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**Research Article** 

# A COMPARATIVE CLINICAL EFFICACY STUDY OF *TRIPHALA KWATH* AND *TRIKATU* CAPSULES IN THE MANAGEMENT OF *MEDOROGA* W.S.R TO DYSLIPIDEMIA

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ABSTRACT

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*Medoroga* is a condition in which there is an excessive accumulation of

Meda Dhatu in the body. Accumulation of Medo Dhatu in different parts of

the body causes blockage of Strotsa which ultimately leads to poor nourishment of other Dhathus. Lack of exercises and Kaphavardhak Ahar Viharar are the two main causes of Medoroga. In modern times, way of life has changed drastically to quick nourishments and inactive tendencies throughout the world. Because of these factors, accumulation of Meda *dhatu* happens immensely. In Ayurveda, dyslipidemia is considered under *Medoroga*. Dyslipidemia is an emerging serious health abnormality associated with co-morbidities including CVD that continues to be the leading cause of death worldwide. It is characterized by an increase in cholesterol, triglycerides, LDL levels, and a decrease in HDL levels. The present study has been designed to compare the efficacy of Triphala Kwath and Trikatu capsules on various clinical parameters in the management of Medoroga w.s.r to dyslipidemia. Twenty patients fulfilling the inclusion criteria were randomly selected for the trial and put into two groups of ten patients each. Trikatu capsules were given to patients in group I and Triphla kwath with Madhu and Trikatu capsules in combination were given

to patients of group II for 8 weeks. Patients were thoroughly assessed on

various scientific parameters during the complete trial period. In group II, a

significant (p < 0.001) improvement was observed in subjective parameters

and serum cholesterol, triglycerides, LDL, VLDL levels while in group I, a

significant (p < 0.001) improvement was observed in HDL only. It may

affirmatively be construed from the study that the best impact of the trial

drugs was observed with Triphala Kwatha with Madhu and Trikatu Capsule

together (i.e. Group II). This combination therapy was most effective in

reducing the overall lipid profile with substantial gains related to

subjective as well as objective parameters without any adverse effects.

**KEYWORDS:** *Medoroga*, Dyslipidemia, *Triphala Kwatha*, *Madhu* &*Trikatu* Capsules.

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### INTRODUCTION

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India is undergoing a rapid epidemiological transition with increasing population, economic wealth and urbanization,<sup>[1]</sup> however increase in adverse lifestyles such as smoking and tobacco use, nutritional habits with intake of unhealthy diet, and increasing sedentary lifestyle ultimately leads to non-communicable diseases, including coronary heart disease, cerebrovascular and peripheral artery diseases<sup>[2,3]</sup>. Even in rural areas of India non-communicable diseases have become the leading

cause of death<sup>[2,3]</sup>. *Medoroga* (Dyslipidemia) is a single contributory factor for many life style diseases. It is characterized by an unusual measure of lipids (triglycerides, cholesterol and fat phospholipids) in the blood. Lipids are the naturally occurring molecules which constitute the structural component of various cell membranes. Lipids have a normal property of *Snehatwa* (lubricity)<sup>[4]</sup>. According to Ayurveda, *Medo Dhatu, Vasa* and *Majja Dhatu* are the three components of out of seven

