

ACUTE RENAL FAILURE TRIAL NETWORK (ATN STUDY)

FORM 01 - SCREENING/ELIGIBILITY

Hospital No.

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Patient ID

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Patient Initials

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Screening Date (mm/dd/yy)

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A. INCLUSION CRITERIA (To randomize the patient, items 1-6 must all be YES)

1. Acute renal failure clinically consistent with a diagnosis of ATN defined as condition (a) plus either condition (b) or (c) below ----- Yes ☐ No ☐

a. Clinical setting of acute ischemic or nephrotoxic injury ----- Yes ☐ No ☐

b. An increase in serum creatinine of ≥ 2 mg/dL for males
or ≥ 1.5 mg/dL for females over a period of ≤ 4 days ----- Yes ☐ No ☐

1. Gender ----- male ☐ female ☐

2. Lowest serum creatinine within 4 days prior to screening -----

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 mg/dL

date obtained -----

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 mm/dd/yy

No value available ----- ☐

3. Serum creatinine at screening -----

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 mg/dL

date obtained -----

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 mm/dd/yy

4. Date of onset of acute renal failure -----

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 mm/dd/yy

c. Oliguria (average urine output < 20 mL/hour for > 24 hours) ----- Yes ☐ No ☐

1. 24-hour urine volume -----

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 mL

2. Clinical need for renal replacement therapy ----- Yes ☐ No ☐

3. Receiving care in critical care unit (e.g., ICU, MICU, SICU, CTICU) ----- ☐ ☐

a. If yes, check one

☐ MICU ☐ SICU ☐ CCU ☐ CTICU ☐ Trauma ☐ Mixed ☐ Other

THIS FORM MUST BE COMPLETED FOR ALL SCREENED PATIENTS

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4. One non-renal organ failure or sepsis; i.e., 1 or more of conditions a-f below is satisfied Yes ☐ No ☐

a. PaO₂/FiO₂ ≤ 300 mmHg Yes ☐ No ☐

If yes, enter values

PaO₂

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 mmHg;FiO₂

	.		
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b. Platelet count ≤ 100,000/mm³ Yes ☐ No ☐

If yes, enter value

			,				/mm ³
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c. Bilirubin ≥ 2.0 mg/dL Yes ☐ No ☐

If yes, enter value

			.		mg/dL
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d. Hypotension requiring pressor support for greater than 1 hour Yes ☐ No ☐

e. Glasgow Coma Scale ≤ 12 Yes ☐ No ☐

i. Patient is on sedation? Yes ☐ No ☐

ii. Best Eye Response (check one):

☐ No eye opening (1)☐ Eye opening to verbal command (3)☐ Eye opening to pain (2)☐ Eyes open spontaneously (4)

iii. Best Motor Response (check one):

☐ No motor response (1)☐ Withdrawal from pain (4)☐ Extension to pain (2)☐ Localizes pain (5)☐ Flexion to pain (3)☐ Obeys commands (6)

iv. Best Verbal Response (check one)

a. Non-Intubated

☐ No verbal response (1)☐ Incomprehensible sounds (2)☐ Inappropriate words (3)☐ Converses/Confused (4)☐ Converses/Orientated (5)

b. Intubated

☐ Generally unresponsive (1)☐ Questionable ability to talk (3)☐ Seems able to talk (5)

v. Record overall score (sum of items ii through iv above)

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f. Sepsis defined as proven or suspected infection associated with one or more organ failures.

Yes ☐ No ☐

If yes,

1. Proven infection Yes ☐ No ☐2. Suspected infection Yes ☐ No ☐3. Site of infection (see Operations manual)

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5. Age \geq 18 years Yes ☐ No ☐6. Informed consent signed? ☐ ☐a. If yes, who signed informed consent (check one)? ☐ Patient ☐ Surrogate

If surrogate signed, be sure to complete Section C of this form.

b. If no, (check one). ☐ Patient refused ☐ Surrogate refused ☐ Surrogate not available**B. EXCLUSION CRITERIA (To randomize the patient, items 1-15 must all be NO)**1. Pre-Morbid serum creatinine $>2\text{mg/dL}$ (males) or $>1.5\text{ mg/dL}$ (females) Yes ☐ No ☐a. Enter pre-morbid serum creatinine

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 mg/dLb. Date obtained

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 mm/dd/yyc. No value available ☐2. Acute renal failure primarily due to an etiology other than ATN Yes ☐ No ☐

If yes,

a. Etiology code (see Ops Manual)

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3. More than 72 hours since BOTH of the following conditions were met. Yes ☐ No ☐

a. Fulfilled definition of ARF.

1. Date definition of ARF first met

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 mm/dd/yyb. BUN $> 100\text{mg/dL}$ 1. BUN at time of screening

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 mg/dL2. BUN 3 days prior to screening

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 mg/dL

VA WEST HAVEN CSP 530
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Yes No

4. More than one hemodialysis treatment or longer than 24 hours since starting CRRT ----- ☐ ☐
5. Prior kidney transplant ----- ☐ ☐
6. Patient pregnant ----- ☐ ☐
7. Patient is a prisoner ----- ☐ ☐
8. Pre-morbid weight > 128.5 kg ----- ☐ ☐
- a. enter pre-morbid weight

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 kg
9. Non-candidacy for acute renal replacement therapy ----- ☐ ☐
10. Moribund state ----- ☐ ☐
11. Patient not expected to survive 28 days because of an irreversible chronic medical condition ----- ☐ ☐
12. Comfort-measures-only status ----- ☐ ☐
13. Participation in a concurrent interventional study ----- ☐ ☐
14. Patient/Surrogate refusal ----- ☐ ☐
15. Physician refusal ----- ☐ ☐

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C. THIS SECTION TO BE COMPLETED IF CONSENT IS BEING PROVIDED BY A SURROGATE

1. Patient deemed by 2 physicians to be unable to provide informed consent ----- Yes ☐ No ☐

If no, surrogate consent cannot be used. Skip to Section E.

If yes, complete remainder of this section.

2. Name of person who signed informed consent

First Name

[illegible]

MI

□

Last Name

[illegible]

3. Relationship of person who signed informed consent to patient (check one)

☐ Spouse/Partner ☐ Parent

☐ Sibling☐ Child☐ Friend☐ Other relative☐ Other, specify:

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4. Basis for using person named in item 2 above as surrogate (check one)

☐ Court-approved legal guardian☐ Durable power of attorney for healthcare

☐ Next of kin

☐ Other, specify:

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D. IS THIS PATIENT ELIGIBLE FOR RANDOMIZATION?

Yes ☐ No ☐

If yes, Complete Randomization Form (Form 03)

Date of Form Completion

□□ / □□ / □□
(mm/dd/yy)

Staff Initials

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