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) January 8, 2018

DARÉ BIOSCIENCE, INC.

(Exact name of registrant as specified in its charter)

Delaware (State or other jurisdiction of incorporation) 001-36395 (Commission File Number) 20-4139823 (I.R.S. Employer Identification No.)

11119 North Torrey Pines Road, Suite 200 La Jolla, CA 92037 (Address of Principal Executive Offices and Zip Code)

Registrant's telephone number, including area code (858) 926-7655

Not Applicable (Former name or former address, if changed since last report.)

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 (see General Instruction A.2. below):

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 $\ oxtimes$

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.

On January 8, 2018, Daré Bioscience, Inc. (the "Company") posted a corporate presentation to its website, which presentation the Company plans to present at various investor and analyst meetings. The presentation is attached hereto as Exhibit 99.1.

The information disclosed under this Item 7.01, including Exhibit 99.1 hereto, is being furnished and shall not be deemed "filed" for purposes of Section 18 of the Securities Exchange Act of 1934, as amended, nor shall it be incorporated by reference into any registration statement or other document pursuant to the Securities Act of 1933, as amended, except as expressly set forth in such filing.

Item 9.01 Financial Statements and Exhibits.

The following exhibit is furnished with this report:

Exhibit Number

Description

99.1 <u>Corporate Presentation prepared as of January 8, 2018.</u>

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.

DARÉ BIOSCIENCE, INC.

Dated: January 8, 2018

By: Name: Title:

/s/ Sabrina Martucci Johnson Sabrina Martucci Johnson President and Chief Executive Officer





THIS PRESENTATION IS FOR INFORMATIONAL PURPOSES ONLY AND IS NOT AN OFFER TO SELL OR A SOLICITATION OF AN OFFER TO BUY ANY SECURITIES OF DARÉ BIOSCIENCE, INC. ("DARE" OR THE "COMPANY"). THIS PRESENTATION INCLUDES CERTAIN INFORMATION OBTAINED FROM TRADE AND STATISTICAL SERVICES, THIRD PARTY PUBLICATIONS, AND OTHER SOURCES. DARÉ HAS NOT INDEPENDENTLY VERIFIED SUCH INFORMATION AND THERE CAN BE NO ASSURANCE AS TO ITS ACCURACY. FURTHER, THIS PRESENTATION SPEAKS AS OF THE DATE SET FORTH HEREIN, AND THE DELIVERY OF THIS PRESENTATION SHALL NOT, UNDER ANY CIRCUMSTANCES, CREATE AN IMPLICATION THAT THERE HAS BEEN NO CHANGE IN DARÉ'S AFFAIRS AFTER THE DATE SET FORTH HEREIN.

THIS PRESENTATION CONTAINS FORWARD LOOKING STATEMENTS WITHIN THE MEANING OF THE PRIVATE SECURITIES LITIGATION REFORM ACT OF 1995 AND OTHER FEDERAL SECURITIES LAWS. IN SOME CASES, YOU CAN IDENTIFY FORWARD LOOKING STATEMENTS BY TERMS SUCH AS "MAY," "WILL," "EXPECT," "PLAN," "ANTICIPATE," "STRATEGY," "DESIGNED," "CURRENTLY," "COULD," "INTEND," "BELIEVE," "ESTIMATE," OR "POTENTIAL" OR THE NEGATIVE OF THESE TERMS OR OTHER SIMILAR EXPRESSIONS. THE FORWARD LOOKING STATEMENTS OF THIS PRESENTATION ARE ONLY PREDICTIONS AND ARE SUBJECT TO A NUMBER OF RISKS, UNCERTAINTIES AND ASSUMPTIONS, INCLUDING, WITHOUT LIMITATION RISKS AND UNCERTAINTIES RELATING TO: THE OUTCOME OR SUCCESS OF DARÉ'S CLINICAL TRIALS; DARÉ'S ABILITY TO MAINTAIN AND PROTECT ITS INTELLECTUAL PROPERTY; AND DARÉ'S ABILITY TO DEVELOP OVAPRENE® AT THE COST AND IN THE TIMELINE SET FORTH HEREIN. THE FORWARD LOOKING STATEMENTS IN THIS PRESENTATION REPRESENT DARÉ'S VIEWS ONLY AS OF THE DATE OF THIS PRESENTATION, AND DARÉ UNDERTAKES NO OBLIGATION TO UPDATE OR REVIEW ANY FORWARD LOOKING STATEMENT, WHETHER AS A RESULT OF NEW INFORMATION, FUTURE DEVELOPMENTS OR OTHERWISE, EXCEPT AS REQUIRED BY LAW.





