EUROPEAN MEDICINES AGENCY
SCIENCE MEDICINES HEALTH

Curriculum Vitae

PERSONAL INFORMATION

Sigurros Sigmarsdottir

WORK EXPERIENCE

November 2020- Present

Assessor

Icelandic Medicines Agency (Iceland)

Assessment of bioequivalence studies and writing of relevant assessment reports. Involvement in scientific advice procedures.

October 2018-June 2020

Regulatory Affairs Associate, Regulatory Affairs

Teva Pharmaceutical (Iceland)

Providing clients from all over the world with regulatory documents to maintain their marketing authorisation for products they sourced from Teva Pharmaceutical via Medis.

July 2017-September 2018

Regulatory Affairs Associate, Regulatory Affairs

Teva Pharmaceutical (Iceland)

Providing dossiers and other relevant regulatory documents to Teva´s local regulatory contacts within growth markets for submissions.

February 2014-June 2017

Regulatory Affairs Scientis/Manager, Clinical R&D

Teva Pharmaceutical/Actavis Group PTC (Iceland)

Delegating and responding to clinical deficiency letters received from agencies within EU for Teva´s bioequivalence studies.

September 2006-January 2014

Scientist, Clinical R&D

Actavis Group PTC (Iceland)

Planning, designing and handling bioequivalence studies to support the registration of Actavis products within EU, US, Australian and other markets. Maintaining and completing Trial master files.

June 2005-August 2006

Assistant Pharmacist/Pharmacist

Lyfjalausnir (Iceland)

Pharmacist at a computerised drug dispensing company.

EDUCATION AND TRAINING

September 2001-May 2006

M.Sc. Pharmacy

University of Iceland (Iceland)

ADDITIONAL INFORMATION

Expertise

Publications

Projects

Memberships

Member of the Pharmaceutical Society of Iceland.

Other Relevant Information