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(54) **METHODS OF TREATING
DERMATOLOGICAL DISORDERS AND
INDUCING INTERFERON BIOSYNTHESIS
WITH SHORTER DURATIONS OF
IMIQUIMOD THERAPY**

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(76) Inventor: **Jefferson J. Gregory**, Bristol, TN
(US)

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Correspondence Address:
EDWARDS ANGELL PALMER & DODGE LLP
P.O. BOX 55874
BOSTON, MA 02205 (US)

(57) **ABSTRACT**

Pharmaceutical formulations and methods for the topical and/or transdermal delivery of imiquimod, including creams, ointments and pressure-sensitive adhesive compositions to treat dermatological disorders, namely, viral infections, such as Type I or Type II Herpes simplex infections and genital warts, actinic keratosis and superficial basal cell carcinoma, and to induce interferon biosynthesis, with shorter durations of therapy, than currently approved for imiquimod by the Food & Drug Administration (“FDA”).

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(60) Provisional application No. 60/900,550, filed on Feb. 8, 2007.

**METHODS OF TREATING
DERMATOLOGICAL DISORDERS AND
INDUCING INTERFERON BIOSYNTHESIS
WITH SHORTER DURATIONS OF
IMIQUIMOD THERAPY**

**CROSS-REFERENCE TO RELATED
APPLICATIONS**

[0001] This present application claims the benefit of priority under 35 U.S.C. 119(e) to U.S. Provisional Patent Application No. 60/900,550 filed on Feb. 8, 2007, the disclosure of which is incorporated herein by reference in its entirety.

FIELD OF THE INVENTION

[0002] This invention pertains to methods and pharmaceutical formulations for the topical or transdermal delivery of 1-isobutyl-1H-imidazo[4,5-c]-quinolin-4-amine, i.e., imiquimod. More particularly, it pertains to creams, ointments, foams, gels, lotions, pressure sensitive adhesive coatings and adhesive-coated sheet materials, which contain imiquimod, that enhance skin penetration of drugs to treat dermatological disorders, namely, viral infections, such as Type I or Type II Herpes simplex infections and genital warts, actinic keratosis and superficial basal cell carcinoma, and to induce interferon biosynthesis, with shorter durations of therapy, than currently approved for imiquimod by the Food & Drug Administration ("FDA").

BACKGROUND

[0003] The compound 1-isobutyl-1H-imidazo[4,5-c]-quinolin-4-amine, known as imiquimod and commercially marketed in the U.S. under the brand name Aldara®, is disclosed in U.S. Pat. No. 4,689,338 and described therein as an antiviral agent and as an interferon inducer, which is incorporated herein by reference in its entirety. A variety of formulations for topical administration of imiquimod are also described therein. This U.S. Pat. No. 4,689,338 is incorporated herein by reference in its entirety.

[0004] U.S. Pat. No. 4,751,087 discloses the use of a combination of ethyl oleate and glyceryl monolaurate as a skin penetration enhancer for nitroglycerine, with all three components being contained in the adhesive layer of a transdermal patch, wherein this U.S. patent is incorporated herein by reference in its entirety.

[0005] U.S. Pat. No. 4,411,893 discloses the use of N,N-dimethyldodecylamine-Noxide as a skin penetration enhancer in aqueous systems, wherein this U.S. patent is incorporated herein by reference in its entirety.

[0006] U.S. Pat. No. 4,722,941 discloses readily absorbable pharmaceutical compositions that comprise a pharmacologically active agent distributed in a vehicle comprising an absorption-enhancing amount of at least one fatty acid containing 6 to 12 carbon atoms and optionally a fatty acid monoglyceride. Such compositions are said to be particularly useful for increasing the absorption of pharmacologically active bases, wherein this U.S. patent is incorporated herein by reference in its entirety.

[0007] U.S. Pat. No. 4,746,515 discloses a method of using glyceryl monolaurate to enhance the transdermal flux of a transdermally deliverable drug through intact skin, wherein this U.S. patent is incorporated herein by reference in its entirety.

[0008] U.S. Pat. No. 5,238,944 discloses topical formulations and transdermal delivery systems containing 1-isobutyl-1H-imidazo[4,5-c]-quinolin-4-amine, wherein this U.S. patent is incorporated herein by reference in its entirety.

SUMMARY OF THE INVENTION

[0009] The present invention provides a substantially non-irritating pharmaceutical formulation for topical and/or transdermal administration of the imiquimod, which formulation comprises:

[0010] a) 1-isobutyl-1H-imidazo[4,5-c]-quinolin-4-amine, i.e., imiquimod, in an amount of about 2 percent to about 4 percent by weight based on the total weight of the formulation; and

[0011] b) a pharmaceutically acceptable vehicle for imiquimod, which vehicle comprises a fatty acid, such as isostearic acid, linoleic acid, oleic acid, super purified oleic acid (an oleic acid having low polar impurities such as peroxides) and a combination thereof, in a total amount of about 3 percent to about 45 percent by weight based on the total weight of the formulation. The formulation is further characterized in that when tested in the hairless mouse skin model described in U.S. Pat. No. 5,238,944, the formulation provides a penetration of the agent of at least about 10% (and preferably at least about 15%) of the total amount of the agent contained in the formulation in 24 hours.

[0012] The salient elements of a pharmaceutical formulation according to the invention are (a) imiquimod and (b) a fatty acid, e.g., isostearic, linoleic, super purified oleic acid and mixtures thereof. A pharmaceutical formulation of the invention can be in any form known to the art, such as a cream, an ointment, a foam, a gel, a lotion or a pressure-sensitive adhesive composition, each form containing the necessary elements in particular amounts and further containing various additional elements.

[0013] A cream of the invention preferably contains about 2 percent to about 4 percent by weight of imiquimod based on the total weight of the cream; about 5 percent to about 25 percent by weight of fatty acid, based on the total weight of the cream; and optional ingredients such as emollients, emulsifiers, thickeners, and/or preservatives.

[0014] An ointment of the invention contains an ointment base in addition to imiquimod and fatty acid. An ointment of the invention preferably contains about 2 percent to about 4 percent by weight imiquimod; about 3 percent to about 45 percent, more preferably about 3 percent to about 25 percent by weight fatty acid; and about 40 percent to about 95 percent by weight ointment base, all weights being based on the total weight of the ointment. Optionally, an ointment of the invention can also contain emulsifiers, emollients and thickeners.

[0015] A pressure-sensitive adhesive composition of the invention contains imiquimod, fatty acid, and an adhesive. The adhesives utilized in a pressure sensitive adhesive composition of the invention are preferably substantially chemically inert to imiquimod.

