



## Main Information

**Primary registry identifying number**

LBCTR2019100038

**Protocol number**

asdasdasdsadsd

**MOH registration number**

24234234234234

**Study registered at the country of origin**

No

**Study registered at the country of origin: Specify**

**Type of registration**

**Type of registration: Justify**

N/A

**Date of registration in national regulatory agency**

**Primary sponsor**

**Primary sponsor: Country of origin**

**Date of registration in primary registry**

14/10/2019

**Date of registration in national regulatory agency**

**Public title**

**Acronym**

**Scientific title**

**Acronym**

**Brief summary of the study: English**

**Brief summary of the study: Arabic**

**Health conditions/problem studied: Specify**

N/A

**Interventions: Specify**

N/A

**Key inclusion and exclusion criteria: Inclusion criteria**

**Key inclusion and exclusion criteria: Gender**

**Key inclusion and exclusion criteria: Specify gender**

**Key inclusion and exclusion criteria: Age minimum**

**Key inclusion and exclusion criteria: Age maximum**

**Key inclusion and exclusion criteria: Exclusion criteria**



**Type of study**

**Type of intervention**

N/A

**Type of intervention: Specify type**

N/A

**Trial scope**

N/A

**Trial scope: Specify scope**

N/A

**Study design: Allocation**

N/A

**Study design: Masking**

N/A

**Study design: Control**

N/A

**Study phase**

N/A

**Study design: Purpose**

N/A

**Study design: Specify purpose**

N/A

**Study design: Assignment**

N/A

**Study design: Specify assignment**

N/A

**IMP has market authorization**

**IMP has market authorization: Specify**

**Name of IMP**

**Year of authorization**

**Month of authorization**

**Type of IMP**

**Pharmaceutical class**

**Therapeutic indication**

**Therapeutic benefit**

**Study model**

N/A

**Study model: Explain model**

N/A

**Study model: Specify model**

N/A

**Time perspective**

N/A

**Time perspective: Explain time perspective**

N/A

**Time perspective: Specify perspective**

N/A



<b>Target follow-up duration</b>	<b>Target follow-up duration: Unit</b>
<b>Number of groups/cohorts</b>	
<b>Biospecimen retention</b> N/A	<b>Biospecimen description</b> N/A
<b>Target sample size</b>	<b>Actual enrollment target size</b>
<b>Date of first enrollment: Type</b>	<b>Date of first enrollment: Date</b>
<b>Date of study closure: Type</b>	<b>Date of study closure: Date</b>
<b>Recruitment status</b>	<b>Recruitment status: Specify</b>
<b>Date of completion</b>	
<b>IPD sharing statement plan</b>	<b>IPD sharing statement description</b>
<b>Additional data URL</b>	
<b>Admin comments</b>	
<b>Trial status</b> Approved	

## Secondary Identifying Numbers

No Numbers



## Sources of Monetary or Material Support

No Sources

## Secondary Sponsors

No Sponsors

## Contact for Public/Scientific Queries

No Contacts

## Centers/Hospitals Involved in the Study

No Centers/Hospitals

## Ethics Review

No Reviews

## Countries of Recruitment

No Countries

## Health Conditions or Problems Studied

No Problems Studied



## Interventions

No Interventions

## Primary Outcomes

No Outcomes

## Key Secondary Outcomes

Name	Time Points	Measure
aSas	sS	ASDdD



## Trial Results

**Summary results**

**Study results globally**

**Date of posting of results summaries**

**Date of first journal publication of results**

**Results URL link**

**Baseline characteristics**

**Participant flow**

**Adverse events**

**Outcome measures**

**URL to protocol files**