Evaluation of Spermatogenic Activity of Panchamrita Parpati in Albino Rats

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ABSTRACT

Most of Rasashastra texts have mentioned Panchamrita parpati having vrushya property expected to contribute positive results in infertility and impotency so an attempt was made in this regard to establish and provide scientific data for reference claimed with spermatogenic activity study. Panchamrita parpati was prepared as per reference of Bhaishiya Ratnavali and subjected for evaluation of its spermatogenic activity in Wister strain albino rats Three samples of Panchamrita parpati (pp) low dose (4.5mg) GI, Moderate dose (13.5mg) GII, & High dose (18 mg) GIII, & Wister strain albino rats formed the materials for study. Test drug doses were administered with Ghee in three Groups & orally Ghee in control group. Analysis of cholesterol, Protein, Glycogen Histometric studies were followed .All the samples of test drug have shown significant spermatognic activity but effect of GIII PP (High dose) was significant than GI, GII, GIV. Panchamrita parpati may be used in different condition of male infertility i.e. in total spermatogenic activity and improve motility, cell division production of sex hormones and spermatocytes to spermatozoa. The effect might be due to balya brumhana, rasayana and vrushya properties and chemical components of Panchamrita parpati.

KEYWORDS: Spermatogenic activity, Panchamrita parpati

INTRODUCTION

Rasashashtra is the study of metal and minerals in which great importance has been given to Rasa. The four basic parada yogas in Rasashashtra are: a) Khalwi Rasayana Kalpa, b) Parapati Rasayana Kalpa, c) Kupipakwa Rasayana Kalpa, d)Pottali Rasayana Kalpa.

Among these Parapati Rasayana Kalpa is one of the pharmaceutical forms of Rasoushadhi. It possesses flake like appearance and can be powdered. Preparation and use of the parpati has been first mentioned by Chakrapani in his commentary Chakradatta during 11th century in the treatment of Grahani. Panchamrita parpati, one of the Rasoushadhi possesses vrushya property as per classical text like Bhaishajya Ratnavali¹, Rasendra sara Sangraha², Rasa Kaamdhenu³ etc explained about the vrushya property of panchamrita parapati.

Reproduction is the unique property and vital process of the living being. Although infertility may not be a public health priority in many countries, it is the *How to cite this paper:* Dr. Rajashri. L. Wathakar "Evaluation of Spermatogenic Activity of Panchamrita Parpati in

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central issue in the lives of the individuals who suffer from it. It is a source of social and psychological suffering for both men and women and can place great pressures on the relationship within the couple⁴

Drastic changes of lifestyle, diet, environment, workrelated stress, etc in modern era, have negatively affected physical and mental health, including increased incidence of infertility and impotency. In 2001 W.H.O Report of a meeting on ' Current Practices and Controversies Assisted in Reproduction', it is mentioned that infertility affects more than 80 million people worldwide.⁴ According to W.H.O, 51.2% couples are infertile because of a male factor.⁵ W.H.O also recommended that infertility should be recognized as a public health issue worldwide, including developing countries.⁶

In India primary and secondary infertility figures as given in W.H.O studies are 3% and 8%. The Ninthfive year plan (1997-2002) document of the Government Of India had included infertility in the comprehensive reproductive and child health package⁷.

According to Ayurvedic classics, aim of Ayurveda is to promote the positive health and protect the health of healthy individuals and to cure the disease of sick⁸. Aims or pursuits of the life are to achieve four purusharthas namely Dharma, Artha, Kaama, and Moksha⁹. Out of which Kaama is interpreted as desire for sexual enjoyment, which is essentially important for procreation as well as recreation and relaxation. In ancient times it was believed that without having child one cannot attain moksha.

Vajeekarana, one of the eight branches of Ayurvedha has mentioned several drugs having vrushya property, which helps one in having progeny, increase sexual desire, improve quality and quantity of semen and enable to satisfy partner.

Almost all the medical systems have indicated drugs and treatment modalities for infertility and increasing libido. But most of them turn to be not so effective as claimed or not cost effective for middle and lower economic class of society.

