

TIME _____

DATE _____

SUBJECT NUMBER _____

PASSKI Research Note

Patient was enrolled into PASSKI study by _____ on _____ at _____
(name) (date)
_____. The study was explained to the patient / LAR in full; patient / LAR was allowed to read
(time)
through consent form and all questions were answered before consenting.

Patient had:

- A creatinine rise from _____ to _____ or
(pre Cr) (post Cr)
- Increase of Creatinine \geq 150% to 200% from baseline _____ or
(specify)
- Sustained oliguria (mean output of <0.5 cc/kg/hr for 6 hours)

In addition patient had a granular cast score of _____ or FENa of _____.
(I-IV)

Patient did not meet exclusion criteria. No study protocol procedures were performed prior to obtaining consent. Patient or LAR received a copy of consent form and one was put in the patient chart.

Patient was not in capacity to give consent, so consent was received from a legally authorized representative (LAR).
