

Clinical Trial Protocol

Iranian Registry of Clinical Trials

10 Jun 2026

Autologous Serum Skin Test as a Diagnostic Aid in Chronic Idiopathic Urticaria

Protocol summary

Summary

Background: Chronic urticaria is defined as urticaria persisting daily for more than six weeks, a significant number of them had autoimmune basis where autologous serum skin test is widely used for detection of chronic autoimmune urticaria. Objectives: To estimate the frequency of autoimmune urticarial in Iraqi patients utilizing the autologous serum skin test and to evaluate its results with the variable clinical features of chronic idiopathic urticaria. Methods: In this prospective study, 54 patients with chronic idiopathic urticaria will be investigated with autologous serum skin test where its results were examined with the different clinical parameters of chronic autoimmune urticaria.

General information

Acronym

IRCT registration information

IRCT registration number: **IRCT2013030912758N1**

Registration date: **2013-03-25, 1392/01/05**

Registration timing: **registered_while_recruiting**

Last update:

Update count: **0**

Registration date

2013-03-25, 1392/01/05

Registrant information

Name

Amar Hameed

Name of organization / entity

Baghdad medical college

Country

Iraq

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Recruitment status

Recruitment complete

Funding source

Iraqi Board for Medical specializations

Expected recruitment start date

2009-11-01, 1388/08/10

Expected recruitment end date

2013-11-01, 1392/08/10

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

Scientific title

Autologous Serum Skin Test as a Diagnostic Aid in Chronic Idiopathic Urticaria

Public title

Investigation for chronic Idiopathic Urticaria

Purpose

Diagnostic

Inclusion/Exclusion criteria

Inclusion criteria: All patients with chronic idiopathic urticaria who have active urticaria at time of presentation and the current attack started at least six week before the test. Exclusion criteria: children younger than 15 years of age and pregnant women were excluded from the study

Age

No age limit

Gender

Both

Phase

N/A

Groups that have been masked

No information

Sample size

Target sample size: 54

Randomization (investigator's opinion)

Not randomized

Randomization description

Blinding (investigator's opinion)

Not blinded

Blinding description

Placebo

Used

Assignment

Parallel

Other design features

Secondary Ids

empty

Ethics committees

1

Ethics committee

Name of ethics committee

Iraqi Board for Medical specializations

Street address

Medical Collection Office, P.O. Box 61080 Postal Code 12114, Baghdad, Iraq.

City

Baghdad

Postal code

12114

Approval date

2009-10-09, 1388/07/17

Ethics committee reference number

1976

Health conditions studied

1

Description of health condition studied

urticaria

ICD-10 code

L50.1

ICD-10 code description

Idiopathic urticaria

Primary outcomes

1

Description

wheal and flare

Timepoint

30 minutes after intradermal injection of 0.05 ml of the patient's serum into patient's forearm

Method of measurement

wheal and flare at the tested site of a diameter (measured by taking the mean of the largest two diameters) at least 1.5 millimeters more than the wheal and flare induced at the control site

Secondary outcomes

1

Description

wheal and flare

Timepoint

45 minutes after intradermal injection of the patient's serum into patient's forearm

Method of measurement

wheal and flare at the tested site of a diameter (measured by taking the mean of the largest two diameters) at least 1.5 millimeters more than the wheal and flare induced at the control site

Intervention groups

1

Description

The Autologous Serum Skin Test (ASST) was performed by drawing 2cubic centimeters of venous blood and stored in a plane tube(not heparinized) and allowed to clot at room temperature ,after that the serum is separated by centrifuging the blood in a centrifuge for 10 minutes at 2000 RPM (Revolution per Minute). 0.05 ml of the patient's serum is injected intradermally on volar aspect of the patient's forearm at site not affected by a wheal in the last 24 hours

Category

Diagnosis

2

Description

At the same time a control test is done by injecting the same volume of normal saline at least 5 centimeters away from the serum injection site

Category

Diagnosis

Recruitment centers

1

Recruitment center

Name of recruitment center

Baghdad Teaching Hospital

Full name of responsible person

Dr.Ammar F.Hameed

Street address

Medical city 61106

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Sponsors / Funding sources

1

Sponsor

Name of organization / entity

Iraqi Board for Medical specializations
Full name of responsible person
prof.Khalifa sharquie
Street address
Medical Collection Office, P.O. Box 61080
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Grant name
Grant code / Reference number
Is the source of funding the same sponsor organization/entity?
Yes
Title of funding source
Iraqi Board for Medical specializations
Proportion provided by this source
100
Public or private sector
empty
Domestic or foreign origin
empty
Category of foreign source of funding
empty
Country of origin
Type of organization providing the funding
empty

Person responsible for general inquiries

Contact

Name of organization / entity
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Person responsible for updating data

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Sharing plan

Deidentified Individual Participant Data Set (IPD)
empty
Study Protocol
empty
Statistical Analysis Plan
empty
Informed Consent Form
empty
Clinical Study Report
empty
Analytic Code
empty
Data Dictionary
empty