*Dhatus* that comprises human body. These additionally show same property of *Snehatwa* thus lipids can be corresponded with these three elements. They show regular highlights however each one of those are available at destinations and perform various capacities. Sneha offers chubbiness, lubricates every cell of the body and provides strength, energy to the body when in normal state<sup>[5]</sup>, when it is in an upset state it causes Rasagata Snehavriddhi and leads to Medodushti (Dyslipidaemia)<sup>[5]</sup>. The depiction of *Medoroga* has been given in different classical texts. It results from various etiological factors, for example, defective eating regimen, doing no physical exercise, getting a charge out of day rest, and taking Kapha inciting diet and sugar rich eating routine. The etiological factors are Kapha and Meda Sadharmi Ansha in overabundance hence produce Kapha Bhuista Dosha Virddhi<sup>[6]</sup>. That Dosha Vridhi because of its very nature influences the Agni and producesAgni Vikriti. As an outcome, the Ama goes straight forward to *Medodhatu* and blends with *Kapha* at the tissue level and causes an increase in Meda Dhatu. This results in *Srotasavrodha*, channels of *Dosha* are blocked and thus Vata in the Kostha causes Jatharagni Sandhukashan, which results in food cravings and leads to over-intake of food consequentially leading to *Medoroga*<sup>[6]</sup>. "Infections and intricacies that happen in patients of *Medoroga* are difficult to treat, than those of patients who don't have *Medoroga*"<sup>[7]</sup>. There is no straight forward reference in Ayurveda that can be associated with dyslipidemia. Still, based on description of *Medoroga*, it can be correlated with Medoroga. Dyslipidemia incorporates hyperlipidemia and hyper lipoproteinemia which are caused because of abnormally raised degrees of lipoproteins in the blood<sup>[8]</sup>. These lipids incorporate phospholipids, cholesterol, cholesterol esters, and triglycerides. Lipoproteins are partitioned into 5 classes based on thickness as (a) HDL (Highthickness lipoprotein), (b) LDL (Low-thickness lipoprotein), (c) IDL (Intermediate thickness lipoprotein), (d) VLDL (Very low thickness lipoprotein), (e) chylomicrons. Dyslipidemia is one of the significant issues, which is characterized by the increased levels of either cholesterol, LDL cholesterol or triglycerides in the serum or both LDL and triglycerides and decrease in HDL in the serum that adds to atherosclerosis<sup>[8]</sup>. Atherosclerosis is the major risk factor for morbidity and mortality associated with Cardio-vascular Disease (CVD)<sup>[2,3]</sup>. Keeping in view the prevalence and intricacies of *Medoroga*, the present study has been designed to compare the efficacy of Triphala Kwath

and *Trikatu* capsules on various clinical parameters in the management of *Medoroga* w.s.r to dyslipidemia.

## **MATERIALS AND METHODS**

To fulfill all the aims and objectives of the study, the following materials and methods were used.

# Selection of the patients

Patients were randomly selected from OPD/IPD of R.G.G.P.G. Ayurvedic College and Hospital, Paprola, Dist. Kangra (H.P.). A detailed history was obtained, physical examination was conducted and relevant investigations were carried out before enrolment of the study subjects.

Total 20 patients were selected for the clinical trial of this research work.

# **Grouping of Patients**

Twenty patients fulfilling the inclusion criteria were randomly selected for the trial and put into two groups of ten patients each.

Group I patients were administered *Trikatu* capsules.

Group II patients were administered *Triphla kwath* with *Madhu* and *Trikatu* capsules.

# Dose of Drugs and Route of Administration

Triphla Kwatha 20ml b.i.d with one tsf Madhu

*Trikatu* capsules 500mg b.i.d with water.

Route of Administration- Oral

Duration of Trial- 8 weeks

## **Inclusion Criteria**

- 1. Diagnosed and confirmed cases of dyslipidemia based on investigations i.e.
- (a) Serum Cholesterol>200mg/dl and <300mg/dl
- (b) Serum Triglycerides>150mg/dl and <450mg/dl
- (c) Serum LDL>100mg/dl and <200mg/dl
- (d) Serum VLDL>30mg/dl and <100mg/dl
- (e) Serum HDL<40mg/dl and <20mg/dl
- 2. Patients between the age group of 40-60 years of either sex who have fulfilled the criteria of objective and subjective parameters.

## **Exclusion Criteria**

- 1. Patients not willing for the trial.
- 2. All *Medo Rogi* presenting with complications like *Prameha Pidika*.
- 3. Drug induced dyslipidemia.
- 4. Patients having illness like TB, carcinoma, renal and liver disorders.
- 5. Patients having history of MI and unstable angina.
- 6. Patients below 40 and above 60years of age.

7. Patients with Serum Cholesterol>300mg/dl, Serum Triglycerides>450mg/dl, Serum HDL< 20mg/dl, Serum LDL>200mg/dl, Serum VLDL> 100mg/dl.

## **Criteria for evaluation**

## Assessment criteria

Clinical results were assessed based on subjective and objective parameters. Scoring system was adopted to assess subjective signs and symptoms. A special proforma was designed for assessment based on Ayurvedic principles and modern scientific knowledge. Patients were also inquired about any growing feeling of physical and mental well-being after the therapy.

