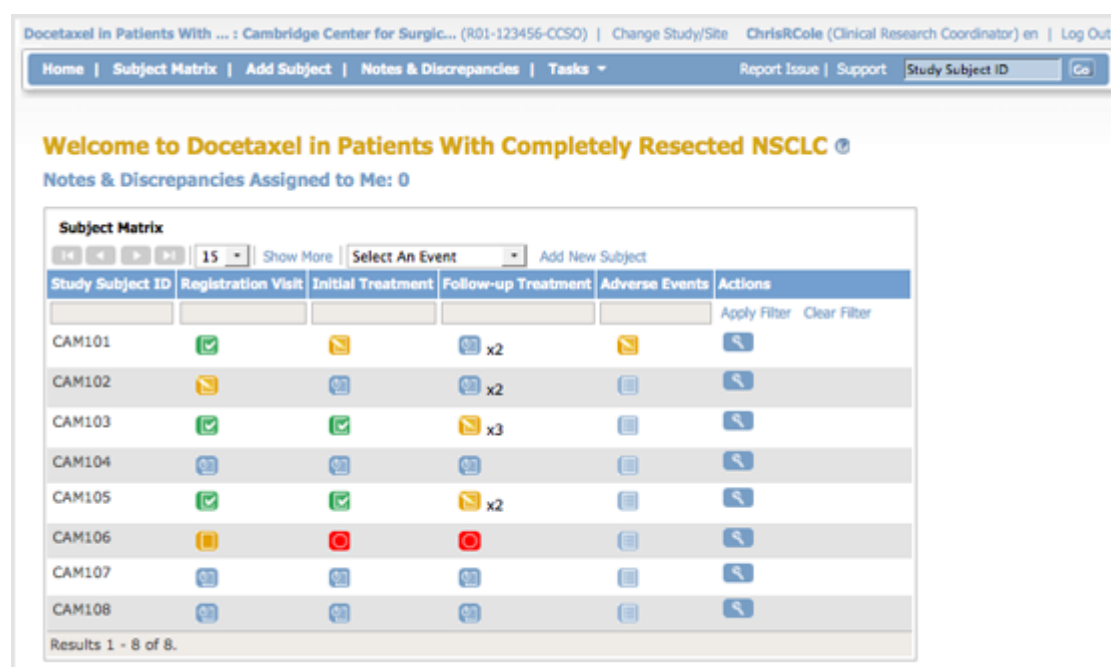


16.3.6 Clinical Research Coordinator Home Page

The body of the home page for a user whose role is Clinical Research Coordinator shows the Subject Matrix for their Site. The Subject Matrix is a table with Event information for all Subjects in a Study. You can view, enter, and change information for Subjects and their Events in the Study. There is one Subject per row, with the Study Subject ID in the first column. The other columns are for each Event Definition in the Study. Each cell contains an icon that identifies the status of Event(s) for the Study Subject. Move the cursor over an icon in the matrix to view and access a Subject's data and actions for that Event. Refer to the Icon Key in the sidebar for icon descriptions. See more details about the [Subject Matrix](#).



The screenshot shows the OpenClinica interface for a Clinical Research Coordinator. The top navigation bar includes links for Home, Subject Matrix, Add Subject, Notes & Discrepancies, and Tasks. The main heading is "Welcome to Docetaxel in Patients With Completely Resected NSCLC". Below this, a table titled "Subject Matrix" displays data for 8 subjects (CAM101 to CAM108). The table has columns for Study Subject ID, Registration Visit, Initial Treatment, Follow-up Treatment, Adverse Events, and Actions. Each cell contains an icon representing the status of the event. For example, CAM101 has a green checkmark for Registration Visit, a yellow envelope for Initial Treatment, and a blue document icon for Follow-up Treatment. The bottom of the table shows "Results 1 - 8 of 8."

Study Subject ID	Registration Visit	Initial Treatment	Follow-up Treatment	Adverse Events	Actions
CAM101	✓	✉	📄 x2	✉	🔍
CAM102	✉	📄	📄 x2	📄	🔍
CAM103	✓	✓	✉ x3	📄	🔍
CAM104	📄	📄	📄	📄	🔍
CAM105	✓	✓	✉ x2	📄	🔍
CAM106	✉	🚫	🚫	📄	🔍
CAM107	📄	📄	📄	📄	🔍
CAM108	📄	📄	📄	📄	🔍

Tasks Available to a Clinical Research Coordinator

Modules (groups of Tasks) available for a Clinical Research Coordinator are:

- Submit Data

Some Tasks a Clinical Research Coordinator can perform are:

- View All Subjects
- Add Subjects
- Add New Study Events
- View Study Events
- Enter CRF Data

This page is not approved for publication.