

Research identification

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|-------------|--|
| Study title | |
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|------------------------|--|
| GHK Reference No. | |
| Protocol No. | |
| Principal investigator | |

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|----------------------|-------------|
| Study start date | ___/___/___ |
| Anticipated end date | ___/___/___ |

Final Report

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|---------------------------|--|---------------------|----------------------------|
| Final Report sequence no. | | Report period | ___/___/___ to ___/___/___ |
| Planned sample size | | No. completed study | |
| No. recruited | | No. withdrew | |

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|---------------------|--|
| Withdrawal reasons: | |
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Summary of Serious Adverse Events (SAE)

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Does the SAE affect the study, and how?

Yes No

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Summary of Complaints from Subjects

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Study Duration

According to schedule

Extended

Premature termination

Summary of Study Outcomes

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Public disclosure of this Clinical Trial*

*Defined by World Health Organization (WHO) as any experimental study which prospectively allocates humans to a medical intervention. For reporting purposes, it includes non-randomised Phase I trials, randomised Phase I, II or III trials, and Phase IV or post-licensure trials of health products if they involve prospective designs (with or without randomisation).

Source: Statement on Public Disclosure of Clinical Trial Results <http://www.who.int/ictrp/results/reporting/en/>

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|---|---|
| Is this study a Clinical Trial? | <input type="checkbox"/> Yes (Proceed to answer the next question) <input type="checkbox"/> No (no need to answer the following questions) |
| This clinical trial is registered in a publicly available, free to access, searchable clinical trial registry | <input type="checkbox"/> Yes (Proceed to answer the next question) <input type="checkbox"/> No (no need to answer the following questions) |
| Name of clinical trial registry | |
| Registry identifier code/number for this clinical trial | |
| Main findings are to be submitted for publication in a peer reviewed journal within 12 months after study completion; OR made available publicly within 24 months of study completion | <input type="checkbox"/> Yes <input type="checkbox"/> No |
| Key outcomes are to be made available publicly within 12 months of study completion by posting to the results section of the primary clinical trial registry, or a free-to-access, publicly available and searchable institutional website where the registry used does not constitute a result database for key outcomes posting | <input type="checkbox"/> Yes <input type="checkbox"/> No |

Report by Principal Investigator

| Name | Signature | Date |
|------|-----------|------|
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