STARRT-AKI Investigator CRF Attestation Form

<table>
<thead>
<tr>
<th>Study Title:</th>
<th>STandard versus Accelerated initiation of Renal Replacement Therapy in Acute Kidney Injury</th>
</tr>
</thead>
<tbody>
<tr>
<td>Site Name:</td>
<td></td>
</tr>
<tr>
<td>Investigator Name:</td>
<td></td>
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<tr>
<td>No. of patients randomized:</td>
<td></td>
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Note: The PI signature on this form should be obtained after all patients randomized at the site have reached their Day 90 follow-up, and ALL the data for patients randomized at the site has been completed on Medidata RAVE.

By signing this form, I attest to having reviewed and verified the data entered for all patients randomized in STARRT-AKI study at this site, and certify that the below mentioned data entered on Medidata RAVE for all patients recruited at this site is accurate, complete, and consistent with the parameters described in the Manual of Operations V4.0 [02APR2018].

- Eligibility criteria
- Randomization
- Baseline
- Daily data
- RRT Initiation data
- Adverse Events/Serious Adverse Events
- Hospital and ICU discharge data
- Resource Utilization through day 28
- Hospital Re-admissions
- Day 90 outcome data
- Study completion data

Principal Investigator Signature: ________________________ Date: __________