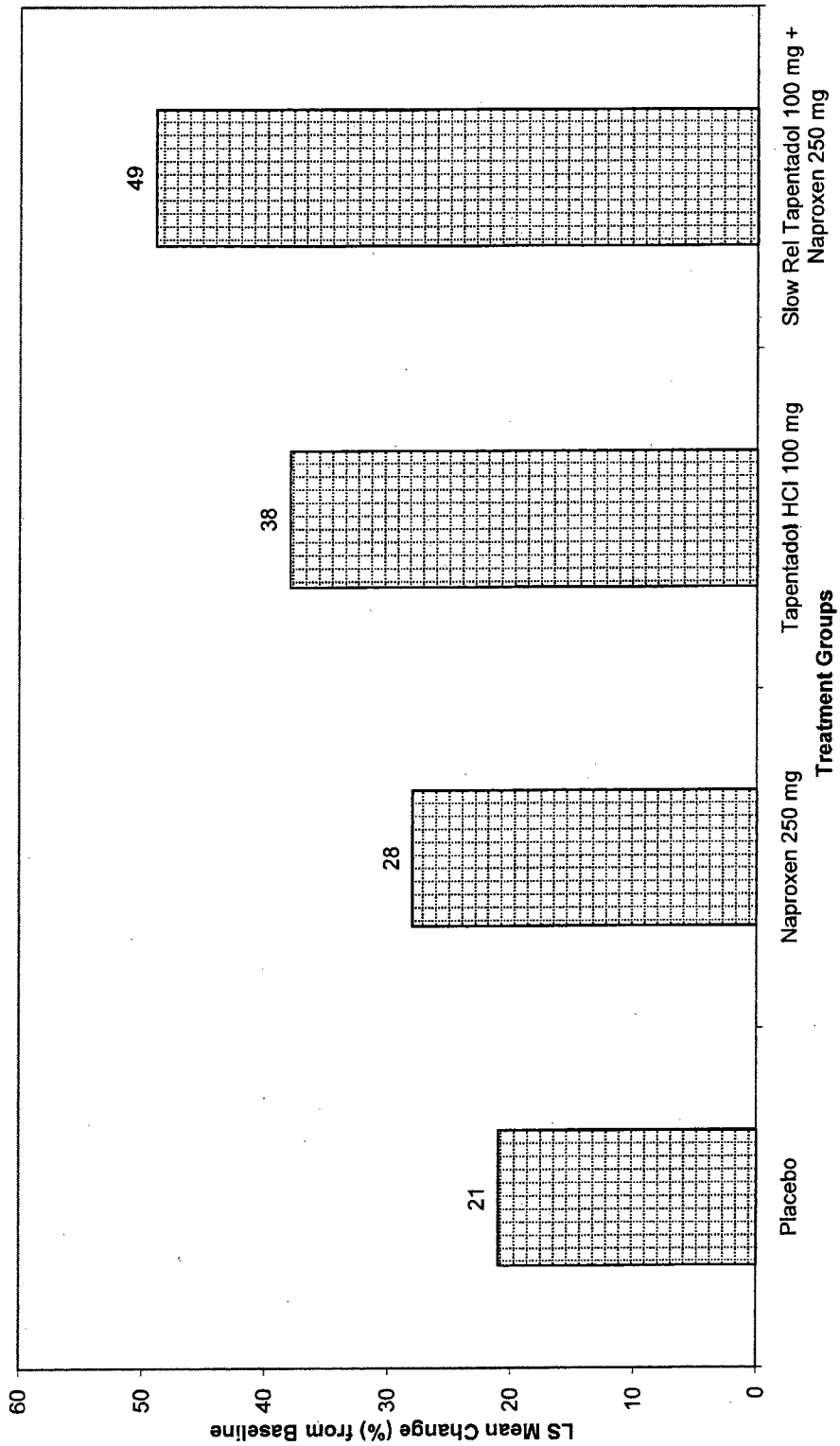
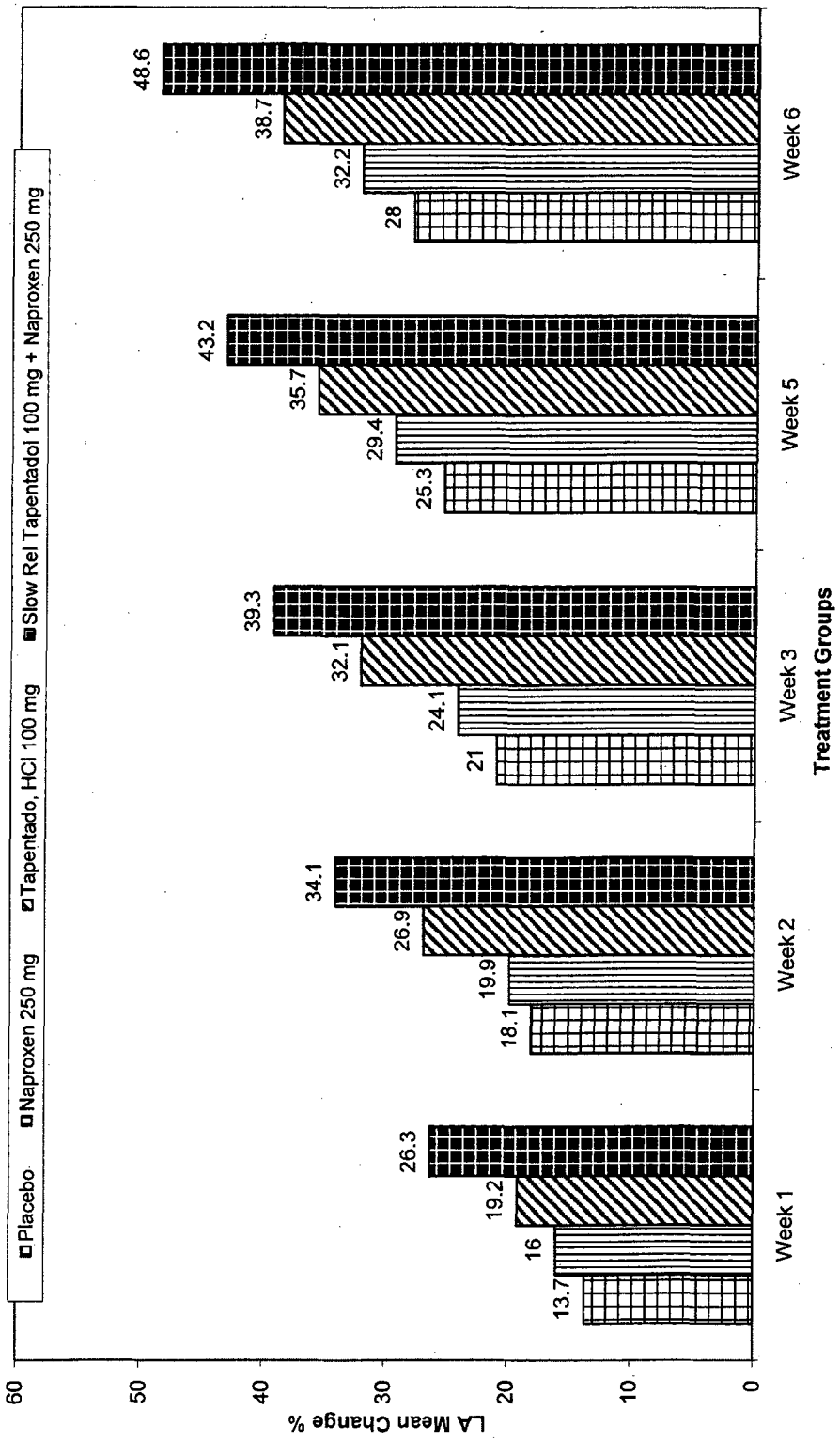
