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

Form 6-K

REPORT OF FOREIGN PRIVATE ISSUER PURSUANT TO SECTION 13a-16 OR 15d-16 UNDER THE SECURITIES EXCHANGE ACT OF 1934

December 4, 2017

001-36686
(Commission file number)

Forward Pharma A/S

(Translation of registrant's name into English)

Østergade 24A, 1st Floor
1100 Copenhagen K, Denmark
(Address of principal executive office)

Indicate by check mark whether the registrant files or will file annual reports under cover of Form 20-F or Form 40-F.

Form 20-F [X] Form 40-F [ ]

Indicate by check mark if the registrant is submitting the Form 6-K in paper as permitted by Regulation S-T Rule 101(b)(1): Yes [ ] No [X]

Indicate by check mark if the registrant is submitting the Form 6-K in paper as permitted by Regulation S-T Rule 101(b)(7): Yes [ ] No [X]

Indicate by check mark whether by furnishing the information contained in this Form, the registrant is also thereby furnishing the information to the Commission pursuant to Rule 12g3-2(b) under the Securities Exchange Act of 1934. Yes [ ] No [X]

If "Yes" is marked, indicate below the file number assigned to the registrant in connection with Rule 12g3-2(b): N/A

Item 1. Company Presentation

On December 4, 2017, Forward Pharma A/S (the "Company") made available on its website a copy of an investor presentation which was also used in meetings with investors at the Global Mizuho Investor Conference 2017. A copy of this presentation is filed herewith as Exhibit 99.1

The fact that this presentation is being made available and filed herewith should not be deemed an admission as to the materiality of ny information contained in the materials. The information contained in the presentation is being provided as of December 4, 2017 and the Company does not undertake any obligation to update the presentation in the future or to update forward-looking statements to reflect subsequent actual results.

SIGNATURE

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, thereunto duly authorized.

Date: December 4, 2017

Forward Pharma A/S

By: /s/ Claus Bo Svendsen

Name: Claus Bo Svendsen

Title: Chief Executive Officer

EXHIBIT INDEX

99.1	Investor Presentation dated December 4, 2017

Exhibit 99.1

FORWARD PHARMA

Forward Pharma (*NASDAQ:FWP*) Corporate Update

December 4, 2017

Claus Bo Svendsen, MD, PhD
Chief Executive Officer

Forward-Looking Statements

FORWARD PHARMA

Certain statements in this presentation may constitute "forward-looking statements" of Forward Pharma A/S within the meaning of the Private Securities Litigation Reform Act of 1995. Forward-looking statements include, but are not limited to, statements which contain language such as "believe", "expect", "anticipate", "estimate", "would", "may", "plan" and "potential". Forward-looking statements are predictions only, which involve known and unknown risks, uncertainties and other factors that may cause actual results to be materially different from those expressed in such statements. Many such risks, uncertainties and other factors are taken into account as part of our assumptions underlying these forward-looking statements and include, among others, risks related to the following: the satisfaction of certain conditions, and the accuracy of certain representations of the Company, in the Settlement and License Agreement entered into with subsidiaries of Biogen Inc. and certain other parties thereto; our ability to obtain, maintain, enforce and defend issued patents with royalty-bearing claims; our ability to prevail in the interference proceeding after all appeals and obtain issuance of the '871 application; our ability to prevail in or obtain a favorable decision in the '355 European opposition proceedings, after all appeals; the expected timing for key activities and an ultimate ruling in such legal proceedings; the issuance and term of our patents; future sales of Tecfidera®, including impact on such sales from competition, generic challenges, regulatory involvement and pricing pressures; the scope, validity and enforceability of our intellectual property rights in general and the impact on us of patents and other intellectual property rights of third parties; and our ability to generate revenue from product sales in the U.S. directly or through an assignee of our U.S. co-exclusive license rights in the event Biogen does not obtain an exclusive license from us in the U.S. Certain of these and other risk factors are identified and described in detail in certain of our filings with the United States Securities and Exchange Commission, including our Annual Report on Form 20-F for the year ended December 31, 2016. We are providing this information as of the date of this presentation and do not undertake any obligation to update any forward-looking statements contained in this presentation as a result of new information, future events or otherwise.

