[redacted]

13 End Of Treatment: 11-gen-2016

Event Status: completed

Study Subject: **10001**

CRF Status: data entry complete

[redacted]

Completion Status - V01

Section Title: End of Neoadjuvant Treatment Completion Status and Reason of Discontinuation

PAINED PAGE

Was Trastuzumab treatment completed as per protocol?: * ◉ Yes ○ No

If No, please specify the reason of discontinuation: * ○ (select one) ○ Adverse event ○ Consent withdrawn ○ Progressive disease ○ Patient refused to continue trastuzumab ○ Other

Specify: *

Was Pertuzumab treatment completed as per protocol?: * ◉ Yes ○ No

If No, please specify the reason of discontinuation: * ○ (select one) ○ Adverse event ○ Consent withdrawn ○ Progressive disease ○ Patient refused to continue pertuzumab ○ Other

Specify: *

Was Palbociclib treatment completed as per protocol?: * ○ Yes ◉ No

If No, please specify the reason of discontinuation: * ○ (select one) ◉ Adverse event ○ Consent withdrawn ○ Progressive disease ○ Patient refused to continue palbociclib ○ Other

Specify: *

Was Fulvestrant treatment completed as per protocol?: * ◉ Yes ○ No ○ Not Done (Cohort B)

If No, please specify the reason of discontinuation: *

Specify: *

Completion Status V01 — 10001

▼ **CRF Header Info**

Click the flag icon next to an input to enter/view discrepancy notes. Please note that you can only save the notes if CRF data entry has already started.

Exit

SCREEN SNAPSHOT

DSDISC (12/12)

Title: End of Neoadjuvant Treatment Completion Status and Reason of Discontinuation

Was Trastuzumab treatment completed as per protocol?: (•) Yes ( ) No *

If No, please specify the reason of discontinuation: * Specify: *
- (•) (select one)
- ( ) Adverse event
- ( ) Consent withdrawn
- ( ) Progressive disease
- ( ) Patient refused to continue trastuzumab
- ( ) Other

Was Pertuzumab treatment completed as per protocol?: (•) Yes ( ) No *

If No, please specify the reason of discontinuation: * Specify: *
- (•) (select one)
- ( ) Adverse event
- ( ) Consent withdrawn
- ( ) Progressive disease
- ( ) Patient refused to continue pertuzumab
- ( ) Other

Was Palbociclib treatment completed as per protocol?: ( ) Yes (•) No *

If No, please specify the reason of discontinuation: * Specify: *
- ( ) (select one)
- (•) Adverse event
- ( ) Consent withdrawn
- ( ) Progressive disease
- ( ) Patient refused to continue palbociclib
- ( ) Other

Was Fulvestrant treatment completed as per protocol?: (•) Yes ( ) No *

If No, please specify the reason of discontinuation: * Specify: *
- (•) (select one)
- ( ) Adverse event
- ( ) Consent withdrawn
- ( ) Progressive disease
- ( ) Patient refused to continue fulvestrant
- ( ) Other