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**PARTICIPANT CONSENT FORM**

**Title: Clinical and microbiological evaluation of one-stage full mouth disinfection in conjunction with systemically administered azithromycin: a Randomised Controlled Clinical Trial**

*I...................................................................................................................................................have read the information provided and any questions I have asked have been answered to my satisfaction. I agree to participate in this activity, I acknowledge that I may withdraw at any time without reason and without prejudice.*

*I understand that all identifiable (attributable) information that I provide is treated as strictly confidential and will not be released by the investigator in any form that may identify me. The only exception to this principle of confidentiality is if documents are required by law.*

*I have been advised as to what data is being collected, the purpose for collecting the data, and what will be done with the data upon completion of the research.*

*I agree that research data gathered for the study may be published provided my name or other identifying information is not used.*

*\_\_\_\_\_\_\_\_\_\_\_\_\_\_   \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_  
 Participant                 Date*

***Approval to conduct this research has been provided by the University of Western Australia, in accordance with its ethics review and approval procedures. Any person considering participation in this research project, or agreeing to participate, may raise any questions or issues with the researchers at any time.***

***In addition, any person not satisfied with the response of researchers may raise ethics issues or concerns, and may make any complaints about this research project by contacting the Human Ethics Office at the University of Western Australia on (08) 6488 3703 or by emailing to***[***humanethics@uwa.edu.au***](mailto:humanethics@uwa.edu.au)

***All research participants are entitled to retain a copy of any Participant Information Form and/or Participant Consent Form relating to this research project.***