

## IMPLANT REMOVAL NOTE

### PATIENT INFORMATION

1. LAST NAME	2. FIRST NAME	3. TODAY'S DATE ____/____/____
4. DOB ____/____/____	5. AGE (YEARS)	6. TIME ____:____ (24hr)
7. GRAVIDITY	8. PARITY	9. DATE OF INSERTION ____/____/____

### ALLERGIES

No Known Allergies     Yes (specify below)

<u>ALLERGY</u>	<u>REACTION</u>

### REASONS FOR REMOVAL

**Chief Complaint:**

- |   |   |  |
|---|---|--|
| <input type="checkbox"/> Vaginal bleeding   | <input type="checkbox"/> Hair loss      | <input type="checkbox"/> Desires pregnancy |
| <input type="checkbox"/> Irregular bleeding | <input type="checkbox"/> Pain; cramping | <input type="checkbox"/> Mood swings       |
| <input type="checkbox"/> Heavy bleeding     | <input type="checkbox"/> Headache       | <input type="checkbox"/> Acne              |
| <input type="checkbox"/> Weight gain        | <input type="checkbox"/> Depression     | <input type="checkbox"/> Breast pain       |
| <input type="checkbox"/> Other: _____       |   |  |

**Secondary Complaints:** \_\_\_\_\_  
 \_\_\_\_\_

Removal Approved By: \_\_\_\_\_

The patient's R / L arm was palpated and the implant device located. The area was prepped with Betadine / Hibiclens. The distal end of the device was palpated and \_\_\_\_ cc of 1% lidocaine with /without epinephrine was injected. A \_\_\_\_ mm incision was made. Any fibrotic tissue was carefully dissected away using blunt and/or sharp dissection. The device was removed in an intact manner. Steri-strips and a sterile dressing were applied to the incision. The patient tolerated the procedure well.

New contraceptive method: \_\_\_\_\_  
 \_\_\_\_\_  
 \_\_\_\_\_

**Clinician signature:** \_\_\_\_\_

**Date:** \_\_\_\_/\_\_\_\_/\_\_\_\_