

1.0 Study Identification. IRB: Newark

This is the first step in your Research Application. You will be automatically guided to the appropriate forms needed to complete your submission.

1.0 * Select the type of submission for your project:

- Research Protocol/Study**
- Research [Tissue or Data] Bank
- Facilitated Review or NCI-CIRB Independent Review
- Emergency Permission
- Humanitarian Use Device (HUD)
- Quality Assurance/Quality Improvement Only
- Western IRB (WIRB)

2.0 Full Title of Project: (If Research [Tissue or Data] bank, Describe where the research [Tissue or Data] bank is physically located (i.e. Institution / Department / Building / Floor / Room)

*** Full Title:**

Evaluating the Effect of Expanded Child Welfare Nursing Services on the Health Care Outcomes of Children in Foster Care.

3.0 * Short Title (Study Name):

The Effect of Nursing Services on Foster Children's Health Care Outcomes.

4.0 * Principal Investigator / Repository Administrator:

[Nina Colabelli](#)

5.0 Study Coordinator / Contact Person:

6.0 Co-Investigators:

Last Name	First Name	Department/Division	Institutional Status	On Probation
Guild	Susan	School of Nursing	Rutgers Paid Staff	no

7.0 Other Staff :

Name	Department	Role	Interaction or access to individuals	Institutional Status	On Probation	Date Modified
Peijia Zha	School of Nursing	Capstone Committee Member, assist with	no	Rutgers Paid Faculty	no	10/9/2014

		data analysis		
Mercedes Echevarria	School of Nursing	Capstone Chair, assist no with data analysis	Rutgers Paid no Faculty	9/15/2014

NOTE:

- **How do I add study team members not in the pick list?**
 - If your study team member does not appear in the pick list for a particular role, then they may not have an account in eIRB. Your first step is to have the study team member register at <https://eIRB.rutgers.edu>. Registration is required during initial logon. An account will be created within 1 to 2 business days.
- All individuals responsible for or working on this project for whom a Rutgers IRB is the IRB of record must be listed here, including individuals who will have responsibility for the consent process, interactions or interventions with subjects, data collection, etc., or who will have access to identifiable private information for research purposes.
- All personnel whose role involves the interaction with living individuals or access to their identifiable information require CITI training.
 - Detailed information for CITI Training can be found [here](#).

View: Add Other Study Staff Member

Study Staff: Designation of Responsibilities

Instructions:

- Use this form to add additional personnel to the team
- Do not add Co-Investigators or the primary Study Coordinator here
- You may add multiple people by clicking the 'OK Add Another' button

* **Other Study Team Member** - click on the button, find the person you want to add:
Peijia Zha

* **Role** - enter a description of the role this person will be performing:
Capstone Committee Member, assist with data analysis

* **Designated Tasks:**

Data Analysis

* **Will the staff member interact with living individuals or have access to their identifiable information:**
 Yes No

View: Add Other Study Staff Member

Study Staff: Designation of Responsibilities

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- Use this form to add additional personnel to the team
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- You may add multiple people by clicking the 'OK Add Another' button

* **Other Study Team Member** - click on the button, find the person you want to add:
Mercedes Echevarria

* **Role** - enter a description of the role this person will be performing:
Capstone Chair, assist with data analysis

*** Designated Tasks:**

Data Analysis

Student Advisor

*** Will the staff member interact with living individuals or have access to their identifiable information:**

Yes No