- Mission: Identify, develop and bring to market a diverse portfolio of novel therapies that expand treatment options, improve outcomes and facilitate convenience for women.
- · Deliver innovation by filling the development gap
 - Fragmented market segment between early and late stage development and the for-profit and non-profit sectors — promising candidates available for licensing, some with existing proof-of-concept human clinical data.
- Initial areas of focus: contraception, sexual & vaginal health, fertility
- Women's health therapeutics global market is expected to reach \$22 billion by 2018.⁽¹⁾
- Global contraception market alone is expected to reach \$33 billion by 2023.⁽²⁾

(1) BCC Research

(2) Global Market Insights





Ovaprene™: convenient non-hormonal contraceptive method

- Unique combination approach seeks to generate effectiveness in the range of the birth control pill
- Enrollment to begin in early 2018 for Postcoital Test (PCT) Clinical Trial; trial completion expected in 2nd half 2019.
- 12 issued patents worldwide (9 U.S.); key patent's last claims expire August 2028, potential extension to 2033

Opportunities for growth and expansion

 Actively identifying and evaluating potential product candidates in women's health to license or develop in addition to Ovaprene™

· Partnering strategy to maximize efficiencies and drive value

- Ideal co-development partners are non-profit organizations with prior U.S. regulatory successes
- Commercialization via an established pharma partner
- Experienced management team with deep expertise in women's health





All about hormones

- Advances beyond "the pill" have largely focused on new combinations, doses, durations and modes of administration
- · Successful hormonal commercial brands, easy to use
 - NuvaRing®
 - Merck's hormonal monthly vaginal ring (2016 revenue: \$777 million)
 - Mirena®
 - Bayer's hormonal intrauterine system (2016 revenue: \$1.13 billion)
- What's lacking?
 - Advances in easy-to-use and effective non-hormonal alternatives

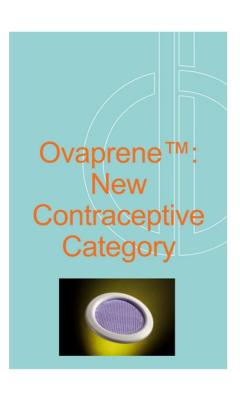
The current contraceptive method mix still fails to meet the needs of all women.





- The global market for contraception was \$19 billion in 2015 and is estimated to grow over 6% to \$33 billion in 2023.
- 40 million women of reproductive age in the U.S. are currently using a contraceptive method, about 62% of the population. Fifty percent of them are using a short-acting reversible method. (2)
- 4 in 10 women are not satisfied with their current method. (3)
- Twenty-two percent of women currently use non-hormonal short-acting methods as their primary form of birth control despite their limitations. (2)
- Condom remains the most widely used non-hormonal option.
- (1) Global Market Insights, Inc. released in February, 2017
- www.gutmacher.org, contraceptive fact sheet
- (3) Ersek, J, Matern Child Health J (2011) 15:497–506





Provides monthly contraceptive protection without hormones

- Silicone reinforced ring with a soft, absorbable scaffolding that encircles a fluid-permeable barrier
- Non-braided, multi-filament mesh functions as a physical barrier to sperm
- Contraceptive-loaded silicone matrix releases non-hormonal ingredients, ferrous gluconate and ascorbic acid, that act together to create a spermiostatic environment.

Prevents advancement of viable sperm

 In a previous human postcoital test clinical study, Ovaprene™ prevented sperm from advancing into the cervical mucus.

No intervention required at the time of intercourse

 Intravaginal ring (IVR) whose monthly usage is similar to that of NuvaRing®. If approved, it will require a physician Rx.



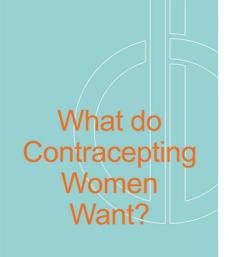


Contraceptive Effectiveness		
Contraceptive Method	Typical Use	Perfect Use
No Method	15.0%	15.0%
Fertility awareness-based methods	76.0%	95.0%
Withdrawal	78.0%	96.0%
Male Condom	82.0%	98.0%
Female condom	79.0%	95.0%
Diaphragm	88.0%	94.0%
Spermicides	72.0%	82.0%
Sponge	88.0%	91.0%
Pill	91.0%	99.7%
Transdermal Patch (OrthoEvra)	91.0%	99.7%
Vaginal ring (NuvaRing)	91.0%	99.7%
Injection (Depo-Provera)	94.0%	99.8%
Implant (Implanon)	100.0%	100.0%
Copper T IUD (ParaGard)	99.2%	99.4%
Levonorgestrel IUD (Mirena)	99.8%	99.8%
Vasectomy	99.9%	99.9%
Female sterilization (tubal ligation)	99.5%	99.5%

A non-hormonal monthly option that is safe and effective, and provides protection over multiple weeks, could fill a current gap.

