



Participant ID:

Participant Initials:

Clinical Center:

Site:

Visit Number:

CRF Date:

RC ID:

CRIC STUDY RE-CONSENT STATUS

RC completes this form to document when a participant provides consent for protocol v.5.x. at the outset of Phase III of the CRIC Study

1. Date of Consent to participate in protocol v5.x: ___ / ___ / ___ (MM/DD/YYYY)