TILLOMED Laboratories Ltd.

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Dear Prescriber,

Welcome to the Cidofovir Exposure Study (**Study**). The Study is sponsored by Tillomed Laboratories Limited (**Tillomed UK**) and its sister companies Tillomed Pharma GmbH and Laboratorios Tillomed Spain SLU (**Sponsors**). It will be conducted by Tillomed UK on behalf of the Sponsors in accordance with a requirement of the European Medicines Agency (**EMA**).

The Study is a prospective, non-interventional, long term exposure, registry study. Its aim is to collect safety data about patients prescribed Tillomed's Cidofovir formulation, Cidofovir Tillomed 75 mg/ml Concentrate for Solution for Infusion (**Cidofovir**) on an observational basis without affecting your patients' standard treatment and care.

The information collected in the registry will be used to understand the medical condition for which you prescribe Cidofovir and the pattern of side effects arising from its use.

Please refer to the synopsis overleaf that provides brief details of the Cidofovir Exposure Study.

Yours sincerely,

Tillomed Laboratories Ltd