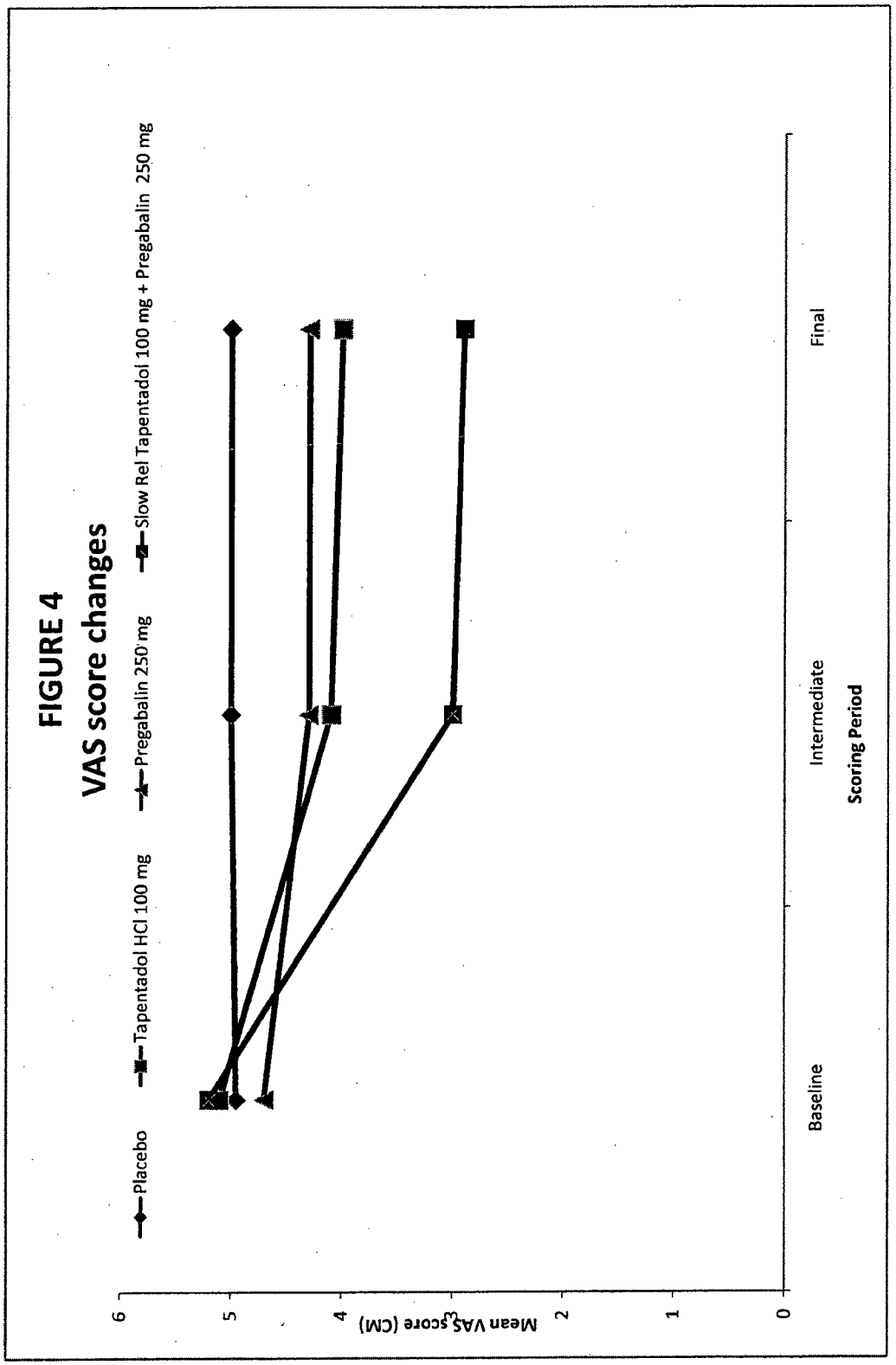