Investment Highlights

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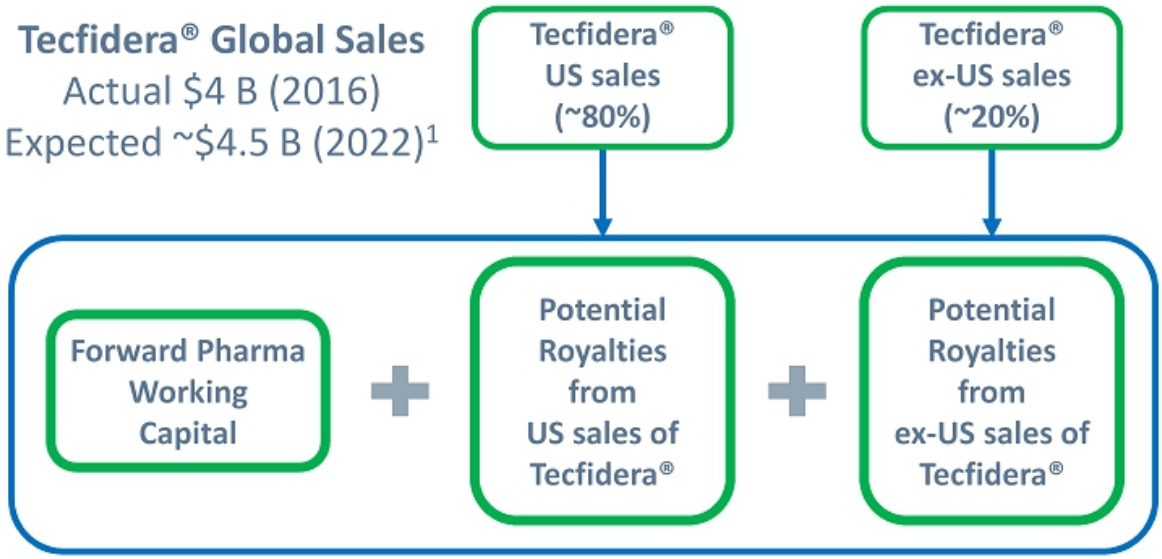
- Tecfidera® (DMF) remains a leading therapy for multiple sclerosis
  - 2022 forecast to $4.46B in global sales
  - US sales ~80%; ex-US sales ~20%
- FWP has IP-gated access to future royalties on Tecfidera® sales
  - Irrevocable license to all DMF IP granted to Biogen in January 2017
  - Potential future royalties on Tecfidera® sales in US and EU dependent on upcoming legal decisions expected in 2018
  - Top legal teams driving appeal of U.S. patent interference decision and European opposition proceeding
- Business optimized to support ongoing IP strategy and continuing obligations per settlement & license agreement
- Capital reduction and shareholder distribution of EUR 917.7 M effected September 2017

