SECTION A: GENERAL STUDY INFORMATION FOR OFFICE USE ONLY:

IDNEW (ID number unrelated to original study ID number) : _____

Cohort (1-4): __

The PFD Investigator verifies that the following inclusion/exclusion criteria were met during a clinical evaluation prior to completion of an Informed Consent for this study.

SECTION B: INCLUSION CRITERIA All answers MUST be YES

ELGB01. Subject has Stage II, III or IV pelvic floor prolapse 1.☐ Yes 2.☐ No

ELGB02. Subject can complete a telephone interview in English 1.☐ Yes 2.☐ No

ELGB03. Subject is willing to participate in urodynamics testing as a planned part of clinical care 1.☐ Yes 2.☐ No

ELGB04. The subject answered PFDI Question 4 – "Do you usually have a sensation of bulging or protrusion from the vaginal area?" 1.☐ Yes 2.☐ No

ELGB05. The subject answered PFDI Question 5 – "Do you usually have a bulge or something falling out that you can see or feel in the vaginal area?" 1.☐ Yes 2.☐ No

ELGB06. Is the answer YES to either B5 or B6 or both? 1.☐ Yes 2.☐ No

The patient offered the following responses to the 6 MESA questions:

DURING THE PAST MONTH:	1.Never	2.Rarely	3.Sometimes	4.Often
ELGB07. Did coughing hard cause you to lose urine?	☐	☐	☐	☐
ELGB08. Did sneezing cause you to lose urine?	☐	☐	☐	☐
ELGB09. Did lifting cause you to lose urine?	☐	☐	☐	☐
ELGB10. Did bending cause you to lose urine?	☐	☐	☐	☐
ELGB11. Did laughing cause you to lose urine?	☐	☐	☐	☐
ELGB12. Did walking briskly/jogging cause you to lose urine?	☐	☐	☐	☐

ELGB13. Was the answer "Sometimes" or "Often" to any of the items in B7-B12? 1.☐ Yes 2.☐ No

SECTION A: GENERAL STUDY INFORMATION FOR OFFICE USE ONLY:

IDnew __ __ __ __ __

SECTION C: EXCLUSION CRITERIA All answers MUST be NO

ELGC01. Subject currently pregnant or within 6 months postpartum 1.☐ Yes 2.☐ No

ELGC02. Subject has neurological disease which might affect voiding function 1.☐ Yes 2.☐ No

ELGC03. Age > 75 1.☐ Yes 2.☐ No

SECTION D: ELIGIBILITY/TREATMENT PLAN (for prolapse/urinary symptoms)

ELGD01. Eligibility status:

1.☐ Eligible 2.☐ Eligible but does not wish to participate in study→ STOP 3.☐ Ineligible → STOP

ELGD02. Date the subject signed informed consent: deleted

ELGD03. Is abdominal sacrocolopexy planned for this subject (following UDS testing)? 1.☐ Yes 2.☐ No

Note: Cohort is the cohort to which the subject was assigned. See the summary for an explanation.