


KLİNİK ÇALIŞMALARIN clinicaltrials.gov kaydı ve NCT numarası alınması

Prof.Dr. Ener Çağrı DİNLEYİCİ

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- Klinik çalışmaların, çalışma başlangıcında kaydedildiği bir bildirim sistemidir.

Bir çok dergi son yıllarda NCT numarası olmayan çalışmaları değerlendirmeye almamaktadır.

clinicaltrials.gov, Amerika Birleşik Devletleri kaynaklı bir çalışma protokol kayıt sistemi olup, özellikle ilaç ve girişimsel çalışmaların kayıtları açısından önemli bir veri tabanıdır.

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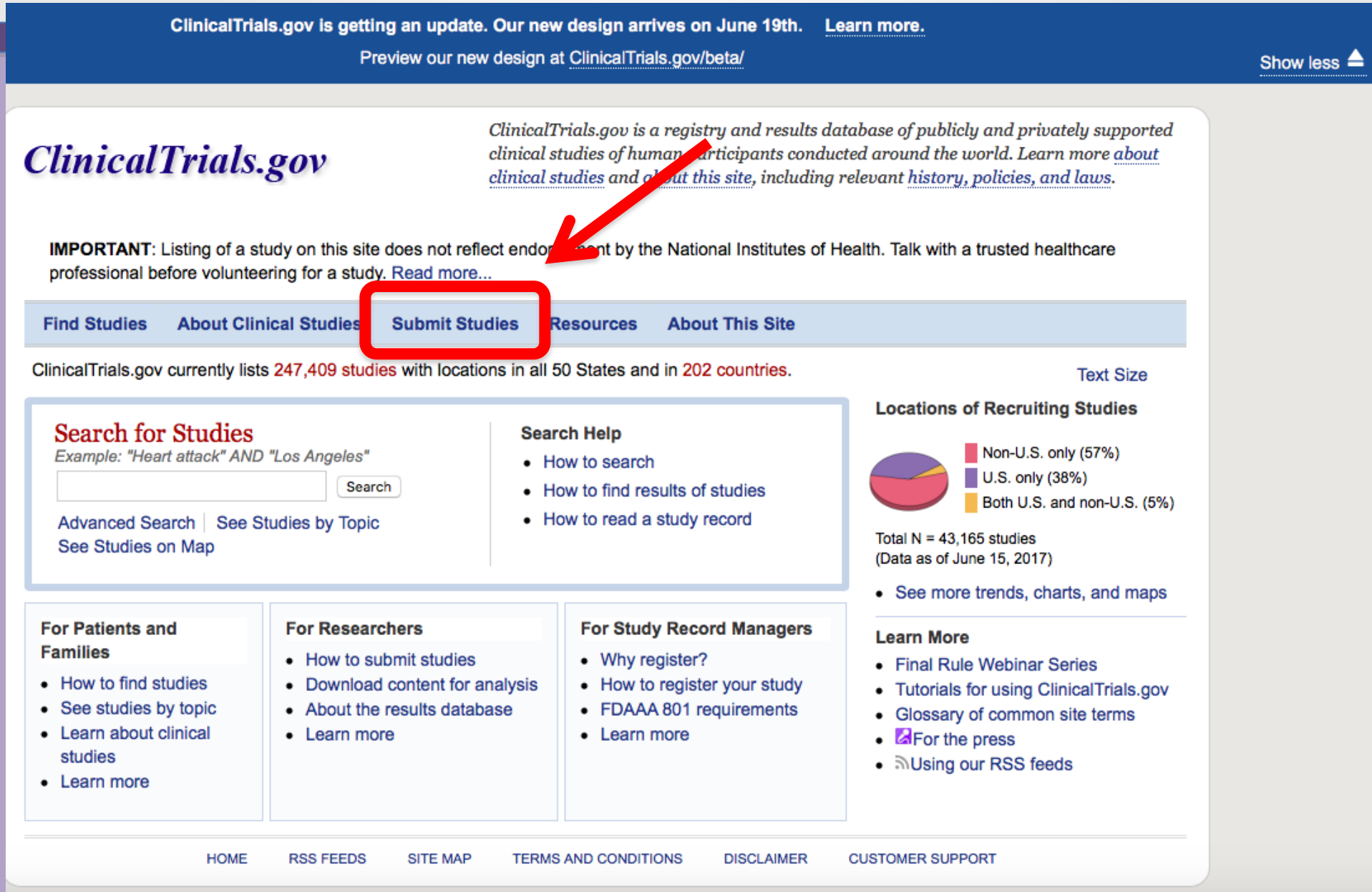

