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**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): **July 6, 2020**

**PACIRA BIOSCIENCES, INC.**

(Exact name of registrant as specified in its charter)

**Delaware**  
(State or other jurisdiction  
of incorporation)

**001-35060**  
(Commission  
File Number)

**51-0619477**  
(IRS Employer  
Identification No.)

**5 Sylvan Way, Suite 300, Parsippany, New Jersey 07054**  
(Address of principal executive offices) (Zip Code)

**(973) 254-3560**  
Registrant's telephone number, including area code

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))

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

<u>Title of each class</u>	<u>Trading symbol</u>	<u>Name of each exchange on which registered</u>
Common Stock, par value \$0.001 per share	PCRX	Nasdaq Global Select Market

Indicate by check mark whether the registrant is an emerging growth company as defined in Rule 405 of the Securities Act of 1933 (CFR §230.405 of this chapter) or Rule 12b-2 of the Securities Exchange Act of 1934 (CFR §240.12b-2 of this chapter).

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.

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**Item 2.02. Results of Operations and Financial Condition.**

On July 6, 2020, Pacira BioSciences, Inc. issued a press release announcing its preliminary unaudited revenue for the second quarter ended June 30, 2020. The full text of the press release issued in connection with the announcement is furnished as Exhibit 99.1 to this Current Report on Form 8-K.

The information in this Item 2.02 of Form 8-K and Exhibit 99.1 attached hereto shall not be deemed “filed” for purposes of Section 18 of the Securities Exchange Act of 1934, as amended (the “Exchange Act”) or otherwise subject to the liabilities of that section, nor shall it be deemed incorporated by reference in any filing under the Securities Act of 1933, as amended, or the Exchange Act, except as expressly set forth by specific reference in such a filing.

**Item 9.01. Financial Statements and Exhibits.**

**(d) Exhibits.**

<b>Exhibit No.</b>	<b>Description</b>
<a href="#">99.1</a>	<a href="#">Press Release dated July 6, 2020</a>
104	Cover Page Interactive Data File (Formatted as Inline XBRL)

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**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.

PACIRA BIOSCIENCES, INC.

Date: July 6, 2020

By: /s/ Kristen Williams

Kristen Williams

*Chief Administrative Officer and Secretary*

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FOR IMMEDIATE RELEASE

NEWS RELEASE

**Pacira BioSciences Reports Preliminary Total Revenue of \$75.5 Million for Second Quarter of 2020**

*-- EXPAREL average daily sales return to year-over-year growth for month of June 2020 --*

**Parsippany, NJ, July 6, 2020** - Pacira BioSciences, Inc. (Nasdaq: PCRX), a leading provider of innovative non-opioid pain management options, today reported preliminary unaudited net revenue of \$75.5 million for the second quarter of 2020, compared to \$102.6 million for the second quarter of 2019. During the second quarter of 2020, average daily sales of EXPAREL® (bupivacaine liposome injectable suspension) were 30 percent, 81 percent and 107 percent of the prior year for the months of April, May, and June, respectively. During the second quarter of 2020, the Company's product sales were negatively impacted by the COVID-19 pandemic, which mandated significant postponement or suspension in the scheduling of elective surgical procedures resulting from public health guidance and government directives. Elective surgery restrictions began to lift on a state-by-state basis in April 2020.

"We are very encouraged by the rapid recovery and consistent uptake in EXPAREL sales and ordering accounts on a weekly basis since the peak of the COVID-19 pandemic impact in April," said Dave Stack, chairman and chief executive officer of Pacira BioSciences. "Ambulatory surgical centers have quickly returned to the operating room to accommodate patients who were required to wait for important surgical interventions during COVID-19-related shutdowns. These customers have been key to our return to year-over-year growth and underscore our investment in working closely with our anesthesia partners to broaden the use of long-acting EXPAREL regional approaches as a cornerstone for multimodal opioid-sparing strategies that continue to enable the shift of inpatient procedures to the 23-hour stay environment."

**Second Quarter Financial Highlights**

- Net product sales of EXPAREL/bupivacaine liposome injectable suspension for the second quarter of 2020 were \$73.8 million, compared to \$99.8 million in the same period in 2019.
  - EXPAREL net product sales for the second quarter of 2020 were \$73.0 million, compared to \$98.9 million in the same period in 2019. Sales of bupivacaine liposome injectable suspension to a third-party licensee for use in veterinary practice for the second quarter of 2020 were \$0.8 million, compared to \$0.9 million in the same period in 2019.
  - iovera<sup>®</sup> net product sales for the second quarter of 2020 were \$1.4 million, compared to \$2.0 million in the same period in 2019. Pacira began recognizing sales of iovera<sup>®</sup> in April 2019 after completing its acquisition of MyoScience, Inc., a privately held medical technology company.
  - Royalty revenue for the second quarter of 2020 was \$0.3 million, compared to \$0.8 million for the same period in 2019.
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The financial information included in this press release is preliminary, unaudited and subject to adjustment. It does not present all information necessary for an understanding of the company's second quarter financial results for 2020. Pacira expects to report its complete financial results for the second quarter and first half of 2020 during the Company's conference call scheduled for August 2020.