Source: Guttmacher (<u>www.guttmacher.org</u>), FDA.gov





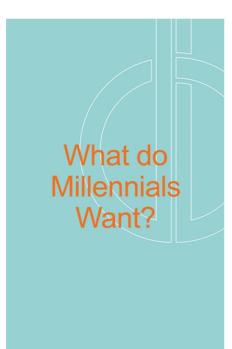
Features desired most in birth control:

- √ Efficacy √ Ease of Use √ Ease of Adherence ✓ No or Low Side Effects √ Easy Reversal
- 67 percent of women report that the vaginal ring has most of the features deemed extremely important(1)
- 85 percent of women would prefer a monthly option with a lower hormone dose than the pill(2)
- 80 percent currently use a non-coital dependent method(3)



⁽¹⁾ Lessard, L, Perspectives on Sexual and Reproductive Health, Volume 44, Number 3,9-2012

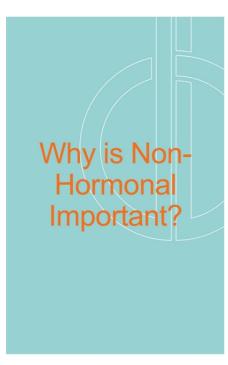
⁽²⁾ Hooper, DJ, Clin Drug Investig. 2010;30(11):74963 (3) Ersek, J, Matern Child Health J (2011) 15:497–506



- A survey of 1,000+ sexually active women aged 18-29 trying to avoid pregnancy revealed⁽¹⁾:
 - 33% are uneasy about what taking the pill might mean for their fertility in the future
 - 26% are scared by the medical warnings they see in the TV commercials
 - 25% say taking something everyday "doesn't work for me"
 - 19% are afraid of a possible side effect like weight gain
 - 17% say they are uncomfortable taking birth control pills because doing so is unnatural







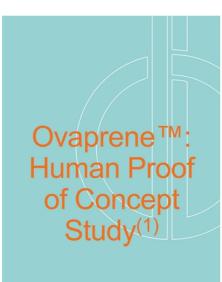
 Many women cannot tolerate hormones – breast tenderness, weight gain, mood swings, decreased sex drive and general discomfort

• Other women are contraindicated for long-term hormone use⁽¹⁾:

- Over 35 years old and smoke or have certain kinds of severe migraine headaches
- Have (or have had) breast cancer or any cancer that is sensitive to female hormones
- Have high blood pressure that medicine can't control
- Have (or have had) blood clots in arms, legs, eyes or lungs
- Have an inherited blood problem that makes it clot more than normal
- Have had a stroke or heart attack
- · Have certain heart valve problems or heart rhythm problems
- · Have diabetes with kidney, eye, nerve or blood vessel damage
- Have liver disease, including liver tumors
- · Have unexplained vaginal bleeding
- Are allergic to hormones used in pills, rings and injections



(1) NuvaRing® hormonal IVR drug label



Postcoital Test (PCT) Clinical Trial Study Design:

 Open-label, single-arm, pilot safety and tolerability study as a barrier contraceptive

Patients:

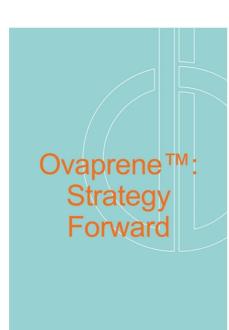
21 women enrolled; all women completed one cycle of use

· Results:

- Postcoital testing revealed no viable sperm in the cervical mucus
- No colposcopic abnormalities, no significant changes in vaginal flora and no serious adverse effects observed

In PCT studies of similar size (Lea's Shield, FemCap, Ortho diaphragm and the SILCS (Caya®) diaphragm), those products with no motile sperm in the cervical mucus during their PCT assessments demonstrated "typical use" contraceptive effectiveness of 88% in pivotal contraceptive studies evaluating pregnancy rates over time.

