**WHAT IS CLAIMED IS:**

1. Intravaginal DHEA for use in the treatment of at least one condition selected from the group consisting of female hypoactive sexual desire disorder, female sexual arousal disorder, female orgasm disorder and female sexual interest arousal disorder in a woman who either (1) is not suffering from symptoms of vulvo-vaginal atrophy and/or (2) is not suffering from moderate to severe dyspareunia.

2. Use of DHEA in the manufacture of an intravaginal medicine for the treatment of at least one condition selected from the group consisting of female hypoactive sexual desire disorder, female sexual arousal disorder, female orgasm disorder and female sexual interest arousal disorder in a woman who either (1) is not suffering from symptoms of vulvo-vaginal atrophy and/or (2) is not suffering from moderate to severe dyspareunia.

3. A pharmaceutical product comprising a container having therein a pharmaceutical composition comprising DHEA and a pharmaceutically acceptable excipient suitable for vaginal administration, said product further including labeling directing the use of said pharmaceutical composition for the treatment of at least one condition selected from the group consisting of female hypoactive sexual desire disorder, female sexual arousal disorder, female orgasm disorder and female sexual interest arousal disorder in a woman who either (1) is not suffering from symptoms of vulvo-vaginal atrophy and/or (2) is not suffering from moderate to severe dyspareunia.

4. The intravaginal DHEA of claim 1 wherein the woman is not suffering from moderate to severe dyspareunia.

5. The intravaginal DHEA of claim 1 wherein the woman is not suffering from severe dyspareunia.

6. The intravaginal DHEA of claim 1 wherein the woman is not suffering from vulvo-vaginal atrophy.

7. The intravaginal DHEA of claim 1 wherein the woman is pre-menopausal.

8. The intravaginal DHEA of claim 1 wherein the woman is post-menopausal.

9. The use of claim 2 wherein the woman is not suffering from moderate to severe dyspareunia.

10. The use of claim 2 wherein the woman is not suffering from severe dyspareunia.

11. The use of claim 2 wherein the woman is not suffering from vulvo-vaginal atrophy.

12. The use of claim 2 wherein the woman is pre-menopausal.

13. The use of claim 2 wherein the woman is post-menopausal.

14. The pharmaceutical product of claim 3 wherein the woman is not suffering from moderate to severe dyspareunia.

15. The pharmaceutical product of claim 3 wherein the woman is not suffering from severe dyspareunia.

16. The pharmaceutical product of claim 3 wherein the woman is not suffering from vulvo-vaginal atrophy.

17. The pharmaceutical product of claim 3 wherein the woman is pre-menopausal.

18. The pharmaceutical product of claim 3 wherein the woman is post-menopausal.

CLAIMS

1. Use of DHEA in the manufacture of an intravaginal medicine for the treatment of at least one condition selected from the group consisting of female hypoactive sexual desire disorder, female sexual arousal disorder, female orgasm disorder and female sexual interest arousal disorder in a woman who either (1) is not suffering from symptoms of vulvo-vaginal atrophy and/or (2) is not suffering from moderate to severe dyspareunia.

2. A pharmaceutical product comprising a container having therein a pharmaceutical composition comprising DHEA and a pharmaceutically acceptable excipient suitable for vaginal administration, when used for the treatment of at least one condition selected from the group consisting of female hypoactive sexual desire disorder, female sexual arousal disorder, female orgasm disorder and female sexual interest arousal disorder in a woman who either (1) is not suffering from symptoms of vulvo-vaginal atrophy and/or (2) is not suffering from moderate to severe dyspareunia.

3. The use of claim 1 wherein the woman is not suffering from moderate to severe dyspareunia.

4. The use of claim 1 wherein the woman is not suffering from severe dyspareunia.

5. The use of claim 1 wherein the woman is not suffering from vulvo-vaginal atrophy.

6. The use of claim 1 wherein the woman is pre-menopausal.

7. The use of claim 1 wherein the woman is post-menopausal.

8. The pharmaceutical product of claim 2 wherein the woman is not suffering from moderate to severe dyspareunia.

9. The pharmaceutical product of claim 2 wherein the woman is not suffering from severe dyspareunia.

10. The pharmaceutical product of claim 2 wherein the woman is not suffering from vulvo-vaginal atrophy.

11. The pharmaceutical product of claim 2 wherein the woman is pre-menopausal.

12. The pharmaceutical product of claim 2 wherein the woman is post-menopausal.