Successful patient enrolment in HAMLET BioPharma's Phase III bladder cancer trial

HAMLET BioPharma is proud to announce that the clinical phase of the Phase III trial has started. Patients with non-muscle invasive bladder cancer are being enrolled and treated according to the study protocol, which is approved by the European and US authorities, EMA and FDA.

This remarkable milestone is reached shortly after the company communicated the successful approval process with the EMA at the end of June, the successful production of Alpha1H drug at Phase III quality and transport to the trial site and the trial site initiation with the clinical team at Motol hospital in Prague, led by Professor Babjuk and his team.

In parallel, the company is moving to start a separate clinical study in patients with severe, therapy resistant cancer *in situ*, at the Department of Urology, University of Iowa.

" Everyone in our extended organization is proud and excited to start the clinical part of the Phase III study. We are extremely grateful for the commitment of the Prague team of bladder cancer experts and the regulatory support by InClino" says Professor Catharina Svanborg, Chairman of the Board.

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