### Subjective criteria

It includes signs and symptoms described in Ayurvedic texts. These are the following:

- Chala- Sphika- Udara -Stana
- Kshudra Shwasa
- Sandhi Shoola
- Kshudha Atimatra
- Pipasa Atiyoga
- Nidradhikya
- Javop rodha
- Swedabadha
- Daurgandhya

Table1: Grading of symptoms									
Symptoms	Grading								
Chal -sphik- udar- stanam (Movement of Body Parts)									
Absence of movement	0								
Little movement after fast activity	1								
Movement after mild activity	2								
Movement even on changing posture	3								
Kshudra Shwasa (Shortness of Breath)									
No shortness of breath	0								
Shortness of breath after hard physical exertion	1								
Shortness of breath after mild physical exertion	2								
Shortness of breath even at rest	3								
Sandhi Shool (Pain in Joints)									
No pain	0								
Mild pain of low intensity causing no disturbance in routine work	1								
Moderate pain occasionally hampering daily routine work	2								
Severe pain causing a definite interruption in routine work	3								
Kshudha Atimatra (Excessive Hunger)									
<2 Chapatis/meal	0								
2-4 Chapatis/meal	1								
4-6 Chapatis/meal	2								
>6 Chapatis/meal	3								
Pipasatiyoga (Excessive Thirst)									
1.5 lit/day of fluid	0								
1.5-2 lit/day of fluid	1								
2-3 lit/day	2								
3 lit/day	3								
Nidradhikya (Excessive Sleep)									
6-8 Hrs/day	0								
8-10 Hrs/day	1								
10-12 Hrs/day	2								
>12 Hrs/day	3								
Javoparodha (Lassitude)									

Table1: Grading of symptoms

Fully active	0
Hesitate to start work but once start, complete it	1
Starts but not complete it	2
Doesn't have the drive, works under compulsion	3
Swedabadha (Excessive Sweating)	
Normal perspiration	0
Mild perspiration after doing exertion	1
Heavy perspiration after doing little exertion	2
Perspiration without exertion	3
Daurgandhya (Bad Body Odour)	
Absence of odour	0
Occasional bad odour	1
Persistent bad odour	2
Persistent bad odour, intolerable to the patient	3

## **Objective Criteria**

- Lipid Profile
- Body Mass Index (BMI),
- Body weight
- Skin Fold Thickness

# **Investigation Based Criteria**

It mainly consists of lipid profile, and patient having deranged lipid value were included in the study i.e.

Serum Cholesterol >200 mg/dl Serum Triglycerides>150 mg/dl Serum LDL>100 mg/dl Serum VLDL>30 mg/dl Serum HDL<40 mg/dl.

# **Other Investigations**

- 1. Hb gm%
- 2. TLC (Total leucocyte count)
- 3. DLC (Differential leucocyte count)
- 4. ESR (Erythrocyte sedimentation rate)
- 5. FBS (Fasting blood sugar)

- 6. Blood urea
- 7. Serum creatinine
- 8. SGOT
- 9. SGPT
- 10. Urine- routine and microscopy

# **Selection of Drug**

Triphala Kwatha and Trikatu Capsules are the important prestigious formulations, which are successfully used from the ancient period. These formulations have been mentioned in various Ayurvedic classics like Chakradatta. Yoga Ratanakara, Sharangadhara Samhita and Vrinda Madhava. The reference of Triphala Kwatha with Madhuused in this trial was selected from Yoga Ratanakara for the management of Medoroga and Trikatu churna from Sushrut Sutrasthan. These drugs were prepared by Pharmacy experts of Charaka Rajkiya Ayurveda Pharmacy, Paprola, Dist Kangra H.P. as per G.M.P. norms.

