XXXX - Essure Case Review

Case Report

Parameter	Findings	Bates Ref	PDF Ref
First Name	XXXX	MR - XXXX - 000001	190
Initial	XXXX		
Last Name	XXXX		
DOB	MM/DD/YYYY		
Preexisting	As on 07/23/2013: Patient denies to being allergic	MR - XXXX Women's	1098
conditions, allergies	to nickel.	Health - 000020	
or contraindications			
for placement of			
Essure			
Prior IUD	07/03/2008: Intrauterine Device (Mirena 20	MR - XXXX Women's	1441,
placement, OB/GYN	mcg/24 hour) -placement.	Health - 7.3.08 to 4.5.12	1433-
medical issues from		-000022, MR - XXXX	1436,
history	08/03/2010: Post coital spotting vaginal	Women's Health - 7.3.08	1428-
	discharge/irritation. Diagnosed with candidiasis of	to 4.5.12 - 000014-	1430,
	vulva and vagina	000017, MR - XXXX	1420-
		Women's Health - 7.3.08	1422,
	10/14/2011: Menstrual cycles are irregular due to	to 4.5.12 - 000009-	1079-
	Mirena. Scant white discharge in vagina. Mild	000011, MR - XXXX	1081
	bleeding in cervix.	Women's Health - 7.3.08	
		to 4.5.12 - 000001-	
	04/05/2012: Menses are currently irregular	000003, MR - XXXX	
	secondary to IUD.	Women's Health -	
		000001-000003	
	03/12/2013: The patient states she has not had		
	menses since having IUD placed. Since the IUD		
	placement, the patient reports non-menstrual		
	bleeding. Also complains of vaginal discharge with		
	foul odor to it. She notices this when she is		
	spotting. Irregular menses, Bacterial vaginitis		
Smoking history	Never smoker	MR - XXXX Women`s	1089-
		Health - 000011-000013	1091
Age at the time of	As on 07/23/2013: 39 years old	MR - XXXX Women`s	1096
Essure Implant		Health - 000018	
Body Mass Index	07/12/2013: BMI: 35.8 kg/m2 (<i>Calculated</i>)	MR - XXXX Women`s	1093,
(BMI) when Essure	Weight: 202 lbs	Health - 000015, 000013	1091
was implanted	Height: 5 feet 3 inches (<i>Taken from visit dated</i>		
	07/08/2013)		
Phase of menstrual	*Reviewer's comment: Her LMP has been limited		
cycle during Essure	to spotting since Depo Provera given on		

Parameter	Findings	Bates Ref	PDF Ref
Placement	07/12/2013 but her urine pregnancy test is negative		
	on the date of Essure placement (07/23/2013)		
Duration between	Prior pregnancy: 04/30/2008	MR - XXXX Women`s	1089-
prior pregnancy and	Essure placement date: 07/23/2013	Health - 000011-000014,	1092,
Essure placement		000020-000021	1098-
	Duration : 5 years 2 months		1099
Warnings given to	Yes	MR - XXXX Women`s	1093
Patient		Health - 000015	
	Essure and bilateral tubal ligation procedures, risks,		
	benefits, and alternatives discussed with patient and		
	she wishes to proceed with Essure.		
	She understands that she will need to use a good		
	method of contraception until hysterosalpingogram		
	confirms bilateral tubal occlusion which is		
	generally 3-6 months after placement of the Essure		
	devices.		
	Essure Medicaid Consent signed on 05/06/2013.		
Date of Essure	07/23/2013	MR - XXXX Women`s	1098-
placement		Health - 000020-000021	1099
Reason for	Reason: Permanent sterilization	MR - XXXX Women`s	1098-
Placement and		Health - 000020-000021	1099
Procedure	Procedure : Hysteroscopic insertion of bilateral		
	Essure fallopian tube micro-inserts		
Essure details	Name of implant: Essure	MR - XXXX Women`s	1098-
	Lot number: 927081	Health - 000020-000021	1099
	Reference number: ESS305		
	*Reviewer's comment: Essure implant label is not		
	available; the above information is taken from the		
	operative report dated 07/23/2013		
Number of attempts	Essure was placed successfully in the first attempt	MR - XXXX Women's	1098-
for successful		Health - 000020-000021	1099
placement of Essure			
Birth control	07/12/2013: Depo Provera IM 150 mg/ml given;	MR - XXXX Women's	1093-
measures for 3	Left Mirena IUD in place to help with	Health - 000015-000016,	1094,
months following	dysmenorrhea and menses	MR - XXXX Women's	1095
placement		Health - 000017	
	*Reviewer's comment: Mirena IUD was removed		
	on 07/23/2013.		
Complications	None		
during Essure			
placement			

Parameter	Findings	Bates Ref	PDF Ref
Complications	07/29/2013: Admits to mild menstrual like	MR - XXXX Women`s	1100-
immediately after	cramping. Patient also complains of increased night	Health - 000022-000023	1101
Essure placement	sweats for one week.		
Diagnostic test	10/28/2013: Hysterosalpingogram (HSG): The	MR - XXXX Women`s	1103-
performed to	Essure implants appear to be in satisfactory	Health - 000025-000026	1104
confirm bilateral	position. There was no intraperitoneal spill of		
tubal blockage	contrast consistent with bilateral occlusion of		
following Essure	fallopian tubes. A small amount of intravasation of		
placement	contrast occurred during the procedure.		
Details regarding	Pregnancy not reported after Essure placement		
pregnancy (If			
reported) following			
Essure placement			
Autoimmune-Like	09/12/2013, 10/31/2013, 01/02/2014, 02/27/2014,	MR - XXXX - 000397-	1962-
Symptoms (Hair	05/01/2014: Numbness and tingling in legs	000401, 000415-000420,	1967,
Loss, Fatigue, Joint		000409-000414, 000421-	1956-
Pain and Rashes)	06/05/2014: Numbness in extremities	000426, 000405-000408,	1961,
		000402-000404	1950-
	06/30/2014: Fatigue	8B7B13342F894C66BF	1955,
		24, XXXX, 145-149	1946-
	07/24/2014, 08/28/2014, 10/30/2014, 12/18/2014:	MR - XXXX - 000393-	1949,
	Numbness in extremities	000396, 000388-000392,	1943-
		000383-000387, 000375-	1945,
	03/19/2015, 05/21/2015: Numbness and tingling in	000379, 000364-000369,	1938-
	legs	000358-000363, 000353-	1942,
		000357	1325-
	07/23/2015: Numbness in extremities	MR - XXXX Women's	1329,
		Health - 000071-000075	1934-
	08/06/2015: Fatigue		1937,
		MR - XXXX Medical	1929-
	03/09/2017: Chronic fatigue	Center - 000039-00004	1933,
		8B7B13342F894C66BF	1924-
	05/30/2017, 03/05/2018, 07/10/2018, 10/16/2018,	24, XXXX, 85-89, 76-	1928,
	02/05/2019, 05/22/2019, 02/19/2020: Fatigue	80, 71-76, 66-71, 52-57,	1916-
		48-52, 19-25, 10-16	1920,
			1905-
			1910,
			1899-
			1904,
			1894-
			1898,
			1149-
			1153, 39-

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	1133,
	1140,
	157,
	675-676,
	1158,
	1159-
	162,
	105-406
10/28/2014, 02/10/2015 : 150 mg of Depo Provera	
given	

Parameter	Findings	Bates Ref	PDF Ref
	03/17/2015: Frothy smooth yellow-colored discharge present. Assessed with vulvovaginitis, discharge of vagina, mild cervicitis. Prescribed Diflucan 150 mg, Flagyl 500 mg		
	05/06/2015: NuSwab + for bacterial vaginosis despite mild changes on Wet prep last time similar to this time therefore will treat with Flagyl. Reports intermittent pain with intercourse and generally bad pain after intercourse. Assessed with bacterial vaginosis, candidal vulvovaginitis. Prescribed Diflucan and Flagyl. 150 mg of Depo Provera given		
	08/06/2015: Dysmenorrhea, hot flashes, and mental instability and pelvic pain. 150 mg of Depo Provera given		
	09/21/2015: Pelvic pain, dyspareunia. Reports increasing vaginal infections beginning approximately 4-6 months ago. Patient desires surgery secondary to pre-menopausal status and sexual activity.		
	10/22/2015: Patient was having menorrhagia with clots and was seen by Urogynecology with a recent Depo-Provera injection		
	11/12/2015: 1 ml of Depo Provera given 12/11/2015: Dyspareunia. Patient desires to		
	proceed with Total Laparoscopic Hysterectomy(TLH)/Bilateral Salpingectomy (BS)		
	12/15/2015 : Dyspareunia		
Details of the Essure Removal Surgery	12/15/2015: Total laparoscopic hysterectomy, bilateral salpingectomy, McCall culdoplasty, and cystoscopy under general anesthesia	MR - XXXX - part 1 - 000172-000173	405-406
Reason for removal and method of removal	Reason for removal: Dyspareunia Method of removal: Bilateral salpingectomy	MR - XXXX - part 1 - 000172-000173	405-406

Parameter	Findings	Bates Ref	PDF Ref
Details of the Essure	Details of procedure:		
removal surgery	The patient's right fallopian tube was identified and		
	elevated at the fimbriated and with the use of the		
	Ligasure device, the mesosalpinx was incised up to		
	the uterine cornu with goad hemostasis noted. The		
	uterine ovarian ligament was then isolated,		
	cauterized, and transected with the LigaSure device		
	Attention was then turned to the left side of the		
	pelvis, at which time, the left fallopian tube, left		
	uterovarian ligament, as well as left round ligament		
	each time were isolated, cauterized, and transected		
	with the Ligasure device.		
Is there a pathology	12/18/2015:	MR - XXXX Women`s	1168-
report from the	Final diagnosis:	Health - 000090-000091	1169
removal surgery?	Fallopian tubes		
	Metallic material within lumen consistent		
	with a prior sterilization		
Condition of the	01/20/2016: Minimal amount of light yellow/brown	MR - XXXX Women`s	1173-
patient post removal	discharge noted in vault secondary to dissolution of	Health - 000095-000097,	1175,
procedure	suture	000101-000103, 000104-	1179-
		000106	1181,
	04/13/2016: Vaginal discharge noted after		1182-
	intercourse. Vaginal/vulvar itching or irritation.	8B7B13342F894C66BF	1184,
	Assessed with candidiasis of vulva and vagina,	24, XXXX, 5-10	1185-
	discharge from the vagina. Prescription for Nystatin		1190
	cream and Diflucan given.		
	11/09/2016: Vaginal discharge. Assessed with		
	Candidiasis vulvovaginitis, Discharge from the		
	vagina. Wet prep positive for yeast. Prescription for		
	Diflucan given.		
	04/07/2020: No abdominal pain. No fatigue. Return		
	to office on 07/13/2020		

Patient History

Past medical history: Hypertension, obesity, stage III colon cancer, anxiety, hyperlipidemia, chronic rhinitis

(PDF ref: 663-664, 127) (Bates ref: A1062A78549849D1B73E, XXXX, 182-183, MR - XXXX Medical Center - 000127)

Past Surgical history: Reduction mammoplasty; Laparoscopic left colectomy with splenic flexure takedown and cholecystectomy 09/2009, repair of ventral incisional hernia with mesh on 10/01/2015, dilation and curettage, lumpectomy

(PDF ref: 665, 663, 558-559, 1084) (Bates ref: A1062A78549849D1B73E, XXXX, 184, 182, 77-78, MR - XXXX Women`s Health - 000006)

Family history: Maternal grandmother: Liver cancer. Maternal cousin, maternal aunt, paternal aunt, maternal niece, paternal niece: Breast cancer. Mother: Arthritis, high cholesterol, type II diabetes, hypertension. Grandmother: Heart disease. Grandfather: Stroke. Father: Myocardial infarction (PDF ref: 663, 131, 525, 127) (Bates ref: A1062A78549849D1B73E, XXXX, 182, MR - XXXX Medical Center - 000131, A1062A78549849D1B73E, XXXX, 44, MR - XXXX Medical Center - 000127)

Social history: No history of alcohol, tobacco or substance abuse (*PDF ref: 663*) (*Bates ref: A1062A78549849D1B73E, XXXX, 182*)

Allergies: Aspirin, which causes bruising and Lipitor which causes aching

(PDF ref: 663) (Bates ref: A1062A78549849D1B73E, XXXX, 182)

Detailed Chronology

DATE	PROVIDER	OCCURRENCE/TREATMENT	BATES REF	PDF REF
07/03/2008	XXXX	Office visit for postpartum examination:	MR - XXXX Women`s	1439- 1440
•	XXXX, M.D. XXXX, CNM	The patient is a G 3 P 2 34-year old African American female who presents for a postpartum exam. She is now 6 weeks out from an uncomplicated vaginal delivery. She has not had any significant problems since her delivery. The patient is not breast feeding. The patient would like to use an Intrauterine Device (IUD) for contraception. Since delivery she has used antihypertensive medication.	Health - 7.3.08 to 4.5.12 - 000020-000021	1440
		Physical examination:		
		Genitourinary:		
		External genitalia: Normal appearance for age, no discharge or inflammatory lesions present		
		Vagina: No discharge present, no inflammatory lesions		
		present, no masses present, normal vaginal vault		
		Cervix: Appearance healthy, no lesions present,		