Rasaoushadhis are having quick and better effect than bottle. herbal medicines with minimum dosage and without unpleasant taste. These medicines can be a better solution for infertility and impotency. Most of the Rasashastra texts have mentioned panchamrita parpati are Materials: having vrushya property. Hence this project op 1. Panchamrita parpati "Evaluation of spermatogenic activity of Panchamrita parpati in albino rats" was undertaken2 to evaluate the spermatogenic activity and to contribute a potent formulation for infertili

Equipments:

Electronic weighing balance Microscope slides with cover slips Centrifuge Microscope with optical meter Spectrophotometer Dissection box Water bath

Glass wares:

Disposable syringes (2 ml) **Pipettes**

Chemicals and Reagents:

1) Carboxyl methyl cellulose 3) 10% formaline 5) Ethanol 7) Sodium Hydroxide 9) Hydrochloric acid 10) Ether 11) Albumin 12) NaCl2 13) CuSO4 15) Sodium Potassium Tartrate 16) Folin-cio-calteu reagent (FCR)

Materials & Method of Phamaceutical study

Preparation of Panchamrita parpat: **Ref:** Bhaishajya Ratnavali

Shuddha Parada 30gm Shuddha Gandhaka-60gm Loha bhasma -15gm Abhraka bhasma - 7.5gm Tamrabhasma _ 3.75gm

Equipments:. Khalwayantra, Loha darvi, Spatula, Agni, Gomaya, Kadali Patra, Samadala pidhanaka, Sneha, Hot Water.

Method:

The drugs should be taken in a Loha darvi, smeared with ghee and heated. On complete heat melting it was should be poured over banana leaf (smeared with ghee) kept on fresh cow dung.

Another banana leaf smeared with ghee was kept over it and pressed gently with fresh cow dung over banana leaf immediately and given pressure gently.

After Swangasheetha, the parpati was collected from the leaf and washed with hot water, dried parpati was converted into powder & preserved in dried glass

MATERIALS **METHODS** AND **EXPRIMENTAL STUDY**

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2. Goghrita 3. Male Albino rats

Weighing machine Neubauer's counting chamber Tissue homogenizer Microtone pH meter Whatman Filter paper No:42 26 No. needle

Test tubes Glass beakers

- 2) Normal Saline 4) Haematoxylene and Eosin stains 6) Sodium Bicarbonate 8) Chloroform 14) Distilled water
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- 17) Trichloroacetic acid (TCA)
- 18) Phenol
- 19) Conc. Sulphuric acid
- 21) Standard cholesterol

20) Acetic anhydride 22) Xylene

23) Paraffin wax Met

Method:

- 1. Healthy adult male albino rats (Wister strain) of 90-100 days old, weighing from 175-200 gm were used for the experiments.
- 2. The animals were maintained under laboratory conditions with controlled environment of temperature, humidity, light and dark cycles.
- 3. The animals were fed with balance pellet diet as prescribed by CFTRI (central food and technological research institute, Mysore) and water and libitum.
- 4. 6 animals were taken in each group for experiment and maximum 3 animals per cage were maintained.
- 5. Animals were selected from animal house of A.V. Samiti's PGCRC Ayurveda Mahavidyalaya Bijapur.

Experimental Protocol:

Sample Size: 24 albino rats were taken for experimental study. 6 albino rats in each group.

Spermatogenic study:

Study Groups:

Group I: Administration of Panchamrita parpati Minimum dose with ghee.

Group II: Administration of Panchamrita parpati Moderate dose with ghee.

Group III: Administration of Panchamrita parpati higher dose with ghee.

Group IV: Administration of Ghee Control

Experimental Protocol

Study design	Group I	Group II	Group III	Group IV
Sample size	6 rats	Rese 6 rats and	6 rats	6 rats
Drug	P.P Low dose with ghee	• P.P Moderate dose with ghee	P.P High dose with ghee	Ghee
Dose108	4.5mg	13.5mg	18mg	0.5ml
Dosage form	Liquid	Liquid	Liquid	Liquid
Route	Oral	Oral	Oral	Oral
Duration of study	30 days	30 days	30 days	30 days
Autopsy	31st day	31st day	31st day	31st day