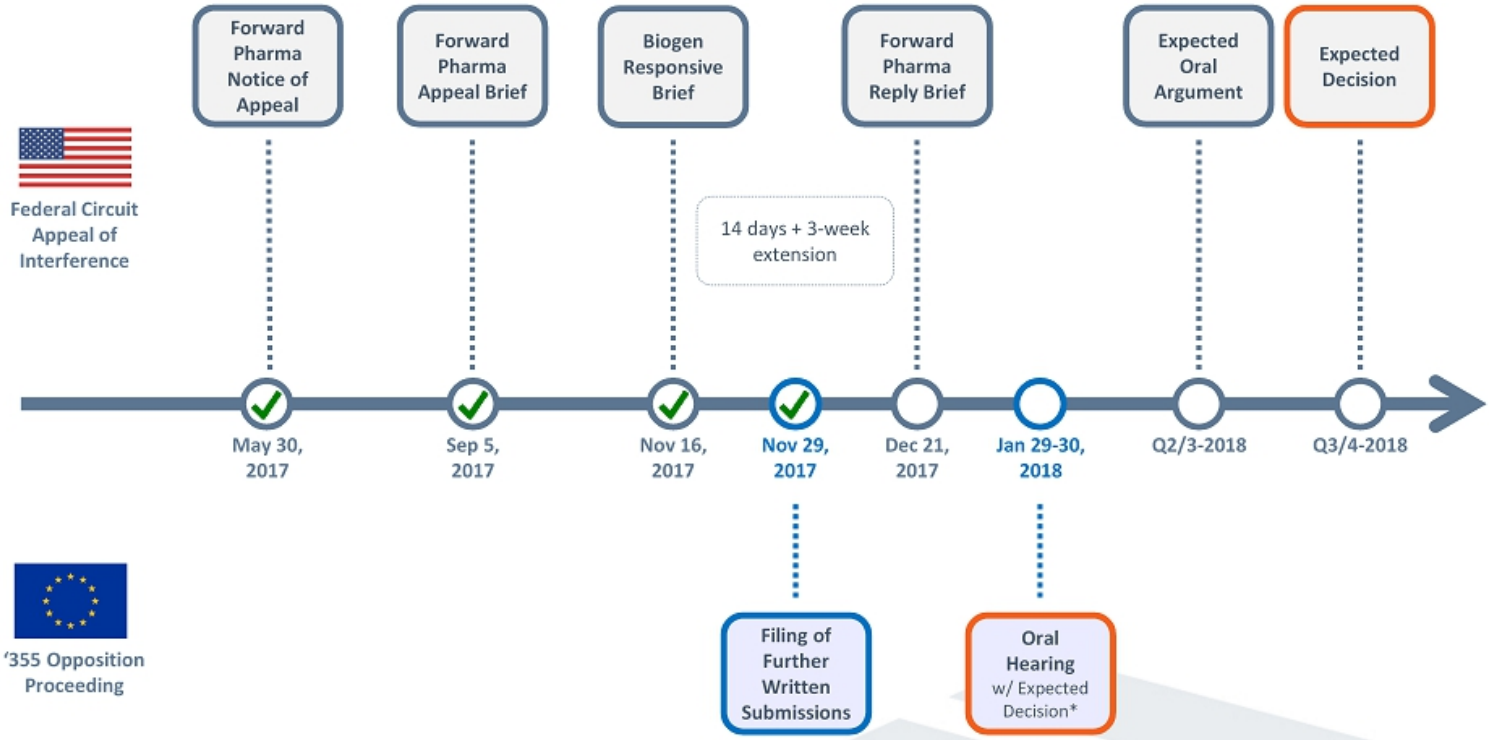
Share Value Drivers under the Settlement and License Agreement