DATE	PROVIDER	OCCURRENCE/TREATMENT	BATES REF	PDF REF
		nontender to palpation, no discharges, no bleeding present, normal midline position, cervix consistency		
		normal		
		Uterus: Nontender to palpation, no masses present,		
		contour; smooth to palpation, position midline/midplane,		
		size normal, shape normal, mobility: normal		
		Adnexa: No adnexal tenderness present, no adnexal		
		masses present		
		Assessment:		
		Post-partum follow-up		
		Contraceptive		
		Plan:		
		Resume all normal activity		
		• Follow-up appointment: One month		
07/03/2008	XXXX	Procedure note for Mirena placement:	MR - XXXX Women`s	1441
	XXXX, M.D.	Preoperative diagnosis: Desires placement of	Health - 7.3.08	
		Intrauterine Device	to 4.5.12 -	
	XXXX, CNM		000022	
		Procedure: Placement of Mirena Intrauterine Device		
		Indication/Consent:		
		Patient is status post vaginal delivery presents today for		
		placement of an IUD. She has been counseled on options		
		for contraception including oral contraceptives, Depo		
		Provera, the contraceptive patch, the contraceptive		
		vaginal ring, as well as permanent sterilization. After		
		careful consideration she has decided to undergo placement of a Mirena IUD. Pregnancy test was		
		indicated. It was negative. The patient was counseled on		
		the risks and benefits of IUD use. She has read and		
	•	signed a detailed patient information and consent		
		brochure on placement and use of the IUD.		
		Description of Procedure:		
		With the patient in the dorsal lithotomy position a		
		speculum was placed in the vagina and the cervix was		
		visualized.		
		There was no significant vaginal or cervical discharge		
		noted. The cervix was cleaned with Betadine. A single-		

DATE	PROVIDER	OCCURRENCE/TREATMENT	BATES REF	PDF REF
		toothed tenaculum was then placed on the anterior lip of the cervix for traction. The uterus was then sounded to 9 cm. Under sterile conditions the IUD was then placed through the cervix into the endometrial cavity without difficulty. The strings were then trimmed to a length of approximately 2-3 centimeters. The tenaculum was the removed from the cervix and good hemostasis was noted. The speculum was then removed from the vagina. Disposition: The patient tolerated the procedure without significant difficulty. She was counseled on refraining from intercourse for approximately 24 hours and to contact the office if she experienced any significant pain, bleeding or fever. Follow-up in 1 month *Reviewer's comment: Interim medical records for the		
00/02/2010		period 07/03/2008-08/03/2010 are not available for review.		1 100
08/03/2010	XXXX, M.D. XXXX, N.P.	Patient's last normal menstrual period is uncertain secondary to irregular bleeding. She presents today for her annual exam. Her last pap was approximately 10/26/2007 ago and with normal results. Her menses are irregular occurring very unpredictably. She describes her menstrual flow as varies and has had occasional spotting. She describes mild cramps of 1-day duration. She is sexually active and uses an IUD for contraception. She reports post coital spotting. She is having significant problems which include vaginal discharge/irritation. Review of systems: Genitourinary: Irregular menses, vaginal discharge, post-coital bleeding Psychiatric: Anxiety, difficulty sleeping Wet mount: Trichomonas: Negative	MR - XXXX Women's Health - 7.3.08 to 4.5.12 - 000014-000017	1433-1436
		Psychiatric: Anxiety, difficulty sleeping		

DATE	PROVIDER	OCCURRENCE/TREATMENT	BATES REF	PDF REF
		Hyphae: Negative		
		Buds: Positive		
		Whiff Test: Negative		
		Assessment: Candidiasis of vulva and vagina		
		Plan: Terazol 3 Vaginal Cream 0.8 %. Return in 1 year		
		for physical examination		
		The patient was also informed to notify her oncologist		
		that she has a Mirena IUD for birth control to see if the		
		progesterone in the IUD will cause problems with treatment or her cancer.		
08/05/2010	XXXX	Labs:	MR - XXXX	1437
00/03/2010		Collected date: 08/03/2010	Women's	1437
		Contested dutes 60/35/2515	Health - 7.3.08	
		Chlamydia trachomatis, Neisseria gonorrhoeae:	to 4.5.12 -	
		Negative	000018	
08/06/2010	XXXX	Cytology report:	MR - XXXX	1438
		Collected date: 08/03/2010	Women's	
	XXXX,		Health - 7.3.08	
	Cytotechnologis	Source: Cervical; Endocervical	to 4.5.12 -	
	t (ASCP)		000019	
		Diagnosis: Negative for intraepithelial lesion and		
		malignancy.		
09/22/2010	XXXX Medical	Office visit for chest pain:	MR - XXXX	127-
	Center		Medical Center -	129
		Sexual activity: Monogamous relationship. Regular	000127-000129	
	XXXX, M.D.	condom use, careful partner selection		
		*Reviewer's comment: Only case relevant information		
		captured from this visit.		
10/01/2010	XXXX Medical	Labs:	A1062A785498	532
	Center	DOGD V :	49D1B73E, XXXX, 51	
1.000.000		POC Pregnancy: Negative	·	10
12/03/2010	XXXX	Follow-up visit intake:	MR - XXXX - 000323-000324	1864- 1865
		Review of systems: Endocrine : Low sex drive	000323-000324	1003
03/24/2011	XXXX Medical	Office visit for annual physical examination:	MR - XXXX	118-
	Center		Medical Center -	120
		Sexual activity: Monogamous relationship. Regular	000118-000120	
	XXXX, N.PC.	condom use, careful partner selection		
1	1	·	1	

DATE	PROVIDER	OCCURRENCE/TREATMENT	BATES REF	PDF REF
		*Reviewer's comment: Only case relevant information captured from this visit.		
03/31/2011	XXXX Medical Center XXXX, N.PC.	Office visit for urinary tract infection: Assessment: Urinary Tract Infection (UTI) - Resolved.	MR - XXXX Medical Center - 000115	115
		PPD is negative Follow-up as needed		
10/14/2011	XXXX	Office visit for annual gynecological examination:	MR - XXXX Women's	1428- 1430
	XXXX, M.D.	Patient's menstrual cycles are irregular due to Mirena. Her last pap was approximately 1.2 years ago and with	Health - 7.3.08 to 4.5.12 -	
	XXXX, CNM	normal results. She describes her menstrual flow as varies and has had occasional spotting. She describes mild cramps of 1-day duration	000009-000011	
		Patient states when she tries to have a bowel movement, when she wipes her vaginal area, she sees blood. She would like to discuss an Essure. Pharmacy-Kerr Drug-		
		Warrenton, or Medical Arts if she is being seen in the office.		
		Medication list: Significant for Mirena IUD 20 mcg/24 hour; Terazol 3 Vaginal Cream 0.8 %		
		Review of systems: Genitourinary: Frequency, irregular menses		
(Physical examination:		
		Vagina: Scant white discharge present Cervix: Mild bleeding present		
		Assessment: Gynecological exam		
		Plan: Return in 1 year for physical examination. Patient given Essure brochure		
10/21/2011	XXXX	Cytology report: Collected date: 10/14/2011	MR - XXXX Women`s	1432

DATE	PROVIDER	OCCURRENCE/TREATMENT	BATES REF	PDF REF
	XXXX, Cytotechnologis t (ASCP)	Source: Cervical; endocervical	Health - 7.3.08 to 4.5.12 - 000013	
		Interpretation:		
		Unsatisfactory for evaluation.		
		Adequacy: Specimen processed and examined, but unsatisfactory for evaluation because of insufficient squamous component. The preparation consists almost		
		entirely of red blood cells		
04/05/2012	XXXX XXXX, M.D.	Office visit regarding permanent sterilization: This is a 38-year old Black/African American female G 3, P 2 whose LMP was 04/2008. She requests permanent sterilization. Her menses are currently irregular secondary to IUD. The patient's past medical and social history are notable for colon cancer. She denies a history of smoking, hypertension, and diabetes. The patient has had Mirena IUD since 2008. She requests Bilateral Tubal Ligation (BTL). Review of systems: Genitourinary: Irregular bleeding Height: 5 feet 3 inches; Weight: 215 lbs; BMI: 38.15 kg/m2	MR - XXXX Women's Health - 7.3.08 to 4.5.12 - 000001-000003	1420- 1422
		Physical examination:		
	10	Genitourinary: External Genitalia: Bloody discharge Cervix: Blood present in os, closed		
		Assessment:		
		Abnormal uterine bleedingContraceptive management		
		Plan:		
		Discussed with patient irregular bleeding not uncommon with the Mirena IUD		
		Discussed with patient that since she is not having any problems with her Mirena IUD would continue until		
		effectiveness has expired.		
04/07/2012	XXXX	Will plan to follow-up as needed. Labs:	MR - XXXX	1423
04/07/2012	ΛΛΛΛ	Laus	ΙνΙΙΧ - ΛΛΛΛ	1423

DATE	PROVIDER	OCCURRENCE/TREATMENT	BATES REF	PDF REF
		Collected date: 04/05/2012	Women's Health - 7.3.08	
		Chlamydia trachomatis, Neisseria gonorrhoeae:	to 4.5.12 -	
		Negative	000004	
06/07/2012	XXXX Medical	ER Follow-up visit for fever and pain:	MR - XXXX	110-
	Center		Medical Center - 000110-000111	111
	XXXX, N.PC.	Sexual activity: Monogamous relationship. Regular condom use, careful partner selection	000110-000111	
	AAAA, N.FC.	condom use, careful partner selection		
		*Reviewer's comment: Only case relevant information		
		captured from this visit.		
10/18/2012	XXXX Medical	Follow-up visit for hypertension:	MR - XXXX	97-
	Center	Complete Management (1) Production	Medical Center - 000097-000100	100
	XXXX, N.PC.	Sexual activity: Monogamous relationship. Regular condom use, careful partner selection		
	777771, 11.1 . C.	condom use, careful parties selection		
		*Reviewer's comment: Only case relevant information		
		captured from this visit.		
02/27/2013	XXXX Medical	Office visit for urinary symptoms:	MR - XXXX Medical Center -	85-86
	Center	Urinary frequency, moderate symptoms, suprapubic	000085-000086	
	XXXX, M.D.	pain/pressure		
	,	Urinalysis: Leukocyte esterase: 2+, Nitrite, blood: 1+		
		Assessment: UTI		
		Plan: Bactrim DS 150 mg, Diflucan 150 mg twice daily		
		Follow-up as needed.		
03/12/2013	XXXX	Office visit for bleeding, vaginal odor:	MR - XXXX Women`s	1079- 1081
	Women's Health	This is a follow-up visit for this 39-year old	Health -	1001
	Ticatui	Black/African American female, Gravida 3 Para 2, who	000001-000003	
	XXXX, M.D.	had an Intrauterine Device (IUD) placed 07/03/2008.		
		Her LMP was 5 years ago. The patient states she has not		
	XXXX, N.P.	had menses since having IUD placed. Since the IUD		
		placement, the patient reports non-menstrual bleeding. She denies dyspareunia, heavy bleeding, and fever. She		
		states she has had spotting off and on for the past two		
		months.		
		The patient also complains of vaginal discharge with		
		foul odor to it.		

DATE	PROVIDER	OCCURRENCE/TREATMENT	BATES REF	PDF REF
		She notices this when she is spotting		
		Medication: Mirena Intrauterine IUD 20 mcg/24 hour		
		Reproductive history: Gravida 3 Para 2 0 2 2 and		
		premenopausal		
		premenopausu		
		Review of systems: Genitourinary: Irregular menses,		
		vaginal odor		
		Physical examination		
		Constitutional: Appearance: Obese		
		Genitourinary External conitalia: Normal apparatus for accura		
		External genitalia : Normal appearance for age, no discharge or inflammatory lesions present		
		Vagina: Normal appearing vaginal vault, well		
		estrogenized, well-rugated, blood-tinged discharge		
		present, without significant rectocele		
		Bladder: Nontender to palpation		
		Urethra		
		Urethral Body: Urethra palpation normal, urethra		
		structural support normal		
		Urethral Meatus: No erythema or lesions present		
		Cervix: Healthy appearance, without abnormal lesions,		
		nontender to palpation, no abnormal discharge, no		
		abnormal bleeding, RID string present		
		Uterus: Nontender to palpation, no masses present, Contour: Smooth to palpation, position		
		midline/midplane, size normal, shape normal, mobility:		
		normal		
•		Adnexa: No adnexal tenderness present, no adnexal		
		masses present		
		Perineum : Perineum within normal limits, no evidence		
		of trauma, no rashes or skin lesions present		
		Anus: Anus within normal limits, no hemorrhoids		
		present		
		T 000		
		In office procedure results:		
		Pregnancy test: Negative Wet Mount:		
		Trichomonas: Negative		
		Clue Cells: Positive		
		Ciac Cons. I ositive		