[0016] A pressure sensitive adhesive composition of the invention preferably contains about 2 percent to about 4 percent by weight imiquimod; about 10 percent to about 40 percent by weight, more preferably of about 15 percent to about 30 percent by weight, and most preferably about 20 percent to about 30 percent by weight of fatty acid; all weights being based on the total weight of the pressure sensitive adhesive composition.

[0017] Optionally, pressure sensitive adhesive compositions of the invention can also contain one or more skin penetration enhancers. The total amount of skin penetration enhancer(s) present in a pressure sensitive adhesive composition of the invention is preferably about 3 percent to about 25 percent by weight, and more preferably about 3 percent to about 10 percent by weight based on the total weight of the pressure sensitive adhesive composition.

[0018] A pressure sensitive adhesive coated sheet material of the invention can be made from a pressure-sensitive adhesive composition of the invention in the form of an article such as a tape a patch, a sheet, or a dressing.

[0019] A formulation of the present invention may be used to topically and/or transdermally administer imiquimod for effectively treating viral infections, for example, Type I or Type II Herpes simplex infections, actinic keratosis and superficial basal cell carcinoma for a shorter duration of time and with the same or increased number of applications per week, as compared to current imiquimod topical therapy.

[0020] For example, a formulation of the present invention containing between greater than about 1% and about 5% imiquimod may be applied from three to seven times per week (once per day) for 8 to 12 weeks to treat viral infections, for example, Type I or Type II Herpes simplex infections, actinic keratosis and superficial basal cell carcinoma. It should be understood that while formulations of the present invention containing between greater than about 1% and about 5% imiquimod are preferred, formulations containing about 2.5%, 2.75%, 3.0%, 3.25%, 3.5%, 3.75%, 4.0%, 4.25% and 4.5% are more preferred and that formulations containing about 2.75%, 3.0%, 3.25%, 3.5%, 3.75% and 4.0% are most preferred.

[0021] As to duration, the present invention contemplates applying an effective amount of imiquimod for a shorter period of time than currently approved by the FDA. More specifically, the present invention contemplates applying an effective amount of imiquimod from three to seven times or more per week to an area in need of imiquimod treatment for about 8 to about 12 weeks, and more preferably between about 4, about 5, about 6 and about 7 times a week for about 8, about 9 or about 10 weeks.

[0022] While the present invention has identified what it believes to be preferred concentrations of imiquimod, numbers of applications per week and durations of therapy, it should be understood by those versed in this art that any effective concentration of imiquimod in a formulation and any numbers of application per week that can accomplish a reduction in therapy duration to effectively treat Type I or Type II Herpes Simplex infections, actinic kurtosis and superficial basal cell carcinoma or induce effective interferon biosynthesis is contemplated by the present invention.

DETAILED DESCRIPTION OF THE INVENTION

[0023] As used in the specification and claims, the phrase "substantially non-irritating" designates formulations that do not cause unacceptable skin irritation in conventional repeat skin irritation tests in albino rabbits such as that described in Raise et al., "Appraisal of the Safety of Chemicals in Food, Drugs and Cosmetics", prepared by the Division of Pharmacology of the Food and Drug Administration, published originally in 1959 by the Association of Food and Drug Officials of the United States, Topeka, Kans. (2nd printing 1965), incorporated herein by reference.

[0024] The present invention provides pharmaceutical formulations such as creams, ointments, foams, gels, lotions and adhesive coatings that contain imiquimod and a fatty acid such as isostearic, linoleic, super purified oleic acid or oleic acid and mixtures thereof. The formulations of the invention provide desirable skin penetrability of the imiquimod.

[0025] The compound imiquimod is a known antiviral agent that is also known to induce interferon biosynthesis. It can be prepared using the method disclosed in U.S. Pat. No. 4,689,338, the disclosure of which is incorporated herein by reference. The compound can be used to treat viral infections such as Type I or Type II Herpes simplex infections and genital warts. Furthermore, the fact that the compound is an interferon inducer suggests that it, and therefore formulations containing it, might be useful in the treatment of numerous other diseases, such as rheumatoid arthritis, warts, eczema, hepatitis B, psoriasis, multiple sclerosis, essential thrombocythaemia, and cancer, such as basal cell carcinoma and other neoplastic diseases. The amount of imiquimod present in a formulation of the invention will be an amount effective to treat the targeted disease state to prevent the recurrence of such a disease or to promote immunity against such a disease. The amount is preferably about 0.5 percent to about 9 percent by weight based on the total weight of a formulation, more preferably between greater than about 1% and about 5% imiquimod, and more preferably between about 2.5%, 2.75%, 3.0%, 3.25%, 3.5%, 3.75%, 4.0%, 4.25% and 4.5%, and most preferred between about 2.75%, 3.0%, 3.25%, 3.5%, 3.75% and 4.0%.

[0026] A fatty acid such as isostearic acid, linoleic acid, super purified oleic acid, oleic acid or a mixture thereof is incorporated into a formulation of the invention. The total amount of fatty acid present in a formulation is preferably about 3 percent to about 45 percent by weight based on the total weight of a formulation. It should be understood that when oleic acid is selected as a fatty acid, that stability may present issue. Thus, stabilizers, such as anti-oxidants and the like, may be required to preserve pharmaceutical elegance and stability over the life of the oleic formulation.

[0027] A pharmaceutical formulation of the invention can be in a form such as a cream, an ointment, a foam, a gel, a lotion, a pressure-sensitive adhesive composition, or other forms known to those skilled in the art, each particular form containing imiquimod and fatty acid in particular amounts, and optionally containing various additional elements. The preferred amounts of drug and fatty acid, and the amounts and types of optional elements used in formulations of the invention are discussed below with particular reference to creams, ointments and adhesive compositions.

[0028] A cream according to the invention contains 1-isobutyl-1H-imidazo[4,5-c]quinolin-4-amine and fatty acid.

[0029] The amount of 1-isobutyl-1H-imidazo[4,5-c]quinolin-4-amine present in a cream is preferably about 0.5 percent to about 9 percent by weight, and more preferably about 1 percent to about 5 percent by weight, based on the total weight of the cream.

[0030] The total amount of fatty acid present in a cream of the invention is preferably about 3 percent to about 45 percent by weight, and more preferably about 5 percent to about 25 percent by weight, based on the total weight of the cream.

[0031] Optionally, a cream of the invention can contain emollients, emulsifiers, thickeners, and/or preservatives.