FIGURE 3

LS Mean Change (%), in Arthritis Pain Intensity VAS Scores of 1-12 Weeks




FIGURE 4
VAS score changes
Placebo
Tapentadol HCl 100 mg
Pregabalin 250 mg
Slow Rel Tapentadol 100 mg + Pregabalin 250 mg
Mean VAS score (CM)
0
1
2
3
4
5
6
Baseline
Intermediate
Final
Scoring Period

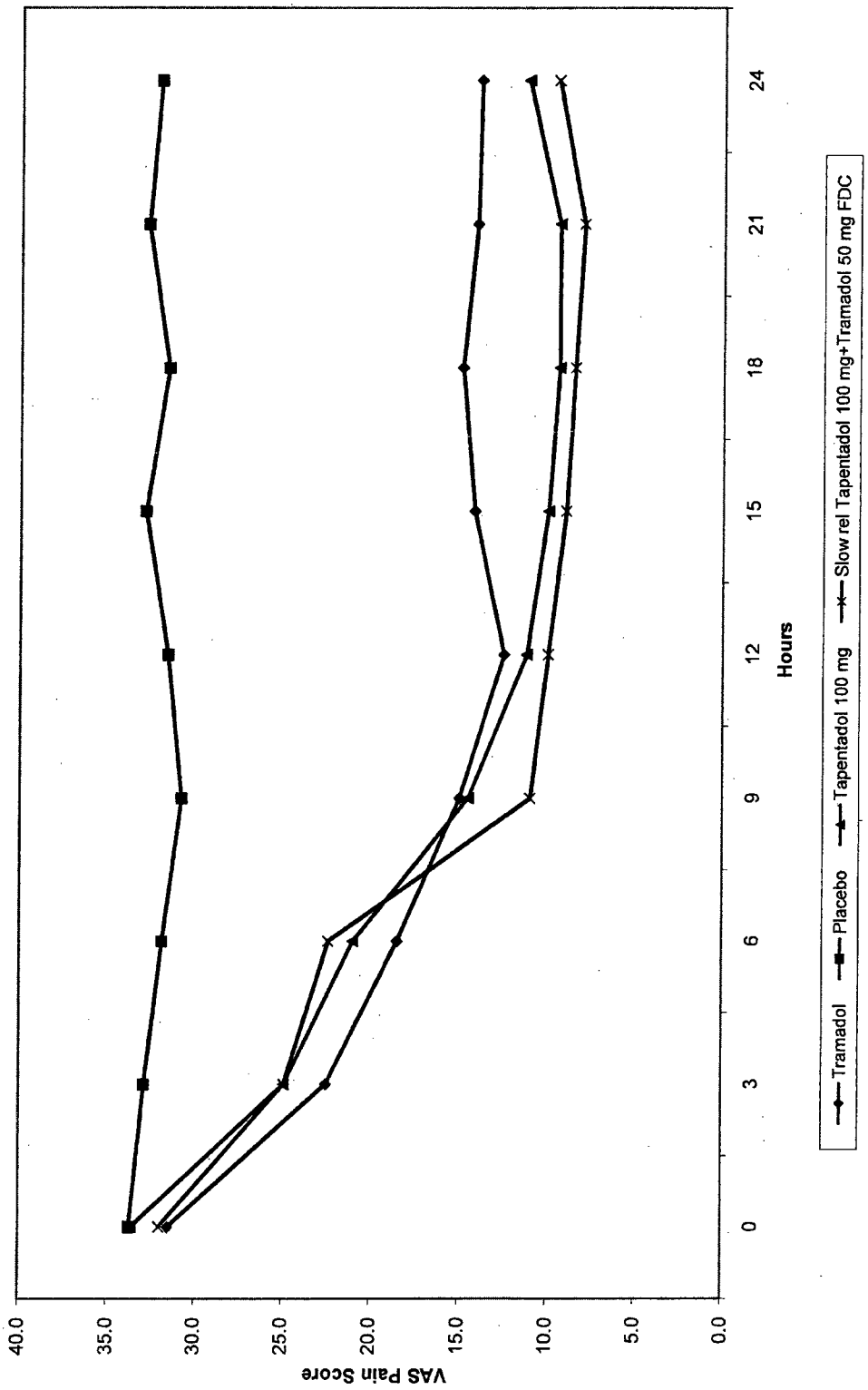
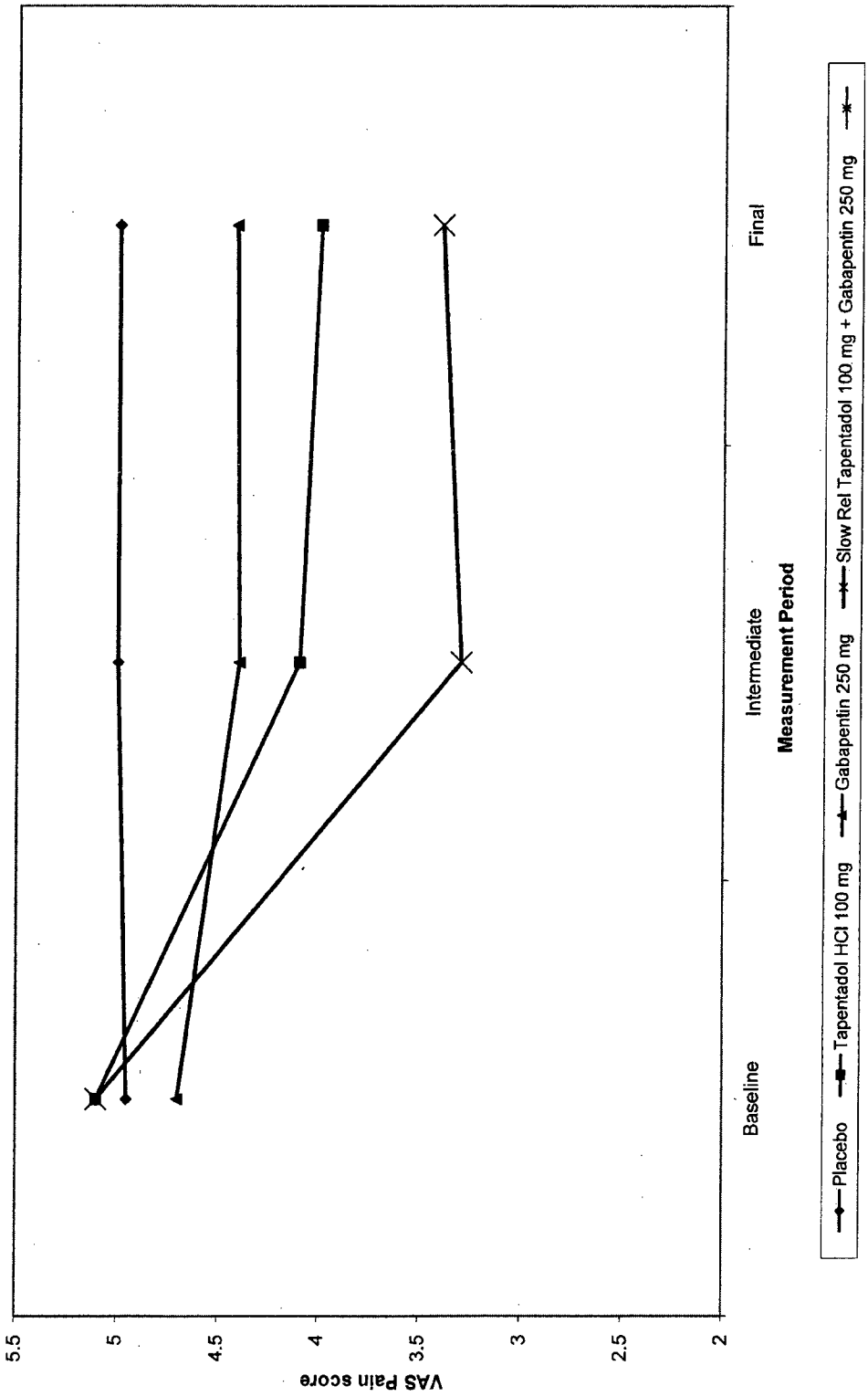


