UNITED STATES SECURITIES AND EXCHANGE COMMISSION Washington, D.C. 20549

FORM 8-K

CURRENT REPORT PURSUANT TO SECTION 13 OR 15(d) OF THE SECURITIES EXCHANGE ACT OF 1934

Date of Report (Date of earliest event reported): September 25, 2019

TherapeuticsMD, Inc.

(Exact Name of Registrant as Specified in its Charter)

001-00100

Nevada (State or Other

Jurisdiction of Incorporation)

(Commission File Number)

87-0233535

(IRS Employer Identification No.)

951 Yamato Road, Suite 220 Boca Raton, FL 33431

(Address of Principal Executive Office) (Zip Code)

Registrant's telephone number, including area code: (561) 961-1900

Not Applicable

(Former name or former address, if changed since last report)

Securities registered pursuant to Section 12(b) of the Act:

		Name of Each Exchange on Which
Title of Each Class	Trading Symbol	Registered
Common Stock, par value \$0.001 per share	TXMD	The Nasdaq Stock Market LLC

Check the appropriate box below if the Form 8-K filing is intended to simultaneously satisfy the filing obligation of the registrant under any of the following provisions:

□ Written communications pursuant to Rule 425 under the Securities Act (17 CFR 230.425)

□ Soliciting material pursuant to Rule 14a-12 under the Exchange Act (17 CFR 240.14a-12)

□ Pre-commencement communications pursuant to Rule 14d-2(b) under the Exchange Act (17 CFR 240.14d-2(b))

□ Pre-commencement communications pursuant to Rule 13e-4(c) under the Exchange Act (17 CFR 240.13e-4(c))

Indicate by check mark whether the registrant is an emerging growth company as defined in Rule 405 of the Securities Act of 1933 (§230-405 of this chapter) or Rule 12b-2 of the Securities Exchange Act of 1934 (§240.12b-2 of this chapter).

Emerging growth company \Box

If an emerging growth company, indicate by check mark if the registrant has elected not to use the extended transition period for complying with any new or revised financial accounting standards provided pursuant to Section 13(a) of the Exchange Act.

Item 7.01. Regulation FD Disclosure.

TherapeuticsMD, Inc. is furnishing as Exhibit 99.1 to this Current Report on Form 8-K an investor presentation which may be used, in whole or in part, and subject to modification, on September 25, 2019 and at subsequent meetings with investors or analysts.

The information in this Current Report on Form 8-K (including the exhibit) is being furnished pursuant to Item 7.01 of Form 8-K and shall not be deemed to be "filed" for the purpose of Section 18 of the Securities Exchange Act of 1934, as amended, or otherwise subject to the liabilities of that section, nor will any of such information or exhibits be deemed incorporated by reference into any filing under the Securities Act of 1933, as amended, or the Securities Exchange Act of 1934, as amended, except as expressly set forth by specific reference in such filing.

Item 9.01. Financial Statements and Exhibits.

(d)	Exhibits	
	Exhibit	Exhibit Index
	<u>Number</u>	Description
	<u>99.1</u> 104	TherapeuticsMD, Inc. presentation dated September 25, 2019. Cover Page Interactive Data File (the cover page tags are embedded within the Inline XBRL document).

SIGNATURES

Pursuant to the requirements of the Securities Exchange Act of 1934, the registrant has duly caused this report to be signed on its behalf by the undersigned hereunto duly authorized.

Date: September 25, 2019

THERAPEUTICSMD, INC.

By: /s/ Daniel A. Cartwright Name: Daniel A. Cartwright

Title: Chief Financial Officer



Forward-Looking Statements

This presentation by TherapeuticsMD, Inc. (referred to as "we" and "our") may contain forward-looking statements. Forward-looking statements may include, but are not limited to, statements relating to our objectives, plans and strategies, as well as statements, other than historical facts, that address activities, events or developments that we intend, expect, project, believe or anticipate will or may occur in the future. These statements are often characterized by terminology such as "believe," "hope," "may," anticipate," "should," "intend, "plan," "will," "expect," "estimate," "project," "positioned," "strategy" and similar expressions and are based on assumptions and assessments made in light of our managerial experience and perception of historical trends, current conditions, expected future developments and other factors we believe to be appropriate.

Forward-looking statements in this presentation are made as of the date of this presentation, and we undertake no duty to update or revise any such statements, whether as a result of new information, future events or otherwise. Forward-looking statements are not guarantees of future performance and are subject to risks and uncertainties, many of which may be outside of our control. Important factors that could cause actual results, developments and business decisions to differ materially from forward-looking statements are described in the sections titled "Risk Factors" in our filings with the Securities and Exchange Commission, including our most recent Annual Report on Form 10-K and Quarterly Reports on Form 10-Q, as well as our current reports on Form 8-K, and include the following: our ability to maintain or increase sales of our products; our ability to develop and commercialize IMVEXXY®, ANNOVERATM, BIJUVA® and our hormone therapy drug candidates and obtain additional financing necessary therefor; whether we will be able to comply with the covenants and conditions under our term loan facility; the potential of adverse side effects or other safety risks that could adversely affect the commercialization of our current or future approved products or preclude the approval of our future drug candidates; the length, cost and uncertain results of future clinical trials; the ability of our licensees to commercialize and distribute our products; our reliance on third parties to conduct our manufacturing, research and development and clinical trials, the availability of reimbursement from government authorities and health insurance companies for our products; the impact of product liability lawsuits; the influence of extensive and costly government regulation; the volatility of the trading price of our common stock and the concentration of power in our stock ownership.

This non-promotional presentation is intended for investor audiences only.

