

SECTION A: GENERAL STUDY INFORMATION FOR OFFICE USE ONLY

A1. Study ID#:

A2. Date Form Completed: ____ / ____ / ____
Month Day Year

A3. Study Staff Initials: ____

A4. Patient's Last Study Visit: ____

SECTION B: FINAL STUDY STATUS

- B1. What was the patient's final study status?
- Completed study 1 → **SKIP TO B5**
 - Lost to follow-up 2 → **SKIP TO B2**
 - Withdrew consent 3 → **SKIP TO B3**
 - Administrative decision 4 → **SKIP TO B1A**
 - Death 5 → **SKIP TO B5 & COMPLETE DEATH FORM**
 - Other 6 ↓

B1a. Specify **administrative decision** or **other**: _____ → **SKIP TO B4**

B2. For patient **lost to follow-up**, date last study data collected: ____ / ____ / ____
Month Day Year

- B2a. Document follow-up efforts below:
- i. _____
 - ii. _____
 - iii. _____

→ **SKIP TO B5**

B3. For patient who **withdrew consent**, date consent withdrawn: ____ / ____ / ____
Month Day Year

B3a. Date last study data collected: ____ / ____ / ____ → **SKIP TO B5**
Month Day Year

B4. For **administrative decision** or **other**, date last study data collected: ____ / ____ / ____ → **SKIP TO B5**
Month Day Year

B5. Additional Comments: _____

→ **IF A4=VBAS OR B1=1, SKIP TO SECTION C**

B6. For randomized patients, did the patient receive any new or continuing treatment for **voiding dysfunction, urge incontinence** or **stress incontinence** since the last study visit for which data was collected?

Yes 1 → **COMPLETE F581**

No 2

SECTION C: PRINCIPAL INVESTIGATOR'S SIGNATURE

I have reviewed and agree with the above-stated information.

Principal Investigator's Signature: _____ Date: ____ / ____ / ____
Month Day Year