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Interpace Biosciences Announces New Draft LCD and Reimbursement for Its Proprietary Thyroid Assay, ThyGeNEXT®

Parsippany, NJ, Dec. 17, 2019 (GLOBE NEWSWIRE) -- Interpace Biosciences, Inc. (Nasdaq: IDXG) today announced that its Medicare Administrative Contractor (MAC) has issued a new draft local coverage determination (LCD) for the Company's ThyGeNEXT® test, representing an increase of approximately \$2,400 per assay over previous reimbursement coverage. This increase in reimbursement rates reflects the expansion of the ThyGeNEXT® panel to aid in identifying the appropriate patients for surgery. In 2018, Interpace processed approximately 12,500 ThyGeNEXT® tests.

Prior to the new LCD code (81455) becoming effective, it was subject to a public comment period, which ended December 15, 2019, and is now subject to an analysis and review period by MAC's Medical Directors. Final approval is expected during the first quarter of 2020. ThyGeNEXT® has been covered by an existing LCD since it was launched in mid-2018 and its predecessor, ThyGenX®, has been covered since 2014.

Jack Stover, President & CEO of Interpace, said, "I am very pleased with the draft local coverage announcement and look forward to the final determination, which when approved will demonstrate the quality of our expanded assay, and most importantly supports continued reimbursement for patients and their families potentially affected by Thyroid cancer."

About ThyGeNEXT® and ThyraMIR®

ThyGeNEXT® utilizes state-of-the-art next-generation sequencing (NGS) to identify more than 100 genetic alterations associated with papillary and follicular thyroid carcinomas, the two most common forms of thyroid cancer, as well as Medullary Thyroid Carcinoma. ThyraMIR® is the first microRNA gene expression classifier. MicroRNAs are small, non-coding RNAs that bind to messenger RNA and regulate expression of genes involved in human cancers, including every subtype of thyroid cancer. ThyraMIR® measures the expression of 10 microRNAs. Both ThyGeNEXT® and ThyraMIR® are covered by Medicare and Commercial insurers, with more than 280 million members covered.

According to the American Thyroid Association, approximately 20% of the 525,000 thyroid fine needle aspirations (FNAs) performed on an annual basis in the U.S. are indeterminate for malignancy based on standard cytological evaluation, and thus are candidates for ThyGeNEXT® and ThyraMIR®.

ThyGeNEXT® and ThyraMIR® reflex testing yields high predictive value in determining the

presence and absence of cancer in thyroid nodules. The combination of both tests can improve risk stratification and surgical decision-making when standard cytopathology does not provide a clear diagnosis.

About Interpace Biosciences

Interpace Biosciences is a leader in enabling personalized medicine, offering specialized services along the therapeutic value chain from early diagnosis and prognostic planning to targeted therapeutic applications.

Interpace Diagnostics is a fully integrated commercial and bioinformatics business unit that provides clinically useful molecular diagnostic tests, bioinformatics and pathology services for evaluating risk of cancer by leveraging the latest technology in personalized medicine for improved patient diagnosis and management. Interpace has four commercialized molecular tests and one test in a clinical evaluation process (CEP): PancraGEN[®] for the diagnosis and prognosis of pancreatic cancer from pancreatic cysts; ThyGeNEXT[®] for the diagnosis of thyroid cancer from thyroid nodules utilizing a next generation sequencing assay; ThyraMIR[®] for the diagnosis of thyroid cancer from thyroid nodules utilizing a proprietary gene expression assay; and RespriDX[®] that differentiates lung cancer of primary vs. metastatic origin. In addition, BarreGEN[®] for Barrett's Esophagus, is currently in a clinical evaluation program whereby we gather information from physicians using BarreGEN[®] to assist us in positioning the product for full launch, partnering and potentially supporting reimbursement with payers.

Interpace Biopharma provides pharmacogenomics testing, genotyping, biorepository and other customized services to the pharmaceutical and biotech industries. The Biopharma business also advances personalized medicine by partnering with pharmaceutical, academic, and technology leaders to effectively integrate pharmacogenomics into their drug development and clinical trial programs with the goals of delivering safer, more effective drugs to market more quickly, and improving patient care.

For more information, please visit Interpace Biosciences' website at www.interpace.com.

Forward-looking Statements

This press release contains forward-looking statements within the meaning of Section 27A of the Securities Act of 1933, Section 21E of the Securities Exchange Act of 1934 and the Private Securities Litigation Reform Act of 1995, relating to the Company's future financial and operating performance. The Company has attempted to identify forward looking statements by terminology including "believes," "estimates," "anticipates," "expects," "plans," "projects," "intends," "potential," "may," "could," "might," "will," "should," "approximately" or other words that convey uncertainty of future events or outcomes to identify these forward-looking statements. These statements are based on current expectations, assumptions and uncertainties involving judgments about, among other things, future economic, competitive and market conditions and future business decisions, all of which are difficult or impossible to predict accurately and many of which are beyond the Company's control. These statements also involve known and unknown risks, uncertainties and other factors that may cause the Company's actual results to be materially different from those expressed or implied by any forward-looking statement. Additionally, all forward-looking statements are

subject to the “Risk Factors” detailed from time to time in the Company's most recent Annual Report on Form 10-K and Quarterly Reports on Form 10Q. Because of these and other risks, uncertainties and assumptions, undue reliance should not be placed on these forward-looking statements. In addition, these statements speak only as of the date of this press release and, except as may be required by law, the Company undertakes no obligation to revise or update publicly any forward-looking statements for any reason.

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