

***KLİNİK ÇALIŞMALARIN
clinicaltrials.gov kaydı ve
NCT numarası alınması***

Prof.Dr. Ener Çağrı DİNLEYİCİ

- Klinik alıřmaların, alıřma bařlangıcında kaydedildiđi bir bildirim sistemidir.

Bir ok dergi son yıllarda NCT numarası olmayan alıřmaları deđerlendirmeye almamaktadır.

clinicaltrials.gov, Amerika Birleřik Devletleri kaynaklı bir alıřma protokol kayıt sistemi olup, zellikle ila ve giriřimsel alıřmaların kayıtları aısından nemli bir veri tabanıdır.

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ClinicalTrials.gov is getting an update. Our new design arrives on June 19th. [Learn more.](#)

Preview our new design at ClinicalTrials.gov/beta/

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ClinicalTrials.gov

ClinicalTrials.gov is a registry and results database of publicly and privately supported clinical studies of human participants conducted around the world. [Learn more about clinical studies](#) and [about this site](#), including relevant [history](#), [policies](#), and [laws](#).

IMPORTANT: Listing of a study on this site does not reflect endorsement by the National Institutes of Health. Talk with a trusted healthcare professional before volunteering for a study. [Read more...](#)

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ClinicalTrials.gov currently lists **247,409 studies** with locations in all 50 States and in **202 countries**.

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Search for Studies

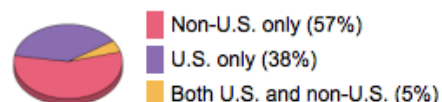
Example: "Heart attack" AND "Los Angeles"

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Locations of Recruiting Studies



Total N = 43,165 studies
(Data as of June 15, 2017)

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For Study Record Managers

- [Why register?](#)
- [How to register your study](#)
- [FDAAA 801 requirements](#)
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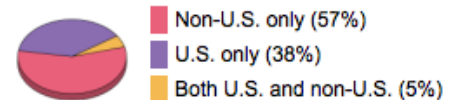
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- [Why Should I Register and Submit Results?](#)
- [FDAAA 801 Requirements](#)
- [How to Apply for an Account](#)
- [How to Register Your Study](#)
- [How to Edit Your Study Record](#)
- [How to Submit Your Results](#)
- [Frequently Asked Questions](#)
- [Support Materials](#)
- [Training Materials](#)

Locations of Recruiting Studies



- Non-U.S. only (57%)
- U.S. only (38%)
- Both U.S. and non-U.S. (5%)

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clinical study information to ClinicalTrials.gov. To avoid duplicate registration, studies should be registered only by the Responsible Party. To help you determine who is responsible for registering a study and submitting results, see the [Elaboration of Definitions of Responsible Party and Applicable Clinical Trial \(PDF\)](#).

If you are the person responsible for registering a study and submitting results, please find out whether your organization has an existing PRS account before applying for one. To apply for an account, follow the instructions below.

[^ TO TOP](#)

Obtaining a PRS Account

Once you have determined that it is appropriate for you to register studies on ClinicalTrials.gov, follow these steps to obtain a PRS account:

1. Check the current [list of organizations with a PRS account](#) to see whether your organization already has a PRS organization account. If so, submit a [PRS Administrator Contact Request Form](#). You will receive contact information for your organization's PRS Administrator(s), whom you can contact directly to request a user login.
2. If your organization does not have a PRS account, you can identify an individual to serve as the PRS Administrator for your organization and that individual can [apply for a PRS organization account](#) on behalf of your organization.
3. If your organization does not currently have anyone who is able to serve as PRS Administrator, you can [apply for a PRS individual account](#); however, this option is not recommended for most organizations. After an individual account is created, an organization is still encouraged to identify appropriate individuals to act as PRS Administrators for the organization.

ClinicalTrials.gov will create a PRS account within 2 business days of receiving your application. Once the account has been created, you will receive an e-mail with instructions for logging in to PRS.

This page last reviewed in December 2014

[^ TO TOP](#)

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Each entity submitting data to ClinicalTrials.gov must adhere to the following terms and conditions, which are intended to ensure the accuracy, currency, and validity of the data:

- Only data for studies that are in conformance with applicable human subjects or ethics review regulations (or equivalent) and applicable regulations of the national (or regional) health authority (or equivalent) may be submitted.
- Notice of changes in recruitment status must be provided as soon as possible, but no later than 30 days after such changes. All other submitted data must be reviewed, verified, and updated as necessary and no less than every 12 months.
- The submitting organization, or individual designated as the [Responsible Party](#), is responsible for the completeness and accuracy of the data submitted to ClinicalTrials.gov.
- Study data must be submitted in English.
- Multiple groups within a single entity (e.g., company, university, government agency) must share a single Protocol Registration and Results System (PRS) organization account.
- Previous versions of study data will be available to the public, although the default view will be the most recent version.