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General Updates and Catalysts

Began IMVEXXY direct to consumer advertising leveraging digital and social media platforms

Expanded payer coverage for IMVEXXY and BIJUVA; awaiting additional coverage decisions

- Expanded commercial coverage for IMVEXXY with the addition of CVS¹
- Awaiting commercial coverage decision from Aetna for IMVEXXY
- The recent addition of Cigna, expanded commercial coverage for BIJUVA to 5 of the top 10 payers1
- Additional commercial and Medicare Part D payer decisions expected 3Q/4Q 2019 for IMVEXXY and BIJUVA

ANNOVERA soft launch begins the week of September 30, 2019

- ANNOVERA has 40% unrestricted coverage in commercial health plans¹
- Added 2 of the top 10 commercial payers for ANNOVERA and several regional plans¹

Recent stock purchases by members of the executive team and board of directors

- CEO purchased approximately \$280k in TXMD stock in 3Q19
- CEO 2019 annual stock-based award of a single stock option to provide more equity to executive team with less dilution to stockholders

MMIT September 2019 and Account insights

Near-Term Events

- Held Satellite Media Tour for Menopause Awareness Month (September)
- IMVEXXY and BIJUVA data will be presented at 2019 Annual Meeting of the North American Menopause Society being held Sept. 25-28, 2019 (2 oral presentations and 5 posters):
 - Results from the REPLENISH Phase 3 trial for BIJUVA that show improvements in several sleep parameters, including sleep disturbances, in menopausal women with moderate to severe vasomotor symptoms taking BIJUVA as compared to placebo
 - Data for BIJUVA showing a consistency of effect for improvements in the frequency and severity of vasomotor symptoms in different subpopulations, regardless of age or body mass index
 - Additionally, two oral presentations will review the growth of the VVA market and patient acceptability data, including patient satisfaction and preference, for IMVEXXY
- ANNOVERA data will be presented at 2019 Annual Meeting of the American Society for Reproductive Medicine (Oct. 12-19, 2019)

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August Launch Metrics

ugust 1-31, 2019)	~44,600
otal patients since launch through August 31, 2019)	~87,400
otal prescribers ² since launch through August 31, 2019)	~14,800

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References:

References: 1. Total prescription data is based on IQVIA prescriber level data plus additional unique patient data identified through utilization of our affordability program. This includes two weeks of estimation for the lag in reporting retail data, which can cause minor fluctuations in historical comparisons. 2. Osphena and intrarosa data sourced from Symphony Health Integrated Dataverse. 3. Vagifiem data sourced from IQVIA National Prescriber Level Data. 4. Market share data based on IQVIA Prescriber Level Data. 4. Market share data based on IQVIA prescriber Level Data.

* Invexy Continued Strong Patient Adherence

Month Initial Prescription Filled	Average # Fills for Those Patients	Maximum Allowable Fills Given the Month of Initial Fill
Aug 2019	1 Fill	1 Fill
Jul 2019	1.9 Fills	2 Fills
Jun 2019	2.4 Fills	3 Fills
May 2019	2.9 Fills	4 Fills
Apr 2019	3.5 Fills	5 Fills
Mar 2019	4.0 Fills	6 Fills
Feb 2019	4.5 Fills	7 Fills
Jan 2019	5.0 Fills	8 Fills
Dec 2018	5.4 Fills	9 Fills
Nov 2018	6.1 Fills	10 Fills
Oct 2018	6.2 Fills	11 Fills
Sep 2018	6.7 Fills	12 Fills
Aug 2018	8.0 Fills	13 Fills
Jul 2018	8.0 Fills	14 Fills

IMVEXXY: 3.8 fills/yr³ (through Aug)

- Vaginal creams: average 1.5 fills/yr⁴
- Vaginal tablets: average 3.5 fills/yr4

each of those patients averaged 6.1 fills from November 2018 through August 2019

Average number of fills per patient is the average number of fills per patient grouped by their initial month on therapy.
 Total prescription data is based on IQVIA prescriber level data plus additional unique patients identified through utilization of our affordability program.
 Average number of fills for all patients is calculated as Total Rx / Total Patients.
 Total Rx/Patient Count.



Commercial Payer Status

			Plans Account for ~71% Imercial Pharmacy Lives
	Plan	% of Lives ¹	Status ²
vew	CVS	15.4%	Adjudicating as of September 2019
	ESI	15.3%	Adjudicating as of 10/1/18
	United	7.5%	Adjudicating as of 3/1/19
	Anthem	7.3%	Adjudicating as of August 2018
	Prime	6.5%	Adjudicating as of 1/1/19
	OptumRx	6.1%	Adjudicating as of 1/1/19
	Kaiser	4.7%	In discussions
	Aetna	4.0%	Awaiting decision ~1.8M of these lives are adjudicating
	Cigna	3.9%	Adjudicating as of 12/15/18
	EnvisionRx	1.8%	Adjudicating as of 1/1/19



Adjudication of claim by payer: IMIVEXXY is on payer formulary as covered product and is being submitted to insurance company for payment by payer to pharmacy.

Plan numbers as of May 2019 2Adjudication status from MMIT September 2019 and Account Insights

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August Launch Metrics

BIJUVA Launch Metrics			
Total paid scripts dispensed to patients ¹ (since launch through August 31, 2019)	~14,200		
Total paid scripts (August 1-31, 2019)	~5,300		
Total patients (since launch through August 31, 2019)	~6,800		
Total prescribers ² (since launch through August 31, 2019)	~3,000		

¹ Total prescription data is based on IQVIA prescriber level data plus additional unique patients identified through utilization of our affordability program. This includes a two week estimation for the lag in reporting retail data, which can cause minor fluctuations in historical comparisons.
² Total Unique Prescribers that have sent a prescription to a pharmacy for at least 1 patient for BIJUVA.

Bijuva (stadiand projestence) capades Coverage Decisions Expected by 4Q19

New

Quicker Process to Payer Coverage than IMVEXXY

- Commercial segment represents vast majority of BIJUVA patients
- Expect 3-4 quarters coverage cycle (from launch) to secure commercial payers
- Amendments to the company's existing payer contracts with little Medicare Part D

Plan	% of Lives ¹	Status ²	
CVS	15.4%	In discussions	
ESI	15.3%	Adjudicating as of 4/19/19	
United	7.5%	Adjudicating as of 8/1/19	
Anthem	7.3%	In discussions	
Prime	6.5%	In discussions	
OptumRx	6.1%	Adjudicating as of 8/1/19	
Kaiser	4.7%	In discussions	
Aetna	4.0%	Adjudicating as of 4/2019	
Cigna	3.9%	Adjudicating as of 9/2019	
EnvisionRx	1.8%	In discussions	

Adjudication of claim by payer: BUUVA is on payer formulary as covered product and is being submitted to insurance company for payment by payer to pharmacy.