FORWARD PHARMA



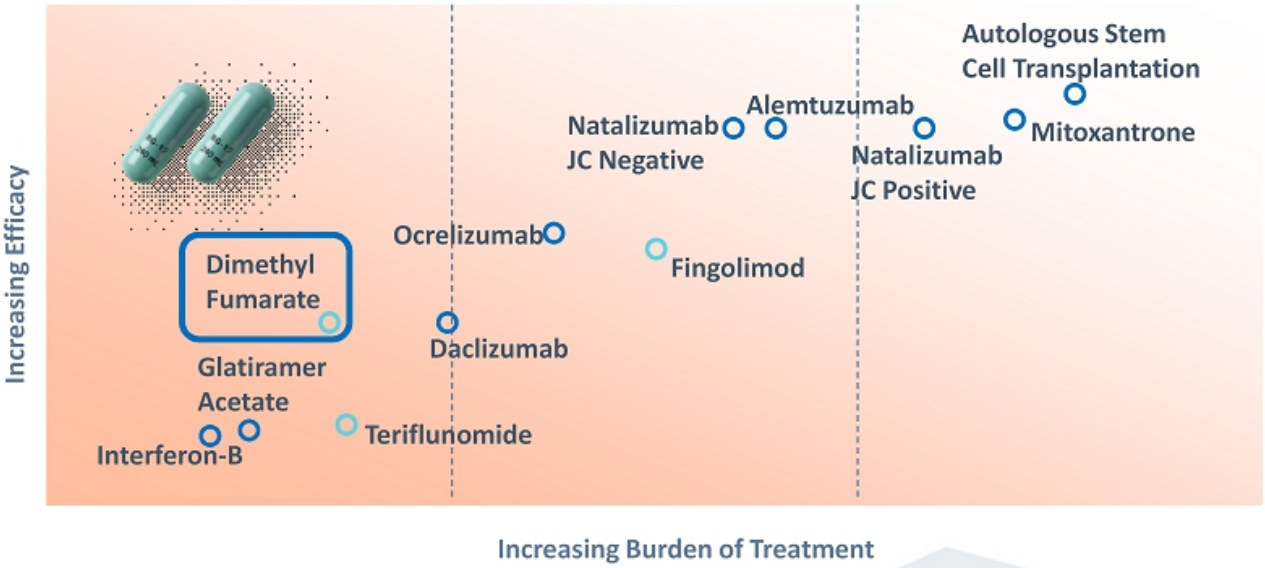
1. Analyst consensus estimates, EvaluatePharma, May 2017. Not meant to be comprehensive; size of boxes is arbitrary and not meant to illustrate comparative differences in amounts. Appropriate risk-adjustment should be applied. Potential investors and current shareholders are strongly urged to read the Settlement and License Agreement in its entirety as well as our Annual Report on Form 20-F for the year ended December 31, 2016, where risk factors are identified and described in detail.

Timeline for IP litigation in U.S. and Europe



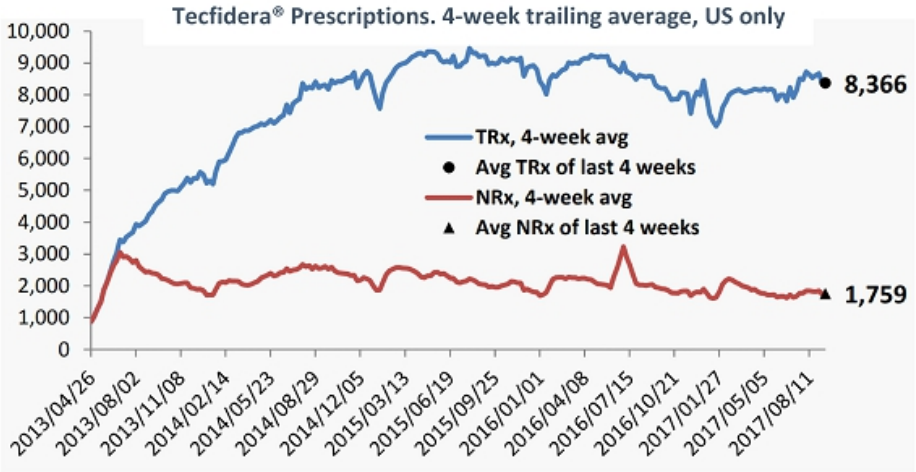
Dates represent current timeline; a green tick mark signifies actual date of completed event.
Documents for the US appeal can be located through https://ecf.cafc.uscourts.gov/ and for the European Opposition through https://register.epo.org/regviewer
* Opposition Division typically issues a decision at the conclusion of the Oral Hearing, with more detailed reasons issued later. Possibility for appeal of the decision of the Opposition Division to the Technical Board of Appeal, with conclusion in an additional 2-3 years.