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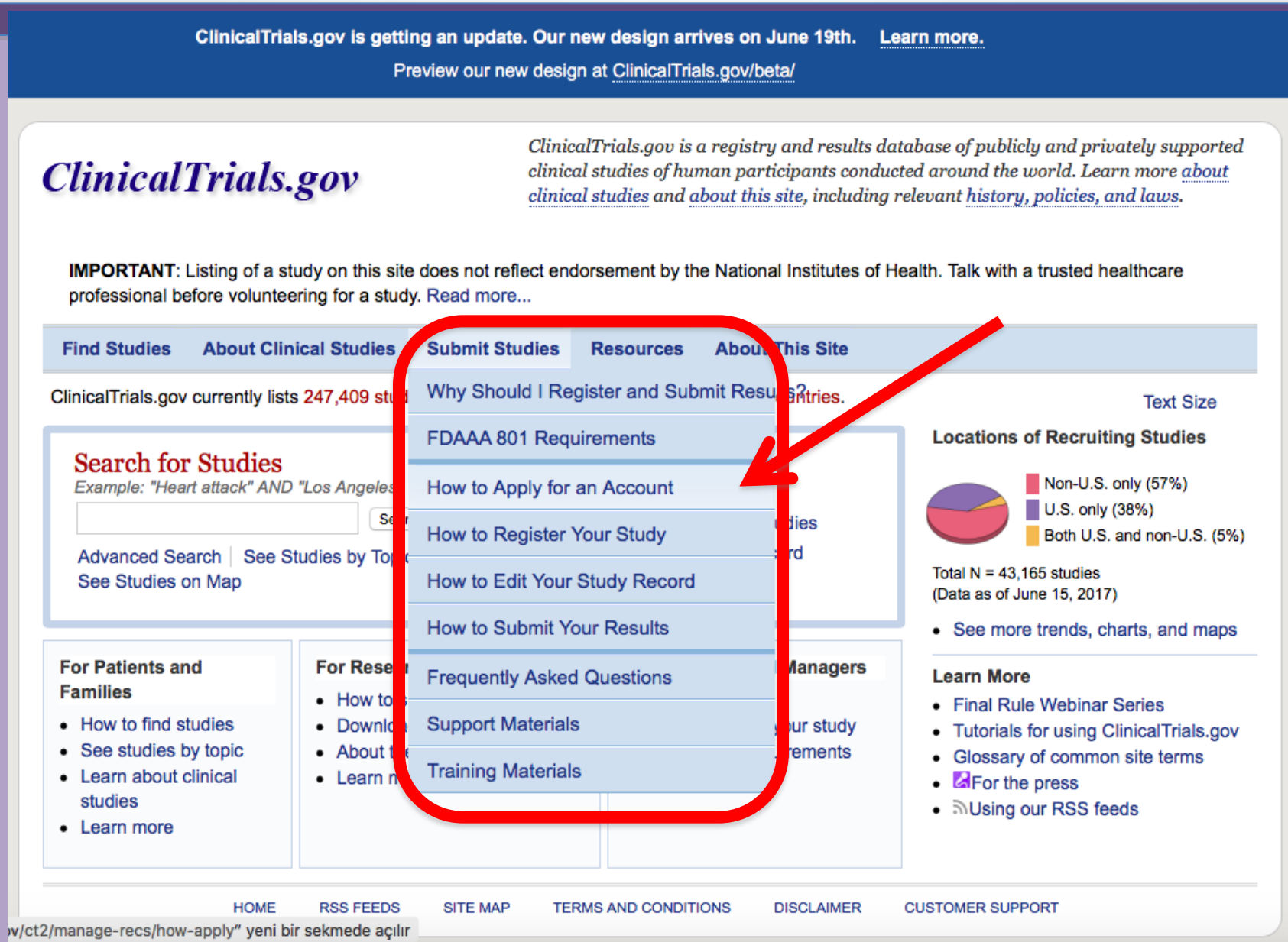

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clinical study information to ClinicalTrials.gov. To avoid duplicate registration, studies should be registered only by the Responsible Party. To help you determine who is responsible for registering a study and submitting results, see the Elaboration 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.