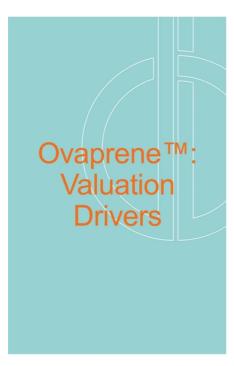


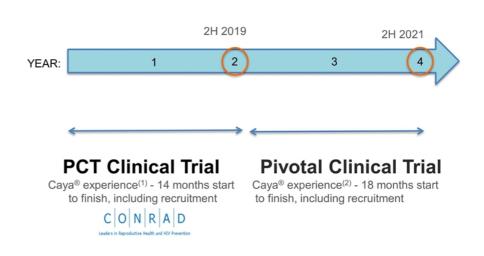


U.S. Regulatory Strategy

- Previous FDA Request for Designation (RFD) determination
- Ovaprene™ will require a PMA with CDRH as lead review division
- Clinical path forward expected to be based on prior CDRH approvals (Example: Caya® diaphragm)
 - **Predictive efficacy** postcoital test (PCT) clinical trial in 15-30 women with 1-3 months of use
 - Proportion of women with an average of fewer than 5 progressively motile sperm per high powered field (HPF) in the cervical mucus 2-3 hours following coitus
 - · Safety assessments, PK, acceptability, fit, and ease of use
 - Single pivotal clinical trial with ~250 completers over 6 months of use
 - Pregnancy probability
 - · Safety, acceptability, fit, ease of use, assessments of vaginal health







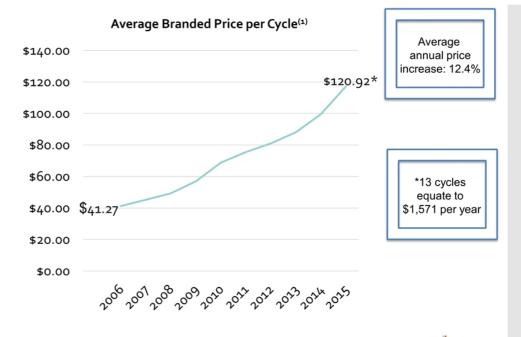
This timeline reflects management's current estimates and constitutes a forward looking statement subject to qualifications elsewhere in the presentation. The actual development timeline may be substantially longer, and Daré is under no obligation to update or review this estimate.

(1) J.L. Schwartz et al. / Contraception 78 (2008) 237-244.

(2) Schwartz JL, Weiner DH, Lai JJ, et al. Contraceptive efficacy, safety, fit, and acceptability of SILCS, a novel single-size diaphragm. 2014.

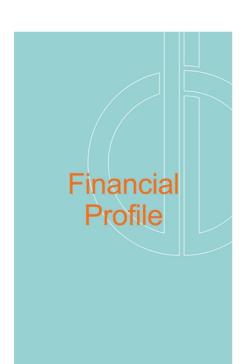
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(1) MediSpan Price Rx Select, January 2016. Average prices of 13 branded contraceptive products.





· Background:

- NASDAQ:DARE
- · Publicly traded via stock purchase agreement (reverse merger), closed July 19, 2017
- Cash at closing sufficient through 2 years, to value inflection point of lead candidate Ovaprene™

· Balance Sheet Data:

- Cash \$8.5 million at 9/30/17
- 6 million shares outstanding
- No debt



Experience	d
Team to	
Move the	
Field Forwar	d

Sabrina Martucci Johnson President and CEO	WCG, Cypress Bioscience, Advanced Tissue Sciences, Baxter Healthcare
Lisa Walters-Hoffert Chief Financial Officer	ROTH Capital Partners, Citicorp Securities, Bank of America, Oppenheimer & Co.
Mark Walters VP, Operations	Pacira, SkyePharma, Alliance, American Home Products
Mike Gendreau, MD, PhD Medical Affairs	Cypress Bioscience, Microprobe, Battelle
Bridget Martell, MD Scientific Affairs	Juniper Pharmaceuticals, Purdue Pharma, Pfizer
Christine Mauck, MD, MPH Clinical Affairs	CONRAD, Population Council, RW Johnson, FDA
Mary Jarosz Global Regulatory Affairs	WCG, Evofem, Abbott Laboratories
Randall Freund Quality Assurance	WCG, Amylin, Cardinal Health, Advantar Labs
Nadene Zack Clinical Operations	Retrophin, Aragon, Cypress Bioscience, Pfizer



Roger Hawley (Chairman)	Zogenix, Alios Biopharma, InterMune, Elan
Susan Kelley, MD	Cerulean, Bayer, BMS, ArQule
William Rastetter, PhD	Cerulean, GRAIL, Receptos, Illumina, IDEC
Robin Steele, JD, LLM	InterMune, Elan, Alvea, Alios Biopharma,
Sabrina Martucci Johnson	WCG, Cypress Bioscience, Advanced Tissue Sciences, Baxter Healthcare



- Ovaprene™: convenient non-hormonal contraceptive method
 - Intended to provide protection over multiple weeks of use, requiring no intervention at the time of intercourse
 - Seeks to generate effectiveness in the range of the birth control pill
 - Near term catalyst enrollment to begin in early 2018 for PCT
- Opportunities for growth and expansion
 - Women's health multi-billion dollar market
 - · Actively evaluating promising candidates available for licensing
 - Daré can fill the gaps in the market by addressing the development gap
- Experienced management team with deep expertise in women's health and success in prior ventures in achieving regulatory approvals and partnering in novel product categories



