

Eli Lilly collaborates with Daiichi Sankyo for commercialisation of migraine drug in Japan

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Eli Lilly Japan will hold a marketing authorization for galcanezumab and Daiichi Sankyo will take charge of its distribution and sales after marketing approval



Eli Lilly Japan K.K. and Daiichi Sankyo Company have concluded an agreement on commercialization collaboration in Japan for galcanezumab (generic name: galcanezumab (genetical recombination)), for which Eli Lilly Japan has submitted a new drug application for the treatment of migraine.

Galcanezumab is a monoclonal antibody with a new mode of action. It binds specifically to calcitonin gene-related peptide (CGRP), which is considered to be associated with migraine, and thereby inhibits binding of CGRP to its receptor. This drug was approved for the prevention of migraine in September 2018 in the United States and is currently used for patients with migraine in 20 countries across the world.

Under the terms of the agreement, Eli Lilly Japan will hold a marketing authorization for galcanezumab and Daiichi Sankyo will take charge of its distribution and sales after marketing approval. Both companies will provide healthcare professionals with information on the drug.

Through the commercialization collaboration with Daiichi Sankyo, which has a solid operating foundation in Japan based on the established position and global network in the fields of central nervous system diseases and pain management, Eli Lilly Japan will strengthen its system to provide healthcare professionals with proper information on galcanezumab after marketing approval.

Daiichi Sankyo will make efforts to improve the quality of life of many more patients with migraine by utilizing its operating foundation in various fields, including the field of central nervous system diseases in Japan, through the partnership with Eli Lilly Japan.

Eli Lilly Japan and Daiichi Sankyo will closely work together through the commercialization collaboration for galcanezumab to