Risk-Reward guides choice of therapy in MS



Adapted from **Coles A, Newer therapies for multiple sclerosis. Ann Indian Acad Neurol 2015;18, Suppl S1:30-4**

Factors Influencing Future Tecfidera® Sales



- **Regulatory Data Exclusivity and Patent Protection**
  - Settlement and License Agreement adds Forward Pharma IP
- **Launch of Ocrevus®**
  - Based on analyst reports and interviews with EU and US KOLs, Ocrevus® is initially converting later stage patients on injectables and adding a Progressive MS market
- **Potential Generic Fingolimod and additional S1P modulators**

Gating Events for Royalties on Tecfidera® Sales

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US

↓

Gating event

If we prevail after all appeals to the Federal Circuit in the Interference Proceeding and as a result thereof obtain issuance of a patent with a claim that covers treatment for MS by orally administering 480 mg per day of DMF

↓

US Royalties as described on next slide

Ex-US

↓

Gating event

If we prevail in the EP'355 Opposition, including any appeals, and as a result thereof obtain issuance of a patent with a claim that covers treatment for MS by orally administering 480 mg per day of DMF

↓

Ex-US Royalties as described on next slide

The summary of the Settlement and License Agreement in this presentation does not purport to be complete and is subject to, and qualified in its entirety by, the full text of the Settlement and License Agreement, which is available on Forward's website. Potential investors and current shareholders are strongly urged to read the Settlement and License Agreement in its entirety as well as our Annual Report on Form 20-F for the year ended December 31, 2016.

Potential Royalty Rates on Tecfidera® Sales

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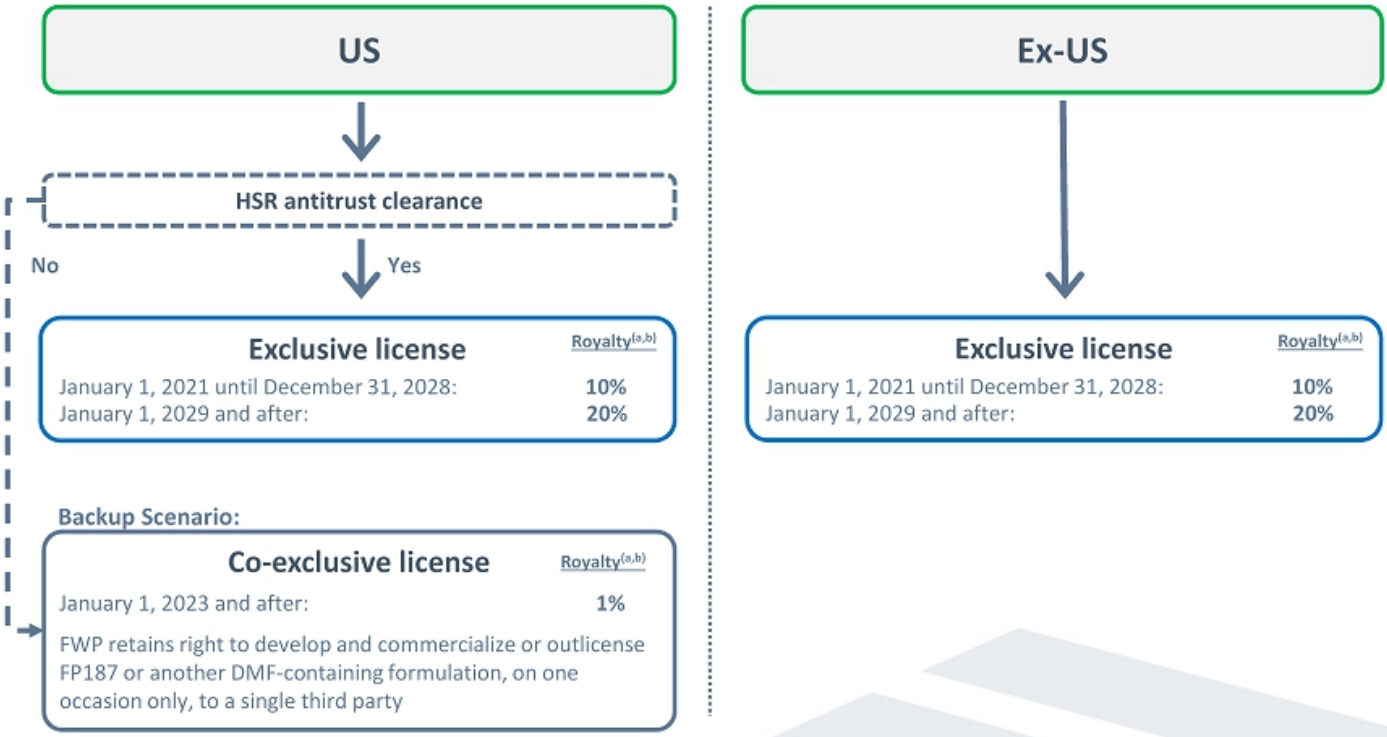