## Table 2: Pharmacological Characters of Ingredients of Triphala Kwath [9]

S.No	Ingredients	Botanical Name	Family	Rasa	Guna	Veerya	Vipaka	Dosha Karma
1.	Haritaki	Terminalia chebula Retz.	Combretaceae	Pancharasa (Alavana Kashaya pradhana)	Laghu, Ruksha	Ushna	Madhura	Tridoshahar
2.	Bibhitaka	Terminalia bellirica (Gaertn.) Roxb.	Combretaceae	Kashaya	Ruksha, Laghu	Ushna	Madhura	Tridoshahar
3.	Amalaki	Emblica officinalis Gaertn.	Euphorbiaceae	Pancharasa (Alavana- Laghu, Amla Ruksha pradhana)		Sheeta	Madhura	Tridoshahar

	Table 5. That mactingical characters of high eulents of Trikutu capsules (										
S.No	Ingredients	Botanical Name	Family	Rasa	Guna	Veerya	Vipaka	Dosha Karma			
1.	Sonth	Zingiber officinale Roscoe	Zingiebraceae	Katu	Laghu Sanigdha	Ushna	Madhur	Kaphvata Shamak			
2.	Marich	Piper nigrum L.	Piperaceae	Katu	Laghu Sheeta	Ushna	Katu	Vatkapha Shamak			
3.	Pippali	Piper longum L.	Piperaceae	Katu	Laghu Sanigha Tikshna	Anushanas heeta	Madhur	Vatakapha Shamak			

Table 3: Pharmacological Characters of Ingredients of *Trikatu* Capsules<sup>[9]</sup>

# **OBSERVATION AND RESULTS**

Among 20 registered patients' maximum patients i.e., 55% were in the age group of 40-50 years followed by 45% patients were in the age group of 50-60 yrs. On gender wise distribution majority of patients were females i.e. 75% and 25% of the patients were male. Housewives were predominant group of study than others i.e. 65%. 20% patients were in Govt job; 5% were in private job and 15% patients were engaged in business.

Majority of the patients (70%) were used to consume mixed diet while rest i.e. 30% patients were taking vegetarian diet. Among 20 registered patients, maximum i.e. 65 % had excessive appetite, 35% had normal appetite. Out of 20 patients, 40% have *Mridu Koshtha*, 50% have *Madhyam Koshtha*, 10% have *Krura Koshtha*. Majority of the patients in both groups i.e. 80% had excessive sleep followed by 20% had sound and adequate sleep. Data showed that patients of *Vatakaphaj Prakriti* were maximum (75%), followed by *Pittakaphaj Prakriti* (25%). The effects of therapy on various subjective and objective parameters are given below from Table 4 to 8 and figure 1 & 2.

S.No	Symptoms	Group	Mean	Score	Sr%	SD <u>+</u>	SE <u>+</u>	'ť	ʻp'	Inter-group
		BT AT Change					Comparison			
1		Group-I	1.375	1	27%	0.518	0.183	2.049	0.080	<i>t</i> = 2.170
	Udara Stana	Group-II	1.4	0.5	64%	0.568	0.18	5.014	<0.001	<i>p</i> = 0.045
2	Kshudra	Group-I	1.375	1	27%	0.518	0.183	2.049	0.080	<i>t</i> =2.573
	Shwasa	Group-II	1.3	0.3	76%	0.667	0.211	4.743	0.001	<i>p</i> =0.020
3	Sandhi Shoola	Group-I	0.25	0.125	50%	0.354	0.125	1.000	0.351	<i>t</i> =-2.342
		Group-II	1.5	0.6	60%	0.568	0.18	5.014	<0.001	<i>p</i> =0.032
4	Kshudha Atimatra	Group-I	1.5	1.25	16.66%	0.463	0.164	1.528	0.170	t=2.399
	Atimutra	Group-II	1.3	0.6	53.84%	0.675	0.213	3.280	0.010	<i>p</i> =0.029
5	E · · ·	Group I	1.375	1.125	18.18%	0.463	0.164	1.528	0.170	<i>t</i> =2.412
	Atiyoga	Group II	1.3	0.4	69.23%	0.568	0.18	5.014	<0.001	<i>p</i> =0.028

**Table 4: Effect of Therapy on Subjective Symptoms** 

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Anjli Sharma et al. Study of Triphala Kwath and	Trikatu Capsules in the Manag	ement of Medoroga w.s.r to Dystibidemia

