

ISSN 2278 - 0211 (Online)

Chemopharmacophoric Aphrodicophore

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Abstract:

Aphrodisiac drugs are the pleasure – oriented pharmaceuticals and have value in sex – therapeutics. We made an attempt to apply new concept of chemobiological mimicry to aphrodisiac chemistry by comparing with endogenous, neurotransmitters and androgens. Dopamine, Norepinepherine, Serotonin and Testosterone were taken as the structural standards for rationalizing the selection of chemobiologic mimetics to design chemopharmacophoric aphrodicophore [CA]. This may be useful in imparting aphrodisiacal specificity at receptor levels and promoting bioenergetic sexual synergy.

1. Introduction

The remarkable growth of aphrodisiac formulations in the recent years for improving the pleasure of love life claimed therapeutical benefits. People are searching such formulative potions at about 80,000 websites of internet. Aphrodisiacs^{1,2} inspire erotic thoughts and desires through triggering the brain neurochemicals. The marketed aphrodisiac formulations³ of botanical and synthetic nature were analyzed at chembiological level for ascertaining structural mimicry.

2. Theoretical Methodology

The aphrodisiac herbology⁴⁻⁷ and synthetic drugs⁸ have notable chemical structures. They are molecular mimics of neurotransmitters and hormones, therefore, structural standards for deriving the chemobiologic entities were selected from two categories.

- Neurotransmitters Dopamine norepinepherine, serotonin
- Hormones androgen- Testosterone

The sexual interest, desire, libido, fantasy, imagination and pleasure are interlocked in the chemistries of neurochemicals, hormones and bioenergetics. Their biological influences involve three types of tropisms – neurotropism, hormonal tropism and bioergotropism.

The aphrodicological structures of the natural products and synthetic compounds were studied. The botanical structures belong to alkaloids, terpenoids, flavonoids, steroids, saponins, glycosides, cordycepins and aromatic aminoacids, whereas the synthetic products (useful in impotence/erectile dysfunction) have heterocyclics

Triazole, imidazole, piperazine indole with acidic and basic functions. The basic framework of chemically oriented aphrodisicophore lies in structural mimicry at chemobiological level^{10, 11}.

3. Discussion and Result

The chemical basis of CA needs pertinent bioactions of the chemobiologic mimics. They are enumerated below.

- Testosterone booster = libido enhancer
- Antioxidant = anti-aging
- Anti-stress = adaptogenic
- Love promoter = releases love chemicals
- Pleasure-enhancer = liberates pleasure chemicals
- Blood flow enhancer = enriches blood flow to genitals

The four types of mimicries were indentified which have aphrodisiacal complementariness. They are

Androgenic mimicry = Testosterone

Catecholic mimicry = Dopamine Norepinephrine

Phenylethylamine mimicry = Dopamine Norepinephrine

Indolyl mimicry = Serotonin

3.1. Androgenic Mimicry¹²⁻¹⁵

Panaxsapogenin Ecdysteroids, curculigol, curculigenin, androstenol, withaferin A Mesterolone sitminoside, simarolide, Quassinoids (partial), Diogenin, Protodiosin and Glycyrrhizinic acid have structural resemblance with androgen at nuclear level. It is assumed that natural aphrodisiacs having such mimicry, may act as substrates for testosterone boosting by direct or indirect actions as listed below:

- Direct stimulation of the leydig cells in the testes
- Stimulation of the pituitary to produce more LH
- Reduction of estradiol levels through competitive receptor binding (anti-estrogens) or decreased production of aromatase enzyme.
- Increase production of enzymes responsible for the biosynthesis of testosterone.

The indirect actions are also capable of impacting testosterone levels

- Reducing prolactin
- Stimulating oxytocin
- Stimulating noradrenaline.

3.2. Catecholic Mimicry

The bioflavonoids having the chrome nuclear structure have catecholic/phenolic functions. Butein, Butrin, Epicatechin, Leucodelphindin, Licochalcone B, Paradol, Resveratol¹⁵ and Rutin are supportive examples. Their antioxidative and anti-ageing bioactions promote physiological youthfulness.

3.3. Phenyethylamine Mimicry

The alkaloids eg Apomorphine mimics dopaminergic features. It is vasodiatory, No donor and PDE inhibitor⁹. All this is favorable for blood flow to penile vasculature.

3.4. Indolyl Mimicry

Yohimbine intensifies libido, parasympathetic activity and blood flow to erectile tissue. It also acts as anti-anxiety. The indolyalkylamine moiety partially mimics serotonin.

