

Testing RG1-VLP Vaccine to Prevent HPV-related Cancers

NCT05985681

Status	RECRUITING
Phase	Phase 1
Sponsor	National Cancer Institute (NCI)
Enrollment	18 participants

Key Eligibility Criteria

Inclusion (18)

- Women, age 18 - 60 years. Because no dosing or adverse event (AE) data is currently available for the use of RG1-VLP in humans, children and adolescents are excluded from this study
 - White blood cell (WBC) between 3000/mm³ - institutional upper limit of normal
 - Hemoglobin (Hgb) between 10 g/dl - institutional upper limit of normal
 - Platelets \geq 100,000/mm³
 - Serum creatinine within institutional normal limits
- ... and 13 more (see full listing online)

Exclusion (21)

- History of any of the following:
 - Prior or current genital warts
 - Treatment for anogenital intraepithelial neoplasia (cervical intraepithelial neoplasia [CIN], anal intraepithelial neoplasia [AIN], vaginal intraepithelial neoplasia [VAIN], vulvar intraepithelial neoplasia [VIN])
 - Systemic cancer treatment within the prior year
 - History of anaphylaxis to vaccines
- ... and 16 more (see full listing online)

Locations (5 total)

University of Alabama at Birmingham Cancer Center, Birmingham, Alabama, United States
Johns Hopkins University/Sidney Kimmel Cancer Center, Baltimore, Maryland, United States
Staten Island University Hospital, Staten Island, New York, United States
... and 2 more locations