Vaccibody is a leader in the rapidly developing field of individualized cancer neoantigen vaccines and is using the Vaccibody technology to generate best-in-class therapeutics to treat cancers with a high unmet medical need. A phase I/IIa neoantigen clinical trial is now enrolling patients with locally advanced or metastatic melanoma, non-small cell lung carcinoma, clear renal cell carcinoma as well as urothelial or squamous cell carcinoma of head and neck.

Vaccibody has a collaboration with Nektar Therapeutics, planning to start testing VB10.NEO in combination with bempegaldesleukin (NKTR-214) in squamous cell carcinoma of head and neck in H2 2019. Vaccibody’s front runner program (VB10.16) is a therapeutic DNA vaccine against HPV16 induced pre-malignancies and malignancies.

The first-in-human study (phase I/IIa), evaluating the safety and immunogenicity of VB10.16 in women with high grade cervical intraepithelial neoplasia (HSIL; CIN 2/3) has published positive 12 months data. Vaccibody has recently started a collaboration with Roche, exploring VB10.16 in combination with their checkpoint inhibitor atezolizumab (Tecentriq™) in patients with advanced or recurrent cervical cancer. First patient is expected to be vaccinated in Q1 2020.

Further information about the company and its drug development programs and capabilities may be found online at [http://www.vaccibody.com](http://www.vaccibody.com/)