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(54) **USE OF MELATONIN FOR INDUCTION OF GENERAL ANESTHESIA**

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(58) **Field of Search** 514/415, 816, 514/817, 818

(56) **References Cited**

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Mallo et al, *Acta Endocrinol, Dec.*, 1988 119(4), 474-80, abstract.*

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(57) **ABSTRACT**

Melatonin (N-acetyl-5-methoxytryptamine), or its biologically active analogues, are used to induce anesthesia.

8 Claims, No Drawings

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USE OF MELATONIN FOR INDUCTION OF GENERAL ANESTHESIA

CROSS REFERENCE TO A RELATED APPLICATION

This application is a provisional of Serial No. 60/233,785 filed Sep. 19, 2000 and claims the benefit of that filing date.

BACKGROUND OF THE INVENTION

In the medical field there is a continuing need for new compounds having demonstrated use for inducing anesthesia. It is not only important to induce beneficial anesthesia, but it must be done in a manner that limits toxicity to patients, and as well, minimizes what is known as "anesthesia hangover".

The pineal hormone melatonin (N-acetyl-5-methoxytryptamine) has several putative functions, including regulation of circadian rhythms, regulation of the reproductive axis and antioxidant activity. Autoradiographic studies and receptor assays have demonstrated the presence of melatonin receptors in various regions of the central nervous system and in other tissues in humans.

Exogenous administration of melatonin has been found by several investigators to facilitate sleep onset and improve quality of sleep. Available data suggest that the sleep-inducing properties of melatonin may differ from those of benzodiazepines. Benzodiazepines decrease duration of REM sleep after single administration of a high dose or long-term administration of low dose. Benzodiazepines also reduce slow-wave sleep, thus negatively influencing sleep quality. In contrast, a single low dose of melatonin produced no suppression of REM sleep. Furthermore, unlike benzodiazepines, melatonin does not induce "hangover" effects.

In a previous publication of the author, *British Journal of Anesthesia* 82 (6):875-80 (1999), low-level dosing of oral melatonin in a sublingual fashion was demonstrated as an effective pre-medication, prior to administering a general anesthetic. Patients who were administered such low-level doses sublingually had a significant decrease in anxiety levels and an increase in levels of sedation before operation. However, as pointed out in that article, the use of melatonin in anesthesia had as of then never been evaluated properly, and to the inventor's present knowledge it has never been used as a general anesthetic.

This invention has as its primary objective the development of pineal hormone melatonin (N-acetyl-5-methoxytryptamine) or its biologically active analogues as a general anesthetic which can be used without any significant anesthetic hangover. The continuing need in the art for meeting such an objective is readily apparent.

SUMMARY OF THE INVENTION

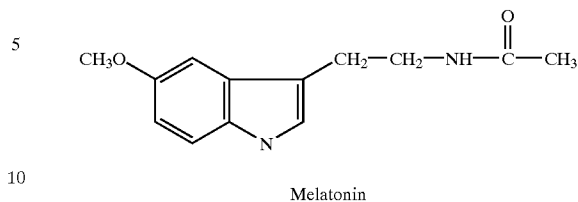
Anesthetic compositions are prepared using a pharmaceutically-acceptable carrier and an anesthetic-inducing effective amount of melatonin or biologically active analogues of melatonin. The invention also relates to the method of administration using the described compositions.

DETAILED DESCRIPTION OF THE PREFERRED EMBODIMENT

N-acetyl-5-methoxytryptamine (melatonin) is synthesized mainly by the pineal gland, and to a lesser extent by

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extra pineal tissues such as the retina, hardierian gland, and gastrointestinal tract. Melatonin has the following structure:



As seen, the chemical formula for melatonin is N-acetyl-5-methoxytryptamine. From time to time in the specification applicant uses the term "N-acetyl-5-methoxytryptamine (melatonin), or its biologically active analogues". As used herein, this phrase refers to the precise compound itself and other compounds having the same general structure, but only differing in minor moieties, and therefore still having the same biological activity of anesthetic-inducing effectiveness. The biologically active compound of the present invention, such as melatonin, may be derived or extracted from the pineal gland, or it can be synthesized from 5-Methoxyindol as a starting material by known routes, Szmuszkovicz et al., *J. Org. Chem.* 25, 857 (1960). Biochemical role of melatonin: *Chem. & Eng. News* 45, 40 (May 1, 1967).

The anesthetic active, i.e., the N-acetyl-5-methoxytryptamine (melatonin), or its biologically active analogues, can be administered with traditionally acceptable pharmaceutical carriers. Examples include Intralipid®, Cyclodextrin, and others, some of which are briefly hereinafter described. However, there is no need for detailed description of suitable anesthetic carriers because they are so well known in the industry.