6	Nidradhikya	Group I	1.625	1.375	15.38%	0.463	0.164	1.528	0.170	<i>t</i> =2.762
		Group II	1.2	0.6	50%	0.699	0.221	2.714	0.024	<i>p</i> =0.014
7	Javoproda	Group-I	0.5	0.125	60%	0.744	0.263	1.426	0.197	<i>t</i> = -2.347
		Group-II	1.7	0.6	64%	0.568	0.18	6.128	< 0.001	<i>p</i> =0.032
8	Swedabadha	Group-I	0.875	0.625	28%	0.463	0.164	1.528	0.170	<i>t</i> = 2.160
		Group-II	0.8	0.1	87.50%	0.675	0.213	3.280	0.010	<i>p</i> =0.046
9	Daurgandhya	Group-I	1.375	1.125	18.18%	0.463	0.164	1.528	0.170	t=2.437
		Group-II	1.3	0.5	61.50%	0.789	0.249	3.207	0.011	<i>p</i> =0.027

# Table 5: Effect Of Therapy on BMI

S.No	Category	ry Group	Mean Score		%	SD±	SE±	ť	ʻp'	Inter-Group
			BT	AT	Change	3DT	JEI	L	р	Comparison
1	DMI	Group-I	31.125	29.75	4.41%	2.106	0.744	2.434	0.045	<i>t</i> =2.138
1.		Group-II	29.8	27.7	7.04%	1.951	0.617	6.034	<0.001	<i>p</i> =0.048

# Table 6: Effect of Therapy on Body Weight

S.No.	Category	Group	Mean Score		%	SD±	SE±	'ť'	(m)	Inter-Group	
			BT	AT	Change	3DT	2FI	L	'p'	Comparison	
1	Pody Woight	Group I	72.7	70.5	2.90%	2.37	0.838	4.968	0.002	<i>t</i> = 2.128	
	1. Body Weight	Group II	72.42	68.3	5.68%	2.063	0.652	6.88	<0.001	<i>p</i> =0.049	

# Table 7: Effect of Therapy on Skin Fold Thickness

C No	Categories	Group	Mean Score		%	SD+	CE .	'ť'	()	Inter group
S.No			BT	AT	Change	5D <u>+</u>	SE <u>+</u>	L	'p'	Comparison
1	Dicon	Group-I	2.95	2.68	9.1%	0.368	0.130	6.251	<0.001	<i>t</i> = 2.1
1	1 Bicep	Group-II	2.89	2.34	19%	0.344	0.108	8.199	<0.001	<i>p</i> = 0.047
2	Tricon	Group-I	3.2	2.987	9.3%	0.470	0.166	4.389	0.002	<i>t</i> =2.516
2	Tricep	Group-II	3.04	2.48	18.42%	0.401	0.127	11.225	<0.001	<i>p</i> = 0.023
3 Supraillia	Suprailliag	Group-I	5.66	4.9	13.4%	1.013	0.358	5.127	0.001	<i>t</i> =2.101
	Supraillac	Group-II	5.63	3.96	29.66%	0.885	0.279	13.19	<0.001	<i>p</i> =0.052

-											
S.No	Lipid Profile	Group	Mean	Score	%	SD <u>+</u>	SE <u>+</u>	'ť	'p'	Inter group	
3.100		aroup	BT	AT	Change	<u>50</u> <u>+</u>	3Ľ <u>+</u>	L	Ψ	Comparison	
1	Serum	Group-I	240.75	211.75	12.04%	36.269	12.823	2.262	0.058	<i>t</i> =10.9	
1	Cholesterol	Group-II	230.7	134.2	41.82%	31.43	9.94	9.708	<0.001	<i>p</i> <0.001	
2	Serum	Group-I	270.375	268.125	0.83%	28.085	9.93	0.227	0.827	<i>t</i> =18.2	
2	<sup>2</sup> Triglyceride	Group-II	296.875	143	51.60%	79	25.2	6.04	<0.001	<i>p</i> <0.001	
3	Serum LDL	Group-I	95.5	87.87	7.9%	30.56	10.807	0.706	0.503	<i>t</i> =5.36	
3	Sel ulli LDL	Group-II	128.4	48.5	62.22%	28.7	9.1	8.70	<0.001	<i>p</i> <0.001	
4	Somum HDI	Group-I	34.25	61.37	44.19%	7.7	2.7	-9.84	<0.001	<i>t</i> =10.7	
4	4 Serum HDL	Group-II	33.2	37.2	10.75%	7	2.2	1.7	0.107	<i>p</i> <0.001	
5	Serum VLDL	Group-I	38.25	36.25	5.2%	3.5	1.2	1.5	0.155	<i>t</i> =9.02	
Э	Seruii VLDL	Group - II	35.2	17.7	49.71%	6.819	2.156	8.1	<0.001	<i>p</i> <0.001	