DATE	PROVIDER	OCCURRENCE/TREATMENT	BATES REF	PDF REF
		Hyphae: Negative		
		Buds: Negative		
		Whiff Test: Positive		
		Assessment		
		 Irregular menses 		
		 Bacterial vaginitis 		
		Plan: Flagyl 500 mg, Provera 10 mg		
		Follow-up in 2 weeks.		
03/15/2013	XXXX Health	Transabdominal ultrasound of the pelvis:	MR - XXXX	1082
			Women's	
	XXXX	History: IUD placement	Health - 000004	
	(Credentials	Impression:		
	unknown)	Echogenic structure consistent with an IUD is		
		demonstrated in the uterus within the mid to		
		upper aspect toward the fundus.		
		 Unremarkable appearance of the ovaries on transabdominal scanning 		
05/06/2013	XXXX	Office visit to discuss about bilateral tubal ligation:	MR - XXXX	1084-
03/00/2013	Women's	office visit to discuss about shateful tusul lighton.	Women's	1085
	Health	This is a 39-year old Black/African American female G	Health -	
		3, P 2 whose LMP was spotting. She requests BTL. Her	000006-000007	
	XXXX, D.O.	menses are currently irregular have normal flow, last for		
		2 days, and are without significant dysmenorrhea. The		
		patient's past medical and social history are		
		unremarkable. She denies a history of smoking. The		
		patient has been using IUD 5 years. She requests BTL.		
		Medication: Mirena IUD 20 mcg/24 hour, Provera oral		
		tablet 10 mg		
		Physical examination:		
		Abdominal Examination: Abdomen nontender to		
		palpation, tone normal without rigidity or guarding, no masses present, healed midline incision present,		
		umbilicus without lesions.		
		Assessment: Sterilization		
		Diana Instructions, Madicaid array day 1 (1)		
		Plan: Instructions: Medicaid papers signed today. Will do when mature		
		do when mature		

DATE	PROVIDER	OCCURRENCE/TREATMENT	BATES REF	PDF REF
05/20/2013	XXXX Medical	Follow-up visit for evaluation:	MR - XXXX	82-84
	Center		Medical Center -	
		Sexual activity: Monogamous relationship. Regular	000082-000084	
	XXXX, N.PC.	condom use, careful partner selection		
		*Reviewer's comment: Only case relevant information		
07/09/2012	VVVV	captured from this visit.	MR - XXXX	1089-
07/08/2013	XXXX Women's	Office visit to discuss permanent sterilization:	Women's	1089-
	Health	Chief complaint: Desires permanent sterilization	Health -	1052
	Ticarui	Cinci complaint. Desires permanent stermzation	000011-000014	
	XXXX, D.O.	This is a 39-year old Black/African American female G3		
	, D.G.	P2022 whose LMP was no cycles due to IUD.		
		Chief complaint:		
		Desires permanent sterilization. Duration:		
		05/06/2013. Other complaints: No problems. She uses		
		Mirena IUD for contraception.		
		Last Pap: 10/14/2011		
		She comes in for discussion of surgery consisting of		
		laparoscopic BTL secondary to desire for sterilization.		
		She will have surgery on 07/10/2013 at XXXX Medical		
		Center.		
		District		
		Discussion of procedure:		
		Patient appears to understand the risks/benefits/alternatives of the procedures and wishes		
		to proceed with the planned procedure. She appears to		
		understand the risk of bleeding that could result in a		
		blood transfusion and the risks of a blood transfusion		
		including, but not limited to: Allergic reaction; and		
		developing Hepatitis, Acquired Immunodeficiency		
		Syndrome (AIDS), or some other blood borne infection.		
		She also appears to understand the risk of infection and		
		cardiac arrest. She also appears to understand the		
		specific risks associated with this/these particular		
		procedures as outlined on the Operative Permit patient		
		appears to understand that sterilization is a permanent		
		procedure. She also appears to understand the failure		
		rate of 1/200 women years (i.e. if 200 women have		
		sterilization this year you can expect on average for 1 of		
		these 200 women each year to get pregnant). In addition,		

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		she appears to understand that should she become		
		pregnant that there is a greater chance than usual of the		
		pregnancy being an ectopic pregnancy and the need for		
		immediate evaluation should she have abnormal		
		menses/bleeding, amenorrhea (i.e. miss her period), or		
		have low abdominal/pelvic pain.		
		Patient also appears to understand that there are methods		
		of contraception (birth control) other than permanent		
		sterilization including, but not limited to, Depo-Provera,		
		birth control pills, contraceptive patch, contraceptive		
		ring, barrier methods of contraception, etc. but she		
		wishes to proceed with permanent sterilization.		
		Medication list: Mirena Intrauterine IUD 20 mcg/24		
		hours, 07/03/2008 place 1 device by intrauterine route		
		once a day for 1825 days; Provera oral tablet 10 mg		
		office a day for 1625 days, 110vera of a tablet 10 mg		
		Reproductive history: Age Menarche: 11; Last		
		menstrual period: 08/01/2007. Menses duration: 3 days.		
		Menopause status: Premenopausal. Flow: Light.		
		Pregnancy summary:		
		Total Pregnancies: 3		
		Full Term: 2		
		Premature: 0		
		Abortion induced: 1		
		Abortion spontaneous: 1		
		Ectopic: 0		
		Multiples: 0		
		Living: 2		
		Pregnancy details: 04/30/2008; birth weight 6 lbs;		
		vaginal delivery, epidural anesthesia. Complications:		
		Chronic HTN, oligo.		
		Weight: 202 lbs.		
		Height: 5 feet 3 inches (Taken from visit dated		
		07/08/2013)		
		BMI: 35.8 kg/m2 (Calculated)		
		Assessment: Sterilization		
		Plan:		
		Abstain from sexual intercourse for 2 weeks		

DATE	PROVIDER	OCCURRENCE/TREATMENT	BATES REF	PDF REF
		Anticipate pain in shoulders radiating to the neck		
		Hospital discharge instructions sheet given to patient		
		Proceed with Laparoscopic Sterilization		
07/12/2013	XXXX	Office visit for discussion of permanent sterilization:	MR - XXXX	1093-
	Women's		Women's	1094
	Health	Patient comes in for discussion of Permanent	Health - 000015-000016	
		Sterilization. Her menses are absent due to IUD. Patient	000013-000016	
	XXXX, M.D.	was seen by Dr. XXX but felt that all her questions were not addressed.		
		She uses Mirena IUD for contraception. Placed on		
		07/03/2008.		
		Her last Pap Smear was 10/14/2011 and was		
		unsatisfactory. She denies a history of abnormal Pap		
		smears. Last mammogram done on 10/27/2011 with		
		negative results at MPMC.		
		Patient appears to understand that Sterilization is a		
		permanent procedure. She also appears to understand the		
		failure rate of 1/200 women years (i.e. if 200 women		
		have sterilization this year you can expect on average for		
		1 of these 200 women each year to get pregnant). In		
		addition, she appears to understand that should she		
		become pregnant that there is a greater chance than usual		
		of the pregnancy being an ectopic pregnancy and the		
		need for immediate evaluation should she have abnormal		
		menses/bleeding, amenorrhea (i.e. miss her period), or		
		have low abdominal/pelvic pain. Patient also appears to		
		understand that there are methods of contraception		
		(Birth control) other than permanent sterilization		
		including, but not limited to, Depo-Provera, birth control		
		pills, contraceptive patch, contraceptive ring, barrier		
		methods of contraception, etc. but she wishes to proceed		
		with permanent sterilization.		
		She reports normal menses without significant		
		dysmenorrhea even when not on hormonal		
		contraception.		
		Essure and BTL Procedures, risks, benefits, and		
		alternatives discussed with patient and she wishes to		
		proceed with Essure.		
		She understands that she will need to use a good method		
		of contraception until HSG confirms bilateral tubal		
		occlusion which is generally 3-6 months after placement		
		of the Essure devices.		

DATE	PROVIDER	OCCURRENCE/TREATMENT	BATES REF	PDF REF
		Essure Medicaid Consent signed on 05/06/2013.		
		Physical examination:		
		Cervix: IUD string present and appears normal length		
		Pregnancy test: Result: Negative		
		Assessment: Essure Counseling		
		Plan:		
		Depo-Provera Intramuscular suspension 150 mg/mL		
		Inject 150 mg by intramuscular route every 3 months for		
		1 day		
		Dispensed: 1 ml vial with 1 refill		
		Xanax oral tablet 2 mg; Ketorolac IM solution 60 mg/2		
		ml, Vicodin oral tablet 5-300 mg		
		Instructions:		
		Schedule Essure procedure		
		Essure instructions for sterilization:		
		Take Ibuprofen 800 mg (e.g. Advil; four 200 mg tablets)		
		by mouth every 8 hours to begin two days prior to the		
		Essure Procedure then every 8 hours after the procedure		
		as needed for pain. I would not expect you to need it for		
		longer than 1-2 days after the procedure.		
		Take Vicodin, 2 tablets by mouth 30 minute prior to		
		appointment then 1 tablet every 4-6 hours as needed for		
		pain after procedure.		
		Have someone drive for you since Vicodin is a narcotic.		
		Take Xanax 1 tablet 2 mg when you take the pain		
		medication.		
		Fill prescription for Toradol and bring to office for injection.		
		Depo today		
		Leave Mirena IUD in place to help with menses and		
		dysmenorrhea even though we can't depend on it for		
		contraception		
07/12/2013	XXXX	Procedure note:	MR - XXXX	1095
	Women's		Women's	
	Health	The patient was injected with Depo Provera on	Health - 000017	
		07/12/2013 using sterile procedure.		
	XXXX, M.D.	Dose: 1 ml		

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		Site: Right Gluteal IM		
	XXXX, L.P.C.	Manufacturer: Greenstone		
		Lot Number: 645413.		
		Expiration date: 01/31/2016		
		Patient was monitored and there were no adverse effects.		
		Injection Given By: A. Milton, LPN		
		Assessment: Initiation Depo-Provera		
		Return to clinic for Essure placement		
07/23/2013	XXXX	Procedure note for Ketorolac injection:	MR - XXXX	1097
	Women's		Women's	
	Health	The patient was injected with Ketorolac on 07/23/2013	Health - 000019	
		using sterile procedure.		
	XXXX, M.D.	Dose: 2 ml		
		Site: Right Gluteal IM		
	XXXX, L.P.C.	Manufacturer: Wockhardt		
		Lot Number: DN10751.		
		Expiration Date: 01/31/2015		
		Patient was monitored and there were no adverse effects.		
		Assessment: Sterilization		
07/23/2013	XXXX	Procedure note for IUD removal:	MR - XXXX	1096
	Women's		Women's	
	Health	Indications:	Health - 000018	
		Patient presents today for IUD removal. Her current IUD		
	XXXX, M.D.	was placed 07/03/2008. She has not had any problems.		
		She requests removal of the IUD because she wants		
		permanent sterilization with Essure. The IUD removal		
		procedure was discussed with the patient and her		
,		questions were answered.		
		Procedure:		
		The patient was placed in a dorsal lithotomy position		
		and appropriately draped. A speculum exam was		
		performed, and the cervix was visualized. The IUD		
		string was visualized. Using ring forceps, the string was		
		grasped, and the IUD removed intact.		
		Post procedure status:		
		The patient tolerated the procedure well with minimal		
		bleeding or pain. Patient was discharged in stable		

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		condition.		
		Assessment: IUD removal		
		Plan: Removal of intrauterine device (IUD) (58301) - 07/23/2013		
		Instructions		
		Call if bleeding, pain or fever occur		
		Birth control counseling given		
07/23/2013	XXXX	Procedure report for Essure implantation:	MR - XXXX	1098-
	Women's		Women's	1099
	Health	This G 3 P 2 0 2 2, LMP has been limited to spotting	Health - 000020-000021	
		since Depo Provera given on 07/12/2013 presents for	000020-000021	
	XXXX, M.D.	office hysteroscopic insertion of bilateral Essure		
		fallopian tube micro-inserts for permanent sterilization.		
		Patient denies to being allergic to nickel. All consents		
		have been signed.		
		Education and consent:		
		She has read the Essure pamphlet and has seen the		
		Essure patient instructional DVD. All her questions have been answered to her satisfaction. She has confirmed to		
		me that she understands this procedure is intended to		
		provide permanent and irreversible sterilization, yet she		
		is also aware that there is a small failure rate that could		
		result in either an intrauterine and/or an ectopic		
		pregnancy. She has indicated to me that she does not		
		wish to ever become pregnant again under any		
		circumstances or contingencies of life. Patient confirmed		
		to me that she understands that the Essure micro-inserts		
		should not be relied on for contraception until she has		
		undergone a Hysterosalpingogram (HSG) which		
		demonstrates bilateral tubal occlusion and satisfactory		
		location of the micro-inserts. She understands that this		
		generally occurs about 3 months after insertion of the		
		micro-insert devices. She also understands that if		
		Bilateral tubal occlusion has not occurred by this 3-		
		month HSG that she will undergo another HSG about 6		
		months after the initial insertion of the micro-insert		
		devices to demonstrate bilateral tubal occlusion and		
		satisfactory location of the micro-inserts. The patient has		
		also confirmed to me that she understands that if the		

