# 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): June 15, 2017

Skyline Medical Inc.

(Exact Name of Registrant as Specified in Charter)

Delaware

(State or Other Jurisdiction of Incorporation)

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001-36790 (Commission File Number) 33-1007393 (I.R.S. Employer Identification Number)

2915 Commers Drive, Suite 900, Eagan, Minnesota 55121 (Address of Principal Executive Offices) (Zip Code)

(651) 389-4800

(Registrant's telephone number, including area code)

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:

] 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 (17 CFR §230.405) or Rule 12b-2 of the Securities Exchange Act of 1934 (17 CFR §240.12b-2). Emerging growth company [ ]

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 8.01. Other Events.

On June 15, 2017, Skyline Medical Inc. issued a press release announcing that the STREAMWAY System meets all requirements and can now be affixed with the CE mark and marketed in 32 European countries. A copy of the press release is attached as Exhibit 99.1 to this report and incorporated herein by reference.

### Item 9.01. Financial Statements and Exhibits.

(d) Exhibits.

# Exhibit No. Description/Exhibit

99.1 Press Release of Skyline Medical Inc. dated June 15, 2017

## 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 hereunto duly authorized.

## Skyline Medical Inc.

Date: June 22, 2017

By: <u>/s/ Bob Myers</u> Bob Myers Chief Financial Officer

#### Skyline Medical Announces CE Mark for the STREAMWAY® System

MINNEAPOLIS, June 15, 2017 (GLOBE NEWSWIRE) -- Skyline Medical Inc. (NASDAQ:SKLN) ("Skyline" or "the Company"), producer of the FDA-approved STREAMWAY® System for automated, direct-to-drain medical fluid disposal, announces that its European Authorized Representative, Emergo Europe B.V., has notified the Dutch Competent Authority that the Company's STREAMWAY System meets all requirements and can now be affixed with the CE mark and marketed in 32 European countries.

"Receipt of the CE mark is an important milestone for Skyline Medical and we are very excited to be able to expand the reach of our STREAMWAY System to Europe," said Dr. Carl Schwartz, chief executive officer of Skyline Medical. "We have already held preliminary discussions with a few European distributors, and with the CE mark in hand we can now advance those negotiations while beginning dialogues with others.

"We are optimistic that we can record our first product sale under the CE mark before year-end, and ramp up STREAMWAY's availability throughout Europe by early 2018. Europe presents a rich market opportunity we are eager to pursue," Dr. Schwartz added.

#### About the STREAMWAY System

Skyline's revolutionary, FDA-cleared STREAMWAY System is the first true direct-to-drain fluid disposal system designed specifically for medical applications, such as radiology, endoscopy, urology and cystoscopy procedures. It connects directly to a facility's plumbing system to automate the collection, measurement and disposal of waste fluids. As of March 31, 2017, Skyline Medical customers have installed 103 STREAMWAY Systems in 52 facilities across 20 states, and in Canada.

The STREAMWAY minimizes human intervention for better safety and improves compliance with Occupational Safety and Health Administration (OSHA) and other regulatory agency safety guidelines. It also provides unlimited capacity for increased efficiency in the operating room, which leads to greater profitability. Furthermore, the STREAMWAY eliminates canisters to reduce overhead costs and provides greater environmental stewardship by helping to eliminate the approximately 50 million potentially disease-infected canisters that go into landfills annually in the U.S. For a demonstration please visit www.skylinemedical.com or call 855-785-8855.

### **About Skyline Medical**

Skyline Medical produces a fully automated, patented, FDA-cleared waste fluid disposal system that virtually eliminates staff exposure to blood, irrigation fluid and other potentially infectious fluids found in the healthcare environment. Antiquated manual fluid handling methods that require hand carrying and emptying filled fluid canisters present an exposure risk and potential liability. Skyline Medical's STREAMWAY System fully automates the collection, measurement and disposal of waste fluids and is designed to: 1) reduce overhead costs to hospitals and surgical centers; 2) improve compliance with OSHA and other regulatory agency safety guidelines; 3) improve efficiency in the operating room, and radiology and endoscopy departments, thereby leading to greater profitability; and 4) provide greater environmental stewardship by helping to eliminate the approximately 50 million potentially disease-infected canisters that go into landfills each year in the U.S. For additional information, please visit www.skylinemedical.com.

#### **Forward-looking Statements**

Certain of the matters discussed in this announcement contain forward-looking statements that involve material risks to and uncertainties in the Company's business that may cause actual results to differ materially from those anticipated by the statements made herein. Such risks and uncertainties include, among other things, current negative operating cash flows and a need for additional funding to finance our operating plan; the terms of any further financing, which may be highly dilutive and may include onerous terms; unexpected costs and operating deficits, and lower than expected sales and revenues; uncertain willingness and ability of customers to adopt new technologies and other factors that may affect further market acceptance, if our product is not accepted by our potential customers, it is unlikely that we will ever become profitable; adverse economic conditions; adverse results of any legal proceedings; the volatility of our operating results and financial condition; inability to attract or retain qualified senior management personnel, including sales and marketing personnel; our ability to establish and maintain the proprietary nature of our technology through the patent process, as well as our ability to possibly license from others patents and patent applications necessary to develop products; the Company's ability to implement its long range business plan for various applications of its technology; the Company's ability to enter into agreements with any necessary marketing and/or distribution partners; the impact of competition, the obtaining and maintenance of any necessary regulatory clearances applicable to applications of the Company's technology; and management of growth and other risks and uncertainties that may be detailed from time to time in the Company's reports filed with the Securities and Exchange Commission, which are available for review at www.sec.gov. This is not a solicitation to buy or sell securities and does not purport to be an analysis of the Company's financial position. Se

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