

Clinical Trial Details (PDF Generation Date :- Sat, 03 Jun 2023 10:51:59 GMT)

CTRI Number Last Modified On Post Graduate Thesis

Type of Trial Type of Study

Study Design Public Title of Study

Scientific Title of Study

Secondary IDs if Any

CTRI/2020/05/024969 [Registered on: 01/05/2020] - Trial Registered Prospectively 28/09/2020 No Interventional Homeopathy Randomized, Parallel Group, Placebo Controlled Trial Homoeopathy as adjuvant in management of Covid-19 infection Effect of adjuvant homoeopathy with standard treatment protocol in management of covid-19: a

randomised, open label, placebo controlled, parallel group study Secondary ID Identifier

NIL

Details of Principal Investigator or overall **Trial Coordinator** (multi-center study)

Details of Principal Investigator		
Name	Prof Dr Pradeep Kumar Gupta	
Designation	Principal	
Affiliation	Naiminath Homoeopathic Medical College, Hospital and Research Centre	
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Details Contact Person (Scientific Query)

Details Contact Person (Scientific Query)		
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Details Contact Person (Public Query)

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Details Contact Person (Public Query)	
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Source of Monetary or Material Support

Source of Monetary or Material Support > Naiminath Homoeopathic Medical College, Hospital and Research Centre

Primary Sponsor

Primary Sponsor Details		
Name	Naiminath Homoeopathic Medical College Hospital and Research Centre	
Address	N.H.19, Firozabad Road, Nawalpur, Etmadpur, Agra, Uttar Pradesh – 283202	
Type of Sponsor	Private medical college	

Details of Secondary Sponsor

Name	Address
NIL	NIL

Countries of Recruitment

List of Countries India

Sites of Study

Name of Principal Investigator	Name of Site	Site Address	Phone/Fax/Email
Prof Dr Pradeep Kumar Gupta	and Hospital	Railway over bridge,	9765270885 nhmcagra@gmail.com

Details of Ethics Committee

Name of Committee	Approval Status	• •	Is Independent Ethics Committee?
Institutional Ethics	Approved	20/04/2020	Yes
Committee-F H Medical			
College			

Regulatory Clearance Status from DCGI

Status	Date
Not Applicable	No Date Specified

Health Condition / Problems Studied

Health Type	Condition
	Coronavirus as the cause of diseases classified elsewhere

Intervention / Comparator Agent

Туре	Name	Details
Intervention	Homoeopathic medicine	Homoeopathic medicines will be given to this group patients as an add-on to standard treatmet. Following Individualization medicines will be selected keeping in view pathological aspect of disease. Dose and potency will be according to the frequency, intensity and duration of signs and symptoms. Repetitions will vary from cae to case basis cosidering disease state and vitality of patient. Range of potency from 30, 200, 1M etc will be used.
Comparator Agent	Placebo	Placebo group patients will receive identical placebo



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			(globules moistened with dispensing alcohol) as an add on to standard protocol treatment. Dose repetition will be a in similar pattern to medicine group.	
Inclusion Criteria		Inclusion	n Criteria	
	Age From	18.00 Year(s)		
	Age To	80.00 Year(s)		
	Gender	Both		
		infection. o Bo	PCR confirmed cases of bth sexes or/> o Age between 18 years to 80 Willing to give signed written informed br/>	
Exclusion Criteria	Exclusion Criteria			
	Details 1. Patients requiring ventilatory support 2. Patients with compromised immunity 3. Severe heart, lung, kidney, brain, blood diseases or other important systemic diseases 4. Subjects unable to complete the study, or not suitable for the study by researchers. 5. Women during pregnancy 6. Lactating mothers		npromised immunity ng, kidney, brain, blood nportant systemic to complete the study, or study by regnancy	
Method of Generating Random Sequence Method of Concealment	Computer generated randomiz Not Applicable	ation		
Blinding/Masking	Open Label			
Primary Outcome	Outcome		Timepoints	
·	Clinical outcome in terms of re or requirement of life support (Ever 24 hr.	
Secondary Outcome	Outcome			
	Outcome		Timepoints	
	i. Time to progression to criticalii. Time to fever clearance iii. Time to resolution of pneum iv. Change in severity of sympolinical syndromes associated infection. v. Time to recovery in terms of Covid-19 vi. Period of hospital stay vii. Requirement of convention clinical improvement	nonia toms as per with COVID - 19 f 2 negative test for	Ever 24 hr.	
Target Sample Size	i. Time to progression to criticalii. Time to fever clearance iii. Time to resolution of pneumiv. Change in severity of sympolinical syndromes associated infection. v. Time to recovery in terms of Covid-19 vi. Period of hospital stay vii. Requirement of convention clinical improvement Total Sample Size=100 Sample Size from India=100 Final Enrollment numbers as	nonia toms as per with COVID - 19 f 2 negative test for hal medication till	Ever 24 hr.	
Target Sample Size Phase of Trial	i. Time to progression to criticalii. Time to fever clearance iii. Time to resolution of pneumiv. Change in severity of sympolinical syndromes associated infection. v. Time to recovery in terms of Covid-19 vi. Period of hospital stay vii. Requirement of convention clinical improvement Total Sample Size=100 Sample Size from India=100 Final Enrollment numbers as	nonia toms as per with COVID - 19 f 2 negative test for hal medication till	Ever 24 hr.	
	i. Time to progression to critical ii. Time to fever clearance iii. Time to resolution of pneum iv. Change in severity of symp clinical syndromes associated infection. v. Time to recovery in terms of Covid-19 vi. Period of hospital stay vii. Requirement of convention clinical improvement Total Sample Size=100 Sample Size from India=100 Final Enrollment numbers ac Final Enrollment numbers ac	nonia toms as per with COVID - 19 f 2 negative test for hal medication till	Ever 24 hr.	

CLINICAL TRIALS REGISTRY - INDIA

ICMR - National Institute of Medical Statistics



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Enrollment (Global)	
Estimated Duration of Trial	Years=0 Months=3 Days=0
Recruitment Status of Trial (Global)	Not Applicable
Recruitment Status of Trial (India)	Not Yet Recruiting
Publication Details	
Brief Summary	The results of the study will be able to predict the role of add on homoeopathic treatment for management of Covid-19 infected. In the absence any known anti-viral and vaccine the positive results of this study will decrease the morality and morbidity due to this pandemic. Further the cost-effectiveness of homoeopathy will enable developing countries in controlling such deadly disease effectively.