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		Essure micro-inserts cannot be placed bilaterally, then		
		she will not rely on this method of sterilization, as		
		Essure has not been proven to be effective when it is		
		placed unilaterally. She has also confirmed to me that		
		she understands that the Essure product is intended to		
		prevent pregnancy and does not protect against either		
		HIV infection or other sexually transmitted diseases.		
		Urine pregnancy test:		
		A urine pregnancy test was performed today, and the		
		result was negative.		
		A "Time-Out" was undertaken to confirm that this was		
		indeed patient and that the intended procedure was a	•	
		voluntary placement of an Essure Permanent Birth		
		Control System under paracervical block and oral		
		analgesics.		
		Pre-operative medications given to patient included:		
		Ibuprofen 800 mg every 6 hours for 48 hours pre-		
		procedure, Toradol 60 mg IM, Vicodin 5/500, and		
		Xanax 1 mg.		
		Procedure:		
		All equipment was checked to ensure that there was no		
		damage and no missing parts. Patient was placed in the		
		lithotomy position and draped appropriately. A metal		
		bivalve speculum was introduced into her vagina and the		
		cervix was prepped with Betadine. A paracervical block		
		was placed with a total of 20 cc of 1% Lidocaine (plain)		
		using a 22-gauge spinal needle.		
	X V	Approximately 25 minutes were allowed to elapse for		
		the paracervical block to take maximal effect. During		
		that time, final preparations were undertaken to connect		
		the camera, light source, sealing cap, fluid inflow and		
		outflow tubing to the operative sheath of the 3-mm rigid		
		Storz hysteroscope. The hysteroscope was focused, a		
		white balance was performed, and the inflow/outflow		
		functions of the sheath were checked. All bubbles were		
		flushed out of the equipment.		
		A single tooth tenaculum was placed on the anterior lip		
		of the cervix. Charlene XXXX was then told that the		
		hysteroscope was about to be inserted into her cervix,		

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		and she was invited to view the procedure on the video		
		monitor. I described the pertinent anatomy to her.		
		Cervical dilatation was not needed. Body temperature		
		0.9% normal saline solution was used to gently distend		
		the uterus with use of a BP Cuff (Maximum of 150 mm		
		Hg) over the bag of distension medium. A total of 700		
		cc was used to distend the uterus during the procedure.		
		We retrieved 250 cc resulting in a loss of less than 450		
		cc of distension medium for the procedure. The		
		procedure lasted about 20 minutes.		
		A thorough hysteroscopic uterine cavity assessment was		
		carried out. Both tubal ostia were seen clearly, giving the		
		expectation of successful bilateral placement of the		
		Essure micro-inserts. The Valved, DryFlow Introducer		
		the sealing cap on the hysteroscope operating channel.		
		The Essure delivery catheter was inserted through the		
		introducer and advanced through the operating channel.		
		The Essure delivery catheter was then advanced into the		
		proximal left fallopian tube with gentle, constant		
		forward movement in an effort to minimize tubal spasm.		
		The catheter was then advanced until the black		
		positioning marker reached the fallopian tube ostium,		
		indicating that the Essure micro-insert was spanning the		
		intramural and the proximal isthmic segments of the		
		fallopian tube, with the outer coil spanning the		
		uterotubal junction. The handle of the Essure micro-		
		insert was then stabilized against the hysteroscope and		
		camera to prevent inadvertent forward movement of the		
		micro-insert during retraction of the delivery catheter.		
`		After once again visually confirming that the positioning		
		marker was at the tubal ostium, I rotated the thumb-		
		wheel on the handle back towards myself at a rate of one		
		click per second until the wheel stopped to withdraw the		
		delivery catheter. The black positioning marker moved		
		away from the tubal ostium and disappeared out of view		
		into the hysteroscope operating channel.		
		At this point the Cold Band notch was seen to be located		
		just outside the tubal ostium and the Green Release		
		Catheter was in view. While continuing to stabilize the		
		handle against the camera and hysteroscope, the button		
		on the handle was then depressed to initiate deployment		

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		of the micro-insert device. The thumbwheel was then		
		once again rotated back to a Hard Stop withdrawing the		
		Green Release Catheter which allowed the outer coils of		
		the Essure micro insert to expanded and detach from the		
		Essure Delivery Catheter. The delivery system was		
		gently withdrawn from the micro-insert by pulling the		
		handle backwards. Once the delivery system was		
		withdrawn, the position of the Essure micro-insert was		
		assessed.		
		The number of expanded coils that appeared trailing into		
		the uterine cavity was 3.		
		The identical steps were undertaken to insert the Essure		
		micro-insert in the opposite tube.		
		The number of expanded coils that appeared trailing into		
		the uterine cavity was 5.		
		She tolerated the procedure well, stating she had only		
		mild cramps at most.		
		Post procedure information:		
		Patient was given the patient ID card provided by the		
		Conceptus company showing the micro-insert reference		
		#ESS305 and Lot #927081. She was asked to carry it		
		with her at all times and show it to other physicians		
		involved in her present or future care. She was also		
		given personalized "Post-operative Guidelines," which		
		are scanned into this chart.		
		We once again reiterated to the patient that she must use		
		an alternative form of contraception until bilateral tubal		
		occlusion and satisfactory location of the micro-inserts		
		could be demonstrated by the HSG. She plans to use		
		Depo Provera for contraception during this time period. I		
		also warned her that there is theoretically an increased		
		risk of ectopic pregnancy during this time period,		
		therefore compliance with a good contraceptive regimen		
		is crucial.		
		She was advised that the IUD/IUS could become		
		entangled with the micro-inserts resulting in failure of		
		tubal occlusion, but she still wishes to use this method of		
		contraception until tubal occlusion is demonstrated.		
		She is being scheduled for HSG to be done in three		
		months. We have discussed that only if micro-insert		
		location is satisfactory and there is evidence of bilateral		

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		tubal occlusion will we then instruct her to discontinue use of her alternative contraception and rely on the Essure micro-inserts for pregnancy prevention. Mirena IUD was blocking bilateral tubal ostia and had to be removed to insert the devices.		
		Assessment: Sterilization with Essure		
		Plan: HCG urine, Surgical hysteroscopy with bilateral fallopian tube cannulation 07/23/2013. Anesthesia paracervical block 4 injection, Ketorolac Tromethamine, per 15 mg Instructions: Depo-Provera until HSG confirms tubal occlusion Return to clinic in 1 week		
		HSG will be ordered in 3 months.		
07/29/2013	XXXX Women's Health XXXX, M.D.	Patient reports for follow-up of Essure on 07/23/2013. Her LMP was 08/01/2007. She is premenopausal. She uses Depo-Provera for contraception. She reports irregular menses since Depo Provera. Patient has been spotting intermittently. Patient states doing well since Essure procedure. Patient denies fever and chills. Admits to mild menstrual like cramping. Patient also complains of increased night sweats for one week. Urinalysis: Protein: Trace; Blood: 2+ Assessment: • Sterilization with Essure • Microscopic hematuria Plan: Urina culture	MR - XXXX Women's Health - 000022-000023	1100-1101
		Urine culture HSG in 3 months then follow-up appointment afterwards		
09/12/2013	XXXX Institute	Office visit for lumbar pain:	MR - XXXX - 000421-000426	1962- 1967
	XXXX, M.D.	Review of systems: Neuro: Numbness and tingling in		

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		legs. Psych: Anxiety, depression		
		*Reviewer's comment: Only case relevant information		
10/22/2012	*****	captured from this visit.	NO MANA	1100
10/23/2013	XXXX	Procedure note:	MR - XXXX Women`s	1102
	Women's	The noticest was injected with Dane Provens on	Health - 000024	
	Health	The patient was injected with Depo Provera on 10/23/2013 using sterile procedure.		
	XXXX, M.D.	10/23/2013 using sterile procedure.		
	MAMA, M.D.	Dose: 150 mg		
	Felicia Spruill,	Site: Right gluteal IM		
	C.M.A.	Manufacturer: Greenstone		
		Lot Number: G41971		
		Expiration Date: 02/2016		
		Patient was monitored and there were no adverse effects.		
		Injection given by: F. Spruill, CMA		
		Assessment: Contraception-follow-up Depo-Provera		
10/28/2013	XXXX Medical	Hysterosalpingogram:	MR - XXXX	1103-
	Center		Women`s Health -	1104
	******	History: For Essure placement check	000025-000026	
	XXXX	D	000020	
	(Credentials	Procedure : Timeout was taken at 0933 to confirm the		
	unknown)	patient's identity, site and procedure. The procedure was performed by Dr Spargo.		
		The speculum was inserted, and the cervix was		
		visualized. Cervix was swabbed ×3 with Betadine. The		
		hysterosalpingogram catheter was inserted and a		
		retention balloon inflated without difficulty. Isovue 300		
		was then injected into the endometrial cavity. The		
		endometrial cavity is normal in size and configuration.		
		No abnormal filling defects are seen. Some intravasation		
		of contrast occurred during the injection. There was no		
		reflux of contrast to either fallopian tube.		
		The Essure implants appear to be in satisfactory		
		position. No additional findings are noted on the post		
		drainage film		
		Impression: The Essure implants appear to be in		
		satisfactory position. There was no intraperitoneal spill		
		of contrast consistent with bilateral occlusion of		
		fallopian tubes. A small amount of intravasation of		

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		contrast occurred during the procedure.		
10/31/2013	XXXX Institute	Follow-up visit for lumbar pain:	MR - XXXX - 000415-000420	1956- 1961
	XXXX, M.D.	Review of systems: Neuro: Numbness and tingling in		
		legs. Psych: Anxiety, depression		
		*Reviewer's comment: Only case relevant information captured from this visit.		
11/11/2013	XXXX	1 0	MR - XXXX	1106-
11/11/2013	Women's	Office visit for urinary frequency:	Women's	1106-
	Health	Patient presents with a history of urinary frequency,	Health - 000028-000030	1100
	XXXX, M.D.	dysuria, and a change in urine color. The problem is described as moderately severe and began 1 day ago.		
	AAAA, W.D.	She reports no additional symptoms. There are no		
	XXXX, N.P.	alleviating factors. There are no aggravating factors. She		
		reports she has not had any recent urinary tract		
		infections. She has not been evaluated for her current		
		complaints. She does not have a history of recurrent		
		urinary tract infections. The patient admits using a		
		sexual enhancement oil yesterday and feels that irritated		
		her. She denies vaginal discharge.		
		Weight: 204 lbs 2 oz		
		Review of systems:		
		Admits: Frequency, dysuria, change in urine color		
		Denies: Vaginal discharge		
		Physical examination:		
	\wedge	Vagina: Normal vaginal vault without central or		
		paravaginal defects, white-colored discharge present, no		
		inflammatory lesions present, no masses present		
		Urinalysis: Blood: 2+; Leukocytes: 2+; Protein: Trace		
		Assessment:		
		Urinary frequency		
		Dysuria		
		Plan: Cipro 500 mg, Phenazopyridine 200 mg		
11/13/2013	XXXX	Urine culture report:	MR - XXXX	1105
		Collected date: 11/11/2013	Women's	

DATE PROVIDE	ER OCCURRENCE/TREATMENT	BATES REF	PDF REF
	Final: Escherichia coli, greater than 100,000 CFU per ml	Health - 000027	
01/02/2014 XXXX Instit	tute Follow-up visit for lumbar pain:	MR - XXXX - 000409-000414	1950- 1955
XXXX, M.D	Anxiety, depression		
	*Reviewer's comment: Only case relevant information captured from this visit.		
01/23/2014 XXXX Women's Health XXXX, M.E XXXX, M.E	Patient presents for the follow-up of HSG done. She uses Essure system for contraception. Her last Pap Smear was 04/25/2013 and was normal per Heather at WCHD.	MR - XXXX Women's Health - 000032-000034	1110-1112

DATE	PROVIDER	OCCURRENCE/TREATMENT	BATES REF	PDF REF
		Repeat HSG because of minimal extravasation of dye at		
		last HSG		
		Note that patient reports she is getting her Annual BPP		
		at Health Departments		
		Return to clinic in 3 months for Depo injection		
01/23/2014	XXXX	Procedure note:	MR - XXXX	1109
	Women's		Women's	
	Health	The patient was injected with Depo Provera on	Health - 000031	
		01/23/2014 using sterile procedure.		
	XXXX, M.D.			
		Dose: 1 ml		
		Site: Right Gluteal TM		
		Manufacturer: Greenstone		
		Lot Number: H94054.		
		Expiration Date: 08/31/2016		
		Patient was monitored and there were no adverse effects.		
		Injection Given By: A. Milton, LPN		
		Assessment: Contraception-follow-up Depo-Provera		
		Plan		
		Depo-Provera non-contraception - 01/23/2014		
		Return to clinic in 3 months for next injection.		
02/27/2014	XXXX Institute	Office visit for lumbar pain:	MR - XXXX -	1946-
			000405-000408	1949
	XXXX, M.D.	Review of systems: Neuro: Numbness and tingling in		
		legs		
		Psych: Anxiety, depression		
		*Reviewer's comment: Only case relevant information		
		captured from this visit.		
04/23/2014	XXXX	Office visit for urinary frequency:	MR - XXXX	1114-
	Women's		Women's	1116
	Health	Medication list: Depo-Provera intramuscular	Health -	
		suspension 150 mg/ml	000036-000038	
	XXXX, D.O.			
		Weight: 212 lbs 4 oz		
	XXXX, M.D.			
		Assessment:		
		Urinary frequency		
		Hematuria		