The composition may be administered by conventional administration methods for anesthetics, i.e., oral administration, nasal respiratory administration, bolus injection, intravenous administration by repeated doses or by continuous infusion, rectal, vaginal, sublingual, cutaneous and slow release routes. It may be, and often is preferred, that it be administered in two or more ways, such as by bolus injection followed by continuous intravenous administration.

The liquid forms in which the novel compositions of the present invention may be incorporated for administration orally or by injection include aqueous solutions, suitably flavored syrups, aqueous or oil suspensions, and flavored emulsions with edible oils such as cottonseed oil, sesame oil, coconut oil or peanut oil, as well as elixirs and similar pharmaceutical vehicles. Suitable dispersing or suspending agents for aqueous suspensions include synthetic and natural gums such as tragacanth, acacia, alginate, dextran, sodium carboxymethylcellulose, methylcellulose, polyvinylpyrrolidone or gelatin.

Preferred compositions for administration by injection include those comprising a melatonin biologically active analogue as the active ingredient, in association with a surface-active agent (or wetting agent or surfactant) or in the form of an emulsion (as a water-in-oil or oil-in-water emulsion).

Suitable surface-active agents include, in particular, non-ionic agents, such as polyoxyethylenesorbitans (e.g. Tween™ 20, 40, 60, 80 or 85) and other sorbitans (e.g. Span™ 20, 40, 60, 80 or 85). Compositions with a surface-active agent will conveniently comprise between 0.05 and

5% surface-active agent, and preferably between 0.1 and 2.5%. It will be appreciated that other ingredients may be added, for example mannitol or other pharmaceutically acceptable vehicles, if necessary.

Suitable emulsions may be prepared using commercially available fat emulsions, such as Intralipid™, Liposyn™, Infonutrol™, Lipofundin™ and Lipiphysan™. The active ingredient may be either dissolved in a pre-mixed emulsion composition, or alternatively it may be dissolved in an oil (e.g. soybean oil, safflower oil, cottonseed oil, sesame oil, corn oil or almond oil) and an emulsion formed upon mixing with a phospholipid (e.g. egg phospholipids, soybean phospholipids or soybean lecithin) and water. It will be appreciated that other ingredients may be added, for example glycerol or glucose, to adjust the tonicity of the emulsion. Suitable emulsions will typically contain up to 20% oil, for example, between 5 and 20%. The fat emulsion will preferably comprise fat droplets between 0.1 and 1.0 μm , particularly 0.1 and 0.5 μm , and have a pH in the range of 5.5 to 8.0.

Particularly preferred emulsion compositions are those prepared by mixing an active compound with Intralipid™ or the components thereof (soybean oil, egg phospholipids, glycerol and water).

Compositions for inhalation or insufflation include solutions and suspensions in pharmaceutically acceptable, aqueous or organic solvents, or mixtures thereof, and powders. The liquid compositions may contain suitable pharmaceutically acceptable excipients as set out above. Preferably the compositions are administered by the oral or nasal respiratory route for local or systemic effect. Compositions in preferably sterile pharmaceutically acceptable solvents may be nebulised by use of inert gases. Nebulised solutions may be breathed directly from the nebulising device, or the nebulising device may be attached to a face mask, tent or intermittent positive pressure breathing machine. Solution, suspension or powder compositions may be administered, preferably orally or nasally, from devices which deliver the formulation in an appropriate manner.

The anesthetic may be used alone or often in combination with other anesthetics simultaneously administered. Put another way, it will be appreciated that when using any combination described herein, both the compound of melatonin or its analogue and the other active agent(s) can be administered to a patient, within a reasonable period of time. It may indeed act synergistically with other anesthetic drugs. The compounds may be in the same pharmaceutically acceptable carrier and therefore administered simultaneously. They may be in separate pharmaceutical carriers such as conventional oral dosage forms which are taken simultaneously. The term "combination" also refers to the case where the compounds are provided in separate dosage forms and are administered sequentially. Therefore, by way of example, one active compound may be administered as a tablet and then, within a reasonable period of time, the second active component may be administered either as an oral dosage form such as a tablet or a fast-dissolving oral dosage form. By a "fast dissolving oral formulation" is meant, an oral delivery form which, when placed on the tongue of a patient, dissolves within about 10 seconds.

The dosage will vary depending upon the deepness of the anesthesia desired, but based upon limited studies to date, it is believed that the dosage most effective will be within the range of 0.001 mg/kg of body weight to about 500 mg/kg of body weight, more predictably preferred is the range of 5 mg/kg of body weight to about 350 mg/kg of body weight.

When patients are administered N-acetyl-5-methoxytryptamine (melatonin) or its biologically active analogues, there is a noticeable decrease in anesthetic hang-over. It is believed that this occurs because melatonin itself is a naturally-occurring hormone synthesized in the body by the pineal gland.