Figure 5, Slow Release Tapentadol + Tramdol Combination, VAS Pain Score

Figure 6, VAS Pain Scrore (Example 11)



REFERENCES CITED IN THE DESCRIPTION

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Az EP 2 596 784 lajstromszámú európai szabadalom igénypontjainak magyar fordítása:

1. Gyógyszerkészítmény, amely tartalmaz lassú hatóanyag-leadású tapentadolt és legalább egy gyógyászatilag elfogadható hordozóanyagot, és egy második hatóanyagot, ahol a második hatóanyag pregabalin vagy gyógyászatilag elfogadható sója.

2. Az 1. igénypont szerinti gyógyszerkészítmény, ahol a tapentadol tapentadol-hidroklorid.

3. Az 1. vagy 2. igénypont szerinti gyógyszerkészítmény, ahol a pregabalin azonnali hatóanyag-leadású formában, nyújtott (szabályozott) hatóanyag-leadású formában vagy késleltetett hatóanyag-leadású formában van jelen gyógyászatilag elfogadható hordozóanyaggal együtt.

4. Az 1-3. igénypontok bármelyike szerinti gyógyszerkészítmény, amely tartalmaz azonnali hatóanyag-leadású bevonatot, ahol az azonnali hatóanyag-leadású bevonat tartalmazza a pregabalint.