DATE	PROVIDER	OCCURRENCE/TREATMENT	BATES REF	PDF REF
		Plan: Urine culture. Levaquin 250 mg. Increase fluid intake		
04/23/2014	XXXX Women's Health XXXX, D.O.	Procedure note: The patient was injected with Depo Provera on 04/23/2014 using sterile procedure. Dose: 150 mg	MR - XXXX Women`s Health - 000039	1117
	XXXX, C.M.A.	Site: Left Gluteal IM Manufacturer: Greenstone Lot Number: H47434. Expiration date: 12/2016		
04/25/2014	XXXX	Urine culture report: Collected date: 04/23/2014 Result: Mixed urogenital flora 25,000-50,000 colony forming units per mL	MR - XXXX Women`s Health - 000035	1113
05/01/2014	XXXX Institute XXXX, M.D.	Follow-up visit for lumbar pain: Review of systems: Neuro: Numbness and tingling in legs Psych: Anxiety, depression *Reviewer's comment: Only case relevant information captured from this visit.	MR - XXXX - 000402-000404	1943- 1945
05/21/2014	XXXX Women's Health XXXX, M.D.	Office visit for annual examination and Depo: Patient complains of Right Lower Quadrant (RLQ) pain that has been present for a while and comes in goes in spells. She had a recent MRI that showed a right adnexal multisepated cyst. Her LMP was no cycles secondary to Depo and has had Essure 7/2013 by Dr. XXXX. She denies pain and bleeding associated with intercourse. She uses Essure since 07/2013. She uses Depo every 3 months to help with dysmenorrhea for contraception. Her last Pap Smear was 04/25/2013 and was normal. She denies dyspareunia and dysuria. She admits to dysmenorrhea and RLQ pain. She denies smoking. Medication: Depo-Provera intramuscular suspension	MR - XXXX Women`s Health - 000041-000044	1119-1122

DATE	PROVIDER	OCCURRENCE/TREATMENT	BATES REF	PDF REF
		150 mg/ml Review of systems: Genitourinary: Dysmenorrhea, RLQ pain		
		Weight: 212 lbs 0.4 oz; Height 5 feet 3 inches; BMI: 37.56 kg/m2		
		Physical Examination:		
		Constitutional: Appearance: Obese		
		Uterus: Retroflexed, uterine size-8 weeks 9 weeks,		
		globular shape, soft,		
		Assessment:		
		Ovarian cyst		
		Uteromegaly		
		Plan:		
		Get pelvic Ultrasonogram (USG) for uteromegaly and		
		cystic structure seen on right ovary on recent MRI		
		Return to clinic 4-5 days after pelvic USG		
05/26/2014	XXXX	Cytology report:	MR - XXXX	1118
		Collected date: 05/21/2014	Women's Health - 000040	
	XXXX,		11eaiui - 000040	
	Cytotechnologis	Source: Cervical; Endocervical		
	t (ASCP)	Interpretation: Negative for intraepithelial lesion and		
		malignancy		
05/30/2014	XXXX Medical	Ultrasound pelvis complete with transvaginal:	MR - XXXX	1123-
	Center	History: Right-sided pain.	Women's Health - 000045-000046	1124
	XXXX (Credentials unknown)	Comparison : 03/15/2013.		
	unknown)	Findings:		
		Transvaginal : Essure prostheses are seen in the fundus		
		bilaterally expected location fallopian tube origins.		
		Impression:		
		Normal uterus and endometrium.		
		 Normal ovaries. 		
		Small amount of free fluid likely physiologic.		

DATE	PROVIDER	OCCURRENCE/TREATMENT	BATES REF	PDF REF
		Small echogenic foci seen in the fundus on each		
		side consistent with Essure prostheses.		
06/02/2014	XXXX	Follow-up visit of USG:	MR - XXXX	1125-
	Women's		Women`s Health -	1127
	Health	Her LMP was 08/01/2007. Her menstrual periods are	000047-000049	
		abnormal secondary to Depo. She uses Essure for	000017 000019	
	XXXX, M.D.	contraception. Her last Pap Smear was 05/21/2014 and was normal.		
		Medication list: Depo-Provera intramuscular		
		suspension 150 mg/ml		
		Weight: 215 lbs; Height: 5 feet 3 inches; BMI: 38.09		
		kg/m2		
		Physical examination:		
		Constitutional: Appearance: Obese		
		Assessment:		
		Abdominal pain, lower abdomen		
		Ovarian cyst-resolved		
		Follow-up as needed		
06/05/2014	XXXX Institute	Follow-up visit for lumbar pain:	MR - XXXX -	1938-
			000397-000401	1942
	XXXX, M.D.	Review of systems: Neuro: Numbness in extremities		
		*Reviewer's comment: Only case relevant information		
		captured from this visit.		
06/30/2014	XXXX	Office visit for joint pain:	8B7B13342F89	1325-
30,20,2014	Associates	Anna tor Journ Kanna	4C66BF24,	1329
		Review of systems: Endocrine: Fatigue	XXXX, 145-149	
	XXXX, M.D.			
		*Reviewer's comment: Only case relevant information		
		captured from this visit.		
07/24/2014	XXXX	Procedure note:	MR - XXXX	1128
	Women's		Women's	
	Health	The patient was injected with Depo Provera on	Health - 000050	
	*******	07/24/2014 using sterile procedure.		
	XXXX, M.D.	Dose: 1 ml		
	VVVV CMA	Site: Right Gluteal IM Manufacturer: Greenstone		
	XXXX, C.M.A.	wanuracturer: Greenstone		

DATE	PROVIDER	OCCURRENCE/TREATMENT	BATES REF	PDF REF
		Lot Number: J27858.		
		Expiration Date: 02/2017		
		Weight 212.6 lbs		
		Patient was monitored and there were no adverse effects.		
		Assessment: Contraception-Follow-up Depo-Provera		
		Return to clinic in 3 months for next injection		
07/24/2014	XXXX Institute	Follow-up visit for lumbar pain:	MR - XXXX - 000393-000396	1934- 1937
	XXXX, M.D.	Review of systems:		
		Neuro: Numbness in extremities		
		Psych : anxiety, depression, insomnia		
		*Reviewer's comment: Only case relevant information		
		captured from this visit.		
08/17/2014	XXXX Health	CT scan of abdomen and pelvis:	MR - XXXX	165-
			Medical Center - 000165-000166	166
	XXXX	Indications: Periumbilical pain with nausea and	000103-000100	
	(Credentials	vomiting. History of colonic resection for colonic cancer		
	unknown)	and cholecystectomy		
		Comparison : Comparison was made with prior from		
		08/15/2012		
		00/10/2012		
		Findings:		
		CT pelvis: There are Essure coil devices identified in the		
		fallopian tubes bilaterally		
		Impression:		
		Findings suggestive of early or mild small bowel A structure with transition point poor the		
		obstruction with transition point near the		
		umbilicus, likely representing adhesions from prior periumbilical surgery. No evidence of free		
		air.		
		Free fluid in the mesenteric folia and		
		dependently within the pelvis, likely reactive.		
		No loculated fluid collections to indicate		
		abscess.		
		Normal appendix.		
		Status post cholecystectomy		

DATE	PROVIDER	OCCURRENCE/TREATMENT	BATES REF	PDF REF
08/26/2014	XXXX Medical Center XXXX, M.D.	Office visit for urinary symptoms: Patient presents with 1 day of urinary symptoms. States she has pressure in the lower stomach and the very painful at the end, peeing more and small volume. Tried AZO OTC on yesterday. Urinary frequency: Yes (Severe) Urgency: Yes Small volume voids: Yes (Moderate) Symptom severity: Severe Suprapubic pain/pressure: Yes (Severe)	MR - XXXX Medical Center - 000070-000071	REF 70-71
		Physical examination: Suprapubic: Mildly tender Dip urinalysis: Leukocyte esterase: 2+, Blood: 3+ Assessment: UTI Supportive care: Increase fluids. Follow-up: 1 week		
08/28/2014	XXXX Institute XXXX, M.D.	Follow-up visit for lumbar pain: Review of systems: Neuro: Numbness Psych: Anxiety, depression *Reviewer's comment: Only case relevant information captured from this visit.	MR - XXXX - 000388-000392	1929- 1933
08/26/2014 & 10/20/2014	XXXX Associates XXXX, M.D.	Follow-up visits for joint pain Review of systems: No fevers, no hair loss. Abdominal pain is much better. No muscle weakness *Reviewer's comment: Only case relevant information captured from these visits.	8B7B13342F89 4C66BF24, XXXX, 141-145 8B7B13342F89 4C66BF24, XXXX, 132-136	1321- 1325, 1312- 1316
10/28/2014	XXXX Women's Health XXXX, M.D.	Procedure note: The patient was injected with Depo Provera on 10/28/2014 using sterile procedure. Dose: 150 mg Site: Right Gluteal IM	MR - XXXX Women's Health - 000051	1129

DATE	PROVIDER	OCCURRENCE/TREATMENT	BATES REF	PDF REF
	XXXX, LPN	Manufacturer: Greenstone		
		Lot Number: 381511		
	XXXX	Expiration Date: 05/2017		
	(Credentials	Weight 209.8 lbs		
	unknown)	Patient was monitored and there were no adverse effects.		
		Injection given by XXXX, LPN		
		Assessment: Contraception-follow-up Depo-Provera		
		Instructions: Return to clinic in 3 months for next		
		injection		
10/30/2014	XXXX Institute	Follow-up visit for lumbar pain:	MR - XXXX - 000383-000387	1924- 1928
	XXXX, M.D.	Review of systems:		
	7171711, WI.D.	Neuro: Numbness in extremities		
		Psych: Anxiety, depression, insomnia		
		*Reviewer's comment: Only case relevant information		
		captured from this visit.		
12/18/2014	XXXX Institute	Follow-up visit for lumbar pain:	MR - XXXX -	1916-
			000375-000379	1920
	XXXX, M.D.	Review of systems:		
		Neuro: Numbness in extremities		
		Psych : Anxiety, depression, insomnia		
		*Reviewer's comment: Only case relevant information		
		captured from this visit.		
01/18/2015	XXXX Health	CT of the abdomen and pelvis with contrast:	MR - XXXX	159-
			Medical Center -	160
	XXXX	History : Diffuse abdominal pain. Vomiting. History of	000159-000160	
	(Credentials	small bowel obstruction. History of colon cancer. Status		
	unknown)	post partial colectomy. Rule out obstruction.		
		Findings:		
		CT pelvis: Essure implants are noted in the fallopian		
		tubes		
		Impuggion		
		Impression:		
		No obstruction. The state of the state		
		• There is a suture line in the colon in the area of		
		the splenic flexure consistent with the history of		
		a partial colectomy.		

DATE	PROVIDER	OCCURRENCE/TREATMENT	BATES REF	PDF REF
		Status post cholecystectomy.		
		 Essure implants are noted in the fallopian tubes. 		
		There are multiple small metallic densities along		
		the anterior peritoneal surface at just above the		
		level of the umbilicus consistent with repair of a		
		ventral hernia.		
02/10/2015	XXXX	Procedure note:	MR - XXXX	1130
	Women's		Women's	
	Health	The patient was injected with Depo Provera on	Health - 000052	
		02/10/2015 using sterile procedure.		
	XXXX, M.D.	Dose : 150 mg		
		Site: Right Gluteal IM		
	XXXX, LPN	Manufacturer: Greenstone		
		Lot Number: L22781		
	XXXX	Expiration Date: 09/2017		
	(Credentials	Weight 209.8 lbs		
	unknown)	Patient was monitored and there were no adverse effects.		
	,	Injection given by XXXX, LPN		
		Assessment: Contraception-follow-up Depo-Provera Return to clinic in 3 months for next injection		
03/17/2015	XXXX	Office Visit for possible prolapse:	MR - XXXX	1132-
03/17/2013	Women's	Office visit for possible prolapse:	Women's	1132-
	Health	Patient complains of a month history of patient states "a	Health -	1133
	Health		000054-000057	
	XXXX, M.D.	ball of tissue coming out of her vagina." She states when		
	$\Lambda\Lambda\Lambda\Lambda$, WI.D.	she was able to punch it back up. She also states bleeding when she pushed it		
		bleeding when she pushed it		
		W. 1-14-206 lb - 4		
		Weight: 206 lbs 4 oz		
		Physical everyination		
		Physical examination: Vagina: Frothy smooth yellow-colored discharge		
		present; mild, cystocele present-grade 1, mild prolapse, grade 1 rectocele present, uterine prolapse present-grade		
		2; cervix reaches vaginal introitus with strain		
		Cervix: Mild friability		
		In-office procedures:		
		Wet mount		
		BV Whiff test vagina QI: Negative		

DATE	PROVIDER	OCCURRENCE/TREATMENT	BATES REF	PDF REF
		Clue Cells XXX QI Wet Prep: Few KOH Prep XXX: Neg T vaginalis Ag Genital QI: Negative Yeast Budding # Ur Comp Assist: Negative Assessment		
03/19/2015	XXXX Institute	Follow-up visit for lumbar pain:	MR - XXXX - 000364-000369	1905- 1910
	XXXX, M.D.	Review of systems: Neuro: Numbness and tingling in legs Psych: Anxiety, depression *Reviewer's comment: Only case relevant information captured from this visit.		
03/20/2015	XXXX	Labs: Collected date: 03/17/2015	MR - XXXX Women`s Health - 000053	1131
		Source: Vaginal		