[0032] Emollients such as long chain alcohols, e.g., cetyl alcohol, stearyl alcohol and cetearyl alcohol; hydrocarbons such as petrolatum and light mineral oil; or acetylated lanolin can be included in a cream of the invention. A cream can contain one or more of these emollients. The total amount of emollient in a cream of the invention is preferably about 5 percent to about 30 percent, and more preferably about 5 percent to about 10 percent by weight based on the total weight of the cream.

[0033] Emulsifiers such as nonionic surface active agents, e.g., polysorbate 60 (available from ICI Americas), sorbitan monostearate, polyglyceryl-4 oleate, and polyoxyethylene(4) lauryl ether or trivalent cationic a cream of the invention. A cream can contain one or more emulsifiers. Generally the total amount of emulsifier is preferably about 2 percent to about 14 percent, and more preferably about 2 percent to about 6 percent by weight based on the total weight of the cream.

[0034] Pharmaceutically acceptable thickeners, such as Veegum.TM.K (available from R. T. Vanderbilt Company, Inc.), and long chain alcohols (i.e. cetyl alcohol, stearyl alcohol or cetearyl alcohol) can be used, A cream can contain one or more thickeners. The total amount of thickener present is preferably about 3 percent to about 12 percent by weight based on the total weight of the cream.

[0035] Preservatives such as methylparaben, propylparaben and benzyl alcohol can be present in a cream of the invention. The appropriate amount of such preservative(s) is known to those skilled in the art.

[0036] Optionally, an additional solubilizing agent such as benzyl alcohol, lactic acid, acetic acid, stearic acid or hydrochloric acid can be included in a cream of the invention.

[0037] If an additional solubilizing agent is used, the amount present is preferably about 1 percent to about 12 percent by weight based on the total weight of the cream.

[0038] Optionally, a cream of the invention can contain a humectant such as glycerin, skin penetration enhancers such as butyl stearate, and additional solubilizing agents.

[0039] It is known to those skilled in the art that a single ingredient can perform more than one function in a cream, i.e., cetyl alcohol can serve both as an emollient and as a thickener.

[0040] Generally, a cream consists of an oil phase and a water phase mixed together to form an emulsion. Preferably, the amount of water present in a cream of the invention is about 45 percent to about 85 percent by weight based on the total weight of the cream.

[0041] The oil phase of a cream of the invention can be prepared by first combining the 1-isobutyl-1H-imidazo[4,5-c]quinolin-4-amine and the fatty acid (if the cream contains benzyl alcohol it can also be added at this point) and heating with occasional stirring to a temperature of about 50° C. to 85° C. When the 1-isobutyl-1H-imidazo[4,5-c]quinolin-4-amine appears to be completely dissolved, the remaining oil phase ingredients are added and heating is continued until dissolution appears to be complete.

[0042] The water phase can be prepared by combining all other ingredients and heating with stirring until dissolution appears to be complete.

[0043] The creams of the invention are generally prepared by adding the water phase to the oil phase with both phases at a temperature of about 65° C. to 75° C. The resulting emulsion is mixed with a suitable mixer apparatus to give the desired cream.

[0044] An ointment of the invention contains an ointment base in addition to 1-isobutyl-1H-imidazo[4,5-c]quinolin-4-amine and fatty acid.

[0045] The amount of 1-isobutyl-1H-imidazo[4,5-c]quinolin-4-amine present in an ointment of the invention is preferably about 0.5 percent to about 9 percent, and more preferably about 0.5 percent to about 5 percent by weight based on the total weight of the ointment.

[0046] The total amount of fatty acid present in an ointment of the invention is preferably about 3 percent to about 45 percent, and more preferably about 3 percent to about 25 percent based on the total weight of the ointment.

[0047] A pharmaceutically acceptable ointment base such as petrolatum or polyethylene glycol 400 (available from Union Carbide) in combination with polyethylene glycol 3350 (available from Union Carbide) can be used. The amount of ointment base present in an ointment of the invention is preferably about 60 percent to about 95 percent by weight based on the total weight of ointment.

[0048] Optionally, an ointment of the invention can also contain emollients, emulsifiers and thickeners. The emollients, emulsifiers, and thickeners and the preferred amounts thereof described above in connection with creams are also generally suitable for use in an ointment of the invention.

[0049] An ointment according to the invention can be prepared by combining 1-isobutyl-1H-imidazo[4,5-c]quinolin-4-amine with fatty acid and heating with occasional stirring to a temperature of about 65° C. When the 1-isobutyl-1H-imidazo[4,5-c]quinolin-4-amine appears to be completely dissolved, the remaining ingredients are added and heated to about 65° C. The resulting mixture is mixed with a suitable mixer while being allowed to cool to room temperature.

[0050] A pressure-sensitive adhesive composition of the invention contains 1-isobutyl-1H-imidazo[4,5-c]quinolin-4-amine, fatty acid, and a pressure sensitive adhesive polymer.

[0051] The amount of 1-isobutyl-1H-imidazo[4,5-c]quinolin-4-amine present in a pressure sensitive adhesive composition of the invention is preferably about 0.5 percent to about 9 percent by weight, and more preferably about 3 percent to about 7 percent by weight based on the total weight of the adhesive composition. The amount of fatty acid present is preferably about 10 percent to about 40 percent by weight, more preferably about 15 percent to about 30 percent by weight, and most preferably about 20 percent to about 30 percent by weight, based on the total weight of the adhesive composition.

[0052] Preferably, the adhesive polymer utilized in a pressure sensitive adhesive composition of the invention is substantially chemically inert to 1-isobutyl-1H-imidazo[4,5-c]quinolin-4-amine. The adhesive polymer is preferably present in an amount of about 55 percent to about 85 percent by weight based on the total weight of the composition. Suitable adhesive polymers include acrylic adhesives that contain, as a major constituent (i.e., at least about 80 percent by weight of all monomers in the polymer), a hydrophobic monomeric acrylic or methacrylic acid ester of an alkyl alcohol, the alkyl alcohol containing 4 to 10 carbon atoms. Examples of suitable monomers are those discussed below in connection with the "A Monomer". These adhesive polymers can further contain minor amounts of other monomers such as the "B Monomers" listed below.

[0053] Preferred adhesives include acrylic pressure-sensitive adhesive copolymers containing A and B Monomers as follows: Monomer A is a hydrophobic monomeric acrylic or

methacrylic acid ester of an alkyl alcohol, the alkyl alcohol containing 4 to 10 carbon atoms, preferably 6 to 10 carbon atoms, more preferably 6 to 8 carbon atoms, and most preferably 8 carbon atoms. Examples of suitable A Monomers are n-butyl, n-pentyl, n-hexyl, isoheptyl, n-nonyl, n-decyl, iso-hexyl, 2-ethyloctyl, isooctyl and 2-ethylhexyl acrylates. The most preferred A Monomer is isooctyl acrylate.

[0054] Monomer B is a reinforcing monomer selected from the group consisting of acrylic acid; methacrylic acid; alkyl acrylates and methacrylates containing 1 to 3 carbon atoms in the alkyl group; acrylamide; methacrylamide; lower alkyl-substituted acrylamides (i.e., the alkyl group containing 1 to 4 carbon atoms) such as tertiary-butyl acrylamide; diacetone acrylamide; n-vinyl-2-pyrrolidone; vinyl ethers such as vinyl tertiary-butyl ether; substituted ethylenes such as derivatives of maleic anhydride, dimethyl itaconate and monoethyl formate and vinyl perfluoro-n-butylate. The preferred B Monomers are acrylic acid, methacrylic acid, the above-described alkyl acrylates and methacrylates, acrylamide, methacrylamide, and the above-described lower alkyl substituted acrylamides. The most preferred B Monomer is acrylamide.

[0055] In one embodiment of a pressure-sensitive adhesive composition of the invention, the pressure-sensitive adhesive copolymer containing A and B Monomers as set forth above preferably contains the A Monomer in an amount by weight of about 80 percent to about 98 percent of the total weight of all monomers in the copolymer. The A Monomer is more preferably present in an amount by weight of about 88 percent to about 98 percent, and is most preferably present in an amount by weight of about 91 percent to about 98 percent. The B Monomer in such a copolymer is preferably present in the pressure-sensitive adhesive copolymer in an amount by weight of about 2 percent to about 20 percent, more preferably about 2 percent to about 12 percent, and most preferably 2 to 9 percent of the total weight of the monomers in the copolymer.

[0056] In another embodiment of a pressure-sensitive adhesive composition of the invention, the adhesive copolymer comprises about 60 to about 80 percent by weight (and preferably about 70 to about 80 percent by weight) of the above-mentioned hydrophobic monomeric acrylic or methacrylic acid ester of an alkyl alcohol (i.e., Monomer A described above) based on the total weight of all monomers in the copolymer; about 4 to about 9 percent by weight based on the total weight of all monomers in the copolymer of a reinforcing monomer selected from the group consisting of acrylic acid, methacrylic acid, an alkyl acrylate or methacrylate containing 1 to 3 carbon atoms in the alkyl group, acrylamide, methacrylamide, a lower alkyl-substituted acrylamide, diacetone acrylamide and N-vinyl-2-pyrrolidone; and about 15 to about 35 percent by weight (and preferably about 15 to about 25 percent by weight) of vinyl acetate based on the total weight of all monomers in the copolymer. In this embodiment the preferred acrylic or methacrylic acid ester is isooctyl acrylate and the preferred reinforcing monomer is acrylamide.

[0057] The above described adhesive copolymers are known, and methods of preparation therefor are well known to those skilled in the art, having been described for example, in U.S. Pat. No. 24,906 ((Arich), the disclosure of which is incorporated herein by reference. The polymerization reaction can be carried out using a free radical initiator such as an organic peroxide (e.g., benzoylperoxide) or an organic azo compound (e.g., 2,2'-azobis(2,4-dimethylpentanenitrile), available under the trade designation "Vazo 52" from DuPont).

[0058] Since pressure-sensitive adhesives such as those described above are inherently rubbery and tacky and are suitably heat and light stable, there is no need to add tackifiers or stabilizers. However, such can be added if desired.

[0059] Optionally, a pressure sensitive adhesive composition of the invention can also contain one or more skin penetration enhancers such as glyceryl monolaurate, ethyl oleate, isopropyl myristate, diisopropyl adipate and N,N-dimethyldodecylamine-N-oxide, either as a single ingredient or as a combination of two or more ingredients. The skin penetration enhancer(s) preferably form a substantially homogeneous mixture with the pressure sensitive adhesive polymer or copolymer. The total amount of skin penetration enhancer(s) present in a pressure sensitive adhesive composition of the invention is preferably about 3 percent to about 25 percent by weight, more preferably about 3 percent to about 10 percent by weight based on the total weight of the adhesive composition.

[0060] When the skin penetration enhancer is a single ingredient, it is preferably a skin penetration enhancer such as isopropyl myristate, diisopropyl adipate, ethyl oleate, or glyceryl monolaurate.

[0061] When a combination skin penetration enhancer is used, it is preferably a combination such as: ethyl oleate with glyceryl monolaurate; ethyl oleate with N,N-dimethyldodecylamine-N-oxide; glyceryl monolaurate with N,N-dimethyldodecylamine-N-oxide; and ethyl oleate with both glyceryl monolaurate and N,N-dimethyldodecylamine-N-oxide.

[0062] A pressure-sensitive adhesive composition of the invention can be prepared by combining dry adhesive, 1-isobutyl-1H-imidazo[4,5-c]quinolin-4-amine, fatty acid, and skin penetration enhancer(s) with an organic solvent. The preferred organic solvents are methanol and ethyl acetate. The total solids content of the adhesive coating is preferably in the range of about 15 percent to about 40 percent, and more preferably in the range of about 20 to about 35 percent based on the total weight of the adhesive coating. The resulting mixture is shaken or mixed for a period of about 20 to 72 hours. When this method is used it is preferred that the 1-isobutyl-1H-imidazo[4,5-c]quinolin-4-amine be in micronized form (i.e., particle size of 1-2 microns in diameter). Optionally, the mixture can be heated during shaking.

[0063] In a preferred method, the 1-isobutyl-1H-imidazo-4,5-c]quinolin-4-amine is combined with the fatty acid and shaken at 40° C. until there appears to be complete dissolution. The remaining ingredients are added and the mixture is shaken for a period of about 20 to 72 hours.

[0064] The pressure-sensitive adhesive compositions described above are preferably coated onto one surface of a suitable backing of sheet material, such as a film, to form a pressure-sensitive adhesive coated sheet material. A pressure-sensitive adhesive coated sheet material of the invention can be prepared by knife coating a suitable release liner to a predetermined uniform thickness with a wet adhesive formulation. This adhesive coated release liner is then dried and laminated onto a backing using conventional methods. Suitable release liners include conventional release liners comprising a known sheet material, such as a polyester web, a polyethylene web, or a polystyrene web, or polyethylene-coated paper, coated with a suitable silicone-type coating such as that available under the trade designation Daubert 164Z, from Daubert Co. The backing can be occlusive, non-occlusive or a breathable film as desired. The backing can be any of the conventional materials for pressure-sensitive adhesive tapes, such as polyethylene, particularly low density polyethylene, linear low density polyethylene, high density

polyethylene, randomly-oriented nylon fibers, polypropylene, ethylene-vinylacetate copolymer, polyurethane, rayon and the like. Backings that are layered, such as polyethylene-aluminum-polyethylene composites are also suitable. The backing should be substantially non-reactive with the ingredients of the adhesive coating. The presently preferred backing is low density polyethylene.

