

LABORATORY CORP OF AMERICA HOLDINGS  
Form 8-K  
November 15, 2017

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

November 15, 2017  
(Date of earliest event reported)

LABORATORY CORPORATION OF  
AMERICA HOLDINGS

(Exact Name of Registrant as Specified in its Charter)

Delaware 1-11353 13-3757370  
(State or other jurisdiction of Incorporation) (Commission File Number) (I.R.S. Employer Identification No.)

358 South Main Street,  
Burlington, North Carolina 27215 336-229-1127  
(Address of principal executive offices) (Zip Code) (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 communication 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))

Item 7.01 Regulation FD Disclosure

LabCorp (NYSE: LH) a leading global life sciences company, today announced the U.S. availability of the PD-L1 IHC 28-8 pharmDx assay as a complementary diagnostic for two newly approved indications in connection with the use of Bristol-Myers Squibb's OPDIVO® (nivolumab) to treat patients with metastatic urothelial carcinoma, also referred to as bladder cancer, and squamous cell carcinoma of the head and neck. The PD-L1 IHC 28-8 pharmDx assay was developed by Agilent's Dako pathology division. While OPDIVO is approved for these indications without use of the test, the test provides physicians with important information about those patients who are most likely to respond positively to OPDIVO. LabCorp's Center for Molecular Biology and Pathology laboratory performed testing for the clinical studies that supported approval of the new indications for the assay.

Exhibit Index [Exhibit 99.1](#)



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.

LABORATORY CORPORATION OF AMERICA HOLDINGS  
Registrant

By: /s/ F. SAMUEL EBERTS III  
F. Samuel Eberts III  
Chief Legal Officer and Secretary

November 15, 2017