5. Az 1-3. igénypontok bármelyike szerinti gyógyszerkészítmény, ahol a készítmény kétrétegű készítmény, amely tartalmaz lassú hatóanyag-leadású tapentadol-réteget és egy réteget, amely tartalmazza a pregabalint.

6. Az 1-5. igénypontok bármelyike szerinti gyógyszerkészítmény, ahol a készítmény olyan in vitro kioldódási profillal rendelkezik, hogy 2 óra eltelte után a tapentadol körülbelül 0 tömeg%-tól körülbelül 30 tömeg%-ig terjedő mennyiségét adja le, 4 óra eltelte után a tapentadol körülbelül 5 tömeg%-tól körülbelül 22 tömeg%-ig terjedő mennyiségét adja le, 6 óra eltelte után a tapentadol körülbelül 15 tömeg%-tól körülbelül 30 tömeg%-ig terjedő mennyiségét adja le, és 8 óra eltelte után a

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tapentadol körülbelül 40 tömeg%-nál nagyobb mennyiségét adja le, amikor a USP szerinti kosaras eljárással mérjük 75 percenkénti fordulatszám (rpm) mellett 900 ml 0,1 N HCl-ban, 37 °C-on.

7. Az 1-6. igénypontok bármelyike szerinti gyógyszerkészítmény, ahol a készítmény bevont tabletta, ahol a bevonat tartalmaz legalább egy vízben oldhatatlan, vizet áteresztő filmképző polimert, legalább egy lágyítószert és legalább egy vízben oldódó polimert, és tartalmazza a második hatóanyagot.

8. Az 1-7. igénypontok bármelyike szerinti gyógyszerkészítmény, ahol a vízben nem oldódó, vizet áteresztő filmképző polimer részaránya a bevonat száraz tömegének körülbelül 20 %-a – körülbelül 90 %-a, a lágyítószer részaránya a bevonat száraz tömegének körülbelül 5 %-a – körülbelül 30 %-a, és a vízben oldódó polimer részaránya a bevonat száraz tömegének körülbelül 10 %-a – körülbelül 75 %-a.

9. A 8. igénypont szerinti gyógyszerkészítmény, ahol a vízben nem oldódó, vizet áteresztő filmképző polimer etilcellulóz, a vízben oldódó polimer polivinilpirrolidon, és a lágyítószer dibutil-szebacát és/vagy ahol a mag tartalmaz kenőanyagot, kötőanyagot, csúsztatóanyagot vagy tartalmazza ezek bármilyen kombinációját és/vagy ahol a hordozóanyag tartalmaz adjuvánst, tartósítószert, antioxidánst, sűrítőanyagot, kelátképző szert, gomba elleni szert, antibakteriális szert, izotóniás szert, ízesítőszert, édesítőszert, habzásgátló szert, színezőanyagot, hígítóanyagot, nedvesítő szert, parietális sejt-aktivátort vagy tartalmazza ezek kombinációját.

10. Az 1-9. igénypontok bármelyike szerinti gyógyszerkészítmény, amelyben a tapentadol körülbelül 5 mg-tól körülbelül 400 mg-ig terjedő mennyiségben van jelen.

11. Az 1-10. igénypontok bármelyike szerinti gyógyszerkészítmény, ahol a pregabalin körülbelül 5 mg-tól körülbelül 500 mg-ig terjedő mennyiségben van jelen.

12. Az 1-11. igénypontok bármelyike szerinti gyógyszerkészítmény fájdalom kezelésére való alkalmazásra.

13. A 12. igénypont szerinti gyógyszerkészítmény neuropátiás fájdalom, csont-ízületi gyulladás, reumatoid ízületi gyulladás, fibromialgia, és hát, csontváz-izomzati fájdalom, ízületi merevséget okozó csigolyagyulladás, fiatalkori reumatoid ízületi gyulladás, diabéteszes neuropátia, migrének, fogfájás, alhasi fájdalmak, iszkémiás fájdalom, műtét utáni fájdalom, vagy érzéstelenítő vagy sebészeti beavatkozás okozta zúzódás (contrition) miatti fájdalom kezelésére való alkalmazásra.

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Meghatalmazott: