Clinical trial information

Registration Item	Details of registration Item	Contents
Title of the study	Title of the study	Phase I, Multiple Oral Dose Study to Evaluate the Tolerability and Pharmacokinetics of Favipiravir in Healthy Volunteers
	Primary sponsor	Toyama Chemical Co., Ltd
	Study Type	interventional (treatment)
	Summary	The purpose of this study is to evaluate the safety, tolerability, and pharmacokinetics of multiple oral doses of favipiravir in healthy male volunteers.
Details of study	Interventional drug name	favipiravir
	Target illness	Healthy Volunteers
	Classification name (code) of the investigational drug	625：antiviral
	Administration route	oral
	Control drug name (code/td)	Placebo
	Classification name (code) of the investigational drug	-
	Administration route	oral
	Objectives of the study	Pharmacokinetics, safety, and tolerability
	Study phase	Phase I study
	Study design	Randomized, double-blind, placebo-controlled
	Inclusion Criteria	・Healthy adult male volunteers ・Aged between 20 and 39 years (inclusive)
	Outcome	Safety, pharmacokinetics
	Study status	Completed
	Duration of the study	Study Completed
	Region	Japan
Contact information	Organization	Toyama Chemical Co., Ltd.
	Division	Data Science and Administration Department
	Contact	https://www.toyama-chemical.co.jp/en/form/general/input.php?id=TCClinicalEn