Plan numbers as of May 2019 "Adjudication status from MMIT September 2019 and Account Insights

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Key Payer Updates

ANNOVERA has 40% Unrestricted Coverage in Commercial Health Plans¹

- Anthem adjudicating ANNOVERA at Tier 3 with no copay as of August 2019¹
- Kaiser Washington ACA drug list covers ANNOVERA at no copay¹
- Cigna adjudicating ANNOVERA at Tier 3 as of August 2019¹
- Starting on January 1, 2020, New York state insurance law requires coverage for all contraceptives, including ANNOVERA, with no copay²
- Currently with the addition of New York, 19 states, plus Washington, D.C., require insurance plans to cover all contraceptives that do not have a generic equivalent

Test and Learn Market Introduction Starting Week of September 30, 2019

MMIT September 2019 and Accounts Insights ²https://www.nysenate.gov/legislation/bills/2019/s659/amendment/a

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Commercial Payer Update Coverage Decisions Expected to be finalized 1Q20

			Plans Account for ~71% Imercial Pharmacy Lives	
	Plan	% of Lives ¹	Status ²	
	CVS	15.4%	In discussions	
	ESI	15.3%	In discussions	
	United	7.5%	In discussions	
w	Anthem	7.3%	Adjudicating at T3, no copay as of August 2019	
	Prime	6.5%	In discussions	
	OptumRx	6.1%	In discussions	
2	Kaiser	4.7%	In discussions Kaiser Washington covering at no copay	
	Aetna	4.0%	In discussions	
N	Cigna	3.9%	Adjudicating at T3 as of August 2019	
	EnvisionRx	1.8%	In discussions	



Adjudication of claim by payer: ANNOVERA is on payer formulary as covered product. and is being submitted to insurance company for payment by payer to pharmacy.

Plan numbers as of May 2019. Adjudication status from .MMIT September 2019 and Account Insights.



Commercial Payer Update

Coverage Decisions Expected to be finalized 1Q20

	Selected Regional Plan Coverage				
	Plan	% of Lives ¹	Status ²		
	MC-Rx (ProcareRx)	0.64%	Adjudicating as of April 2019		
lew	Magellan Rx	0.47%	Adjudicating at no copay as of August 2019		
	BCBS of Massachusetts	0.47%	Adjudicating at no copay as of August 2019		
lew	Excellus	0.27%	Adjudicating as of September 2019		
lew	EmblemHealth	0.25%	Adjudicating at no copay as of September 2019		
lew	Weilmark	0.23%	Adjudicating as of August 2019		
	Harvard Pilgrim	0.18%	Adjudicating at no copay as of August 2019		
	Independent Health Association	0.06%	Adjudicating as of August 2019		
lew	BC of Idaho	0.00%	Adjudicating at no copay as of September 2019		
lew	Summacare	0.00%	Adjudicating at no copay as of September 2019		
lew	Clear Script PBM	0.00%	Adjudicating as of August 2019		
lew	Univera Healthcare	0.00%	Adjudicating as of August 2019		

. . . .

^{1P}ion numbers as of May 2019 ³MMIT September 2019 and Account Insights and is being submitted to insurance company for payment by payer to pharmacy.





BIRTH CONTROL STATE LAWS REGARDLESS OF ACA MANDATES

8 STATES REQUIRE COVERAGE WITH COPAY REGARDLESS OF ACA DECISION (~27 Million women in these states)





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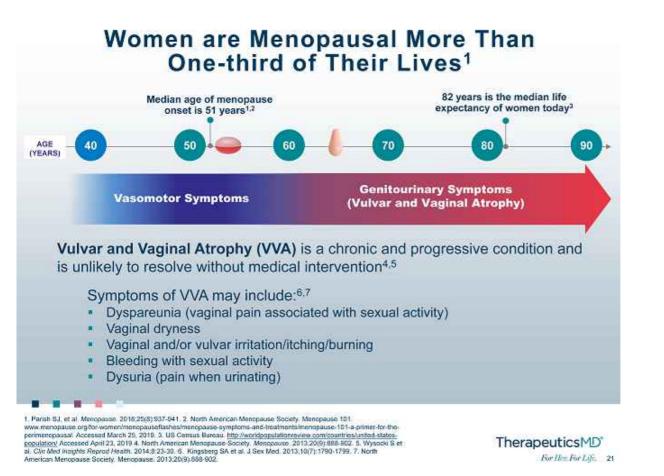
Conditions, and Eligibility Criteria apply.

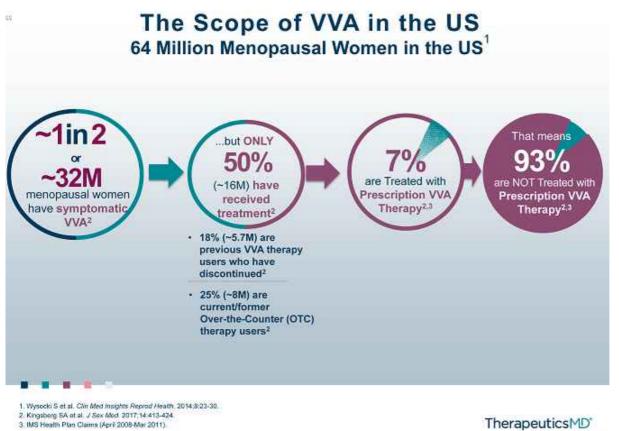
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	IOF WOR	nen throughout their life	cycles
	* Invexy estadd vagrel neerts 4may 10 reg	Bijuva regimer Istradul and progesterineli capitales	Annovera- (segeteure actue and string to any constraints and been to segvent actue
Key Value Proposition	Easy to use, lowest approved dose, designed to support patient compliance	First and only FDA-approved bio- identical combination product	First and only long-lasting (one year/13 cycles), procedure-free, patient-controlled, reversible birth control product
Affected US Population	32 million women ^{1,2}	36 million women ⁴	43 million women ⁶
US TAM Opportunity	>\$20B3	>\$25B ^{3,5}	\$5B ⁷
Status	Approved May 29, 2018 Launched August 2018	Approved October 28, 2018 Launched April 2019	Approved August 10, 2018 Test & Learn Introduction: 4Q19 Full scale launch expected: 1Q20

The Power of A Women's Health Portfolio







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IMVEXXY Product Characteristics Compare Favorably¹⁻⁹

