

Rasha 123

11/08/2025 20:57:27

/lai				

Primary registry identifying number

LBCTR2018090151

MOH registration number

20555/2018

Study registered at the country of origin

Yes

Type of registration

Prospective

Date of registration in national regulatory agency

15/09/2018

Primary sponsor

Rasha

Date of registration in primary registry

22/04/2020

Public title

Rasha 123

Scientific title

Rasha 123

Brief summary of the study: English

Great Job

Brief summary of the study: Arabic

السجل الوطني للابحاث السريرية الاول من نوعه

Health conditions/problem studied: Specify

Heart Failure

Interventions: Specify

New medication

Key inclusion and exclusion criteria: Inclusion criteria

all adults

Key inclusion and exclusion criteria: Gender

Both

Key inclusion and exclusion criteria: Age minimum

Protocol number

Rasha123

Study registered at the country of origin: Specify

Type of registration: Justify

N/A

Primary sponsor: Country of origin

Lebanon

Date of registration in national regulatory agency

15/09/2018

Acronym

RH

Acronym

RH

Key inclusion and exclusion criteria: Specify gender

Key inclusion and exclusion criteria: Age maximum

75

Key inclusion and exclusion criteria: Exclusion criteria

less than 18

Type of study

Observational

Type of intervention

Pharmaceutical

Trial scope

Therapy

Study design: Allocation

Non-randomized controlled trial

Study design: Control

Uncontrolled

Study design: Purpose

Treatment

Study design: Assignment

Other

IMP has market authorization

No

Name of IMP

In Love

Type of IMP

Others

Pharmaceutical class

heart diseases

Therapeutic indication

to treat heart failure

Therapeutic benefit

increase ejection fraction

Study model Case-Control

Study model: Specify model

N/A

Time perspective

Prospective

Time perspective: Specify perspective

N/A

Type of intervention: Specify type

pharmaceutical with behavioral

Trial scope: Specify scope

N/A

Study design: Masking

N/A

Study phase

1 to 2

Study design: Specify purpose

Study design: Specify assignment

single arm

IMP has market authorization: Specify

USA

Year of authorization Month of authorization

2014

Study model: Explain model

OK

Time perspective: Explain time perspective



Lebanon Clinical Trials Registry

OK

Months

Target follow-up duration

30

Number of groups/cohorts

4

Biospecimen retention

Samples without DNA

Biospecimen description

Target follow-up duration: Unit

DNA saliva samples

Target sample size

1000

Date of first enrollment: Type

Anticipated

Date of study closure: Type

Anticipated

Recruitment status

Complete

Date of completion

24/04/2020

IPD sharing statement plan

No

Actual enrollment target size

1

Date of first enrollment: Date

20/10/2018

Date of study closure: Date

20/10/2020

Recruitment status: Specify

IPD sharing statement description

I do not want to share

Additional data URL

later

Admin comments

Trial status

Approved

Secondary Identifying Numbers				
Full name of issuing authority	Secondary identifying number			
KSA	25325			
NO one	55555			

Sources of Monetary or Material Support

Name

King faisal hospital

Secondary Sponsors

Name

NA

Contact for Public/Scientific Queries

Contact type	Contact full name	Address	Country	Telephone	Email	Affiliation
Public	Rasha Hamra	Beirut	Iceland	0255.0000	rasha@gmail	МОН
Scientific	Marie Ok	Beirut	Lebanon	565540000 0	marie@yahoo	МОН
Public	NN	Beirut	Belize	5555	r@yahoo	МОН

Centers/Hospitals Involved in the Study Principles investigator Center/Hospital name Name of principles investigator Ethical approval speciality AUB RHOncologist Approved RH USJ Oncologist NA



Ethics Review						
Ethics approval obtained	Approval date	Contact name	Contact email	Contact phone		
American University of Beirut Medical Center King faisal hospital	11/10/2018	RH	@gmail	565458		
Makassed General Hospital	11/12/2019	RH	@yahoo	3336661		

Countries of Recruitment Name Lebanon

Health Conditions or Problems Studied				
Condition	Code	Keyword		
heart failure	Heart failure (I50)	RF		

Interventions					
Intervention	Description	Keyword			
pharmaceutical and behavioral	new medication with lifestyle changes	RH and forever			
New treatment	New	RH			

Primary Outcomes				
Name	Time Points	Measure		
increast EF	in 6 months	EF		

Key Secondary Outcomes				
Name	Time Points	Measure		
I dont have	later	Ok		



Lebanon Clinical Trials Registry

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	