DATE	PROVIDER	OCCURRENCE/TREATMENT	BATES REF	PDF REF
		Result: NuSwab Vaginitis Plus (VG+) Atopobium vaginae High - 2 Score BVAB2 Low-0 score Megasphaera 1. High - 2 Score Candida albicans, NAA: Negative Candida glabrata, NAA: Negative Trichomonas vaginalis by, NAA: Negative Chlamydia trachomatis, NAA: Negative Neisseria gonorrhoeae, NAA: Negative		
05/06/2015	XXXX Women`s Health	Office visit for pelvic pressure and pelvic organ prolapse: Patient complains of a 2-3 months history of pelvic	MR - XXXX Women's Health - 000059-000062	1137- 1140
	XXXX, M.D.	pressure and pelvic pain and discomfort, increased pelvic pressure when sitting down, and urine frequency. She states the symptoms are not accentuated by standing, lifting or straining. The symptoms are not relieved by lying down. The patient also reports urinary frequency.		
		NuSwab + for Bacterial vaginosis (BV) despite mild changes on Wet prep last time similar to this time therefore will treat with Flagyl. Reports intermittent pain with intercourse and generally bad pain after intercourse		
	\Q	Weight: 200 lbs 4 oz Physical examination:		
		Vagina: Grade 1 cystocele present, grade 1 rectocele present, grade 2 uterine prolapse present Wet mount:		
		Bacterial Vaginosis (BV) Whiff test Vagina QI: Negative Clue Cells XXX QI Wet Prep: Positive KOH Prep XXX: Negative. T vaginalis Ag Genital QI: Negative		
		Yeast Budding # Ur Comp Assist: Positive. Assessment		

DATE	PROVIDER	OCCURRENCE/TREATMENT	BATES REF	PDF REF
		 Uterovaginal prolapse, incomplete Microhematuria Bacterial Vaginosis (BV) Candidal vulvovaginitis 		
		Plan: Urine culture Diflucan 150 mg, Flagyl 500 mg Refer to Duke Urogyencology for evaluation and treatment of uterovaginal prolapse		
05/06/2015	XXXX Women`s Health XXXX, M.D. K. Griffin, MA	Procedure note: The patient was injected with Depo Provera on 5/6/2015 using sterile procedure. Dose: 150 mg Site: Left Gluteal IM Manufacturer: Greenstone Lot Number: L58524. Expiration date: 12/2017 Weight 200.4. lbs Patient was monitored and there were no adverse effects. Injection given by K Griffin, MA Assessment: Contraception-follow-up Depo-Provera	MR - XXXX Women`s Health - 000063	1141
05/08/2015	XXXX Medical Center	Return to clinic in 3 months for next injection Urine culture: Collected date: 05/06/2015 Result: No growth	A1062A785498 49D1B73E, XXXX, 401	882
05/21/2015	XXXX Institute XXXX, M.D.	Follow-up visit for lumbar pain: Review of systems: Neuro: Headache, numbness and tingling in legs Psych: Anxiety	MR - XXXX - 000358-000363	1899- 1904
07/23/2015	XXXX Institute	*Reviewer's comment: Only case relevant information captured from this visit. Follow-up visit for lumbar pain:	MR - XXXX - 000353-000357	1894- 1898
	XXXX, M.D.	Review of systems: Neuro: Numbness in extremities Psych: Anxiety, depression, insomnia		

DATE	PROVIDER	OCCURRENCE/TREATMENT	BATES REF	PDF REF
		*Reviewer's comment: Only case relevant information		
08/06/2015	XXXX Women's XXXX, M.D.	*Reviewer's comment: Only case relevant information captured from this visit. Office visit for annual gynecological examination: Patient complains of extreme hot flashes, night sweats, irritability, anxiety, and pelvic pain. Her LMP was no cycles secondary to Depo and has had Essure 07/2013 by Dr. XXXX. She denies bleeding associated with intercourse. She uses Essure for contraception. She denies dyspareunia and dysuria. She admits to dysmenorrhea, hot flashes, and mental instability and pelvic pain Patient reports that Urogyn told her that she had bladder spasms and put her on Oxybutynin that helped with symptoms. Review of systems: Gastrointestinal: Abdominal pain Genitourinary: Dysmenorrhea Psychiatric: Anxiety, depression Weight: 199 lbs 4 oz; height 5 feet 3.5 inches; BMI: 34.74 kg/m2 Physical examination: Vagina: Normal vagina with atrophy, no discharge present, no inflammatory lesions present, no masses present, grade 1 uterine prolapse present Mental status: Mood and Affect: Depressed, flattened Patient teary-eyed when talking about her medical problems and symptoms; decreased estrogenicity and rugations Urinalysis: Hemoglobin strip: 3+	MR - XXXX Women's Health - 000071-000075	1149-1153
		Assessment: Hematuria Ovarian cyst Uteromegaly Fatigue		

DATE	PROVIDER	OCCURRENCE/TREATMENT	BATES REF	PDF REF
		Menopausal symptoms		
		Plan:		
		Get pelvic USG for uteromegaly and cystic structure		
		seen on right ovary on recent MRI		
		Return to clinic 4-5 days after pelvic USG		
08/06/2015	XXXX	Procedure note:	MR - XXXX	1148
	Women's		Women's Health - 000070	
	Health	The patient was injected with Depo Provera on	11cartii - 000070	
	WWW MD	08/06/2015 using sterile procedure.		
	XXXX, M.D.	Dose: 150 mg		
	XXXXX CMA	Site: Right Gluteal IM		
	XXXX, CMA	Manufacturer: Greenstone		
		Lot Number: L50623.		
		Expiration date: 01/2018 Weight 199.4 lbs		
		Patient was monitored and there were no adverse effects.		
		Injection given by T. Wilson/RMA		
		injection given by 1. wilson/kiviA		
		Assessment: Contraception-follow-up Depo-Provera		
		Assessment. Contraception-10110w-up Depo-110vera		
		Return to clinic in 3 months for next injection		
08/07/2015	XXXX Medical	Office visit for weakness:	MR - XXXX	54-55
00/07/2015	Center	Older And Total Medianicos	Medical Center -	0.00
		Patient presents in office complains of feeling very bad	000054-000055	
	XXXX, M.D.	and weak. Patient states that her heart rate dropped to 44		
		for about 3 days in a row. Patient states that she has been		
		feeling agitated		
		Assessment:		
		Early menopause		
		Anxiety		
		,		
		Plan:		
		Continue to hold the Systolic, continue to take the		
		Lisinopril		
		Check BP and pulse at least 3 times a week and extra if		
		you feel off		
		Follow-up as needed		
08/09/2015	XXXX Medical	Urine culture:	A1062A785498	758-
	Center	Collected date: 08/06/2015	49D1B73E, XXXX, 277-278	759

DATE	PROVIDER	OCCURRENCE/TREATMENT	BATES REF	PDF REF
		Final: 10,000 - 50,000/m Escherichia coli		
08/12/2015	XXXX XXXX Pathology	Cytopathology report: Collected date: 08/06/2015 Cytologic diagnoses:	MR - XXXX Women`s Health - 000069	1147
	Laboratory Associates	 Satisfactory for evaluation. Negative for intraepithelial lesion or malignancy 		
	Qing Cai, SCT (ASCP)			
09/21/2015	XXXX Women's Health XXXX, M.D.	Office visit for pelvic pain, dyspareunia, pelvic prolapse: Chief complaint: Pelvic Pain Pelvic prolapse The patient is 41-year old African American/Black female G 4 P 2 0 2 2 who reports today for further evaluation of dyspareunia and pelvic pain. She has a known history of grade 2-3 uterine prolapse. She reports increasing symptoms and notes at times the need to push the cervix back into the vagina after very strenous activity. She has also noted increasing dyspareunia and pelvic pain especially with standing. She also reports increasing vaginal infections beginning approximately 4-6 months ago. She was evaluated by Duke UroGyn who at that time did not recommend surgery. She presents today inquiring about surgical treatment options. Her LMP was 08/01/2007 due to Depo Provera. She is premenopausal. She uses Essure for contraception. Her PMHx is incorporated in separate enclosed PMHx form. Her last Pap Smear was 08/06/2015 and was normal.	MR - XXXX Women's Health - 000076-000079	1154-1157
		Medication list: Depo-Provera 150 mg/mL intramuscular suspension Review of systems: Genitourinary: Amenorrhea, complete emptying of bladder on voiding, dyspareunia,		

DATE	PROVIDER	OCCURRENCE/TREATMENT	BATES REF	PDF REF
		weight: 199 lbs 6 oz Assessment: Dyspareunia Chronic pelvic pain in female Uterine prolapse Plan: Grade 3 uterine prolapse: Patient with grade 3 prolapse on exam with Valsalva. She is currently symptomatic with symptoms of dyspareunia and pelvic pain. Treatment options re-addressed at this time including pessary versus surgery. Patient at this time desires surgery secondary to pre-menopausal status and currently sexually activity. Surgery also recommended given increasing symptomatology. Discussed laparoscopic hysterectomy given multiple abdominal surgeries with high probability of extensive adhesive disease. Surgery request submitted today for Total Laparoscopic Hysterectomy (TLH), Bilateral Salpingectomy (BS), and cystoscopy. Will scheduled with general surgery for abdominal entry and adhesiolysis as needed. Follow up for preoperative		
10/22/2015	XXXX Medical Center XXXX, M.D. XXXX, N.P.	Follow-up visit status post hemicolectomy: Patient was having menorrhagia with clots and was seen by Urogynecology with a recent Depo-Provera injection. Plan: She is scheduled for a hysterectomy for menorrhagia in 12/2015 and she notes that Dr. XXXX has been asked to assist her Gynecologist in view of surgical scars from colon surgery. She is likely in perimenopause with hot flashes. Ovaries will remain in situ per a reported GYN plan. We discussed the use of Effexor for hot flashes that her Gynecologist had also advised; at this point, she does not want to initiate a new medication. I have encouraged her to discuss this again with her Gynecologist should she become more symptomatic	A1062A785498 49D1B73E, XXXX, 194-195	675- 676

DATE	PROVIDER	OCCURRENCE/TREATMENT	BATES REF	PDF REF
		following her surgery. She will return to our clinic in 1		
		year, to which she agrees.		
11/12/2015	XXXX	Procedure note:	MR - XXXX	1158
	Women's		Women's	
	Health	The patient was injected with Depo Provera on	Health - 000080	
		11/12/2015 using sterile procedure.		
	XXXX, M.D.	Dose: 1 ml		
		Site: Right Gluteal IM		
	XXXX, L.P.N.	Manufacturer: Greenstone		
		Lot Number: M40837		
		Expiration date: 04/30/2018		
		Weight 212.2 lbs		
		Patient was monitored and there were no adverse effects.		
		Injection given by: A. Milton, LPN		
		Assessment: Contraception-follow-up Depo-Provera		
		Return to clinic in 3 months for next injection		
12/03/2015	XXXX	Office visit back, leg neck and arm pain:	MR - XXXX -	1620-
			000079-000081	1622
	XXXX, M.D.	Review of systems: Endocrinology: Low sex drive		
		*Reviewer's comment: Only case relevant information		
		captured from this visit.		
12/11/2015	XXXX	Office visit for preoperative evaluation:	MR - XXXX	1159-
	Women's		Women's	1162
	Health	Patient uses Essure for contraception. She comes in for	Health - 000081-000084	
		discussion of Total Laparoscopic	000061-000064	
	XXXX, M.D.	Hysterectomy(TLH)/Bilateral Salpingectomy		
		(BS)/cystoscopy secondary to grade 2 symptomatic		
		uterine prolapse and dyspareunia. Her surgery is		
		scheduled for 12/15/2015 at XXXX Hospital		
		Discussion for procedure:		
		She appears to understand the risks/benefits/alternatives		
		of the procedures and wishes to proceed with the		
		planned procedure. She appears to understand the risk of		
		bleeding that could result in blood transfusion and the		
		risk of a blood transfusion including allergic reaction,		
		developing hepatitis, AIDS, or some other blood borne		
		infection. She also appears to understand the risk of		
		infection and cardiac arrest. She also appears to		
		understand the specific risks associated with this/these		

DATE	PROVIDER	OCCURRENCE/TREATMENT	BATES REF	PDF REF
		particular procedures as outlined on the operative permit		
		Medication list: Depo Provera 150 mg/ml intramuscular suspension		
		Review of systems: Genitourinary: Amenorrhea, complete emptying of bladder on voiding, voiding normal amounts of urine.		
		Weight: 213 lbs 2 oz		
		Assessment:		
		Risks, benefits and alternatives to procedure discussed.		
		Patient desires to proceed. Postoperative pain medication		
		prescribed. Follow-up in 1 week for routine		
		postoperative evaluation.		
12/11/2015	XXXX Medical	Consent for total laparoscopic hysterectomy, bilateral	MR - XXXX -	394-
	Center	salpingectomy, cystoscopy, possible laparotomy	part 1 - 000161- 000162	395
12/11/2015	XXXX Medical	Preoperative interview:	MR - XXXX -	400
	Center		part 1 - 000167	
	VVVV	Planned procedure: Total laparoscopic hysterectomy,		
	(Credentials	bilateral salpingectomy, cystoscopy		
	unknown)	Surgical diagnosis: Uterine prolapse, dyspareunia		
	unitio (vii)	Surgicul ulugnosis eterme prompse, aj spureumu		
		Height: 64 inches; Weight: 96.6 kg; BMI: 36.1 kg/m2		
12/15/2015	XXXX Medical	Labs:	MR - XXXX	1167
	Center		Women's	
		POC pregnancy: Negative	Health - 000089	
12/15/2015	XXXX Medical	Operative report for total laparoscopic hysterectomy,	MR - XXXX - part 1 - 000172-	405- 406
	Center	bilateral salpingectomy, McCall culdoplasty, and	000172-	400
	XXXX, M.D.	cystoscopy:		
	(Surgeon)	Pre and postoperative diagnosis: Grade 2-3		
		symptomatic uterine prolapse, dyspareunia.		
	Dr. XXXX			
	(Assistant)	Anesthesia: General		