Estrogens			Non-estrogens			
	Estrace® Cream (estradiol vaginal cream, USP, 0.01%)!	Premarin® (conjugated estrogens) Vaginal Cream ²	Vagifem® (estradiol vaginal inserts) ⁴	IMVEXXY® (estractiol vaginal inserts) ⁶	Intrarosa/8) (prasterone) vaginal inserts7	Osphena® (ospentifene) tablets, for oral use
Product	Designed and the second			innessa .	P. Marmora	
	- () Allergon	Pfizer	50	Therapeutics%07	🙈 amag	DUCHERNAY USA
FDA approval	1984	1978	1999	2018	2016	2013
Rx MSB Dollars of Brand & Generic 2018 ⁴	\$540,000,000	\$462,226,000	\$420,030,000	\$44,000,000	\$35,001,000	\$73,908,000
2018 Total Units ³	1,902,000	1,220,000	1,500,000	205,500 (10 months)	169,000	218,000
Method of administration	Vaginal cream	Vaginal cream	Vaginal insert	Vaginal Insert	Vaginal insert	Oral tablet
Application	Reusable vaginal applicator- cream	Reusable vaginal applicator- cream	Disposable vaginal applicator- tablet	No applicator needed- softgel vaginal insert	Disposable vaginal applicator- bullet insert	Oral daily table
Active ingredient	100 mcg estradiol	625 mcg/g conjugated equine estrogens	10 mcg estradiol	4 mcg or 10 mcg estradiol	6,500 mcg prasterone	60,000 mcg ospemifene
Average maintenance dose	100 mcg 2x/week	312.5 mcg 2x/week	10 mcg 2x/week	4 mcg or 10 mcg 2x/week	6,500 mcg daily	60,000 mcg daily
WAC package price (2019) ¹⁰	\$344.78 (42.5-g tube)	\$373.56 (30-g tube)	\$170.16 (8 tablets)	\$180.00 (8 softgel capsules)	\$202.00 (28 inserts)	\$648.00 (90 tablets)
WAC 28-day supply (2019) ¹⁹	\$97.35	\$130.75	\$170.16	\$180.00	\$202.00	\$201.60

References: 1. Ustroce Vaginni Cream (package insert), Initias, CA. Allenger USA, Inc. 2017. 2. Ptemarin Vaginal Cream (package insert), Protatepha, PA. Wyeth Pharmasourizata Inc., a subsidiary of Pfazer Inc. 2017. 3. Esting (package insert), New Yook, NY: Pharmasia & Uputin Company LLC, a subsidiary of Pfazer Inc. 2017. 4. Vagine (package insert), Watham, NA: MAO, 97. 5. MVEXXY (package Insert), Bock Raton, FL. TherapoutisMD, Inc. 2019. 7. Intrainas (package insert), Watham, MA: MAO, 97. 5. MVEXXY (package Insert), Bock Raton, FL. TherapoutisMD, Inc. 2019. 7. Intrainas (package insert), Watham, MA: MAO, 97. 5. MVEXXY (package Insert), Bock Raton, FL. TherapoutisMD, Inc. 2019. 9. Therapouties (Control And Control And

There have been no head-to-head trus between IMVEXXY and any of the products labed above. All indemarks are the property of their respective owners. Abbreviations: WWC, wholesan application cost. TherapeuticsMD*

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* Invexy stadd vaginal inserts) The VVA Market Exceeds \$1.7B Gross Revenue Annually

Product	12 Months through June 2019 Total Units ¹	Gross Dollars for 2018 ¹
Estrace® Cream Brand & Generics	2,000,000	\$554,450,000
Premarin®	1,190,000	\$460,760,000
Vagifem® Brand & Generics	1,500,000	\$454,550,000
Estring®	259,000	\$114,360,000
Osphena®	217,000	\$75,910,000
Intrarosa®	209,000	\$46,940,000
2018 Value of the VVA N	larket	\$1,700,000,000

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1) Symptrony Health Solutions PHAST Data powered by IDB; MBS dollars All trademarks are the property of liber respective owners.



Value of Additional Fills

Average Net Revenue / Unit	25%	35%	45%	55%
\$80	\$184M	\$257.6M	\$331.2M	\$404.8
\$100	\$230M	\$322M	\$414M	\$506
Р	ercent of market	based on patient	Count of 2.3M an	d 5 fills pe
Average Net Revenue / Unit	25%	35%	45%	55%
\$80	\$230M	\$322M	\$414M	\$506N
\$100	\$287.5M	\$402.5M	\$517.5M	\$632.5
e P	ercent of marke	t based on patient	count of 2.3M an	d 6 fills pe
Average Net Revenue / Unit	25%	35%	45%	55%
\$80	\$276M	\$386.4M	\$496.8M	\$607.2
\$100	\$345M	\$483M	\$621M	\$7591

Market opportunity is calculated by multiplying the number of patients on products annually times the market share times the average number of fills per patient per year times the average potential net revenue per unit. At \$100 average net revenue, the value per fill ranges from \$57M to \$126M, depending on market share.

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Commercial Average Non Preferred Copay

- Commercial Average Non Preferred Copay is \$59
- IMVEXXY co-pay card offer can bring this down to \$35

	Average Copayment	Average Coinsurance
Plans With Three or More Tiers		
First Tier	\$11	19%
Second Tier	\$33	26%
Third Tier	\$59	36%
Fourth Tier	\$105	31%
Plans With Two Tiers		
First Tier	\$11	NSD
Second Tier	\$31	28%
Plans With the Same Cost Sharing For All Covered Drugs	1992	5200 F
First Tier	NSD	20%
NOTE: Number of tiers refers to the number of tiers NSD: Not Sufficient Data	excluding those specifically for specialty o	drugs.
SOURCE: KFF Employer Health Benefits Survey, 20	18	

Source: 2018 Employer Health Benefits Survey, Section 9: Prescription Drug Benefits (KFF, Oct. 3, 2018), https://www.kff.org/report.section/2018-employer-health-benefits-survey-section-9-prescription-drug-benefits/ (accessed June 5, 2019).