### **About Pacira BioSciences**

Pacira BioSciences, Inc. (Nasdaq: PCRX) is a leading provider of non-opioid pain management and regenerative health solutions dedicated to advancing and improving outcomes for health care practitioners and their patients. The company's long-acting local analgesic, EXPAREL<sup>®</sup> (bupivacaine liposome injectable suspension) was commercially launched in the United States in April 2012. EXPAREL utilizes DepoFoam<sup>®</sup>, a unique and proprietary product delivery technology that encapsulates drugs without altering their molecular structure, and releases them over a desired period of time. In April 2019, Pacira acquired the iovera<sup>®</sup> system, a handheld cryoanalgesia device used to deliver precise, controlled doses of cold temperature only to targeted nerves. To learn more about Pacira, including the corporate mission to reduce overreliance on opioids, visit [www.pacira.com](http://www.pacira.com).

### **About EXPAREL<sup>®</sup>**

EXPAREL (bupivacaine liposome injectable suspension) is indicated for single-dose infiltration in adults to produce postsurgical local analgesia and as an interscalene brachial plexus nerve block to produce postsurgical regional analgesia. Safety and efficacy have not been established in other nerve blocks. The product combines bupivacaine with DepoFoam, a proven product delivery technology that delivers medication over a desired time period. EXPAREL represents the first and only multivesicular liposome local anesthetic that can be utilized in the peri- or postsurgical setting. By utilizing the DepoFoam platform, a single dose of EXPAREL delivers bupivacaine over time, providing significant reductions in cumulative pain scores with up to a 78 percent decrease in opioid consumption; the clinical benefit of the opioid reduction was not demonstrated. Additional information is available at [www.EXPAREL.com](http://www.EXPAREL.com).

### **Important Safety Information for Patients**

EXPAREL should not be used in obstetrical paracervical block anesthesia. In studies where EXPAREL was injected into the wound, the most common side effects were nausea, constipation, and vomiting. In studies where EXPAREL was injected near a nerve, the most common side effects were nausea, fever, and constipation. EXPAREL is not recommended to be used in patients younger than 18 years old or in pregnant women. Tell your healthcare provider if you have liver disease, since this may affect how the active ingredient (bupivacaine) in EXPAREL is eliminated from your body. EXPAREL should not be injected into the spine, joints, or veins. The active ingredient in EXPAREL: can affect your nervous system and your cardiovascular system; may cause an allergic reaction; may cause damage if injected into your joints.

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## About iovera<sup>o</sup><sup>®</sup>

The iovera<sup>o</sup> system is used to destroy tissue during surgical procedures by applying freezing cold. It can also be used to produce lesions in peripheral nervous tissue by the application of cold to the selected site for the blocking of pain. It is also indicated for the relief of pain and symptoms associated with osteoarthritis of the knee for up to 90 days. In one study, the majority of the patients suffering from osteoarthritis of the knee experienced pain and system relief beyond 150 days.<sup>1</sup> The iovera<sup>o</sup> system's "1×90" Smart Tip configuration (indicating one needle which is 90 mm long) can also facilitate target nerve location by conducting electrical nerve stimulation from a separate nerve stimulator. The iovera<sup>o</sup> system is not indicated for treatment of central nervous system tissue.

## Important Safety Information

The iovera<sup>o</sup> system is contraindicated for use in patients with the following: Cryoglobulinemia; Paroxysmal cold hemoglobinuria; cold urticaria; Raynaud's disease; open and/or infected wounds at or near the treatment line. Potential complications: As with any surgical treatment that uses needle-based therapy, there is potential for temporary site-specific reactions, including but not limited to: bruising (ecchymosis); swelling (edema); inflammation and/or redness (erythema); pain and/or tenderness; altered sensation (localized dysesthesia). Typically, these reactions resolve with no physician intervention. Patients may help the healing process by applying ice packs to the affected sites, and by taking over-the-counter analgesics.

## Forward-Looking Statements

*Any statements in this press release about the company's future expectations, plans, outlook, projections and prospects, and other statements containing the words "believes," "anticipates," "plans," "estimates," "expects," "intends," "may," "will," "would," "could," "can" and similar expressions, constitute forward-looking statements within the meaning of The Private Securities Litigation Reform Act of 1995. Actual results may differ materially from those indicated by such forward-looking statements as a result of various important factors, including risks relating to: the impact of the worldwide COVID-19 (Coronavirus) pandemic and related global economic conditions; the success of the company's sales and manufacturing efforts in support of the commercialization of EXPAREL; the rate and degree of market acceptance of EXPAREL; the size and growth of the potential markets for EXPAREL and the company's ability to serve those markets; the company's plans to expand the use of EXPAREL to additional indications and opportunities, and the timing and success of any related clinical trials; the ability to realize anticipated benefits and synergies from the acquisition of MyoScience; the ability to successfully integrate iovera<sup>o</sup> and any other future acquisitions into the company's existing business; the commercial success of iovera<sup>o</sup> and other factors discussed in the "Risk Factors" of the company's most recent Annual Report on Form 10-K and in other filings that the company periodically makes with the SEC. In addition, the forward-looking statements included in this press release represent the company's views as of the date of this press release. Important factors could cause actual results to differ materially from those indicated or implied by forward-looking statements, and as such the company anticipates that subsequent events and developments will cause its views to change. However, while the company may elect to update these forward-looking statements at some point in the future, it specifically disclaims any obligation to do so. These forward-looking statements should not be relied upon as representing the company's views as of any date subsequent to the date of this press release.*

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<sup>1</sup>Radnovich, R. et al. "Cryoneurolysis to treat the pain and symptoms of knee osteoarthritis: a multicenter, randomized, double-blind, sham-controlled trial." *Osteoarthritis and Cartilage* (2017) p1-10.

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**Investor Contact:**

Susan Mesco, (973) 451-4030  
susan.mesco@pacira.com

**Media Contact:**

Coyne Public Relations  
Alyssa Schneider, (973) 588-2270  
aschneider@coynepr.com

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