



PANHPVAX study – vaccination against human papillomaviruses **Healthy volunteers wanted**

The clinical phase I study is assessing the safety, tolerability and immunogenicity of a novel vaccine (PANHPVAX antigen with cyclic di-AMP as booster) against human papillomaviruses in healthy volunteers.

Study period 7 to 8 months

15 visits at the study center: Time commitment

> 3 x all day for vaccination, approx. 8 hrs each 1x screening and 1x follow-up, approx. 1.5 hrs each

10 x check-ups, approx. 30 min. each

Times indicated are approximate. The study days will be

planned flexibly.

Total of three vaccinations with the novel vaccine PANHPVAX Study procedure

Control checks (blood samples, physical examinations)

Participants Healthy men and women

Age 18 to 45 years

No regular use of medication (except for oral contraceptives/ Requirements

"the pill", thyroid hormones)

Reliable contraception methods used

Not yet vaccinated against human papillomaviruses (HPV) No known chronic HPV infection (e.g. chronic genital warts)

Compensation EUR 2,000

Principal investigator/ PD Dr. med. Antje Blank

lead physician

Heidelberg University Hospital | Medical Clinic

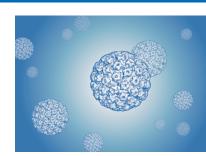
Dept. of Clinical Pharmacology and Pharmacoepidemiology

Early Clinical Trial Unit Pharmacology (KliPS) Im Neuenheimer Feld 410 | 69120 Heidelberg

Are you interested? Then call in for a face-to-face chat!

To arrange an appointment, please contact the Early Clinical Trial Unit Pharmacology (KliPS) of Heidelberg University Hospital:

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