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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

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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.

ŞİFRE VE KULLANICI ADI GELDİKTEN SONRA

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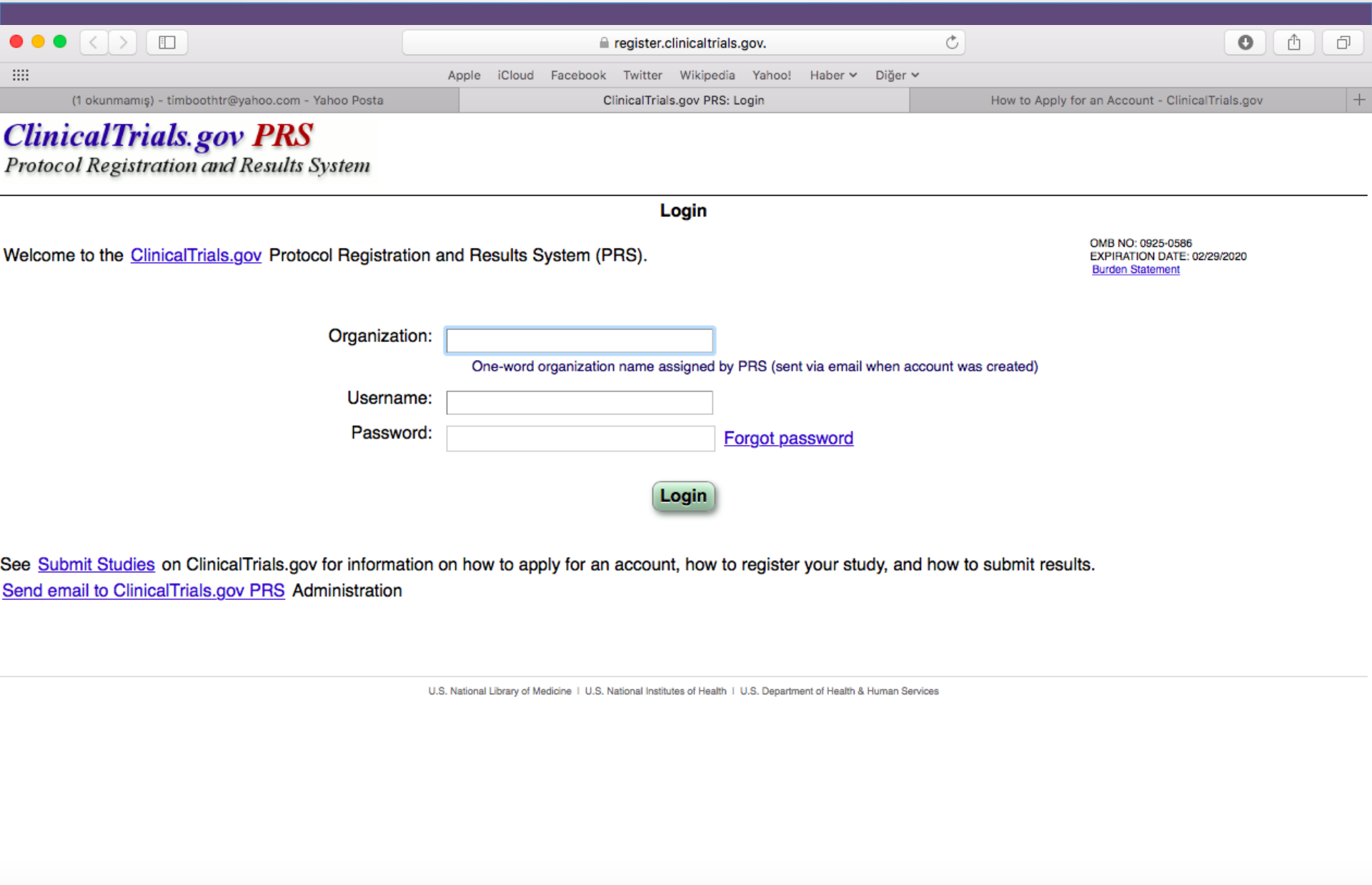