(a) Subject to, among other things, expiration or invalidation of the patents or impact of generic entry on a country-by-country basis, as defined in the License Agreement
(b) Royalties payable on a country-by-country basis on DMF-containing products indicated for MS that, but for the License Agreement, would infringe a Forward licensed patent

USPTO Interference Proceeding

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- **Interference declared April 13, 2015**

  A patent interference is an administrative proceeding at the USPTO used to determine which party is the first to invent a common invention claimed by both parties.

- **Forward awarded "Senior Party" status**

  The Senior Party has the earliest effective filing date to the common invention; entitled to the presumption that it is the first inventor.

- **On March 31, 2017, the USPTO PTAB ruled in favor of Biogen**

  Without addressing which party was the first to invent the common invention claimed by both parties, the PTAB concluded that the Forward patent application did not have sufficient written description support for the claimed invention.

- **Forward is appealing the ruling to the U.S. Court of Appeals for the Federal Circuit**

  Specialist team led by Kathleen Sullivan from Quinn Emanuel Urquhart & Sullivan, LLP.

  Forward Pharma Reply Brief due on December 21, 2017.

  Should the Forward appeal be successful, the interference will be returned to the USPTO to resume the interference proceeding. After completion of the interference proceeding, a further appeal to the U.S. Court of Appeals is possible.

European EP'355 Opposition Proceedings

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- **EP2801355 patent granted by European Patent Office (EPO) on May 20, 2015**
- **Subject to several oppositions filed with the EPO by third parties (including Biogen)**
- **The first instance hearing of the Opposition Proceedings in the EPO is scheduled for January 29-30, 2018**

  Opposition Division typically issues a decision at the conclusion of the Oral Hearing, with more detailed reasons issued later.

  Possibility for appeal of decision of Opposition Division to the Technical Board of Appeal, with conclusion in an additional 2-3 years.

Capital Reduction Executed in September

FORWARD PHARMA

- Decision was based on a careful evaluation of the most appropriate capital allocation strategy after receipt of the non-refundable $ 1.25 billion cash fee from Biogen

- **EUR 917.7 M in total returned to shareholders**

- The capital reduction is the final step of the organizational transformation to align the amount of working capital with the adjusted business activities following the Settlement and License Agreement with Biogen.

Balance Sheet and Operating Results

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- Capital Reduction completed in September 2017
  - EUR 917.7 M distributed to shareholders
  - Current number of issued shares is 94,367,998, of which 24.34% are listed as ADSs on NASDAQ (1 ADS represents 2 shares)