## AYUSHDHARA, 2020;7(5):2864-2874 Table 8: Effect of Therapy on Serum Lipid Profile

Figure 1: Effect of Therapy on BMI, Body Weight and Skin Fold Thickness Effect of Therapy on BMI, Body Weight & Skin Fold Thickness

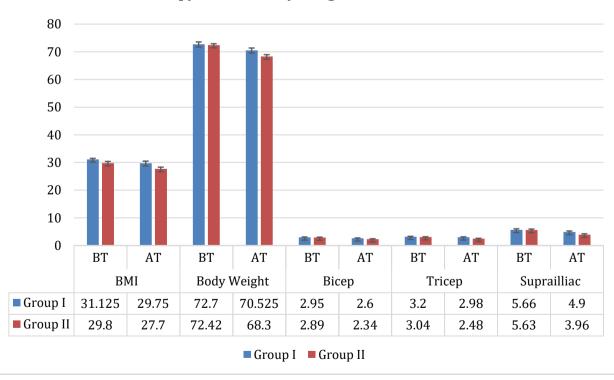


Figure1.\*BT (Before Treatment), AT (After Treatment)

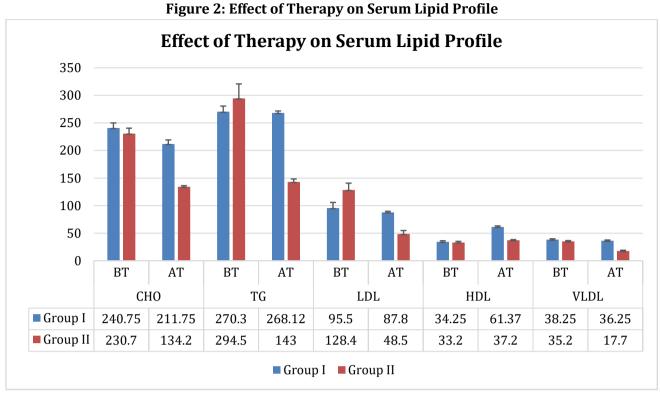


Figure2.\*BT (Before Treatment), AT (After Treatment), CHO (Cholesterol), TG (Triglycerides), LDL (Low Density Lipoproteins), HDL (High Density Lipoproteins), VLDL (Very Low-Density Lipoproteins) **DISCUSSION** 

In subjective parameters a highly significant (*P*<0.001) improvement was found in *Chala Sphika Udara Stana, Sandhi Shoola, Pipasa Ati Yoga, Pipasa Ati Yoga, Javoprodha* and significant improvement was found in *Kshudra Shwasa, Kshudha Atimatra, Nidradhikya, Swedabadha* and *Daurgandhya* in patients of Group II. While in Group I, insignificant improvement was found in the above mentioned subjective parameters as shown in Table 4.

There was a statistically highly significant decrease (p<0.001) in BMI by 7.04% in group II. In group in only statistically significant (p=0.045) improvement i.e., 4.41% was observed as shown in Table 5.

On intergroup comparison, statistically significant difference was found between the results of both the groups (p=0.048). There was a highly significant decrease (p<0.001) in weight by 5.68% in group II. In group I, statistically significant (p=0.002) improvement i.e., 2.90% was observed as shown in Table 6.

On intergroup comparison, a significant difference was found between the results of both the groups (p=0.049).

There was a highly significant decrease (p<0.001) in Skinfold Thickness of biceps by 19% in group II. In group I, only 9.1% decrease in Skinfold thickness of biceps was observed after the therapy

which was statistically significant (p<0.001). On intergroup comparison, a significant difference between the results of both the groups (p=0.047). There was highly significant decrease (p<0.001) in Skinfold thickness of Triceps by 18.42% in group II. In group I, only 9.3% decrease in Skinfold thickness of Triceps was observed after the therapy which significant (p=0.002). intergroup was On comparison, a significant difference was found between the results of both the groups (p=0.023). There was a highly significant decrease (p<0.001) in skinfold thickness of suprailiac by 29.66% in group II. In group I, only 13.4% decrease in skinfold thickness of suprailiac was observed after the therapy which was significant (p=0.001). On intergroup comparison, there was no significant difference between the results of both the groups (p=0.052) as shown in Table 7 and fig. no. 1.

