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AcuCort receives renewed marketing authorisation for Zeqmelit® in Norway

The pharmaceutical company AcuCort has received a renewed marketing authorisation for its medicinal product Zeqmelit® from the Norwegian Medical Products Agency (Direktoratet for medisinske produkter), Norway's equivalent of the Swedish Medical Products Agency. The decision grants approval without a time limitation.

AcuCort was first granted a marketing authorisation for Zeqmelit® in Norway in 2022. That approval was time-limited, which is standard for new medicines. Following an application for renewal, the Norwegian Medical Products Agency has now granted an extended marketing authorisation for the Norwegian market, this time without a time restriction.

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About AcuCort AB (publ)

AcuCort has developed and is commercializing Zeqmelit®, a new rapidly dissolving oral film placed on the tongue, based on the well-known cortisone substance dexamethasone. The drug is a smart product in a new, innovative, patented, and user-friendly administration form primarily for the treatment of severe and acute allergic reactions, croup in children, nausea and vomiting during chemotherapy, and for the treatment of patients with COVID-19 requiring supplemental oxygen therapy. Zeqmelit® is approved in Sweden, Denmark, Norway, and Finland. AcuCort (ticker: ACUC) is listed on the Spotlight Stock Market. Visit www.acucort.se for more information.