RAVIMIAMET

F. Hoffmann-La Roche AG
Grenzacherstrasse 124
4058 Basel
SWITZERLAND

18.07.2024 nr RKU-4/48

DECISION

to grant authorisation for clinical trial on the basis of Regulation (EU) No 536/2014 of the European Parliament and of the council

State Agency of Medicines has received the application from sponsor F. Hoffmann-La Roche AG on 05.06.2024 to conduct a clinical trial under the conditions stipulated in Regulation (EU) No 536/2014 art 5 (1) and § 99^1 (1) of Estonian Medicinal Products Act (MPA).

Based on art 8 of Regulation (EU) No 536/2014, considering the aspects covered by Part I and Part II of the assessment report, on the basis of § 99^6 section 1 p 1 and (3) of Medicinal Product Act

State Agency of Medicines has decided to give the approval to conduct the clinical study protocol no MN43964 under the following conditions:

protocol no: MN43964 (Version 2, dated August 29, 2023)

full title of the trial: A Multicenter, Single-arm, Open-label, Extension, Rollover Study To Evaluate The Long-term Safety And Efficacy Of Ocrelizumab In Patients With Multiple Sclerosis

sponsor of the trial: F. Hoffmann-La Roche AG

number of subjects in Estonia: 7

principal investigators and study locations:
- Dr Katrin Gross-Paju, Lääne-Tallinna Keskhaigla AS, Paldiski mnt 68, 10617 Tallinn, Estonia
- Dr Sulev Haldre, Tartu University Hospital, L. Puusepa tn 8, 50406 Tartu, Estonia

The addressee may file a challenge with State Agency of Medicines within 30 days as of the day when the addressee became or should have become aware of the decision as prescribed in Code of Administrative Procedure § 71(1). If the addressee wishes to have the decision to annulled by the administrative court, he or she may submit a complaint to the Tartu Administrative Court as prescribed in § 7(1) and § 46(1) of the Code of Administrative Court Procedure within 30 days after the date on which the decision was notified to the applicant.

(digitally signed)

Ott Laius
Deputy Director General
on duties of Director General