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Medicare Part D Median Preferred Copay

Medicare Part D Median Preferred Copay is \$40

	Preferred generics		Generics		Preferred brands*		Non-preferred drugs		Special	ty drugs
Name of PDP	2018	2019	2018	2019	2018	2019	2018	2019	2018	2019
Median for all PDPs	\$7	\$1	\$6	\$5	\$37/21%	\$40/2016	40%	-40%	26%	26%
Top 10 PDPs										
SilverScript Choice	\$3	\$3	\$14	\$13	\$42	342	46%	45%	33%	33%
AARP MedicareRx Preferred	\$5	\$5	\$12	\$10	\$37	\$40	40%	40%	3346	33%
Humana Walmart Rx	\$1	\$1	44	\$4	23%	2096	3596	35%	25%	25%
Humana Preferred Rx	\$0	\$0	:\$1	51	2091	25%	3590	3796	25%	259
AARP MedicareRx Saver Plus	\$5	51	53	\$6	533	\$25	30%	3396	25%	25%
Aetna Medicare Rx Saver	\$1	\$1	\$2	\$2	\$30	\$30	35%	35%	26%	279
WellCare Classic	\$0	\$5	51	52	\$35	\$37	4291	41%	25%	25%
Humana Enhanced	\$3	\$5	\$7	\$10	542	\$47	44%	50%	3346	33%
AARP MedicareRx Waigreens	\$0	50	56	-55	\$21	\$30	32%	32%	25%	25%
Aetna Medicare Rx Value Plus	\$3	51	\$2	\$2	\$47	547	50%	4795	33%	33%

Source: Juliette Cubanski, Anthony Damico, and Tricia Neuman, Medicare Part D: A First Look at Precription Drug Plans in 2019 (Kff, Oct. 16, 2018). https://www.kff.org/report-section/medicare-part-d-a-first-look-at-prescription-drug-plans-in-2019tables/ (accessed June 5, 2019).

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Medicare Part D Payer Status

Decisions Expected 3Q/4Q19

Top 6 Plans Account for ~78% of all Medicare Part D Pharmacy Lives				
Plan	% of Lives1	Status ²		
United	20.7%	Adjudicating as of 2/1/19		
Humana	17.9%	Decision expected 3Q/4Q19		
CVS Caremark	14.1%	Decision expected 3Q/4Q19		
Wellcare with Aetna lives	13.6%	Decision expected 3Q/4Q19		
Express Scripts/ Cigna	8.5%	Decision expected 3Q/4Q19 ~1M of these lives are adjudicating as of June 2019		
Kaiser	3.6%	Adjudicating maintenance pack as of 10/1/18 and starter pack as of 3/1/19		



Adjudication of claim by payer: IMVEXXY is on payer formulary as covered product and is being submitted to insurance company for payment by payer to pharmacy.

¹Plian numbers as of July 2019 and is b ²Adjudication status from MMIT September 2019 and Account Insights

* Invexy How Adjudication Rate* Will Change When Payer Cycle Completes

2Q 2019 Actuals

	Column A	Column B	Column C
IMVEXXY	No Insurance	Commercial Insurance	Medicare Eligible Patients
% of Business	3%	62%	35%
% Adjudicated	0%	50%	8%
Contribution to Overall Adjudication Rate	0%	31%	3%
Overall Adjudication Rate	34%	6 (up from 27% i	n 1Q19)

Target Adjudication as Contracting is Finalized

	Column A	Column B	Column C	
IMVEXXY	No Insurance	Commercial Insurance	Medicare Eligibl Patients	
% of Business	6%	70%	24%	
% Adjudicated	0%	75%	65%	
Contribution to Overall Adjudication Rate	0%	53%	17%	
Overall Adjudication Rate		70%		

Chart 2

Chart 1

"Adjudication Rate= Percent of Business multiplied by percent of claims being covered.

IMVEXXY Model Different Than Typical Pharmaceutical Launch

Pharmacy Discounts Payer Rebates Returns, Allowances & Other Accruals	Patient Copay Assistance	Where We Focused
Payer Rebates Returns, Allowances & Other Accruals et Revenue Cost of Sales ross Margin Sales & Marketing Cost Copay Assistance substituted	Wholesale Costs	
Returns, Allowances & Other Accruals et Revenue Cost of Sales ross Margin Sales & Marketing Cost Copay Assistance substituted	Pharmacy Discounts	
et Revenue Cost of Sales ross Margin Sales & Marketing Cost Copay Assistance substituted	Payer Rebates	
Cost of Sales ross Margin Sales & Marketing Cost Copay Assistance substituted	Returns, Allowances & Other Accru	uals
Copay Assistance substituted	et Revenue	
Sales & Marketing Cost Copay Assistance substituted	Cost of Sales	
Sales & Marketing Cost Copay Assistance substituted for Marketing Cost	ross Margin	
	Sales & Marketing Cost	Copay Assistance substituted for Marketing Cost

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Example: How a Prescription is Paid & the Impact on Manufacturer

	Column A Column B Patient's Commercial Insurance Insurance Used Doesn't Cover w/ Patient Product Yet Deductible Not Yet Met & High Deductible Plans		Column C Commercial Insurance Used w/ Average Copay	Column D Medicare Part D Insurance Used w/ Average Copay	
Payment from Copay Card (cost to Manufacturer)	\$200	\$200	\$40	\$0	
Payment from Insurance Company	\$0	\$0	\$160	\$195	
Payment from Patient	\$ 35	\$ 35	\$ 35	\$ 40	
Total Amount Received by Pharmacy	\$235	\$235	\$235	\$235	

- For columns A and B, the copay card covers most of the cost of the product for the patient
- · For columns C and D, the insurance company pays most of the cost of the product for the patient

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IMVEXXY Catalysts





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Product Development Rationale

- 2002 Women's Health Initiative (WHI) study showed that the long-term use of certain synthetic hormones (a combination of medroxyprogesterone acetate and conjugated equine estrogens) increased the risk of breast cancer, stroke, heart attack and blood clots
 - Prior to BIJUVA, all FDA-approved combination hormonal products contain a synthetic progestin and not a bio-identical progesterone
- After WHI, women and healthcare providers shifted to bio-identical hormone therapy as an alternative despite estradiol and progesterone combinations being *unapproved* drugs for use together
- Compounding filled the need for bio-identical hormone therapy

Bijuvama

Astraction and procesteromed capsules.