[0065] The pressure-sensitive adhesive coated sheet material of the invention can be made in the form of an article such as a tape, a patch, a sheet, a dressing or any other form known to those skilled in the art.

[0066] Preferably, an article in the form of a patch is made from an adhesive coated sheet material of the invention and applied to the skin of a mammal. The patch is replaced as necessary with a fresh patch to maintain the particular desired therapeutic effect of the 1-isobutyl-1H-imidazo[4,5-c]quinolin-4-amine.

Inherent Viscosity Measurement

[0067] The inherent viscosity values reported in the Examples below were obtained by the conventional method used by those skilled in the art. The measurement of the viscosity of dilute solutions of the adhesive, when compared to controls run under the same conditions, clearly demonstrates the relative molecular weights. It is the comparative values that are significant; absolute figures are not required. In the examples, the inherent viscosity values were obtained using a Cannon-Fenske #50 viscometer to measure the flow time of 10 ml of a polymer solution (0.2 g polymer/deciliter tetrahydrofuran, in a water bath controlled at 25° C.). The examples and the controls were run under identical conditions. The test procedure followed and the apparatus used are explained in detail in the Textbook of Polymer Science, F. W. Billmeyer, Wiley-

Interscience, 2nd Edition, 1971 under: Polymer chains and their characterization, D. Solution Viscosity and Molecular Size, pp 84-85, the disclosure and textbook of which is incorporated by reference.

[0068] The following examples are provided to illustrate the invention, but are not intended to be limiting thereof. Parts and percentages are by weight unless otherwise specified. Examples of creams, ointments and pressure sensitive adhesive compositions contemplated by the present invention are described in U.S. Pat. No. 4,689,338 and U.S. Pat. No. 5,238,944. Percent modifications for, e.g., imiquimod and vehicle, to generate imiquimod formulations as described herein are likewise contemplated by the present invention. In addition, the formulations described and disclosed in U.S. patent application, Ser. No. 11/276,324, are also contemplated by the present invention. Thus, U.S. patent application, Ser. No. 11/276,324, is incorporated herein by reference in its entirety.

Preparative Method 1

Laboratory Scale Preparation of Isooctylacrylate/Acrylamide Copolymer

[0069] To a 114 gram narrow-mouth glass bottle were added: 18.6 g isooctyl acrylate, 1.4 g acrylamide, 0.04 g benzoyl peroxide, 27.0 g ethyl acetate and 3.0 g methanol. The solution was purged for thirty five seconds with nitrogen at a flow rate of one liter per minute. The bottle was sealed and placed in a rotating water bath at 55° C. for twenty-four hours to effect essentially complete polymerization. The polymer was diluted with ethyl acetate/methanol (90/10) to 23.2 percent solids and had a measured inherent viscosity of 1.26 dl/g in ethyl acetate.

Preparative Method 2

Pilot Plant Scale Preparation of Isooctylacrylate/Acrylamide Copolymer

[0070] 155 kg isooctylacrylate, 11.6kg acrylamide, 209.1 kg ethyl acetate and 23.2 kg methanol were charged to a clean, dry reactor. Medium agitation was applied. The batch was deoxygenated with nitrogen while heating to an induction temperature of 55° C. 114 g Lucidol.TM.70 initiator (available from Pennwalt Corp.) mixed with 2.3 kg ethyl acetate was charged to the reactor. The temperature was maintained at 55° C. throughout the reaction. After 5.5 hours reaction time, 114 g Lucidol.TM.70 mixed with 2.3 kg ethyl acetate were charged to the reactor. After 9.0 hours reaction time, an additional 114 g Lucidol.TM.70 initiator mixed with 2.3 kg ethyl acetate were charged to the reactor. The reaction was continued until the percent conversion was greater than 98 percent as measured by gas chromatographic evaluation of residual monomer concentration. The resulting polymer solution was diluted to 25-28 percent solids with ethyl acetate/methanol (90/10) and had a measured Brookfield viscosity of 17,000-21,000 centipoises using spindle #4 at 12 rpm. The polymer had a measured inherent viscosity of 1.3-1.4 dl/g in ethyl acetate.

[0071] The above procedure was found to provide a pressure-sensitive adhesive that is equivalent in the practice of the present invention to a pressure-sensitive adhesive prepared according to

Preparative Method 1.

[0072] A 25-30 percent solids solution of the isooctyl acrylate:acrylamide (93:7) adhesive copolymer in ethyl acetate/methanol (90:10) was coated onto a two-sided release liner using a knife-coater and coating at 0.5 mm in thickness. The adhesive-coated laminate was dried first at 82° C. for 3 minutes and then at 116° C. for 3 minutes. The dried adhesive coating was then stripped off the release liner and placed in a glass bottle. The foregoing procedure results in a reduction of the amount of any residual monomer in the adhesive copolymer.

Preparative Method 3

Preparation of Isooctyl Acrylate: Acrylamide: Vinyl Acetate (75:5:20) Copolymer

[0073] The procedure of PREPARATIVE METHOD 1 above acrylate, 8.0 g acrylamide; 32.0 g vinyl acetate, 0.32 g benzoyl peroxide, 216.0 g ethyl acetate and 24.0 g methyl alcohol. The resulting polymer was diluted with the ethyl acetate/methyl alcohol mixture to 21.52% solids. The adhesive polymer had a measured inherent viscosity of 1.40 dl/g in ethyl acetate at a concentration of 0.15 g/dl. Its Brookfield viscosity was 2,300 centipoise.

Preparative Method 4

Preparation of Isooctyl Acrylate Acrylamide: Vinyl Acetate (75:5:20) Copolymer

[0074] A master batch was prepared by combining 621.0 g of isooctyl acrylate, 41.4 g of acrylamide, 165.6 g of vinyl acetate, 1.656 g of 2,2'-azobis(2,4-dimethylpentanenitrile) (available from the DuPont Company as Vazo.TM.52), 884.52 g of ethyl acetate and 87.48 g of methanol. A 400 g portion of the resulting solution was placed in an amber quart bottle. The bottle was purged for two minutes with nitrogen at a flow rate of one liter per minute. The bottle was sealed and placed

in a rotating water bath at 45° C. for twenty-four hours to effect essentially complete polymerization. The copolymer was diluted with 250 g of ethyl acetate/methanol (90/10) to 26.05% solids and had a measured inherent viscosity of 1.27 dl/g in ethyl acetate at a concentration of 0.15 g/dl. Its Brookfield viscosity was 5580 centipoise.