Cordycepins¹⁶ have an exceptional non-mimetic structure and behaves as fortifier of aphrodisiac potential. They empower internal energy of body, enhance immunity, protect from free radiacl damage and act as anti-physical stress structure. The synthetic-pharmaceutical^{17,18} are leodopa, Trazodone, Bupropinon, Fenfluramine, Fluoxetine and Paraxetine. They partially

The synthetic-pharmaceutical^{17,18} are leodopa, Trazodone, Bupropinon, Fenfluramine, Fluoxetine and Paraxetine. They partially mimic amphetamine and catechol chemistry of the neurotransmitters. The propyl chain, 6- membered heterocyclic (preferably piperazine piperidine) and strong electron attracing groups induce the structural deviation from the chembiological mimicry, but this is useful as their antidepressive bioactions, synerge the aphrodisiac efficacy.

4. Conclusion

The hormonal tropism, prostate secretion, activation of love chemicals and blood flow to the gentials are the physiological determinants of sexual competency. Therefore suggested chemopharmacophorical approdisicophore of four chemobiomimetic features

- Catecholic
- Phenylethylamine
- Androgenic
- Indolyl amine

Should meet the basic essentials of aphrodisiacs. Hopefully this model may help in developing love- inspiring and pleasure enhancing pharmaceuticals.

5. Acknowledgement

The authors express their thankfulness to Shri Kshitiz Nagrath for helping in the preparation of manuscript.

6. References

- 1. Aphrodisiacs: Love Potions or Poisons? http://www.netasia.net/users/truehealth/aphrodisiacs.htm
- 2. Aphrodisiacs: http://www.naturalnurse.com/naturalaphro.htm
- 3. Anil Ballwin: Natural products as Aphrodisiacs: M.sc Thesis (Pharmchem) submitted to H.N.B. University Garhwal Uttrakhand India 2008.
- 4. http://www.everaldo.com herbal aphrodisiac
- 5. "Plants of Love: The history and aphrodisiac and a guide to their identification and use by Christian Ratsch, Ten speed Press, Berkeley, CA, ISBN: 0-898915-928-8
- 6. S.C. Malhotra and D.P. Sharma (eds.), Pharmacological investigations of certain medicinal plants and compound preparations used in Ayurveda and Siddha. CCr in Ayurveda and Siddha, New Delhi, India (1996)
- 7. N.G. Bisset (ed.), Herbal drugs and Phytopharmaceuticals. CRC Pre boca Raton, U.S. A.(1993) P.C. Sharma, M.B. Yelne and T.J. Dennis (eds.), Databaseon Medicinal Plants used in Ayurveda, Vol.1 (2000) Vol. 2 (2001), Vol. 3(2001) Vol. 4(2002), Vol 6(2003) Central council for research in Ayurveda and Siddha, New Delhi India.
- 8. Evans WO. Chemical aphrodisiacs. Psychopharmacol Bull. 1969;5(2);10-17[Pub Med]
- 9. Kuthe A, Wiedenroth A, Magert HJ, Uckert S, Forssmann WG, Stief CG, et al. Expression of different phosphodiesterase genes in human cavernous smooth muscle. J urol. 2001;165:280-3[Pub Med]
- 10. Aphrodisiacs: Their biological basis.http://sulcus.berkeley.edu/mcb/165_001/papers/manuscripts/_98.html What Some Scientists are saying about Supposed Aphrodisiacs. ABC7.com http://webapp.abclocal.go.com/kabc/health/031805_HS_Supposed _Aphrodisiacs.html
- 11. Designer Aphrodisiac. http://www.naturalnurse.com/naturalaphro.htm
- 12. A selection of prime Ayurvedic Plant Drugs Ancient-Modern Concordance, Sukh Dev, 2006, Anamaya Publishers, F-154/2, Lado Sarai, New Delhi-110030, India
- 13. http://www.medicallook.com/reviews/Mesterolone.html
- 14. http://images.google.co.in/images?imgurl=http://www.mdidea.com/support/shogaol.jpg&imgrefurl=http://www.mdidea.com/support/glossary_phytochemicals_s.html&h=200&w=200&sz=118&hl=en&start=14&tbnid=Cw60MitYVTPAVM:&tbnh=104&tbnw=104&prev=/images%3Fq%3Dthe%2Bchemical%2Bstructure%2Bof%2Bquassinoids%26gbv%3D5226hl%3Den.
- 15. http://images.google.co.in/images?imgurl=http://www.mdidea.com/support/shogaol.jpg&imgrefurl=http://www.mdidea.com/support/glossary_phytochemicals_s.html&h=200&w=200&sz=118&hl=en&start=14&tbnid=Cw60MitYVTPAVM:&tbnh=104&tbnw=104&prev=/images%3Fq%3Dthe%2Bchemical%2Bstructure%2Bof%2Bquassinoids%26gbv%3D5226hl%3Den. Resveratrol[501-36-01]
- 16. Professor P.L. Greenhoff, Mycology Research Laboratory. www.mycologyresearch.com(circa September 2004)
- 17. Burger's medicinal chemistry and drug-discovery sixth edition vol. 6 Nervous system agents edited by Donald J Abraham ISBN-0471-27401-1 (2003) John Wiley sons 484-508
- 18. Lemke Foye's Principles of Medicinal Chemistry 5th Edn (1974) reprinted (2002)