- All the major medical societies and the FDA encourage the prescribing of FDA approved hormones
- > NEED FOR AN FDA-APPROVED COMBINATION BIO-IDENTICAL HORMONE THEREAPY

Hormone Therapy Options for Vasomotor Symptoms Before BIJUVA

After WHI (2002), a majority of women and clinicians shifted to bio-identical hormone therapy1

FDA-APP	ROVED	NOT FDA-APPROVED
Combination <u>Synthetic</u> Estrogens + Progestins ²	Separate <u>Bio-identical</u> Estradiol & Progesterone	Compounded <u>Bio-identica</u> Estradiol + Progesterone
~ 2.5 million total annual prescriptions ³	~ 3.9 million total annual prescriptions (each) ¹	12 - 18 million total annual prescriptions ⁴
Prempro [®] , Activella [®] , Angeliq [®] , Femhrt [®] , Climara Pro [®] , Combipatch [®]	Oral or transdermal estradiol & Prometrium®	Compounded estradiol + progesterone
FDA-approved	Not FDA-approved to be used together	Not FDA-approved
1 copay	2 copays	Often not covered by insurance
Insurance coverage	Insurance coverage	Almost 100% out of pocket

1) Symphony Health Solutions PHAST Data powered by IDV: 12 months as of December 31 2018
2) Products include synthetic progestin with synethetic or bio-identical estragen
3) includes the following drugs Activetatio: FerriHT10: Angelogi, Generic 17b + Progestins, Premproß, Premphase6, Duavea®, Brisdete®
4) Compositio of Flainet, J. QuintlesMBA, PerriHT10: Angelogi, Generic 17b + Progestins, Premproß, Premphase6, Duavea®, Brisdete®
4) Compositio of Flainet, J. QuintlesMBA, PerriHT10: Angelogi, Generic 17b + Progestins, Premproß, Premphase6, Duavea®, Brisdete®
4) Compositio of Flainet, J. QuintlesMBA, PerriHT10: Angelogi, Generic 17b + Progestins, Premproß, Premphase6, Duavea®, Brisdete®
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4) Compositio of Flainet, J. QuintlesMBA, PerriHT10: Angelogi, Generic 17b + Progestins, Premproß, Premphase6, Duavea®, Brisdete®
4) Compositio of Flainet, J. QuintlesMBA, PerriHT10: Progestins, Prempinates, and the property of their respective owners.



BIJUVA is indicated in a woman with a uterus for the treatment of moderate to severe vasomotor symptoms due to menopause

KEY CLINICAL ATTRIBUTES

- First and only bio-identical combination of estradiol to reduce moderate to severe hot flashes combined with progesterone to help reduce the risk to the endometrium
- Strong efficacy and safety data
- Sustained steady state of estradiol
- No clinically meaningful changes in weight or blood pressure
- No clinically meaningful changes in coagulation or lipid parameters
- No clinically meaningful changes in mammograms
- Clinically meaningful improvements in quality of life and sleep disturbance data
- High desired amenorrhea rates (no bleeding)

OTHER KEY ATTRIBUTES

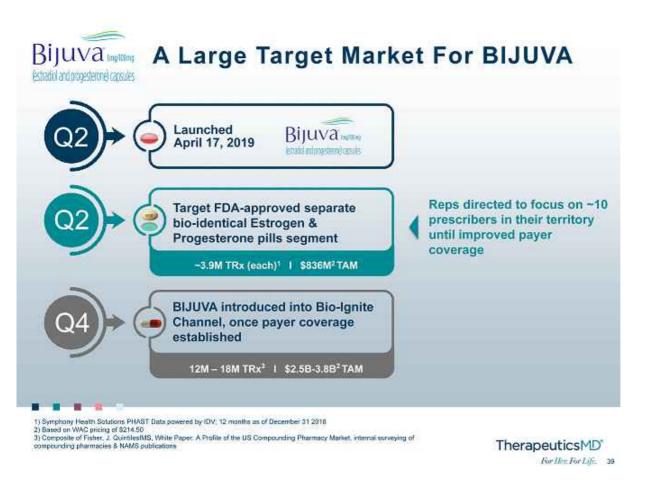
- Once-a-day single oral softgel capsule only continuous combined progesterone and estradiol product
- No peanut oil unlike other FDA-approved progesterone products
- One prescription, one copay
- BIJUVA is available in blister packages containing 30 capsules



References:

BUJUVA (package insert), Boca Raton, FL: TherapouticsMD, Inc. 2019. Labo RA, et al. Obstet Gynecol. 2018;132(1):161-170. Labo RA, et al. North American Menopause Society Annual Meeting. October 3 – 6, 2018, San Diego, CA, USA, abstract number S-2.

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Launch Plan Mirrors IMVEXXY

Focused on Driving Early Behavior Change that Leads to Long Term Adoption



Offer not valid for patients enrolled in Medicare, Medicaid, or other federal or state health care programs
 (including any state pharmaceutical assistance programs). Program Terms, Conditions, and Eligibility Criteria apply.

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Impact of Faster Payer Cycle on Adjudication Rate*



2Q 2019 Actuals

	Column A	Column B	Column C
BIJUVA	No Insurance	Commercial Insurance	Medicare Eligible Patients
% of Business	3.1%	89.1%	7.8%
% Adjudicated	0%	37.6%	7.6%
Contribution to Overall Adjudication Rate	0%	33.4%	0.6%
Overall Adjudication Rate	34% (up from 25% in M	/lay 2019)

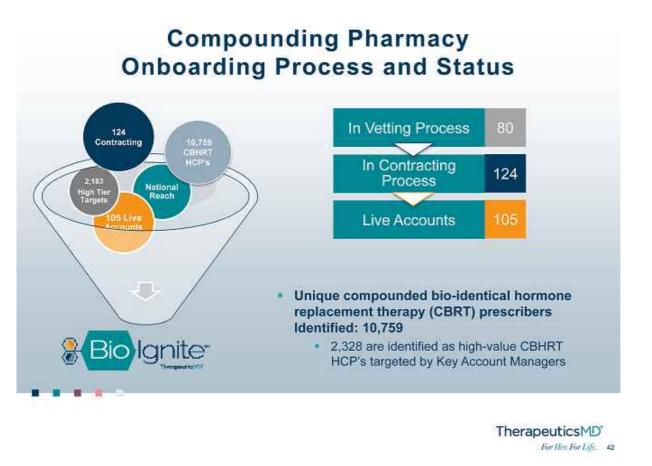


Target at Fully Established Insurance Coverage

	Column A	Column B	Column C
BIJUVA	No Insurance	Commercial Insurance	Medicare Eligible Patients
% of Business	6%	87%	8%
% Adjudicated	0%	75%	65%
Contribution to Overall Adjudication Rate	0%	65%	5%
Overall Adjudication Rate		70%	

*Adjudication Rate= Percent of Business multiplied by % of claims being covered.