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	Dr. XXXX	Procedure performed: Total laparoscopic		
	(Assistant)	hysterectomy, bilateral salpingectomy, McCall		
		culdoplasty, and cystoscopy		
		Findings : Grade 3 uterine prolapse while the patient was		
		anesthetized, normal cervix, normal bilateral fallopian		
		tubes, and ovaries. Small bowel noted to be adhesed to		
		the abdominal ventral mass.		
		Description of procedure: The patient was taken to the		
		Operating Room with preoperative IV fluids as well as		
		prophylactic: Ancef infusing. The patient's general		
		endotracheal anesthesia was obtained without difficulty.		
		She was placed in a dorsal lithotomy position. The		
		vagina and abdomen were prepped and draped in 3		
		normal sterile fashion. A preoperative time-out was		
		called. A weighted speculum and Beaver were inserted		
		into the vagina for visualization of the cervix, which		
		appeared normal and multiparous. The patient was noted		
		to have grade 3 uterine prolapse while anesthetized. The		
		uterus was sounded to 8 cm. The RUMI uterine		
		manipulator was then placed into the uterine cavity		
		successfully. A Foley catheter was inserted and noted to		
		be draining clear urine. At this time, Dr. XXX with		
		General Surgery was present in the room for abdominal		
		entrance, secondary to the patient's history of colonic		
		resection as well as abdominal mesh placement. A 12 x		
		12 cm area was circumscribed around the umbilicus in		
		the area of the previous mesh. A mid right quadrant 5-		
		mm skin incision was made to accommodate a 5-mm		
		trocar, which was advanced under direct visualization		
		atraumatically. A broad inspection of the abdomen		
		revealed several portions of smell bowel adhesed to the		
		ventral mesh. However, the pelvis was noted to be free		
		of any adhesive disease. An infraumbilical port was then		
		placed outside the 12 x 12 cm circumference under		
		direct visualization atraumatically as well with the third		
		5-mm port placed in the left lower quadrant		
		atraumatically and under direct visualization. The pelvis		
		was inspected again and was noted to be free of any		
		adhesive disease. The uterus as well as tubes and ovaries		
		were all noted to be normal. The patient's right fallopian		

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		tube was identified and elevated at the fimbriated and		
		with the use of the Ligasure device, the mesosalpinx was		
		incised up to the uterine cornu with goad hemostasis		
		noted. The uterine ovarian ligament was then isolated,		
		cauterized, and transected with the LigaSure device. The		
		right round ligament was cauterized and transected with		
		the Liga-Sure device as well. The anterior leaf of the		
		broad ligament was then dissected with partial		
		skeletonization of the uterine arteries. The posterior leaf		
		was also partially dissected as well to provide a greater		
		visualization of the uterine arteries. The bladder flap was		
		partially created from the right side of the patient with		
		the use of the Ligasure device. At this time, the uterine		
		arteries were well skeletonized and cauterized and		
		transected with the Ligasure device on the right side of		
		the patient. Good hemostasis was noted at this time.		
		Attention was then turned to the left side of the pelvis, at		
		which time, the left fallopian tube, left uterovarian		
		ligament, as well as left round ligament each time were		
		isolated, cauterized, and transected with the Ligasure		
		device. The uterine arteries and the left were again		
		skeletonized with complete dissection of the bladder		
		from the lower uterine segment. At this time, the uterine		
		arteries on the left were cauterized and transected with		
		the LigaSure device. Given full dissection of the bladder		
		from the lower uterine segment, the anterior colpotomy		
		was made with the L-hook and in a circumferential		
		fashion, the cervix was transected from the vaginal		
		mucosa. The vaginal portion of the case was undertaken.		
		The specimen was removed intact and handed off and		
		sent to Pathology. A weighted speculum and Deaver		
		were inserted into the vagina for visualization of the		
		vaginal cuff.		
		The pelvic peritoneum was grasped with long Allis		
		clamps and with the use of 0 Monocryl, a standard		
		McCall culdoplasty was performed incorporating the		
		bilateral uterosacral. This stitch was left hanging until		
		the end of the case.		
		The bilateral vaginal cuff angles were than grasped with		
		long Allis clamps and closed in a locked running fashion		
		from both cuff angles to the midline. The vaginal cuff		
		sutures were then tied down. The McCall suture was -		

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		was also tied down at this time with good elevation of the vaginal vault noted. The vagina was irrigated copiously and cleared of all clots and debris and the vaginal cuff inspected and found to be hemostatic. Cystoscopy was undertaken at this time. Cystoscopy revealed normal bladder with exception of a 0.5-1 cm cyst that appeared to be benign. There was noted to be bilateral clear urine efflux from bilateral patent ureters without any damage or suture noted to the bladder. The laparoscopic portion of the case was undertaken for the final time. The abdomen was insufflated with gas. All pedicles were inspected and found to be hemostatic. The abdomen was then irrigated copiously and cleared of all clots and debris and partially deflated of gas and all pedicles again remained hemostatic. At this time, the abdomen was completely deflated of gas and all instruments and ports removed. The 3 laparoscopic skin incisions were closed with Dermabond. Sponge, lap, and needle counts were correct ×2 and the patient was extubated and taken to the PACU in stable condition. Specimen removed: Uterus, cervix, as well as bilateral fallopian tubes, all sent no Pathology. Drains: None.		
	10	Estimated blood loss: 50 ml. IV Fluids: 2500 ml. Urine output: 500 ml		
12/15/2015	XXXX Medical Center XXXX, M.D.	Diet: Begin with liquids and tight foods Progress to normal diet if you are not nauseated No alcoholic beverages for 24 hours	MR - XXXX - part 1 - 000152- 000154	385- 387
		Activities: The duration of drowsiness varies with each person. You will recover from these effects by tomorrow		

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		 Do not make important personal/business decisions or sign legal documents for 24 hours Do not drive or operate machinery for 24 hours or until not taking pain medicine Limit your activities for 24 hours. Do not engage in sports, heavy work or heavy lifting until your physician gives you permission May shower in 24 hours Pain take prescription as directed Ice pack Heat therapy Use deep breathing, distraction, repositioning Wound care: Use ice packs as needed for pain and inflammation Keep dressing dry Follow-up with Dr. XXX in 1 week. Keep the wound dry. 		
12/15/2015	XXXX Medical	Hospitalization related records: Medication sheets,	MR - XXXX -	388-
	Center	consent, anesthesia record, orders	part 1 - 000155- 000156, MR - XXXX - part 1 - 000160, MR - XXXX - part 1 - 000164, MR - XXXX - part 1 - 000165-000166, MR - XXXX - part 1 - 000168- 000171, MR - XXXX - part 1 - 000177-000248	389, 393, 397, 398- 399, 401- 404, 410- 481
12/18/2015	XXXX Medical	Pathology report:	MR - XXXX Women`s	1168- 1169
	Center	Collected date: 12/15/2015	Health -	1109
	XXXX, M.D.	Final diagnoses:	000090-000091	
		Cervix, uterus and fallopian tubes, hysterectomy		
		with bilateral salpingectomy:		
		Mild chronic cervicitis with squamous metaplasia and multiple Nabothian cysts.		

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		No evidence of dysplasia in recuts.		
		Negative for malignancy.		
		Uterus:		
		Inactive endometrium with focal evidence of		
		ciliated cell metaplasia.		
		No evidence of endometritis or hyperplasia.		
		Focal superficial adenomyosis noted.		
		Multiple intramural leiomyomata. Subscript I in recommendation		
		Subserosal leiomyoma.Fibrous adhesions with admixed peritoneal		
		inclusion cysts along serosal sur face.		
		No evidence of endometriosis.		
		Negative for atypia and malignancy.		
		3,1		
		Fallopian tubes		
		 Metallic material within lumen consistent with a 		
		prior sterilization		
		Cross description		
		Gross description: Uterus, cervix, bilateral tubes: In formalin is a 136 gram		
		uterus and cervix (8.7 cm cervix to fundus, 5.4 cm cornu		
		to cornu and 4.1 cm anterior to posterior) with attached		
		presumed discontinuous right fimbriated fallopian tube		
		(4.9 cm in length x 0.5 cm in diameter) and a 2.1 cm in		
		length x 0.3 cm in diameter portion left fallopian tube.		
		The serosal surface of the fallopian tubes is purple-gray		
		and grossly unremarkable. The portion of apparent left		
		fallopian tube exhibits silver coiled metallic material within the lumen. The serosal surface of the uterus is		
· ·		tan-purple with several foci of disruption and a 0.2 cm in		
		greatest dimension white-tan subserosal nodule. There is		
		a 0.9 x 0.7 x 0.2 cm collection of unilocular		
		cysts/edematous adhesions on the posterior surface. The		
		cervix (3.9 x 3.6 cm) is purple-gray and locally		
		hemorrhagic. The endometrial cavity (5.4 x 2,4 cm) is		
		lined by retracted brown-yellow tissue (0.1 cm thick).		
		The myometrium (1.1-2.1 cm thick) is pink-tan and		
		exhibits multiple white tan intramural nodules, up to 1.0 cm in greatest dimension.		
12/28/2015	XXXX	Follow-up visit status post TLH and BS:	MR - XXXX	1170-
		1 1	Women's	1172

DATE	PROVIDER	OCCURRENCE/TREATMENT	BATES REF	PDF REF
	Women's Health XXXX, M.D.	Since her surgery she has done well. She reports normal bowel and urinary habits. She denies vaginal bleeding. She is ambulating with minimal difficulty and tolerating a regular diet.	Health - 000092-000094	
		Medication list: Depo-Provera 150 mg/mL intramuscular suspension Weight: 207 lbs		
		Physical examination: Lap sites ×3 healing well without surrounding erythema or induration		
		Assessment: Postoperative exam		
		Instructions: Patient healing well without clinical signs/symptoms of infection. Incision care discussed. Benign pathology and surgical findings discussed today. Patient encouraged to		
01/20/2016	XXXX	slowly increase activity to baseline with exception of heavy lifting and intercourse. Follow-up in 3 weeks for continued post-operative evaluation and pelvic exam. Follow-up visit for continued postoperative	MR - XXXX	1173-
	Women's Health XXXX, M.D.	evaluation: Patient continues to do well. She reports an increase in light brown vaginal spotting without active bright red bleeding. She has resumed all normal activity without complication.	Women's Health - 000095-000097	1175
		Medication list: Depo-Provera 150 mg/mL intramuscular suspension, Metrogel Vaginal 0.75 % vaginal gel		
		Weight: 222 lbs		
		Physical examination: Gastrointestinal: Abdomen: Lap sites ×3 well healed Vagina: Minimal amount of light yellow/brown		

DATE	PROVIDER	OCCURRENCE/TREATMENT	BATES REF	PDF REF
		discharge noted in vault secondary to dissolution of suture		
		Assessment: Postoperative exam		
		Instructions: Patient continuing to heal well without clinical		
		signs/symptoms of infection. Pelvic exam unremarkable		
		today with dissolved suture noted in vault. Reassurance		
		given concerning vaginal discharge, and this is normal with vaginal suture dissolution. Patient encouraged to		
		continue all normal activity with exception of heavy		
		lifting and intercourse.		
		Disposition: Call or return if symptoms worsen or		
02/10/2016	XXXX	persist. Follow-up visit for final postoperative evaluation:	MR - XXXX	1176-
02/10/2010	Women's	Follow-up visit for final postoperative evaluation.	Women's	1178
	Health	Patient continues to do well. She has resumed all normal	Health - 000098-000100	
		activity without complication. She denies any further	000098-000100	
	XXXX, M.D.	episodes of vaginal bleeding.		
		Medication list: Depo-Provera 150 mg/mL		
		intramuscular suspension, Metrogel Vaginal 0.75 %		
		vaginal gel		
		Weight: 223 lbs		
		Physical examination:		
		Gastrointestinal: Abdomen: Laparoscopic sites ×3 well healed		
		Vagina: Vaginal cuff well healed, no suture or abnormal		
		discharge present, non-tender to palpation		
		Assessment: Postoperative exam		
		Plan: Patient completely healed without further need for		
		activity restriction. Patient encouraged to resume all		
		normal activity including heavy lifting and intercourse.		
		Patient advised that she no longer requires pap smears		
		secondary to TLH with negative cervical pathology and no personal history of severe cervical dysplasia. Follow-		
		up in 12/2016 for annual exam or sooner if symptoms		