(Acceptance Required)

Accept

Do Not Accept

If your organization is already registered with ClinicalTrials.gov, provide the following information to request contact with your organization's PRS Administrator.

* Organization:

Please enter the name exactly as it appears on the [list of organizations with a PRS account](#).

EskisehirOU|

Requestor Information

* Name:

* Department or Group:

* Phone:

Please enter a valid phone number, including area code.

* Email:

Questions about this form and the PRS may be sent to register@ClinicalTrials.gov.

* Required

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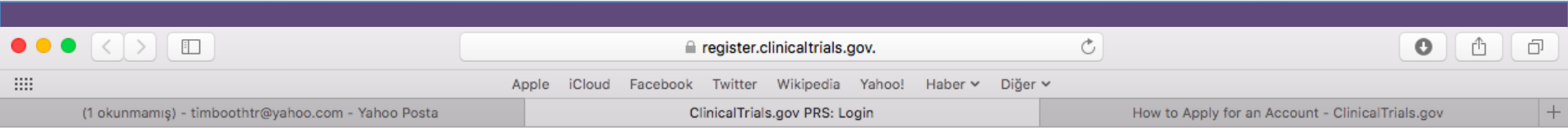
Kişisel bilgilerinizi doldurup admin sisteminden kullanıcı adı ve şifrenizi Beklemeniz gerekmektedir. Şifrenin ulaşması hemen gerçekleşmemektedir.

Questions about this form and the PRS may be sent to register@ClinicalTrials.gov.

* Required

ŞİFRE VE KULLANICI ADI GELDİKTEN SONRA

register.clinicaltrials.gov



ClinicalTrials.gov PRS

Protocol Registration and Results System

Login

Welcome to the [ClinicalTrials.gov](https://www.clinicaltrials.gov) Protocol Registration and Results System (PRS).

OMB NO: 0925-0586
EXPIRATION DATE: 02/29/2020
[Burden Statement](#)

Organization:

One-word organization name assigned by PRS (sent via email when account was created)

Username:

Password:

[Forgot password](#)

Login

See [Submit Studies](#) on ClinicalTrials.gov for information on how to apply for an account, how to register your study, and how to submit results.
[Send email to ClinicalTrials.gov PRS Administration](#)

ClinicalTrials.gov PRS *Protocol Registration and Results System*

Login

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Organization:

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YENİ ÇALIŞMA KAYDI

Quick Links

- [New Record](#)
- [Quick Start Guide](#)
- [Problem Resolution Guide](#)

Records ▾ Accounts ▾ Help ▾

Record List

Showing: 4 records

Search:

[Show/Hide Columns](#)

	Protocol ID	ClinicalTrials.gov ID	Brief Title	Record Status	Last Update	Responsible Party	Problems
Open	PROBAGE006	NCT02765217	Effect of Lactobacillus Reuteri DSM 17938 to Prevent Antibiotic-associated Diarrhea in Children (PEARL)	Public	05/05/2016 16:51	Ener Cagri DINLEYICI timboothtr@yahoo.com	<ul style="list-style-type: none">• Not Recently Updated• Record Has 1 Error
Open	PROBAGE	NCT01927094	Effects of Probiotics and/or Prebiotics on the Duration of Diarrhea and Hospitalization in Children (PROBAGE)	Public	02/12/2015 10:33	Ener Cagri DINLEYICI timboothtr@yahoo.com	
Open	Prob-esity	NCT01927107	Effects of Probiotics in Obese Children (Prob-esity)	Public	08/19/2013 11:29	Ener Cagri DINLEYICI timboothtr@yahoo.com	
Open	VARICOMP	NCT01887496	Varicella-related Hospitalizations in Turkey (VARICOMP)	Public	06/26/2013 07:59	Ener Cagri DINLEYICI timboothtr@yahoo.com	

KEY: Results Delayed Results PRS Review

XML Upload No longer public PRS Review Comments

[Download...](#)

- Klinik alıřmaların kaydında tm detayların girilmesi istenmektedir ve NCT numarası verilmeden nce bir komite tarafından alıřma deęerlendirilmektedir.

Tm sorularınızda bana 2708 numaralı telefonda ulaşabilirsiniz.