The following examples are offered to further illustrate, but not limit the invention disclosed herein.

All experiments were carried out in male Sprague-Dawley rats (300–350 g). Rats were maintained on a 12 hour light/12 hour dark cycle with free access to food and water. All surgical procedures were performed under sterile conditions (skin preparation, sterile field drape, gloves, mask, etc.). All instruments and materials were ethylene oxide sterilized. Non-fasting adult male Sprague Dawley rats (~300 gm) were anesthetized with halothane in oxygen and weighed. The hair over the ventral neck and over the back (between the scapulae) was removed with an electric razor. In the supine position the ventral neck was washed with povidone-iodine, followed by a 3-cm skin incision, just left of mid-line. All bleeding points were cauterized. Both the left jugular vein and left common carotid artery were isolated via blunt dissection. The left jugular vein was cannulated with a heparinized (20 U/ml) saline-filled silastic catheter (0.012-in ID, 0.025-in OD) advanced ~3-cm into the right atrium. The jugular catheter was secured to the vein with 4-O silk at the point of insertion, as well as at the rostral jugular ligature.

After implantation of intravascular catheters, rats were housed in individual stainless steel cages. Studies with melatonin were carried out 5–7 days after surgery. Crystalline Melatonin powder was obtained from Sigma (Sigma Chemical Co. St. Louis, Mo.).

The melatonin was prepared for anesthetic use in the following manner:

100 mg melatonin added to 1 ml of intralipid and 1 ml of Ringer's Lactate (final concentration=50 mg melatonin/ml).

Results

Rat 1: 250 mg/kg followed approximately 1 minute later by 65 mg/kg: the animal was very drugged but did not lose righting reflex.

Rat 2: 250 mg/kg resulted in loss of righting and eyelash reflexes and inability to pull his hind paw in response to pressure applied to it.

Rat 3: 320 mg/kg resulted in loss of righting and eyelash reflexes and inability to pull his hind paw in response to pressure applied to it.

Rat 4: 370 mg/kg resulted in loss of righting and eyelash reflexes and inability to pull his hind paw in response to pressure applied to it.

Additional preparation of melatonin occurred with cyclodextrin as follows:

100 mg melatonin added to 1 ml of cyclodextrin 40% and 1 ml of intralipid (final concentration=50 mg melatonin/ml)

Results

Rat 1: 315 mg/kg: the animal moves slowly but no loss of righting reflex.

Rat 2: 460 mg/kg resulted in loss of righting reflex.

Another group of rats received the solvent alone and did not result in any effect. This demonstrates the anesthetic property of melatonin, and that the invention accomplishes its stated objectives.

Melatonin in 40% Cyclodextrin

Rat 1: 315 mg/kg resulted in ptosis, loss of eye blink response and loss of paw pinch response to a pressure of 60 mmHg using a very low profile load cell (Omega part number LCKD-1KG, measurement range of 0–1 kg) from OMEGA Engineering, INC. One Omega Drive, Stamford, Conn. 06907-0047. The righting reflex was lost for 27 min.

Rat 2: 374 mg/kg resulted in ptosis and loss of eye blink response. The righting reflex was lost for 15 min. The animal responded to a paw pinch response to a pressure of 60 mmHg by pulling his paw without vocalization.

Rat 3: Administration of the solvent (40% cyclodextrin) did not affect the animal behavior and did not result in sedation or hypnosis.

In a recent publication of the inventor concerning this invention, additional data relating to the invention is disclosed, *Anesthesia and Analgesia*, 91:473–479(2000), the disclosure of which is incorporated herein by reference.

The following examples are comparative examples demonstrating the effectiveness of melatonin for induction of general anesthesia in rats and how it compares to other known anesthetics in inducing general anesthesia.

COMPARATIVE EXAMPLES

The goal of these examples was to determine the doses of melatonin, thiopental and propofol needed to induce anesthesia in 50% and 95% of rats and to evaluate the time course of different indices of anesthesia. Rats were randomly assigned to receive three cumulative doses of 6.67 mg/kg i.v. thiopental, 3.3 mg/kg i.v. propofol or 70 mg/kg i.v. melatonin or three cumulative injections of the vehicle in which these drugs were dissolved at intervals of approximately 1 min. After the final cumulative dose, measurements of anesthesia end-points were made at fixed intervals for an additional 20 minutes. Separate groups of rats received a single bolus injection of 20 mg/kg i.v. thiopental, 10 mg/kg i.v. propofol or 275 mg/kg i.v. of melatonin or the vehicle in which these drugs were dissolved. Measurements of anesthesia end-points were made in these rats at fixed intervals for 20 minutes. Righting reflex was scored on a four-point scale (1=immediate/brisk, both feet under the rat; 2=complete, but slower than normal; 3=slow, feet not placed under body; and 4=absent). The threshold pressure (mm Hg) at which the rat withdrew or vocalized after pinch of one hindpaw was measured. For paw pinch, a subminiature, very low profile load cell Omega part number LCKD-1KG, measurement range of 0–1 kg) (OMEGA Engineering, INC., Stamford, Conn.) was used to measure the amount of pressure applied to the rat's paw. The action of compressing the load cell between the faces of the sponge clamp assembly results in a change in impedance within the load cell, and this in turn, was then converted to mm Hg pressure by the monitor. The presence or absence of eyelash reflex was noted on a three-point scale (1=normal, 2=weak; and 3=absent). The strength of grip by the forepaws was determined on a four-point scale (0=absent, 1=weak, 2=moderate and 3=strong).