Observation & result

Effect of drugs on body weight before and after the administration of drug

			n=06
Groups	Before administration	After administration	Before v/s After
			t=8.654
Group-I	1833.33±4.082	240.833±14.289	p=0.0001
		After administration 240.833±14.289 251.66±13.292 258.33±9.832 238.33±1.05	HS
			t=10.542
Group-II (PP)	189.166±5.845	251.66±13.292	p=0.0001
			HS
			t=14.45
Group-III (PP)	187.5±6.892	258.33±9.832	p=0.0001
			HS
Crown W (Control)	177.5 ± 1.12	228 22+1 05	t=97.056
Group-1v (Collirol)	177.J± 1.12	230.33±1.03	p=0.0001

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Showing the effect of drugs on net increase in bodyweight after administration of drugs

				11-00
Group	GI	G II	G III	G IV
Weight gain	52.503±0.71	62.494±0.71	70.83±0.75	60.83±0.54
17.1 16				

Values are Mean ± SEM

Table Showing the effect of drugs on biochemical parameters.

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n - 116	•
11-171	

n_06

Biochemical	After	r Admiı Dr	nistratio ugs	istration of gs GIV/S GII v/S GIII v/S GI v/S G GIV CIV CIV CIV			G I v/s	G II v/s		
Analysis	GI	GII	G III	GIV	GIV	GIV	GIV	GII	GIII	GIII
Cholestrol (mg/gm)	11.533± 0.1633	12.9± 0.894	14.783± 0.318	9.38± 0.204	t=25.133 p=0.0001 HS	t=38.712 p=0.0001 HS	t=34.96 p=0.0001 HS	t=25.733 p=0.0001 HS	t=36.926 p=0.0001 HS	t=38.712 p=0.0001
Glycogen (mg/gm)	2.836± 0.0427	3.4± 0.1265	3.476± 0.0258	2.64± 0.08	t=5.294 p=0.0004 HS	t=12.438 p=0.0001 HS	t=24.362 p=0.0001 HS	t=10.347 p=0.0001 HS	t=31.423 p=0.0001 HS	t=1.442 p=0.1799 NS
Protien (mg/gm)	16.1± 0.1673	18.4± 0.1095	21.25± 0.1643	14.77± 0.120	t=26.871 p=0.0001 HS	t=54.734 p=0.0001 HS	t=78.015 p=0.0001 HS	t=50.820 p=0.0001 HS	t=76.357 p=0.0001 HS	t=35.357 p=0.0001 HS

Values are Mean ± SEM

Table No: Showing the effect of the drugs on Spermatogenic elements **n=06** After Administration of Drugs GI GI **Sperma** GII **G**III GII GI togenic v/s v/s GI **G** III GII **GIV GIV Elements GIV GIV** GII **G**III **G**III t=3.15 t=3.58 t=0.37 t=3.17 t=3.68 t=2.46 Spermat 91.83± 101.0± $102.42 \pm 90.17 \pm$ p=0.717 p=0.025 p=0.016 p=0.025 p=0.014 p=0.033ogonia 7.00 1.100.976 8.35 NS S HS S HS HS t=0.07 t=0.37 t=6.39 t=0.27 t=6.24 t=24.17 Sperma $204.86 \pm 172.8 \pm$ 173.3± p=0.944 p=0.726 p=0.001p=0.80p=0.002 p=0.00 174.67±1.63 tocytes 12.1 2.7912.0 NS HS HS NS NS HS t=0.003 t=3.6 t=6.76 t=3.60 t=6.69 t=9.00 228.29 ± 205.00 217.5± Spermatids 204.83±8.35 p=0.973 p=0.015 p=0.001 p=0.016 p=0.001p=0.00 2.17 2.14 ± 8.20 NS HS HS HS HS HS t=0.86 t=0.17 t=3.66 t=1.28 t=2.47 t=8.17 Spermcount $48.72 \pm$ 46.17± 53.3± $46.50 \pm$ p=0.413|p=0.873|p=0.015|p=0.249|p=0.057|p=0.00(mil/susp) 4.48 1.97 0.906 4.48 NS NS HS NS S HS

