

QUAD

From the NAPWA website at <http://www.napwa.org.au/trials/quad>

Clinical trials information

QUAD

This trial is **no longer enrolling**. This may mean the trial is fully enrolled, or that enrolment in the trial has been halted or suspended. You cannot enrol in this trial.

About this trial

This study will compare the safety and [efficacy](#) (Of a drug or treatment). The maximum ability of a drug or treatment to produce a result regardless of dosage. A drug passes efficacy trials if it is effective at the dose tested and against the illness for which it is prescribed. In the standard procedure, Phase II clinical trials gauge efficacy, and Phase III trials confirm it. of a single tablet taken once daily - the Quad pill which contains fixed doses of elvitegravir (an [experimental](#) (Of a drug) Not licensed for use in humans, or as a treatment for a particular condition. Experimental drugs are studied in clinical trials to determine their safety and efficacy, and are sometimes made available via Special Access Schemes prior to their approval. integrase inhibitor) + emtricitabine and tenofovir (Truvada) + GS-9350 (an experimental boosting agent) - versus a regimen containing ritonavir-boosted atazanavir + Truvada.

Official title:

A Phase 3, Randomized, Double-Blind Study to Evaluate the Safety and Efficacy of Elvitegravir/Emtricitabine/Tenofovir Disoproxil Fumarate/GS-9350 Versus Ritonavir-Boosted Atazanavir Plus Emtricitabine/Tenofovir Disoproxil Fumarate in HIV-1 Infected, Antiretroviral Treatment-Naïve Adults

What is this trial studying?

new_drug – Truvada, elvitegravir (GS-9137), cobicistat (GS-9350)

Start date:

April 2010 (This may be the proposed or expected start date for trials which have not yet started.)

How many participants will this trial enrol?

700 (The exact number of participants may be lower or slightly higher than this. Some trials also have specific quotas for participants from each state, city or clinic.)

How long is this trial planned to go for?

Participants in this trial will be asked to take the trial drugs for 48 weeks. Following the conclusion of the trial, participants will be followed up for a further 48 weeks.

Links to further information:

- http://clinicaltrials.gov/ct2/show/study/NCT01106586?term=HIV+and+elvitegravir&rank=6&show_loc=Y#locn

Who can enrol in this trial?

You *may* be eligible to participate in this trial if you meet the following criteria:

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At least 18 years old
Have never taken HIV treatments
Viral load at least 5000 copies/ml

This is a summary of key inclusion and exclusion criteria for this trial. There may be other criteria which may exclude some people from participation in this trial. Some laboratory tests may also be required. Consult your doctor, or view the trial protocol or informed consent documentation to see the full range of exclusion and inclusion criteria.

Disclaimer

While NAPWA has taken every care to compile the information on this page and to keep it up-to-date, we cannot guarantee its correctness and completeness.

Before making the decision to participate in any clinical research, visit the NAPWA website for background information on participating in clinical research.

Contact NAPWA if you have any questions or comments about this trial.