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) April 26, 2021

BECTON, DICKINSON AND COMPANY

(Exact Name of Registrant as Specified in Its Charter)

			N I			
-			New Jersey (State or Other Jurisdiction of	f Incorporation)		
	001-4802			22-0760120		
	(Commission File Number)			(IRS Employer Identification No.)		
	1 Becton Drive,	Franklin Lakes,	New Jersey	07417-1880		
<u>-</u>	(Address of Principal Executive Offices)			(Zip Code)	_	
	(201) 847-6800					
-	(Registrant's Telephone Number, Including Area Code)					
			N/A			
(Former Name or Former Address, if Changed Since Last Report)						
Check the appropr	riate box below if the Form 8-K Filing	is intended to simultaneously satisf	ry the filing obligation of the registrant under	r any of the following provisions (see General Instruction A.2. below):		
□ V	V-i4	.l. 425 d db . Cisi A .s (15	(CED 220425)			
	Written communications pursuant to Rule 425 under the Securities Act (17 CFR 230425)					
	Soliciting material pursuant to Rule 14a-12 under the Exchange Act (17 CFR 240.14a-12)					
□ P	Pre-commencement communications pursuant to Rule 14d-2(b) under the Exchange Act (17 CFR 240.14d-2(b))					
□ P	re-commencement communications pu	ursuant to Rule 13e-4(c) under the l	Exchange Act (17 CFR 240.13e-4(c))			
Securities registere	ed pursuant to Section 12(b) of the Act	t:				

Title of Each Class	Trading Symbol	which registered
Common stock, par value \$1.00	BDX	New York Stock Exchange
Depositary Shares, each representing a 1/20th interest in a share of 6.00% Mandatory Convertible Preferred Stock, Series B	BDXB	New York Stock Exchange
1.000% Notes due December 15, 2022	BDX22A	New York Stock Exchange
1.900% Notes due December 15, 2026	BDX26	New York Stock Exchange
1.401% Notes due May 24, 2023	BDX23A	New York Stock Exchange
3.020% Notes due May 24, 2025	BDX25	New York Stock Exchange
0.632% Notes due June 4, 2023	BDX/23A	New York Stock Exchange
1.208% Notes due June 4, 2026	BDX/26A	New York Stock Exchange
1.213% Notes due February 12, 2036	BDX/36	New York Stock Exchange

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

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. \Box

ITEM 7.01 Regulation FD Disclosure

On April 26, 2021, Becton, Dickinson and Company ("BD") issued a press release announcing it submitted a 510(k) premarket notification to the U.S. Food and Drug Administration for the BD AlarisTM System.

The 510(k) submission is intended to bring the regulatory clearance for the BD AlarisTM System up to date, implement updated features and address open recall issues, including through a new version of BD AlarisTM System software that will provide clinical, operational and cybersecurity updates.

The press release is attached as Exhibit 99.1. The information in this Item 7.01 and in Exhibit 99.1 to this Current Report on Form 8-K is being furnished and 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 thereof, nor shall it be incorporated by reference into future filings by BD under the Exchange Act or under the Securities Act of 1933, as amended, except to the extent specifically provided in any such filing.

ITEM 9.01 Financial Statements and Exhibits.

Exhibit 99.1 Press release dated April 26, 2021, which is furnished pursuant to Item 7.01.

Exhibit 104 Cover Page Interactive Data File (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.

BECTON, DICKINSON AND COMPANY (Registrant)

By:	/s/ Gary DeFazio
_	Gary DeFazio
	Senior Vice President and Corporate Secretary

Date: April 26, 2021



NEWS RELEASE

FOR IMMEDIATE RELEASE

BD Announces FDA 510(k) Submission for BD Alaris™ System

FRANKLIN LAKES, N.J. (April 26, 2021) – BD (Becton, Dickinson and Company) (NYSE: BDX), a leading global medical technology company, today announced it has submitted a 510(k) premarket notification to the U.S. Food and Drug Administration (FDA) for the BD Alaris™ System, the most widely used infusion pump in acute care hospitals across the United States.

The 510(k) submission is intended to bring the regulatory clearance for the BD AlarisTM System up to date, implement updated features and address open recall issues, including through a new version of BD AlarisTM System software that will provide clinical, operational and cybersecurity updates.

"Today marks an important milestone in our commitment to our customers and patients and Advancing the world of health™," said Michael Garrison, worldwide president of Medication Management Solutions for BD. "The 510(k) submission is the first step in the review process with the FDA, and we look forward to working through the FDA review process to obtain clearance for the updated BD Alaris™ System."

The BD Alaris™ System allows clinicians to deliver medications, fluids and blood products through a single integrated platform that includes large volume pumps, syringe pumps and patient-controlled analgesia (PCA) modules for adult, pediatric and neonatal patients.

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About BD

Forward Looking Statements

This press release contains certain forward-looking statements regarding BD's 510(k) premarket notification to the FDA for the BD Alaris™ System. Forward-looking statements involve risks and uncertainties that could cause actual results to differ materially from those expressed or implied in such statements, and there can be no assurance that BD will obtain 510(k) clearance from the FDA for the BD Alaris™ System or as to the timing of any such clearance. Many of these risks and uncertainties are beyond the company's control, including without limitation, risks relating to regulatory clearance and market acceptance of the BD Alaris™ System, the remediation of our infusion pump business and other factors listed in our 2020 Annual Report on Form 10-K and other filings with the SEC. BD expressly disclaims any undertaking to update any forward-looking statements set forth herein to reflect events or circumstances after the date hereof, except as required by applicable laws or regulations.

Contacts:

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