EXAMPLE 1

[0075] A cream according to the present invention was prepared from the following ingredients:

	% by Weight	Amount
<u>Oil Phase</u>		
1-Isobutyl-1H-imidazo[4,5-c]-quinolin-4-amine	1.0	40.0 g
Isostearic acid	10.0	400.0 g
Benzyl alcohol	2.0	80.0 g
Cetyl alcohol	2.2	88.0 g
Stearyl alcohol	3.1	124.0 g
Polysorbate 60	2.55	102.0 g
Sorbitan monostearate	0.45	18.0 g
<u>Aqueous Phase</u>		
Glycerin	2.0	80.0 g
Methylparaben	0.2	8.0 g
Propylparaben	0.02	0.8 g
Purified water	76.48	3059.2 g

The materials listed above were combined according to the following procedure:

[0076] The glycerin, methylparaben, propylparaben and water were weighed into a 4 liter glass beaker then heated on a hot plate with stirring until the parabens isostearic acid and 1-isobutyl-1H-imidazo[4,5-c]-quinolin-4-amine were weighed into an 8 liter stainless steel beaker and heated on a hot plate until the amine was in solution (the temperature reached 69° C.). The benzyl alcohol, cetyl alcohol, stearyl alcohol, polysorbate 60 and sorbitan monostearate were added to the isostearic acid solution and heated on a hot plate until all material was dissolved (the temperature reached 75° C.). With both phases at approximately the same temperature (65°-75° C.), the water phase was added to the oil phase. The mixture was mixed with a homogenizer for 13 minutes then put into a cool water bath and mixed with a 3 inch propeller for 40 minutes (the temperature was 29° C.). The resulting cream was placed in glass jars.

EXAMPLES 2-9

[0077] Using the general method of Example 1, the cream formulations shown in Tables 1 and 2 were prepared.

TABLE 1

	% by Weight Example			
	2	3	4	5
<u>Oil Phase</u>				
1-Isobutyl-1H-imidazo[4,5-e]quinolin-4-amine	1.0	1.0	1.0	1.0
Isostearic acid	10.0	10.0	5.0	5.0
Benzyl alcohol		2.0	—	

TABLE 1-continued

	% by Weight Example			
	2	3	4	5
Cetyl alcohol		1.7	—	
Stearyl alcohol		2.3	—	
Cetearyl alcohol	6.0	—	6.0	6.0
Polysorbate 60	2.55	2.55	2.55	2.55
Sorbitan monostearate	0.45	0.45	0.45	0.45
Brij .TM. 30 ^a				10.0
<u>Aqueous Phase</u>				
Glycerin	2.0	2.0	2.0	2.0
Methylparaben	0.2	0.2	0.2	0.2
Propylparaben	0.02	0.02	0.02	0.02
Purified water	77.78	77.78	82.78	72.78

Brij .TM. 30 (polyoxyethylene(4) lauryl ether) is available from ICI Americas, Inc.

TABLE 2

	% by Weight Example			
	6	7	8	9
<u>Oil Phase</u>				
1-Isobutyl-1H-imidazo-r4,5-clquinolin-4-amine	1.0	1.0	1.0	1.0
Isostearic acid	10.0	25.0	10.0	6.0
Benzyl alcohol		2.0		2.0
Cetyl alcohol		2.2	1.7	
Stearyl alcohol		3.1	2.3	
Cetearyl alcohol	6.0	—		6.0
Polysorbate 60	2.55	3.4	2.55	2.55
Sorbitan monostearate	0.45	0.6	0.45	0.45
Brij .TM. 30	10.0	—		
<u>Aqueous Phase</u>				
Glycerin	2.0	2.0	2.0	2.0
Methylparaben	0.2	0.2	0.2	0.2
Propylparaben	0.02	0.02	0.02	0.02
Purified water	67.78	60.48	79.78	79.78

EXAMPLE 10

[0078] A cream according to the present invention was prepared from the following ingredients:

	% by Weight	Amount
<u>Oil Phase</u>		
1-Isobutyl-1H-imidazo[4,5-4-quinolin-4-amine	1.0	3.00 g
Isostearic acid	5.0	15.0 g
White petrolatum	15.0	45.0 g
Light mineral oil	12.8	38.4 g
Aluminum stearate	8.0	24.0 g
Cetyl alcohol	4.0	12.0 g
Witconol .TM. 14 ^a	3.0	9.00 g
Acetylated lanolin	1.0	3.0 g
Propylparaben	0.063	0.19 g

-continued

	% by Weight	Amount
Aqueous Phase		
Veegum .TM. K ^b	1.0	3.0 g
Methylparaben	0.12	0.36 g
Purified water	49.017	147.05 g

Witconol .TM. 14 (polyglyceryl4 oleate) is available from Witco Chemical Corp. Organics Division

^bVeegum .TM. K (colloidal magnesium aluminum silicate) is available from R. T. Vanderbilt Company Inc.

The materials listed above were combined according to the following procedure:

[0079] The 1-isobutyl-1H-imidazo[4,5-c]quinolin-4-amine and the isostearic acid were weighed into a glass jar and heated with occasional stirring until the amine was dissolved (the temperature reached 68° C.). To this solution was added, the petrolatum, mineral oil, aluminum stearate, cetyl alcohol, Witconol.TM 14, acetylated lanoline and propylparaben. The mixture was heated to 75° C. In a separate beaker, the methylparaben and water were combined and heated until the paraben dissolved (the temperature reached 61° C.). The Veegum.TM.K was added to the aqueous solution and heated at 75° C. for 30 minutes while mixing with a homogenizer. With both phases at 75° C., the aqueous phase was slowly added to the oil phase while mixing with a homogenizer. Mixing was continued for 30 minutes while maintaining a temperature to about 80° C. The jar was then capped and the formulation was allowed to cool.

EXAMPLE 11

[0080] An ointment according to the present invention was prepared from the following ingredients:

	% by Weight	Amount
1-Isobutyl-1H-imidazo[4,5-c]quinolin-4-amine	1.0	0.20 g
Isostearic acid	5.0	1.00 g
Mineral oil	12.8	2.56 g
White petrolatum	65.2	13.04 g
Cetyl alcohol	4.0	0.80 g
Acetylated lanolin	1.0	0.20 g
Witconol .TM.	143.0	0.60 g
Aluminum stearate	8.0	1.60 g

The materials listed above were combined according to following procedure:

[0081] The 1-isobutyl-1-imidazo[4,5-c]quinolin-4-amine and the isostearic acid were placed in a glass jar and heated with stirring until the amine was dissolved. The remaining ingredients were added and the resulting mixture was heated to 65° C. and then mixed while being allowed to cool to room temperature.