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Market Opportunity

	Percer	nt of Addressable	Market	
Average Net Revenue / Unit	25%	35%	45%	55%
80	\$316M	\$442.4M	\$568.8M	\$695.2M
5100	\$395M	\$553M	\$711M	\$869M
100	\$395M	\$553M	\$711M	\$869M



BIJUVA Catalysts





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ANNOVERA: 2019 Prix Galien USA Award Nominee

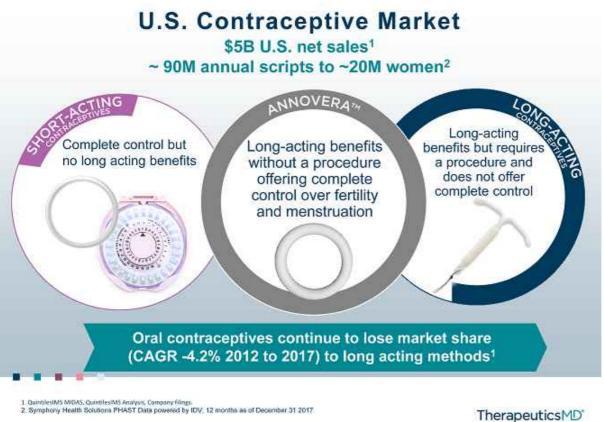


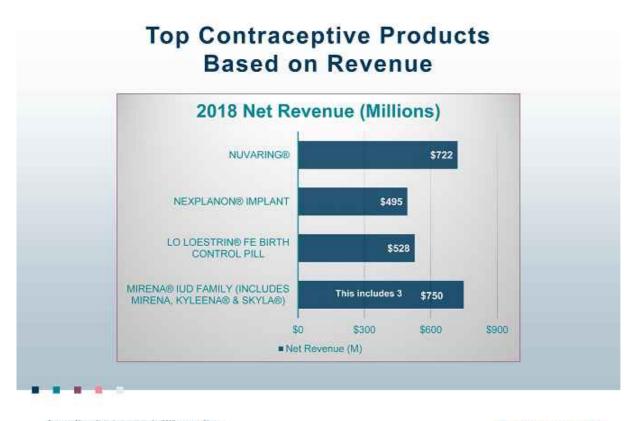
The Prix Galien Award, counted among the global health innovation industry's most prized honors, recognizes outstanding biomedical and medical technology product achievement that improves the human condition

2019 Nominee:	2	018 TRx MBS Dollars ¹
	Ibrance®	\$2,293,000,000
Annovera	IMBRUVICA®	\$2,334,000,000
(segesterme atetate and ethnivi estradici vacinal system) Devena utili anarozzati ana confer	Gleevec®	\$362,000,000
	Januvia®	\$6,237,000,000
	Chantix®	\$1,258,000,000

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Company filings: Net sales as reported in 2018 company filings All trademarks are property of their respective owners.

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Access Attributes

- Market shift to long-acting contraception
- Offer women a long-term birth control option without requiring a procedure for insertion and removal like IUDs or Implants
- Available to all prescribers no special training, equipment, or inventory
- Acceptable for women who haven't had a child (nulliparous) or are not in a monogamous relationship¹
- "Vaginal System" the only product in a potential new category of contraception with potential for \$0 co-pay
- Does not require refrigeration

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Lonv, et al. Use of intrauterine devices in nulliparaus women. Contraception 95 (2017): 529-537

Clinical Attributes



- Only FDA-approved long-lasting reversible birth control that doesn't require a procedure or repeat visit
 - Empowers women to be in control of their fertility and menstruation
 - ANNOVERA is the only user-directed single 12-month birth control product (used in repeated 4-week cycles for 13 cycles)
- Highly effective in preventing pregnancy when used as directed (97.3%)
- High patient satisfaction in clinical trials (phase 3 acceptability study of 905 women)¹
 - -89% overall satisfaction, adherence (94.3%) and continuation (78%)
- Softer and more pliable than NuvaRing®
- Only product with new novel progestin segesterone acetate²
 No androgenic or glucocorticoid effects at contraceptive doses*
- Low rates of discontinuation related to irregular bleeding (1.7%)

Merkatz, Ruh B., Mariena Plagianos, Elena Hoskin, Michael Cooney, Paul C. Hewett, and Barbara S. Mensch. 2014. "Acceptability of the Nestarone®Nethinyl estradiol contraceptive vaginal ing. Development of a model: implications for introduction," Contraception 90(5): 514–521. * Narender Kumar, Samuel S. Keide, Yun-Yen Tsong, and Kalyan Sundaram. 2000. "Nestorone: a Progestin with a Unique Pharmacological Profile," Steroids 65: 829-838

*Based on pharmacological studies in animals and in vitro receptor binding studies. The clinical significance is not known. All trademarks are the property of their respective owners.



High Patient Satisfaction

- Phase 3 acceptability study (n=905 subjects)¹
 - Overall, nearly 90% of women in a global clinical trial were satisfied with ANNOVERA as a form of contraception
 - Most women ranked ANNOVERA highly in characteristics related to:
 - Ease of use
 - Comfort
 - Expulsion
 - Physical effects during sexual activity
- High rates of adherence (94.3%)² and continuation (78%)

	Eas	e of Use	
Ease of inserting (N=905)	Ease of removing (N=905)	Ease of remembering CVS insertion (N=905)	Ease of remembering CVS removal (N=905)
90.8% (n=823)	88.2% (n=798)	87.6% (n=793)	85.2% (n=771)

Merkatz, Ruth B., Marlena Plagianos, Elena Hoskin, Michael Cooney, Paul C. Hewelt, and Barbara S. Mensch. 2014. "Acceptability of the Nestorone@rethinyl estradict contraceptive vaginal ring: Development of a model: implications for introduction," Contraception 90(5): 514–521. "Adherence is defined as fellowing the 21-day-int?-day-out cyclic regimen and not removing the vaginal system >2 hours in the proceeding cycle.