Thiopental was purchased from Abbott Laboratories (Northern Chicago, Ill.). Propofol was purchased from Zeneca Pharmaceuticals (Wilmington, Del.). Melatonin was purchased from Sigma Chemical Co. (St. Louis, Mo.). Thiopental and propofol were dissolved in saline and Intralipid™ respectively. Melatonin was dissolved in a mixture comprising 25% v/v propylene glycol and 25% v/v 1-methyl-2-pyrrolidinone in sterile water. For the cumulative injections, the individual doses were administered in a volume of 0.2 ml and the maximum volume of drug injected did not exceed 0.6 ml. For the bolus injections, the volume of drug injected ranged from 0.6 to 0.75 ml.

Intravenous injection of saline, Intralipid™ or the vehicle for melatonin did not affect righting reflex, grip strength, or

eyelash reflex. Neither saline nor Intralipid™ altered paw withdrawal threshold. However, the vehicle for melatonin produced a significant short-lived increase in paw withdrawal threshold that subsequently decreased to near baseline levels.

Cumulative i.v. injection of divided doses of thiopental caused a progressive loss of righting reflex, grip strength and eyelash reflex with an estimated ED₀₅ (and 95% CI) for the loss of righting reflex of 23.8 (15.4–36.7) mg/kg i.v. Bolus injection of 20 mg/kg thiopental resulted in an immediate loss of righting reflex and grip strength that was maximal at 1 min and resolved within 15 min. These effects were not accompanied by a change in paw withdrawal threshold.

Cumulative i.v. injection of divided doses of propofol caused a progressive loss of righting reflex, grip strength and eyelash reflex with an estimated ED₀₅ (and 95% CI) for loss of righting reflex of 14.9 (6.4–34.9) mg/kg. Bolus injection of 10 mg/kg i.v. propofol caused an immediate loss of righting reflex and grip strength that was maximal for 5 min and resolved within 10 min. These effects were accompanied by a significant increase in paw withdrawal threshold of similar duration.

Cumulative i.v. injection of divided doses of melatonin caused a progressive loss of righting reflex and grip strength, but did not appreciably blunt the eyelash reflex. It also dose-dependently increased paw withdrawal threshold. The estimated ED₀₅ (and 95% CI) of melatonin for loss of righting reflex was 312 (205–476) mg/kg i.v. Bolus injection of 275 mg/kg i.v. melatonin resulted in an immediate loss of righting reflex and grip strength that was maximal for 5 min and resolved to near baseline values by 15 min. This dose of melatonin also increased paw withdrawal threshold as compared to vehicle. The increase in paw withdrawal threshold persisted for at least 20 min, and was apparent at doses of 140 mg/kg or greater.

This data demonstrates that intravenous administration of melatonin can induce general anesthesia. This anesthesia is accompanied by an analgesia that persists after the return of consciousness.

What is claimed is:

1. A method of inducing anesthesia comprising:

administering to a patient an anesthesia inducing effective amount of N-acetyl-5-methoxytryptamine or a biologically active analogue thereof.

2. The method of claim 1 wherein the anesthetic is N-acetyl-5-methoxytryptamine.

3. The method of claim 1 wherein the amount of N-acetyl-5-methoxytryptamine or biologically active analogue thereof administered is a dose of from about 0.001 mg/kg of body weight to about 500 mg/kg of body weight.

4. The method of claim 1 wherein the amount of N-acetyl-5-methoxytryptamine or biologically active analogue thereof administered is from about 5 mg/kg of body weight to about 350 mg/kg of body weight.

5. The method of claim 1 wherein the administering is by a method selected from the group consisting of oral administration, nasal respiratory administration, bolus injection, intravenous administration, by repeated doses or by continuing infusion, rectal, vaginal, sublingual and cutaneous administration.

6. The method of claim 5 wherein the administration is by an initial bolus injection, followed by intravenous administration.

7. The method of claim 1 wherein the administration is in combination with simultaneous administration of another anesthetic.

8. The method of claim 1 wherein N-acetyl-5-methoxytryptamine or biologically active analogue thereof is derived from the pineal gland.