EXAMPLE 12

[0082] Using the general procedure of Example 11 an ointment containing the following ingredients was prepared.

	% by Weight	Amount
1-Tsobutyl-1H-imidazo[4,5-c]-quinolin-4-amine	1.0	0.20 g
Isostearic acid	6.0	1.20 g
Polyethylene Glycol 400	55.8	11.16 g
Polyethylene Glycol 3350	32.6	6.52 g
Stearyl alcohol	4.6	0.92 g

EXAMPLES 13-15

[0083] Creams of the present invention were prepared using the ingredients shown in Table 3. The Example 1 except that benzyl alcohol was used with the isostearic acid to dissolve the 1-isobutyl-1H-imidazo[4,5-c]quinolin-4-amine.

TABLE 3

	Example		
	13	14	15
	% by Weight		
Oil Phase			
1-Isobutyl-1H-imidazo[4,5-c]quinolin-4-amine	5.0	5.0	4.85
Isostearic acid	25.0	25.0	24.3
Benzyl alcohol	2.0	2.0	1.94
Cetyl alcohol	2.2	2.2	1.16
Stearyl alcohol	3.1	3.1	1.75
Petrolatum	3.0		2.91
Polysorbate 60	3.4	3.4	4.13
Sorbitan monostearate	0.6	0.6	0.73
Stearic acid			9.71
Aqueous Phase			
Glycerin	2.0	2.0	1.94
Methylparaben	0.2	0.2	0.19
Propylparaben	0.02	0.02	0.02
Purified water	53.48	56.48	46.39

EXAMPLE 16

[0084] A cream according to the present invention was prepared from the following ingredients:

	% by Weight	Amount
Oil Phase		
1-Isobutyl-1H-imidazo[4,5-c]-quinolin-4-amine	4.0	0.80 g
Isostearic acid	20.0	4.00 g
Benzyl alcohol	2.0	0.40 g
Cetyl alcohol	2.2	0.49 g
Stearyl alcohol	3.1	0.62 g
Polysorbate 60	3.4	0.68 g
Sorbitan monostearate	0.6	0.12 g

-continued

	% by Weight	Amount
Aqueous Phase		
1-Isobutyl-1H-imidazo [4,5-c]-quinolin-4-amine	1.0	0.2 g
Glycerin	2.0	0.4 g
85% Lactic acid	1.0	0.22 g
Methylparaben	0.2	0.04 g
Propylparaben	0.02	0.004 g
Purified water	60.48	12.0 g

The materials listed above were combined according to the following procedure:

[0085] The isostearic acid and 0.8 g of 1-isobutyl-1H-imidazo[4,5-c]quinolin-4-amine were combined in a glass jar and heated with stirring until the amine had dissolved. The remaining oil phase ingredients were added to this solution and the mixture was heated to about 70° C. The aqueous phase ingredients were weighed into a separate beaker and heated with stirring until the amine and the parabens had dissolved. With both phases at about 70° C., the water phase was added to the oil phase and mixed with a propeller until the mixture cooled to room temperature.

EXAMPLE 17

[0086] A mixture of 5.9415 g of the 93:7 isooctyl acrylate:acrylamide adhesive copolymer prepared in PREPARATIVE METHOD 2 above, 1.5126 g isostearic acid, 2.0075 g ethyl oleate, 0.3021 g glyceryl monolaurate, 0.2936 1-isobutyl-1H-imidazo[4,5-c]quinolin-4-amine (micronized) and 23.7 g of 90:10 ethyl acetate:methanol was placed in a small glass jar. The jar was placed on a horizontal shaker and shaken at room temperature for about 13 hours. The formulation was coated at a thickness of 20 mils

onto a 5 mil Daubert 164Z liner. The laminate was oven dried for 3 minutes at 105° F., for 2 minutes at 185° F., and for 2 minutes at 21 0° F. The resulting adhesive coating contained 59.1 percent 93:7 isooctyl acrylate:acrylamide adhesive copolymer, 15.0 percent isostearic acid, 20.0 percent ethyl oleate, 3.0 percent glyceryl monolaurate and 2.9 percent 1-isobutyl-1H-imidazo[4,5-c]quinolin-4-amine. The material was then laminated with 3 mil low density polyethylene backing and die cut into 2.056 cm.sup.2 patches.

EXAMPLES 18-20

Pressure-Sensitive Adhesive Coated Sheet Materials

Prepared Using Unmicronized

[0087] 1-Isobutyl-1H-imidazo[4,5-c]quinolin-4-amine

[0088] Using the general method of Example 17 the formulations shown below were prepared. 1-Isobutyl-1H-imidazo [4,5-c]quinolin-4-amine that had been ground with a mortar and pestle was used. The adhesive was the 93:7 isooctyl acrylate:acrylamide copolymer prepared in PREPARATIVE METHOD 1 above. The solvent was 90:10 ethyl acetate:methanol. All formulations were mixed at room temperature.

	Example		
	18	19	20
1-Isobutyl-1H-imidazo [4,5-c]-quinolin-4-amine	5.0	3.0	3.0
Ethyl oleate	5.1	5.0	8.0
Isostearic acid	10.0	10.0	6.0
Oleic acid	20.0	20.0	13.0
Glyceryl monolaurate	1.5	1.5	1.5
N,N-dimethyldodecylamine-N-oxide	1.0	1.1	3.0
Adhesive	57.4	59.3	65.4

EXAMPLE 21

[0089] A formulation with the same components in the same proportions as Example 18 was prepared using a different method. The 1-isobutyl-1H-imidazo[4,5-c]quinolin-4-amine was combined with the oleic and isostearic acids and shaken at 40° C. until there was complete dissolution of the 1-isobutyl-1H-imidazo-[4,5-c]quinolin-4-amine. The remaining ingredients were added and shaken a 40° C. for 72 hours. Patches measuring 2.056 cm.sup.2 were prepared by the general method of Example 17.

EXAMPLE 22

[0090] A mixture of 2.4734 g 1-isobutyl-1H-imidazo[4,5-c]quinolin-4-amine 3.3315 g isostearic acid and 6.6763 g oleic acid was prepared. To 1.8738 g of the above mixture was added 2.8750 g of the 93:7 isooctyl acrylate:acrylamide adhesive copolymer prepared in PREPARATIVE METHOD 2 above, 0.2548 g of ethyl oleate, 0.0510 g NN-dimethyl-dodecylamine-N-oxide, 0.0820 g glyceryl monolaurate (from Lauricidin, Inc.) and 14.0457 g of 90:10 ethyl acetate:methanol. The above was shaken for 30 hours at room temperature on a horizontal shaker. Transdermal patches were then prepared generally according to the procedures of Example 17.