ANNOVERA Key Attributes

	Oral Contraceptives	Vaginal Ring NuvaRing [®]	Contraceptive Injection	Vaginal System ANNOVERA™	IUDs
Duration of Action	Daily pill intake	1 month (21/7 regimen)	3 months	1 year (21/7 regimen)	3-10 years
Patient Control	Stop at any time	Removable at any time	Stop at any time, but residual effects for 3 months	Removable at any time	Procedure required
Nulliparous Women	Yes	Yes	Yes	Yes	Not universally acceptable
Product Administration	Oral intake	Patient administered Flexible vaginal ring	Physician in-office injection every 3 months	Patient administered Soft and pliable ring- shaped vaginal system	Physician in-office procedure for insertion and removal
Patient Convenience	Daily pill presents compliance and adherence risks; potential increase in unplanned pregnancies	Monthly pharmacy visit	Physician in-office injection, prescriber stocking required	Annual pharmacy visit	Physician in-office procedure, prescriber stocking required
Healthcare Provider Convenience	Filled at pharmacy	Filled at pharmacy; Refrigeration required prior to being dispensed	Prescriber required to hold inventory	Filled at pharmacy; No refrigeration; No inventory or capital outlay	Prescriber required to hold inventory
Yearly WAC	Lo Loestrin® Fe: \$1,829.36	NuvaRing® \$2,114.19	Depo-Provera® \$799.12	\$2,000	Liletta® \$749.40 + \$425.25 for Insertion/removal Plus office visits and screenings

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Patient Types

- Broad-based product a single contraceptive product for most patient and prescriber types
- Supports patient preference
- Amenable to women of all reproductive ages and demographics
- Highly effective
- Self-administered, long-lasting product that is reversible
- Nulliparous women (never had a child before)
- Between children birth spacing
- Women not in monogamous relationships
- Ideal for adolescents of reproductive age who don't want to take a product everyday, but don't want a procedure or nulliparous or non-monogamous
- College women no need for monthly refills
- Women in the military control fertility for 1 year



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Launch Approach the sector of the sector of

Begins Week of September 30, 2019

Pre-Launch A	ctivities	National Launch
WAC Price set at \$2,000/ unit (1 year) Continue dialogue with FDA regarding potential ACA decision designating ANNOVERA as a new method of contraception Start payer discussions	 Test and learn market introduction Production ramps to ~10,000 units for the 4Q19 	 Initial focus on OBGYN target overlap with Menopause Products Early consumer focus given how influential women are in the choice of birth control Full-scale production anticipated 1Q20
3Q 2019	4Q 2019	1Q-3Q 2020

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Market Opportunity

	Percent	of Overall Birth Control N	arket NRx	
Average Net Revenue / Unit	1.0%	1.5%	2.0%	2.5%
\$1,000	\$280M	\$420M	\$560M	\$700M
\$1,500	\$420M	\$630M	\$840M	\$1.05M
\$1,750	\$490M	\$735M	\$980M	\$1.2M
	Addressable NuvaRin	ng Market NRx: 1.2M	1	
	Pe	rcent of NuvaRing Market	NR×	
Average Net Revenue / Unit	25%	35%	45%	55%
\$1,000	\$300M	\$420M	\$540M	\$660M
\$1,500	\$450M	\$630M	\$810M	\$990M
\$1,750	\$525M	\$735M	\$945M	\$1.15M

Market opportunity is calculated by multiplying the annual addressable market times the market share times the average potential net revenue per unit.

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The Power of the Portfolio Multiple Paths to \$1B of Peak Sales

Average Net Revenue / Unit	25%	35%	45%	55%	
\$80	\$184M	\$257.6M	\$331.2M	\$404.8M	7 Invexy
\$100	\$230M	\$322M	\$414M	\$506M	estrado legra rear engi-tis
Tota		essable FDA Market			
12		nt of Addressable N			
Average Net Revenue / Unit	25%	35%	45%	55%	Bijuva
\$80	\$316M	\$442.4M	\$568.8M	\$695.2M	establi and angele sens
\$100	\$395M	\$553M	\$711M	\$869M	
	Total Addressa	ble Birth Control M	arket NRx: 28M		
Average Net Revenue / Unit	1.0%	1.5%	2.0%	2.5%	Annovero
\$1,000	\$280M	\$420M	\$560M	\$700M	Interesting Street
\$1,500	\$420M	\$630M	\$840M	\$1.05M	
\$1,750	\$490M	\$735M	\$980M	\$1.2M	

Diversified risk with 3 FDA-approved products, creating multiple paths to \$1B peak sales opportunity Example: \$230M (IMVEXXY), \$395M (BIJUVA) and \$420M (ANNOVERA) = \$1B peak sales potential Therapeutics MD*



2019 Financial Guidance

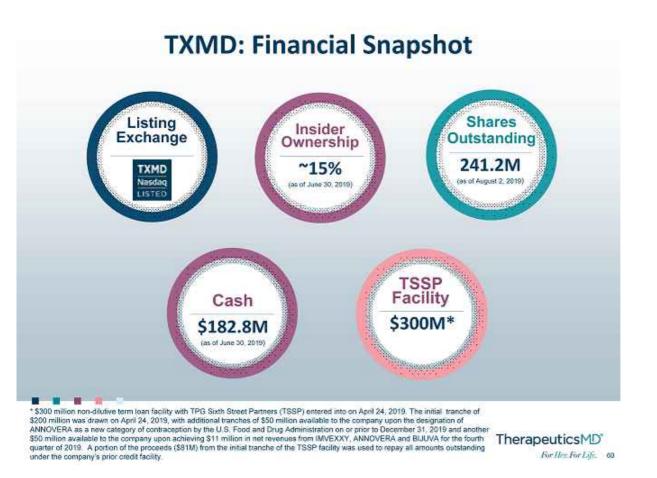
	3Q2019 Estimate	4Q2019 Estimate	FY2019 Estimate
FDA-Approved Products Net Revenue	\$4.50 - 6.50M	\$11.00 - 13.00M	\$20.75 - 24.75M
Prenatal Vitamins Net Revenue	\$2.25 - 2.50M	\$1.75 - 2.25M	\$8.70 - 9.45M
Total TXMD Net Revenue	\$6.75 - 9.00M	\$12.75 - 15.25M	\$29.45 - 34.20M
Important Guidance Notes: As our sales force focus shifts increase for prenatal vitamins, percentage of overall company	we anticipate prenatal vi		

\$300M Non-Dilutive Term Loan Financing Secured

\$200M accessed to date with up to additional \$100M through Specific Company Milestones

	Amount (\$)	TXMD Company Milestone ¹	Anticipated Timing
Tranche 1	\$200 million	Closing of the facility	Completed in April 2019
Tranche 2	\$50 million	Designation of ANNOVERA as a new category of birth control by the U.S. Food and Drug Administration on or prior to December 31, 2019	Second Half of 2019
Tranche 3	\$50 million	Achieving \$11 million in net revenues from IMVEXXY, BIJUVA and ANNOVERA for the fourth quarter of 2019	First Quarter of 2020

 TXMD Company Milestones are draw triggers for additional tranches of funding only and are not affirmative covenants that the company must otherwise meet. Ability to draw additional tranches is also subject to satisfaction (or waiver) of other customary conditions precedent.







IMVEXXY Quarterly Performance

Invexxy (estradol vagral inserts)

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