#### **Serum Lipid Profile**

There was a highly significant decrease (p<0.001) in serum cholesterol by 41.82% in group II. In group I, only 12.04% decrease in serum cholesterol was observed after the therapy which was not significant (p=0.058). On intergroup comparison, a highly significant difference was found between the results of both the groups (p<0.001). There was a highly significant decrease (p<0.001) in serum triglycerides by 51.60% in group II. In group I only 0.83% decrease in serum

triglycerides was observed after the therapy which was not significant (p=0.827). On intergroup comparison, a highly significant difference was found between the results of both the groups (P<0.001). There was a highly significant decrease (p<0.001) in serum LDL by 62.22% in group II. In group I, only 7.9% decrease in serum LDL was observed after the therapy which was not statistically significant (p=0.503). On intergroup comparison, a highly significant difference was found between the results of both the groups (p<0.001). There was a significant increase (p=0.005) in serum HDL by 44.19% in group I. In group II, only 10.75% increase in serum HDL was observed after the therapy which was not significant (p=0.107). On intergroup comparison, a highly significant difference was found between the results of both the groups (p<0.001). There was a highly significant decrease (p<0.001) in serum VLDL by 49.71% in group II. Decrease by 5.2% in serum VLDL was observed in group 1 after the therapy which was not significant (p=0.155). On intergroup comparison, a significant difference was found between the results of both the groups (p<0.001). So, in group II, a highly significant (p<0.001) improvement was found in serum cholesterol, triglycerides, LDL, VLDL while in group I. a statistically highly significant (p < 0.001)improvement was observed in HDL only as shown in Table 8 and figure 2.

Insignificant results were found in the other parameters including TLC, DLC, ESR, FBS, Blood urea, Serum creatinine etc.

The Inter-group comparison showed that Triphala Kwath with Madhu and Trikatu capsules in combination are more effective in dyslipidemia as compare to only single drug i.e., *Trikatu* capsules. The difference was statistically significant. Hence the study proved the efficacy of Triphala Kwath with Madhu and Trikatu capsules in combination with the symptomatology of Medoroga (Dyslipidemia). No side effects of Triphala Kwath with Madhu and Trikatu Capsules were reported as evident by the fact that no patient complained of any untoward symptom after the administration of drug.

An ideal drug is the one that breaks the pathogenesis of the disease without producing any side effects. It is the total effect of all the ingredients that plays a vital role in the treatment of disease. In *Medoroga, Vata* and *Kapha Doshas* are involved along with *Medodhatvagni mandya* and *Srotorodha*. So, drugs used in the present study have *Vatakapha Shamak, Deepana, Pachna, Srotoshodhna,* and *Medohar* properties. Ingredients of present trial drugs i.e., *Triphala Kwath* and *Trikatu* capsules possess most of the above said properties which are desirable in an ideal *Medohara* drug. *Triphala Kwatha* contains drugs which are mainly *Laghu*, *Ruksha Guna*, *Ushna Virya* and *Tridoshhara*. Because of all these properties *Triphala* work as *Medohara*. *Trikatu* being *Katu Rasa* and *Katu Vipaka*, *Ushna Virya* and *Laghu Ruksha Guna* in nature just inverse to *Medodhatu* diminish the amount of *Medodhatu* in body. *Madhu* was used as *Anupaan* in the present study which potentiates the action of the drug. Because of *Kashaya Rasa*, *Guru*, *Ruksha Guna* it acts as *Tridoshshamak* and *Medoghana*.

In the present study, a significant decrease in the total cholesterol, triglycerides, LDL and VLDL in *Triphala Kwath* with *Madhu* treated patients may be due to reduction in the absorption of cholesterol<sup>[10]</sup>. Oral administration of *T. Chebula* is reported to increase gastric emptying, which might be the reason for the decrease in the cholesterol absorption.<sup>[11]</sup> Drugs like Statins reduce LDL cholesterol by inhibiting HMG-CoA reductase in the liver. In our study, we have observed a reduction in the LDL cholesterol after *Triphala Kwath* with *Madhu*administration. This may be due to, at lease in a part, the HMG-CoA reductase inhibitory action of *Triphala*<sup>[10]</sup>.