What is claimed is:

1. A method of topical and/or transdermal administration of imiquimod for treating a dermatological disorder in a mammal, which method comprises.

(a) applying an effective amount of a formulation containing imiquimod on the skin of a mammal once a day at least 3 times a week on 3 different days for at least between about 8 weeks and 12 weeks; and

(b) allowing said imiquimod to remain in contact with the skin for a sufficient time following said application to permit an effective amount of the imiquimod to penetrate the skin to achieve the antiviral effect.

2. A method of claim 1, wherein said formulation is applied for 8 weeks.

3. A method of claim 1, wherein said formulation is applied for 9 weeks.

4. A method of claim 1, wherein said formulation is applied for 10 weeks.

5. A method of claim 1, wherein said formulation is applied for 11 weeks.

6. A method of claim 1, wherein said formulation is applied for 12 weeks.

7. A method of claims 1-6, wherein the formulation is applied once a day at least 4 times a week.

8. A method of claims 1-6, wherein the formulation is applied once a day at least 5 times a week.

9. A method of claims 1-6, wherein the formulation is applied once a day at least 6 times a week.

10. A method of claims 1-6, wherein the formulation is applied once a day at least 7 times a week.

11. A method of claims 1-10, wherein the formulation contains from greater than about 1% to about 5% imiquimod.

12. A method of claim 11, wherein the formulation contains about 1.25% imiquimod.

13. A method of claim 11, wherein the formulation contains about 1.5% imiquimod.

14. A method of claim 11, wherein the formulation contains about 1.75% imiquimod.

15. A method of claim 11, wherein the formulation contains about 2% imiquimod.

16. A method of claim 11, wherein the formulation contains about 2.25% imiquimod.

17. A method of claim 11, wherein the formulation contains about 2.5% imiquimod.

18. A method of claim 11, wherein the formulation contains about 2.75% imiquimod.

19. A method of claim 11, wherein the formulation contains about 3% imiquimod.

20. A method of claim 11, wherein the formulation contains about 3.25% imiquimod.

21. A method of claim 11, wherein the formulation contains about 3.5% imiquimod.

22. A method of claim 11, wherein the formulation contains about 3.75% imiquimod.

23. A method of claim 11, wherein the formulation contains about 4.0% imiquimod.

24. A method of claim 11, wherein the formulation contains about 4.25% imiquimod.

25. A method of claim 11, wherein the formulation contains about 4.5% imiquimod.

26. A method of claims 1-25, wherein the dermatological disorder is a viral disease.

27. A method of claim 26, wherein the viral disease is a Type I Herpes simplex infection.

28. A method of claim 26, wherein the viral disease is a Type II Herpes simplex infection.

29. A method of claims 1-25, wherein the dermatological disorder is genital warts.

30. A method of claims 1-25, wherein the dermatological disorder is actinic keratosis.

31. A method of claims 1-25, wherein the dermatological disorder is superficial basal cell carcinoma.

32. A method of claims 1-31, wherein the formulation further includes a fatty acid.

33. A method of claim 32, wherein the fatty acid is isostearic acid.

34. A method of claim 32, wherein the fatty acid is oleic acid.

35. A method of claim 32, wherein the fatty acid is super purified oleic acid.

36. A method of claim 32, wherein the fatty acid is linoleic acid.

37. A method of claims 1-36, wherein the formulation is selected from the group consisting of a cream, ointment, gel, foam, lotion and adhesive-coated sheet material.

38. A method of topical and/or transdermal administration of imiquimod to induce interferon biosynthesis in a mammal, which method comprises:

- (a) applying an effective amount of a formulation containing imiquimod on the skin of a mammal once a day at least 3 times a week on 3 different days for at least between about 8 weeks and 12 weeks; and
- (b) allowing said imiquimod to remain in contact with the skin for a sufficient time following said application to

permit an effective amount of the imiquimod to penetrate the skin to achieve the antiviral effect.

39. A method of claim 38, wherein said formulation is applied for 8 weeks.

40. A method of claim 38, wherein said formulation is applied for 9 weeks.

41. A method of claim 38, wherein said formulation is applied for 10 weeks.

42. A method of claim 38, wherein said formulation is applied for 11 weeks.

43. A method of claim 38, wherein said formulation is applied for 12 weeks.

44. A method of claims 38-43, wherein the formulation is applied once a day at least 4 times a week.

45. A method of claims 38-43, wherein the formulation is applied once a day at least 5 times a week.

46. A method of claims 38-43, wherein the formulation is applied once a day at least 6 times a week.

47. A method of claims 38-43, wherein the formulation is applied once a day at least 7 times a week.

48. A method of claims 38-47, wherein the formulation contains from greater than about 1% to about 5% imiquimod.

49. A method of claim 48, wherein the formulation contains about 1.25% imiquimod.

50. A method of claim 48, wherein the formulation contains about 1.5% imiquimod.

51. A method of claim 48, wherein the formulation contains about 1.75% imiquimod.

52. A method of claim 48, wherein the formulation contains about 2% imiquimod.

53. A method of claim 48, wherein the formulation contains about 2.25% imiquimod.

54. A method of claim 48, wherein the formulation contains about 2.5% imiquimod.

55. A method of claim 48, wherein the formulation contains about 2.75% imiquimod.

56. A method of claim 48, wherein the formulation contains about 3% imiquimod.

57. A method of claim 48, wherein the formulation contains about 3.25% imiquimod.

58. A method of claim 48, wherein the formulation contains about 3.5% imiquimod.

59. A method of claim 48, wherein the formulation contains about 3.75% imiquimod.

60. A method of claim 48, wherein the formulation contains about 4.0% imiquimod.

61. A method of claim 48, wherein the formulation contains about 4.25% imiquimod.

62. A method of claim 48, wherein the formulation contains about 4.5% imiquimod.

63. A method of claims 38-48, wherein the formulation further includes a fatty acid.

64. A method of claim 63, wherein the fatty acid is isostearic acid.

65. A method of claim 63, wherein the fatty acid is oleic acid.

66. A method of claim 63, wherein the fatty acid is super purified or refined oleic acid.

67. A method of claim 63, wherein the fatty acid is linoleic acid.

68. A method of claims 38-67, wherein the formulation is selected from the group consisting of a cream, ointment, gel, foam, lotion and adhesive-coated sheet material.

* * * * *