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FDA / Center for Biologics Evaluation and Research

After user has logged in using username and password

## Blood Establishment Registration - Select Establishment

Last Session Login: 03/28/2024 11:10:57

Please note: Beginning October 1, 2018, the CBER electronic registration and product listing system (eBER) for manufacturers of blood products and medical devices that are licensed under section 351 of the Public Health Service Act has been updated and expanded to include additional fields for establishment types, blood products, and processes. Additional changes include a requirement to provide a unique facility identifier (see 21 CFR 607.25 (a) and (b)(3)), specified by FDA as your Data Universal Numbering System (DUNS) number (issued by Dun and Bradstreet) and replacement of Form FDA 2830 with a new form entitled *Blood Establishment Registration and Product Listing for Manufacturers of Blood Products and Licensed Devices*.

If your establishment has never registered before:  
Select the **Initial Registration** button below.

If you want to edit the list of establishments you have access to or request access to an existing establishment's registration information: Select the **Edit User Establishments Profile** button below.

If you are updating your establishment registration information: Enter either your FDA Establishment Identifier (FEI) or Central File Number (CFN) in the appropriate box below.  
If you are returning to complete a form started in a previous session enter the Pre-Confirmation Number below.  
Then Select the **Edit This Establishment** button below.

### Frequently Asked Questions

\*FDA Establishment Identifier (FEI):

OR \*Central File Number (CFN):

OR \*Pre-Confirmation Number:

\*Required

Expiration Date:  ?  
Update

OMB Control Number and expiration date visible on page prior to user entering FEI Number associated with their establishment.