Values are Mean ± SE

Table No: Showing the effect of the drugs on micrometric changes of testis

Micrometric Changes of	Aft	er Admi Dr	nistratio ugs	n of	G I v/s	G II v/s		G I v/s	G I v/s	G II v/s
the Testis	GI	GII	G III	GIV	G IV	GIV	V/S G I V	GII	G III	GIII
Diamatar of	7922+	9333.3	10200+	7922+	t=0.00	t=7.48	t=10.38	t=10.45	t=7.33	t=7.54
Dialificter of	1033±	±81.6	10200 ± 200	/033±	p=1.00	p=0.001	p=0.000	p=0.000	p=0.00	p=0.001
Testis (µm)	480		300	484	NS	HS	HS	HS	HS	HS

Values are Mean ± SEM

Table Showing the effect of the drugs on micrometric changes of seminiferous tubule

Micrometric Changes	After Administration of				GI	GII	G III	GI	GI	G II
		Dr	ugs		v/s	v/s	v/s	v/s	v/s	v/s
	GI	GII	G III	G IV	G IV	GIV	G IV	GII	G III	G III
Diamatar of tubulas	182 71	205 67	200 14	192 21	t=0.05	t=4.41	t=5.11	t=4.42	t=5.14	t=4.73
Diameter of tubules	183.7± 12.1	203.07	1.51 ± 1.07	185.5± 12.3	p=0.963	p=0.007	p=0.004	p=0.007	p=0.004	p=0.001
(μπ)		±1.31			NS	HS	HS	HS	HS	HS

Values are Mean ± SEM

n=06

n=06

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Table Showing the effect of the drugs on reproductive accessory organs.											
Effects of drugs	After A	Administ	ration of	f Drugs	GI	GII	G III	GI	GI	GII	
on accessory organs	GI	G II	G III	GIV	v/s G IV	v/s G IV	v/s G IV	v/s G II	v/s G III	v/s G III	
Weight of Testis	1.338± 0.068	1.39± 0.0110	1.4371± 0.0214	1.345± 0.046	t=0.20 p=0.849 NS	t=2.31 p=0.069 NS	t=4.48 p=0.004 HS	t=1.82 p=0.129 NS	t=3.38 p=0.020 S	t=5.10 p=0.001 HS	
Epididymis	0.425± 0.027	0.4566 ±0.005	0.510± 0.020	0.416± 0.0258	t=0.54 p=0.601 NS	t=3.72 p=0.014 HS	t=7.20 p=0.00 HS	t=2.78 p=0.039 S	t=6.30 p=0.00 HS	t=6.00 p=0.00 HS	
Seminal Vesicles	0.2933± 0.0242	0.368± 0.129	0.3429± 0.0243	0.2833 ±0.0207	t=0.77 p=0.461 NS	t=1.59 p=0.172 NS	t=4.77 p=0.001 HS	t=1.40 p=0.221 NS	t=3.67 p=0.0014 HS	t=0.48 p=0.654 NS	
Prostate	0.0533± 0.0082	0.0750 ±0.008	0.0643± 0.00535	0.050± 0.0089	t=0.67 p=0.517 NS	t=5.00 p=0.001 NS	t=3.42 p=0.011 HS	t=4.54 p=0.001 HS	t=2.81 p=0.02 S	t=2.70 p=0.027 S	
Vas Deferens	0.0500 ± 0.0082	0.0566 ±0.0052	0.0683± 0.00408	0.0450± 0.0105	t=0.89 p=0.392 NS	t=2.44 $p=0.044$ S	t=5.00 p=0.002 HS	t=1.58 p=0.153 NS	t=4.57 p=0.004 HS	t=4.34 p=0.002 HS	

Table Showing the effect of the drugs on reproductive accessory organs.

Values are Mean ± SEM

Discussion & Conclusion:

- > All the samples of test drug have shown significant spermatognic activity.
- In total spermatogenic activity and improve motility, cell division production of sex hormones and spermatocytes to spermatozoa.

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Effect of GIII PP (High dose) was significant than S > The effect may be due to Balya, Brumhana, GI, GII, GIV Panchamrita parpati us in different condition of male infertility.
The effect may be due to Balya, Brumhana, Rasayana and Vrushya properties and chemical components of Panchamrita parpati