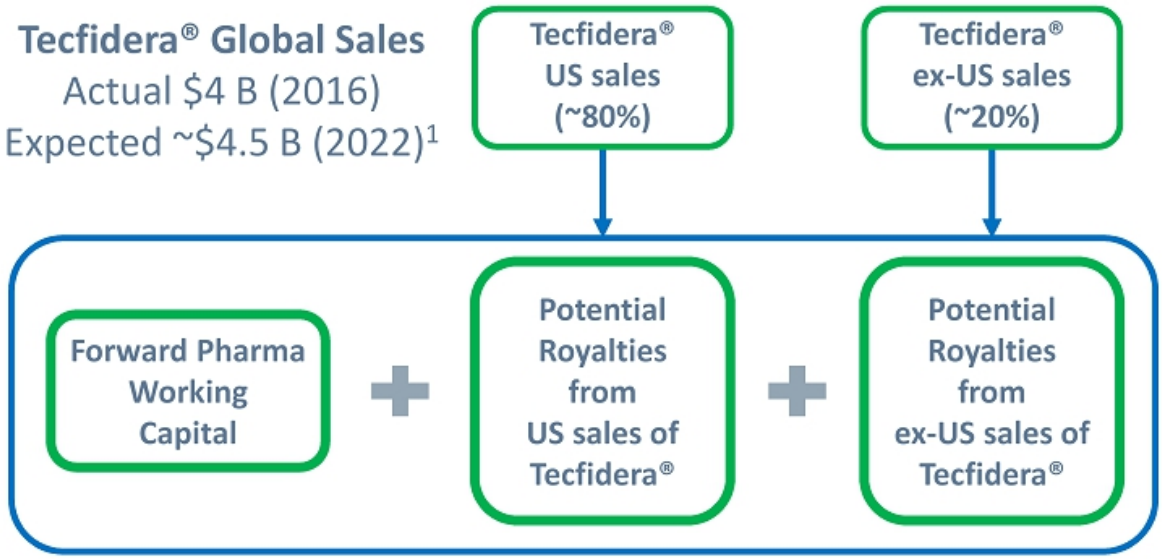
- Stock information (*NASDAQ: FWP*; per November 29, 2017)
  - Closing price: $ 4.20
  - Market Cap : $ 198.2 M

Balance Sheet	**At June 30, 2017** USD '000s
Cash and Investments	1,440,441
Other assets	1,393
Total assets	1,441,834
Total shareholder equity	1,174,295
Total liabilities	267,539
Total shareholder equity and liabilities	1,441,834

Operating Statement	**6 months ended June 30, 2017** USD '000s
Revenue	1,250,000
Operating expenses	(36,406)
Other expenses (net)	(584)
Income tax expense	(271,774)
Net income	941,236

Share Value Drivers under the Settlement and License Agreement

FORWARD PHARMA



1. Analyst consensus estimates, EvaluatePharma, May 2017. Not meant to be comprehensive; size of boxes is arbitrary and not meant to illustrate comparative differences in amounts. Appropriate risk-adjustment should be applied. Potential investors and current shareholders are strongly urged to read the Settlement and License Agreement in its entirety as well as our Annual Report on Form 20-F for the year ended December 31, 2016, where risk factors are identified and described in detail.

Upcoming Events

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- **Upcoming qualifiers for future royalty from Tecfidera® sales**
  - **Appeal of U.S. PTAB interference decision to Federal Circuit**
    - Expected Oral Argument – Q2/3 2018
    - **Expected Decision – Q3/4 2018**
  - **European EP'355 opposition**
    - **Opposition Hearing with expected decision – January 29-30, 2018**
    - Expected Written Decision with arguments – Q2 2018

Contact

Claus Bo Svendsen, MD, PhD
Chief Executive Officer

Forward Pharma Investor Relations
investors@forward-pharma.com

Appendix

US Potential Royalty-Bearing Claims

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4 patent applications are currently pending that, if issued, may contain claims that may be royalty-bearing if Forward obtains a Relevant Patent after all appeals in the Interference Proceeding

	Application Number	Description
US	11/576,871	Treating MS with DMF at 480 mg/day **Interference declared; FWP as Senior Party – April 13, 2015** **PTAB ruled in favor of Biogen – March 31, 2017** **Final Brief in the Appeal to the Federal Circuit due – December 21, 2017**
	15/581,966	Claims to up-titration of DMF to 480 mg/day for the treatment of MS to be filed
	14/212,503	Treating MS with DMF at 480 mg/day to reach certain MMF levels in the bloodstream On appeal from final rejection
	15/728,872	Claims to a pharmaceutical composition comprising DMF in an amount of 50-90% by weight to be filed

A Relevant Patent is a patent that covers treatment for MS by orally administering 480 mg per day of DMF.
If we prevail in the Interference Proceeding after any appeals to the Federal Circuit, we further expect the 11/576,871 application, if ultimately issued, would be entitled to patent term adjustment that would result in an estimated patent expiration in 2029 or later. There is no assurance that patent term adjustment would be obtained to fully compensate for all such time lost.

