

Pediatric Heart Transplant Study

Form 03⁹⁹: Initial Immunosuppressor & Antibiotics

ID# P

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P Institution Code	Sequential Patient Number	Patient Initials
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A. Initial Immunosuppression: (complete and mail 30 days post transplant)

1. Induction Therapy (cytolytic therapy soon after transplant not used to specifically treat known rejection)

Yes (if yes complete this question) No (if no, skip to number 2)

*Induction Agents:

Specifics of Induction; indicate any dose or agent change on a new line:

**OKT3
ALG
ATG.**

If other, please specify.

AGENT*	Pre-Op Dose?	Intra-Op Dose?	Dose/Day & units	Start Date	Stop Date
	<input type="checkbox"/> Yes <input type="checkbox"/> No	<input type="checkbox"/> Yes <input type="checkbox"/> No		____-____-____	____-____-____
	<input type="checkbox"/> Yes <input type="checkbox"/> No	<input type="checkbox"/> Yes <input type="checkbox"/> No		____-____-____	____-____-____
	<input type="checkbox"/> Yes <input type="checkbox"/> No	<input type="checkbox"/> Yes <input type="checkbox"/> No		____-____-____	____-____-____
	<input type="checkbox"/> Yes <input type="checkbox"/> No	<input type="checkbox"/> Yes <input type="checkbox"/> No		____-____-____	____-____-____

2a. Cyclosporine: --Pre-Op Dose Yes No (if yes, specify: _____ mg PO IV)

--Date First Post Op Cyclosporine Dose: ____-____-____. Dose: _____ mg, PO IV

--Brand: Sandimmune Neoral Generic/Other/Study, specify: _____

--Date First Post Op Oral/NGtube Dose (if initial post op dose given IV): ____-____-____

2b. Tacrolimus (Prograf, FK-506): --Pre-Op Dose: Yes No (if yes, specify: _____ mg PO IV)

--Date First Post Op tacrolimus dose: ____-____-____. Dose: _____ mg (PO IV)

3a. Azathioprine (Imuran): --Pre-Op Dose: Yes No (if yes, specify: _____ mg PO IV)

--Date First Post Op azathioprine dose: ____-____-____. Dose: _____ mg (PO IV)

3b. Mycophenolate (Cellcept): --Pre-Op Dose: Yes No (if yes, specify: _____ mg PO IV)

--Date First Post Op mycophenolate dose: ____-____-____. Dose: _____ mg (PO IV)

4. Steroids: Pre-Op: None Prednisone Prednisolone Solumedrol: _____ mg PO IV

Intra-Op: None Solumedrol: _____ mg IV Other, specify: _____ dose: _____

Post-Op: Start Date: ____-____-____ (first post op dose)

Initial Dose: _____ mg PO IV (total dose that day) Type: Prednisone Prednisolone Solumedrol

At 14 Days: _____ mg PO IV (total dose per day) Type: Prednisone Prednisolone Solumedrol

At 30 Days: _____ mg PO IV (total dose per day) Type: Prednisone Prednisolone Solumedrol

6. Other immunosuppression (specify: _____):

a. Pre-Op Dose Yes No If yes, specify: _____ mg PO IV Blinded dose per protocol.

b. Date First Post Op Dose (if any): ____-____-____. Dose: _____ mg, PO IV Blinded dose per protocol

7. List and describe any unusual pre-op or early (1st 30 days) immunosuppression or procedures (including plasmapheresis, photopheresis, immunoabsorption, or radiation (TLI)) with dates:

B. Prophylactic Antibiotics/Antivirals started Pre-op through 30 days post op:

8. Infection Prophylaxis started during first 30 days post transplant (not used to treat a known infection):

<input type="checkbox"/> Ganciclovir (Cytovene)	Date Start: ____-____-____	Days of Therapy Intended: _____
<input type="checkbox"/> Acyclovir (Zovirax)	Date Start: ____-____-____	Days of Therapy Intended: _____
<input type="checkbox"/> Trimethoprim/sulfa	Date Start: ____-____-____	Days of Therapy Intended: _____
<input type="checkbox"/> Antifungal (specify: _____)	Date Start: ____-____-____	Days of Therapy Intended: _____
<input type="checkbox"/> Other (specify: _____)	Date Start: ____-____-____	Days of Therapy Intended: _____
<input type="checkbox"/> Immune Globulin	Date Start: ____-____-____	Days of Therapy Intended: _____
<input type="checkbox"/> Other,specify: _____	Date Start: ____-____-____	Days of Therapy Intended: _____
<input type="checkbox"/> Peri-operative* antibiotics: Specify: _____	Date Start: ____-____-____	Days of Therapy Intended: _____

* Peri-operative includes: pre-operative, intra-operative, and started prophylactically immediately post-operative.

PRINT IN BLACK INK ONLY: USE THIS FORM FOR ALL PATIENTS OR EVENTS FROM January 1, 1999.

2/14/99 RCB

Person Completing this form: _____

Date Original Form Mailed (do not send copy): _____