It is claimed that Amalaki, one of the components of *Triphala* has various biological activities such as improves digestion<sup>[12]</sup>, improves liver function and hepatoprotective<sup>[13]</sup>. *Triphala* has a balancing and rejuvenating effect on Vata, Pitta and Kapha. It is known that colon health is important. Proper regulation of the colon is a key to good health and longevity. Triphala works well as a colon cleanser and as a Rasayana (rejuvenation) for the colon. It is rich in fibers, which helps in and regulation digestion of bowel. Oral administration of Haritaki reported to increase gastric emptying, might be the reason of decreased absorption<sup>[11]</sup>. Most probably in the present study significant decrease total cholesterol, in Triglyceride and VLDL on the administration of Triphala Kwath may be due to reduction in absorption of cholesterol<sup>[14]</sup>, results in improved liver functions<sup>[13]</sup>. and decrease lipid peroxidation<sup>[15]</sup>.

*Madhu* has been used both as food and medicine since ancient times. *Madhu* is a natural product that has been widely used for its therapeutic effects. *Madhu* as *Anupana* was used in the present study in combination with *Triphala Kwath+Trikatu* Capsules. As mentioned in Ayurveda *Anupana* (vehicle) is defined as the *Pana* which is taken immediately after *Oushadhaanga* (part of medicine) and Oushadhi voaa (medicine formulation).<sup>[16]</sup> Anupana is defined as that, which enhances the properties of medicines along with it is taken. The mechanism by which Madhu exerts its effect as a natural product, may be due to its inhibitory effect on HMG-CoA reductase, which is very important in the production of cholesterol in the body, thereby effectively preventing hypercholesterolemia, lowering LDL, triglycerides and increasing HDL levels. These actions effectively reduce the risk of major coronary events, including first and second heart attacks and stroke. in adults with unhealthy cholesterol levels<sup>[17]</sup>.

Trikatu being of Katu, Tikta Rasa and *Katuvipaka* in nature<sup>[1]</sup>, i.e., just opposite of *Medodhatu*, reduce the quantity of *Medodhatu* and also make the channels patent for easy conduction of nutrients for nourishment to following Dhatus. Several popular preparations of *Trikatu* have been mentioned by the Ayurvedic physicians for the treatment of many Kaphavataja diseases, including Agnimandya (poor digestion) and Ama (undigested food and its toxic byproducts). By therapeutic actions such as *Deepana* (appetizer), *Pachana* (digestive), Rukshana (producing dryness), Lekhana (producing sliminess), Karshana (extraction), and Shoshana (absorption), Tikshna (sharp), Laghu (light) and Sukshma (micro in size)<sup>[1]</sup>, as a whole *Trikatu* reduces the quantity of *Meda* (which has the nature of Ama) and also makes the channels patent to carry on the nutrients to subsequent *Dhatus* as per the chronological order mentioned in Ayurveda. These pharmacological actions may be due to its chemical substance piperine which enhances the secretion of digestive juices and might catalyze the functions of enzymes in small intestine too, i.e. it helps in improving the function of Jatharagni (digestive fire). Improvement of Jatharagni function, in total, also helps in a finer disintegration of nutrients, in turn helping maximum absorption for nourishing rest of the *Dhatus*, and thus also facilitates the function of *Bhutagnis* (metabolism). In short, *Trikatu* acts against the deposition of lipids and thus helps clean the eventually blocked channels <sup>[18]</sup>. So, *Trikatu* Capsules probably acts by blocking the formation of different lipids or cholesterol at various stages in the biosynthetic pathway. Because it is a *Rasayana*, it may also cause an increase in the HDL cholesterol [18].

## CONCLUSION

It can be inferred from the present study that the best effects of the Trial drugs were seen with *Triphala Kwatha* with *Madhu* along with *Trikatu* Capsule together (Group II), which is most effective in reducing the overall lipid profile i.e., serum cholesterol, serum triglyceride, LDL, VLDL and increasing the HDL, with substantial gains related to subjective as well as objective parameters and that too, without any adverse effects. It can be concluded that *Triphala Kwath* with *Madhu* and *Trikatu* Capsules are effective in treating *Medoroga* (dyslipidemia) when given in combination.

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