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		arise. Disposition: Call or return if symptoms worsen or persist.		
04/13/2016	XXXX Women's Health	Office visit for vaginal discharge and vulvar itching: Chief complaint: Vaginal discharge; vulvar itching	MR - XXXX Women`s Health -	1179- 1181
	Health XXXX, M.D.	Patient presents today with complaint of vaginal discharge noted after intercourse. She reports the discharge is yellow in character. She describes the discharge as yellow. Significant risk factors include: New chemical or hygienic agent exposure. The chemical/hygienic agent exposure includes deodorant soap. She has had previous treatment for this condition that included Diflucan. Patient stated Monday she tried Monistat, which did not help, also had a prescription for Diflucan which helped but patient still complains of yellow vaginal discharge Review of systems: Genitourinary: Vaginal discharge, vaginal/vulvar itching or irritation Weight: 217 lbs Physical examination: External genitalia: Slight erythema noted Vagina Scant amt of thin white discharge noted in vault In-office procedures: Wet prep BV Whiff test Vagina QI: Negative Clue cells XXX QI wet prep: Negative KOH prep XXX: Positive Trichomonas vaginalis Ag Genital QI: Negative Yeast Budding # Ur Comp Assist: Negative Assessment Candidiasis of vulva and vagina Discharge from the vagina	000101-000103	
		Instructions:		

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		Vaginal discharge: Exam and wet prep positive yeast.		
		Prescription for Nystatin cream and Diflucan sent to		
		pharmacy today. Follow up as needed or sooner if		
		symptoms persist.		
06/10/2016	XXXX Medical	Office visit for urinary symptoms:	MR - XXXX	45-47
	Center	Diagnosis: Acute UTI	Medical Center - 000045-000047	
	XXXX, M.D.			
11/09/2016	XXXX Women's	Office visit for vaginal discharge:	MR - XXXX Women`s	1182- 1184
	Health	Patient complains of a thin, white, and foul-smelling	Health -	1104
	Пеанн	vaginal discharge. She also reports no additional	000104-000106	
	VVVV M D			
	XXXX, M.D.	symptoms. Risk factors: The patient denies risk factors of recently		
		entering into a new sexual relationship, multiple recent		
		sexual partners, chlamydia, gonorrhea, unprotected sex,		
		and frequent yeast infections. The patient reports she has		
		not been exposed to new chemicals or hygienic agents.		
		There are the following aggravating factors: exercise.		
		She has had previous treatment for this condition which		
		included: Diflucan, She states this therapy has resulted		
		in complete relief of symptoms.		
		Review of systems: Genitourinary: Vaginal discharge		
		Weight: 204 lbs 8 oz		
		Physical examination:		
		Vagina: Vaginal cuff normal in appearance with		
		adherent thick clump white discharge noted.		
		In-Office Procedures:		
		Wet prep		
	_	BV Whiff test vaginal QI: Negative		
		Clue Cells XXX QI Wet Prep: Negative		
		KQH Prep XXX: Positive		
		Trichomonas vaginalis Ag Genital QI: Negative		
		Yeast Budding # Ur Comp Assist: Positive		
		Assessment:		
		 Candidiasis vulvovaginitis 		
		 Discharge from the vagina 		

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		Instructions Wet prep positive for yeast today. Prescription for Diflucan sent to pharmacy. Discussed vaginal hygiene and avoidance of harsh soaps or chemicals. Disposition: Call or Return if symptoms worsen or persist.		
03/09/2017	XXXX Medical Center XXXX (Credentials unknown)	Office visit for multiple concerns: Patient with concern for increased fatigue/"just tired which has for a while. Does not feel like she can get out of the bed, just as tired when she gets up as when she goes to bed. Also complains of slight frontal headache, recurrent left neck knot which swelling sometimes which has been going on for about a week Review of systems: Constitutional: Fatigue, energy level Assessment: • Enlarged thyroid • Localized swelling with mass of neck • Chronic fatigue Prescription for Diflucan 150 mg and Amoxicillin 500 mg	MR - XXXX Medical Center - 000039-000042	39-42
04/03/2017	XXXX Medical Center XXXX (Credentials unknown)	Office visit for urinary symptoms: Diagnosis: UTI: Cipro, Fluconazole, scheduled voiding, push fluids, drink plenty of water, hygiene - wipe front to back	MR - XXXX Medical Center - 000036-000038	36-38
05/30/2017	XXXX Associates XXXX, M.D.	Follow-up visit for arthritis: Review of systems: Endocrine: Fatigue	8B7B13342F89 4C66BF24, XXXX, 85-89	1265- 1269
03/05/2018	XXXX Associates XXXX, M.D.	Follow-up visit for arthritis: Review of systems: Endocrine: Fatigue	8B7B13342F89 4C66BF24, XXXX, 76-80	1256- 1260

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07/10/2018	XXXX	Follow-up visit for arthritis:	8B7B13342F89	1251-
	Associates		4C66BF24, XXXX, 71-76	1256
	VVVV MD	Review of systems: Endocrine: Fatigue	XXXX, /1-/0	
07/20/2018	XXXX, M.D. XXXX Medical	Office visit for urinary symptoms:	MR - XXXX	22-25
07/20/2016	Center	Office visit for urmary symptoms.	Medical Center -	22-23
	Conter	Assessment: UTI	000022-000025	
	XXXX, M.D.			
		Supportive care: Increase fluids.		
07/26/2018	XXXX	Office visit for annual gynecological exam:	MR - XXXX	1442-
	Women's		Women's Health - 7.26.18	1445
	WWW MD	The patient is physiologic menopausal. There have been	- 000001 - MR -	
	XXXX, M.D.	no significant changes in her health history since her last	XXXX	
	XXXX, CNM	visit. She complains of constipation. Patient admits she is taking several different medications. She denies	Women's	
	AAAA, CIVIVI	vaginal discharge, abdominal pain, and dysuria. The	Health - 7.26.18 - 000004	
		patient is not taking HRT medication. She is having mild	- 000004	
		vasomotor symptoms.		
		Review of systems: Gastrointestinal: Constipation		
		Physical examination:		
		Genitourinary: Normal		
		And any of a first state of a second state of a		
		Assessment: Encounter for general routine gynecological exam with abnormal findings		
		gynecological exam with abhormal midnigs		
		Plan:		
		Diflucan 150 mg oral tablet, Nystatin 100,000 unit/g		
		topical cream.		
'				
		Return to clinic in 1 year and as needed.		
08/12/2018	XXXX Medical	Urine culture:	A1062A785498	1021
	Center	Collected date: 08/09/2018	49D1B73E, XXXX, 540	
		Popult. 10,000 50,000/ml Each amichic acti	212111, 5TO	
10/16/2018	XXXX	Result: 10,000-50,000/ml Escherichia coli Follow-up visit for arthritis:	8B7B13342F89	1246-
10/10/2018	Associates	ronow-up visit for artiffus;	4C66BF24,	1240-
	11550014105	Review of systems: Endocrine: Fatigue	XXXX, 66-71	
	XXXX, M.D.			
02/05/2019	XXXX	Follow-up visit for arthritis:	8B7B13342F89	1232-
	Associates		4C66BF24,	1237

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	XXXX, M.D.	Review of systems: Endocrine: Fatigue	XXXX, 52-57	
05/22/2019	XXXX Associates	Follow-up visit for arthritis:	8B7B13342F89 4C66BF24,	1228- 1232
	XXXX, M.D.	Review of systems: Endocrine: Fatigue	XXXX, 48-52	
10/11/2019	XXXX Medical Center	Follow-up visit for UTI: Diagnosis: UTI: Follow-up on Monday for a repeat	MR - XXXX Medical Center - 000011-000013	11-13
02/19/2020	XXXX, M.D. XXXX Associates Smith Carl,	Urine culture Follow-up visit for neck and low back pain: Review of systems: Gastrointestinal: Abdominal pain	8B7B13342F89 4C66BF24, XXXX, 19-25	1199- 1205
02/26/2020	M.D. XXXX Associates Smith Carl, M.D.	Endocrine: Fatigue Follow-up visit for neck and low back pain: Review of systems: Gastrointestinal: Abdominal pain Endocrine: Fatigue	8B7B13342F89 4C66BF24, XXXX, 10-16	1190- 1196
03/23/2020	XXXX Medical Center XXXX, M.D.	Office visit for annual wellness visit: Assessment: Routine medical exam Anxiety: Stable no new intervention, continue Ativan 1 mg as needed	MR - XXXX Medical Center - 000001-000007	1-7
04/07/2020	XXXX Associates XXXX, M.D.	Follow-up visit for arthritis: Review of systems: Gastrointestinal: No abdominal pain Genitourinary: No incontinence, hematuria, difficulty urinating or increased frequency Endocrine: No fatigue	8B7B13342F89 4C66BF24, XXXX, 5-10	1185- 1190
09/06/2009 - 05/18/2020	Multiple providers	Return to office on 07/13/2020 Records not relevant to case review: Admission record, after care instructions, anesthesia record, assessment, bilateral digital mammogram, colonoscopy, consent, CT scan of chest, abdomen and pelvis with contrast, discharge instructions, discharge summary, echocardiogram, electrocardiogram, ER		1002- 1020, 101- 109, 1022- 1035, 1038- 1051,

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		record for high bp, ER visit for toothache, jaw pain, fax		1055-
		sheets, follow-up visit status post chemotherapy for		1077,
		colon cancer, headache, Holter monitor report, labs,		1083,
		mammogram, medication sheets, MRI of right and left		1086-
		shoulder, MRI of right shoulder, office visits for		1088,
		arthritis, back pain, bilateral shoulder pain, chest pain,		112-
		constipation, depression, hypertension, finger open		114, 1142-
		wound, hypertension, incisional hernia, joint pain, left		1144,
		heel pain, potassium check, respiratory symptoms,		116-
		shoulder joint pain, throat pain, upper respiratory		117,
		infection, influenza vaccine, operative report for		1163-
				1166,
		laparoscopic repair of ventral incisional hernia with		1197-
		mesh, orders, others, post anesthesia evaluation,		1198,
		preanesthesia, preoperative interview, procedure report,		1206-
		progress notes, referral report, ultrasound, X-ray of		1214, 121-
		abdomen, X-ray of cervical spine, X-ray of chest, X-ray		121-
		of left foot, X-ray of small bowel, X-ray of thoracic		1214-
		spine, assessment, bilateral L3 medial branch block,		1227,
		bilateral L3, L4, L5 radiofrequency thermocoagulation		123-
		of medial branch under fluoroscopic guidance, bilateral		126,
		L4 medial branch block, bilateral L5-S1 facet steroid		1238-
		injection with fluoroscopic guidance, bilateral S1 trans		1246,
		foraminal epidural steroid injections with fluoroscopic		130-
		guidance, labs, left L3 radiofrequency neuroablation,		135, 1260-
		mammogram, MRI of cervical spine without contrast,		1269-
		MRI of lumbar spine without contrast, office visit for		1312,
		neck pain, lower back pain, orders, others, right L3, L4,		1317-
		L5 radiofrequency thermocoagulation of medial branch		1320,
		under fluoroscopic guidance, X-ray of cervical spine		1330-
				1392,
		Bates ref: 8B7B13342F894C66BF24, XXXX, 100-132,		136-
		137-140, 150, 151-171, 17-18, 172-174, 175-177, 178-		141, 1394-
		190, 192-193, 194, 195-197, 201-203, 205-212, 214,		1394-
		215-216, 218-239, 26-47, 58-66, 80-85, 89-100		1398-
		A1062A78549849D1B73E, XXXX, 101-102, 105-106,		1419,
		108-117, 120-133, 135-139, 1-4, 140-154, 157-185, 19- 20, 196-211, 21-22, 212-213, 216-229, 23, 231-243, 24,		14-16,
		20, 190-211, 21-22, 212-213, 210-229, 23, 231-243, 24, 244-253, 25-26, 254-272, 27, 273, 274-276, 279-289,		142-
		28, 29, 290-292, 299, 30, 302-316, 31, 317, 318-319,		147,
		320, 322, 32-33, 323-334, 335-336, 338, 339-353, 34-		149-
		43, 354-355, 356-360, 361-362, 363, 364-365, 366-368,		151,
		369-381, 382, 383-385, 388-398, 399-400, 402-426,		153- 156,
		428-475, 44-46, 47-48, 476-491, 49, 493-518, 521-537,		156, 161-
		52-76, 538-539, 541-554, 557-570, 5-7, 574-596, 77-99		101-

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		MR - XXXX Medical Center - 000008-000010, 000014-		164,
		000016, 000017-000021, 000026-000035, 000043-		167-
		000044, 000048-000053, 000056-000069, 000072-		384,
		000081, 000087-000096, 000101-000117, 000121-		17-21,
		000126, 000130-000147, 000149-000151, 000153-		26-32,
		000156, 000161-000189		396,
		MR - XXXX - 000001 - 000044		409,
		MR - XXXX - part 1 - 000001-000151, 000163, 000176, MR - XXXX Women`s Health - 000005		43-44, 48-50,
		MR - XXXX Women's Health - 000003 MR - XXXX Women's Health - 000008-000010, 000064		482-
		000066, 000085-000088		498,
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		MR - XXXX - 000370-000374, 000325, 000351-000352,		530,
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		000273, 000270-000271, 000268-000269, 000313,		614,
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		000337, 000055-000058, 000253, 000316-000317,		666,
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		000231, 000223-000228, 000217-000222, 000236-		907,
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