Dummy Case Report Form (CRF)

Visit 1 - Demographics

Subject ID: _____

Visit Date (DD-MMM-YYYY): _____

Subject Initials: _____

Date of Birth (DD-MMM-YYYY): _____

Sex (M/F): _____

Race: _____

Ethnicity: _____

Investigator Signature: _____

Date: _____

Visit 1 - Vitals

Subject ID: _____

Visit Date (DD-MMM-YYYY): _____

Height (cm): _____

Weight (kg): _____

Systolic BP (mmHg): _____

Diastolic BP (mmHg): _____

Pulse (bpm): _____

Investigator Signature: _____

Date: _____

Visit 1 - Medical History

Subject ID: _____

Visit Date (DD-MMM-YYYY): _____

Condition: _____

Date of Onset: _____

Ongoing? (Y/N): _____

Treatment Given: _____

Outcome: _____

Investigator Signature: _____

Date: _____

Visit 1 - Concomitant Medications

Subject ID: _____

Visit Date (DD-MMM-YYYY): _____

Medication Name: _____

Indication: _____

Dose: _____

Route: _____

Start Date: _____

End Date: _____

Investigator Signature: _____

Date: _____

Visit 1 - Adverse Events

Subject ID: _____

Visit Date (DD-MMM-YYYY): _____

Event Term: _____

Start Date: _____

End Date: _____

Severity: _____

Related to Study Drug? (Y/N): _____

Investigator Signature: _____

Date: _____

Visit 2 - Demographics

Subject ID: _____

Visit Date (DD-MMM-YYYY): _____

Subject Initials: _____

Date of Birth (DD-MMM-YYYY): _____

Sex (M/F): _____

Race: _____

Ethnicity: _____

Investigator Signature: _____

Date: _____

Visit 2 - Vitals

Subject ID: _____

Visit Date (DD-MMM-YYYY): _____

Height (cm): _____

Weight (kg): _____

Systolic BP (mmHg): _____

Diastolic BP (mmHg): _____

Pulse (bpm): _____

Investigator Signature: _____

Date: _____

Visit 2 - Medical History

Subject ID: _____

Visit Date (DD-MMM-YYYY): _____

Condition: _____

Date of Onset: _____

Ongoing? (Y/N): _____

Treatment Given: _____

Outcome: _____

Investigator Signature: _____

Date: _____

Visit 2 - Concomitant Medications

Subject ID: _____

Visit Date (DD-MMM-YYYY): _____

Medication Name: _____

Indication: _____

Dose: _____

Route: _____

Start Date: _____

End Date: _____

Investigator Signature: _____

Date: _____

Visit 2 - Adverse Events

Subject ID: _____

Visit Date (DD-MMM-YYYY): _____

Event Term: _____

Start Date: _____

End Date: _____

Severity: _____

Related to Study Drug? (Y/N): _____

Investigator Signature: _____

Date: _____

Visit 3 - Demographics

Subject ID: _____

Visit Date (DD-MMM-YYYY): _____

Subject Initials: _____

Date of Birth (DD-MMM-YYYY): _____

Sex (M/F): _____

Race: _____

Ethnicity: _____

Investigator Signature: _____

Date: _____

Visit 3 - Vitals

Subject ID: _____

Visit Date (DD-MMM-YYYY): _____

Height (cm): _____

Weight (kg): _____

Systolic BP (mmHg): _____

Diastolic BP (mmHg): _____

Pulse (bpm): _____

Investigator Signature: _____

Date: _____

Visit 3 - Medical History

Subject ID: _____

Visit Date (DD-MMM-YYYY): _____

Condition: _____

Date of Onset: _____

Ongoing? (Y/N): _____

Treatment Given: _____

Outcome: _____

Investigator Signature: _____

Date: _____

Visit 3 - Concomitant Medications

Subject ID: _____

Visit Date (DD-MMM-YYYY): _____

Medication Name: _____

Indication: _____

Dose: _____

Route: _____

Start Date: _____

End Date: _____

Investigator Signature: _____

Date: _____

Visit 3 - Adverse Events

Subject ID: _____

Visit Date (DD-MMM-YYYY): _____

Event Term: _____

Start Date: _____

End Date: _____

Severity: _____

Related to Study Drug? (Y/N): _____

Investigator Signature: _____

Date: _____