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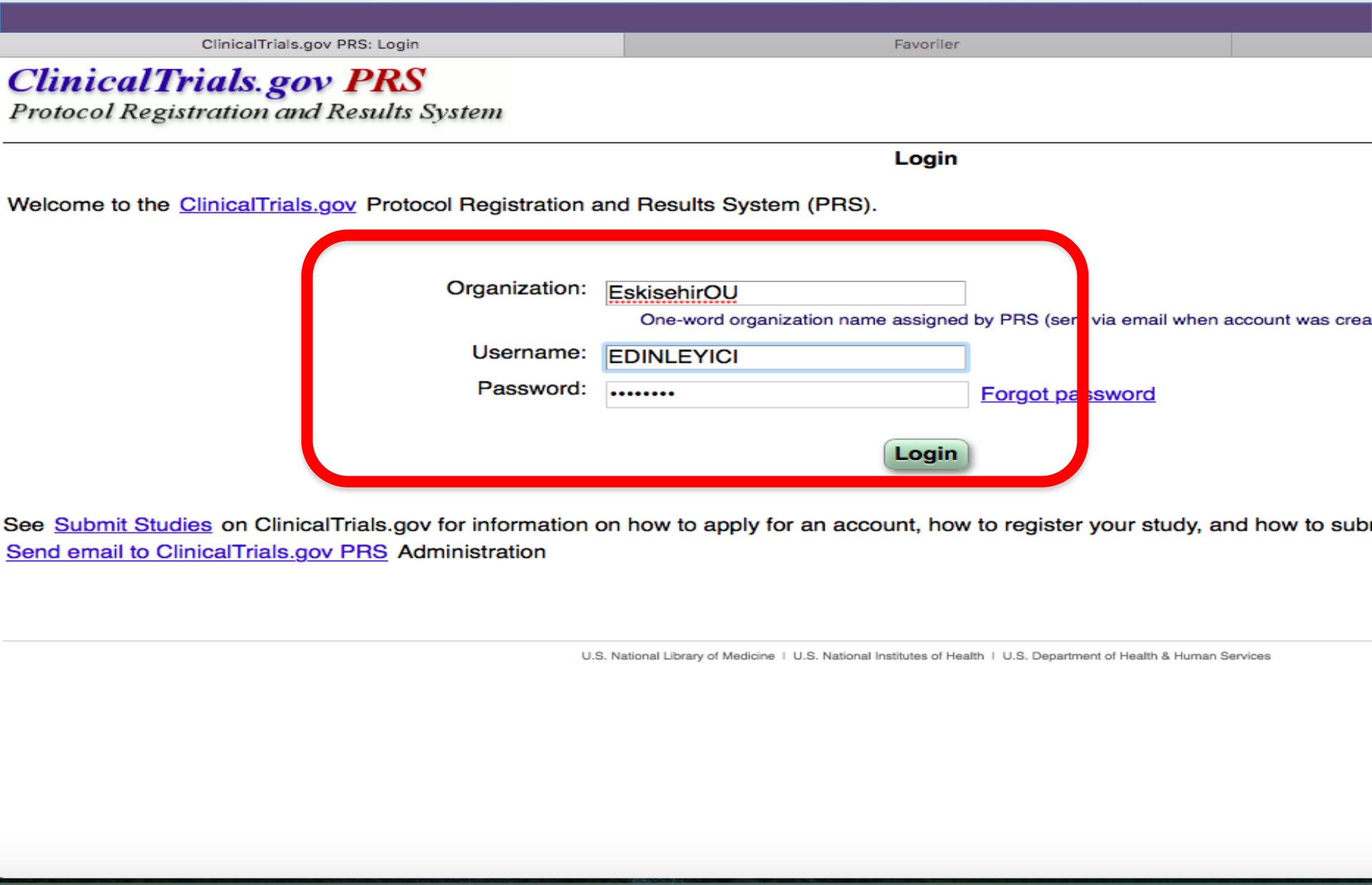
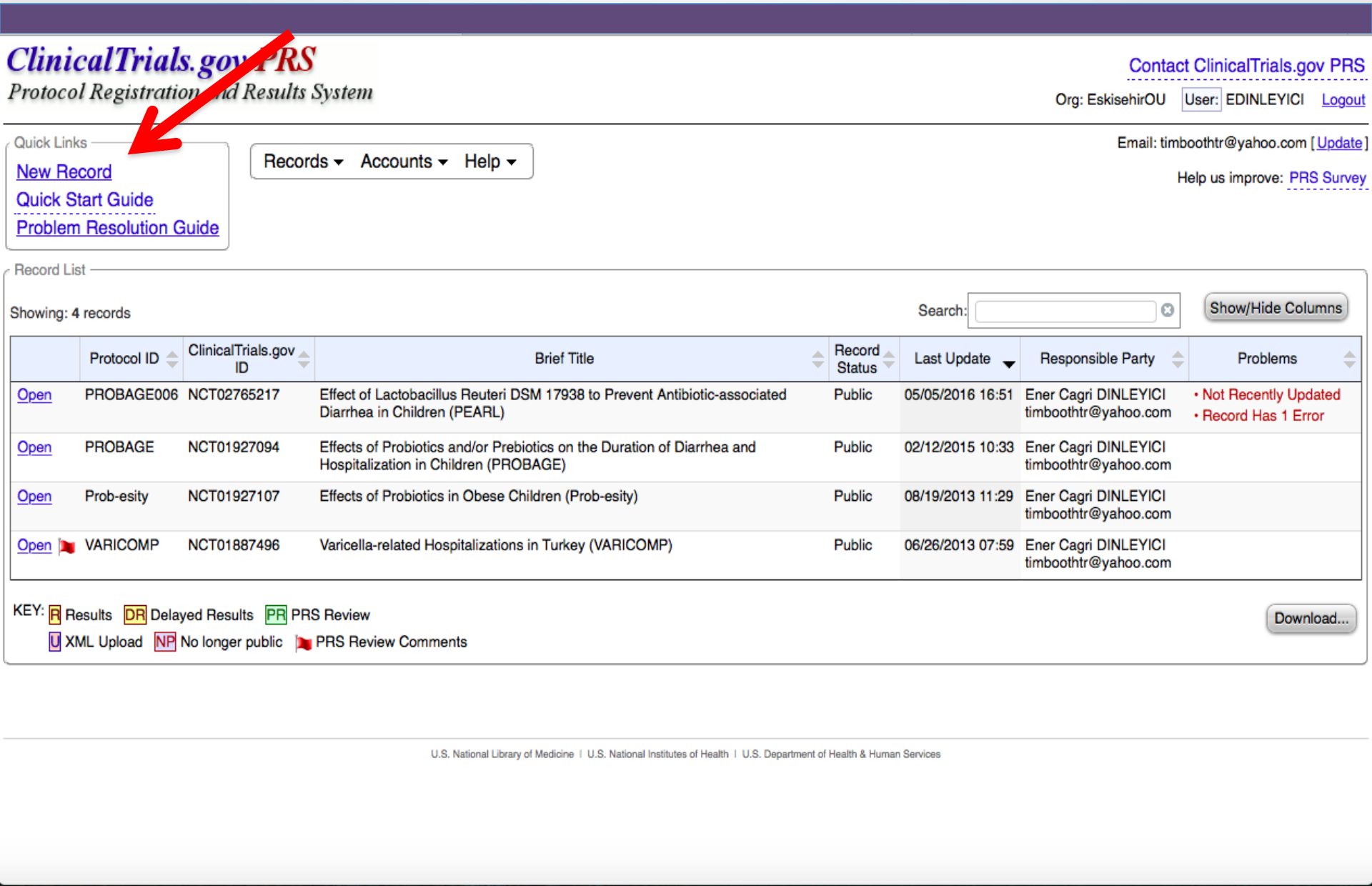


YENİ ÇALIŞMA KAYDI



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- Klinik çalışmaların kaydında tüm detayların girilmesi istenmektedir ve NCT numarası verilmeden önce bir komite tarafından çalışma değerlendirilmektedir.

Tüm sorularınızda bana 2708 numaralı telefondan ulaşabilirsiniz.