

Review Form 1.6

Journal Name:	Journal of Pharmaceutical Research International
Manuscript Number:	Ms_JPRI_77935
Title of the Manuscript:	EVALUATION OF EFFICACY OF NIGELLA SATIVA ON GINGIVAL HEALTH
Type of the Article	Original Research Article

General guideline for Peer Review process:

This journal's peer review policy states that **NO** manuscript should be rejected only on the basis of '**lack of Novelty**', provided the manuscript is scientifically robust and technically sound. To know the complete guideline for Peer Review process, reviewers are requested to visit this link:

(<http://peerreviewcentral.com/page/manuscript-withdrawal-policy>)

PART 1: Review Comments

	Reviewer's comment	Author's comment (if agreed with reviewer, correct the manuscript and highlight that part in the manuscript. It is mandatory that authors should write his/her feedback here)
Compulsory REVISION comments	<p><u>Under Title:</u> Concentration of <i>Nigella sativa</i> to be mentioned in title.</p> <p><u>Under study population:</u> a. Department of Periodontology should be there instead of Periodontics b. Instead of Saveetha Dental College and Hospitals, it should have been hospital. c. The ethical clearance certificate should be produced (Clinical trial registry number from DCGI)</p> <p><u>Under Methodology:</u> Clarify why oral prophylaxis was carried at baseline ? Consider making that as a separate group of study viz. there will be 3 study groups – Oral prophylaxis group, CHX group, Nigella sativa group. KINDLY CHANGE THE METHODOLOGY AS PER THIS & REFRAME THE STUDY TO MAKE ITS CLINICAL RELEVANCE</p> <ul style="list-style-type: none"> - Then compare results of all three modalities on gingivitis 	
Minor REVISION comments		
Optional/General comments		

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PART 2:

	Reviewer's comment	Author's comment <i>(if agreed with reviewer, correct the manuscript and highlight that part in the manuscript. It is mandatory that authors should write his/her feedback here)</i>
Are there ethical issues in this manuscript?	<i>(If yes, Kindly please write down the ethical issues here in details)</i> There has to be a clinical trial registry done under DCGI. Produce a relevant supporting document for the same.	

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