Ex-US Potential Royalty-Bearing Claims

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If Forward obtains a Relevant Patent in the European EP'355 opposition proceedings including all appeals therefrom, and can show on a country-by-country basis that Tecfidera® infringes a valid licensed patent, royalties may be payable. In Europe, there are presently four patents and patent applications with potentially royalty-bearing claims.

	Application Number	Description
Ex-US	EP14172398.1 (Pat. No. EP2801355)	Treating MS with 480 mg/day of DMF wherein the pH controlled release compositions have an enteric coat **Granted – May 20, 2015; Opposition hearing before EPO scheduled for January 29-30, 2018**
	EP15166243.4 (Pat. No. EP2965751)	Treating MS with 480-600 mg fumaric acid esters (including DMF)/day where the composition releases fumaric acid esters depending on pH. Application pending
	EP14172396.5 (Pat. No. EP2792349)	Controlled release composition of DMF for use in treating hyperproliferative, inflammatory or autoimmune disorders other than psoriasis with 480 mg/day **EPO issued intention to grant – September 13, 2017**
	EP16001391.8 (Pat. No. EP3093012)	Controlled release pharmaceutical composition comprising DMF in an amount of 50-90% by weight **EPO issued intention to grant – May 8, 2017**

A Relevant Patent is a patent that covers treatment for MS by orally administering 480 mg per day of DMF.

Key IP Overview:

Core Composition and Erosion Matrix Patent Families[1]

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Patent / Application	Patent Family	Status
U.S. App. 11/576,871	Core Composition	Pending and involved in an interference proceeding. A decision was issued by the PTAB on March 31, 2017 in favor of Biogen. We have filed an appeal of the decision to the Federal Circuit.
U.S. App. 15/581,966	Core Composition	Pending.
U.S. App. 14/212,503	Core Composition	On appeal from final rejection.
U.S. App. 15/728,872	Core Composition	Pending.
EP2801355	Core Composition	Granted. Under opposition with EPO.
EP1799196	Core Composition	Granted. Under opposition with EPO.
EP2801354	Core Composition	Granted. Under opposition with EPO.
EP2316430	Core Composition	Revoked by decision of July 10, 2015; under appeal. Appeal hearing currently scheduled for May 2018.
EP3093012	Core Composition	Pending. Intention to grant issued.
EP2965751	Core Composition	Pending.
EP2792349	Core Composition	Pending. Intention to grant issued.
U.S. Patent No. 8,906,420	Erosion Matrix	Granted.
U.S. App. 14/561,010	Erosion Matrix	On appeal from final rejection.
EP2379063	Erosion Matrix	Granted; opposition rejected; appeal pending.
EP2564839	Erosion Matrix	Granted. Under opposition with EPO.
JP5788331	Erosion Matrix	Granted as JP2012-514624.

1. Beyond the core composition patent and erosion matrix patent families, other patent families include U.S. Patent Application Nos. 14/419,031, 15/584, 439 and 14/914,025 and European Patent Application Nos. EP2879672, EP3038606 and EP3038605. As a result of the corporate restructuring that was completed pursuant to Appendix D of the Settlement and License Agreement, the intellectual property of Forward Pharma that is the subject of the Settlement and License Agreement was ultimately transferred to FWP IP ApS, a Danish limited liability company, and the capital stock of FWP IP ApS was transferred to a newly formed independent Danish foundation. For more information regarding this restructuring and transfer, see our Form 6-K and press release dated November 22, 2017.

Date of preparation: November 22, 2017