A Phase 1/2 Study of EG-70 as an Intravesical Administration to Patients with BCG-Unresponsive NMIBC

Status: Recruiting

Eligibility Criteria

Sex: Male or Female
Age Group: 18 years and over
This study is NOT accepting healthy volunteers

Inclusion Criteria:

- BCG-unresponsive NMIBC with carcinoma in situ (CIS - ineligible for or have elected not to undergo cystectomy - women of childbearing potential must be willing to use highly effective birth control methods; Males are required to use a condom for the duration of the study treatment through 3 months post-dose - at least walking and capable of all selfcare but unable to carry out any work activities; up and about more than 50% of waking hours - see link to clinicaltrials.gov for complete inclusion criteria

Exclusion Criteria:

- other active cancer that required treatment in the past two years - history of partial cystectomy - history of severe asthma or other respiratory diseases - human immunodeficiency virus, Hepatitis B, or Hepatitis C infection - see link to clinicaltrials.gov for complete exclusion criteria

Conditions & Interventions

Conditions:
Cancer
Keywords:
Bladder Cancer With Carcinoma in Situ, CIS, Clinics and Surgery Center (CSC), NMIBC

More Information

Description: This study will evaluate the safety and tolerability of EG 70, a gene therapy, which is given inside the bladder. This study will measure its effectiveness on eliminating bladder tumors in participants with NMIBC who have failed or did not receive adequate Bacillus Calmette- Guérin (BCG) therapy. The study drug will be given into the lining of the bladder. This may cause an immune response inside the bladder and kill the cancer cells.
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