

Biocept Provides Update on COVID-19 Testing with More than 350,000 Samples Received to Date

April 14, 2021

SAN DIEGO--(BUSINESS WIRE)--Apr. 14, 2021-- <u>Biocept. Inc.</u> (Nasdaq: BIOC), a leading provider of molecular diagnostic assays, products and services, announces that since the Company began offering SARS-CoV-2 testing services in June 2020, it has now received more than 350,000 samples for testing. These samples are processed using Biocept's RT-PCR technology at its CLIA-certified, CAP-accredited high-complexity molecular laboratory in San Diego.

"We are exceptionally proud to continue supporting our community with SARS-CoV-2 testing and providing healthcare providers with rapid turnaround of valuable information to help contain viral spread," said Michael Nall, President and CEO of Biocept. "RT-PCR testing for COVID-19 currently remains an important component of our business while we focus on our core business, including the opportunity to serve patients diagnosed with advanced cancer through commercializing our differentiated CNSideTM neuro-oncology offerings."

About Biocept

Biocept, Inc. develops and commercializes molecular diagnostic assays that provide physicians with clinically actionable information for treating and monitoring patients diagnosed with a variety of cancers. In addition to its broad portfolio of blood-based liquid biopsy assays, Biocept has developed the CNSide™ cerebrospinal fluid assay that detects cancer that has metastasized to the central nervous system. Biocept's patented Target Selector™ technology captures and quantitatively analyzes CSF tumor cells for tumor-associated molecular markers, using technology first developed for use in blood. Biocept also is leveraging its molecular diagnostic capabilities to offer nationwide COVID-19 RT-PCR testing to support public health efforts during this unprecedented pandemic. For more information, visit www.biocept.com. Follow Biocept on Facebook, LinkedIn and Twitter.

Forward-Looking Statements Disclaimer Statement

This release contains forward-looking statements that are based upon current expectations or beliefs, as well as a number of assumptions about future events. Although we believe that the expectations reflected in the forward-looking statements and the assumptions upon which they are based are reasonable, we can give no assurance that such expectations and assumptions will prove to have been correct. Forward-looking statements are generally identifiable by the use of words like "may," "will," "should," "could," "expect," "anticipate," "estimate," "believe," "intend," or "project" or the negative of these words or other variations on these words or comparable terminology. To the extent that statements in this release are not strictly historical, including without limitation statements regarding our ability to provide physicians with clinically actionable information for treating and monitoring patients diagnosed with a variety of cancers, such statements are forward-looking, and are made pursuant to the safe harbor provisions of the Private Securities Litigation Reform Act of 1995. The reader is cautioned not to put undue reliance on these forward-looking statements, as these statements are subject to numerous risk factors as set forth in our Securities and Exchange Commission (SEC) filings. The effects of such risks and uncertainties could cause actual results to differ materially from the forward-looking statements contained in this release. We do not plan to update any such forward-looking statements and expressly disclaim any duty to update the information contained in this press release except as required by law. Readers are advised to review our filings with the SEC, which can be accessed over the Internet at the SEC's website located at www.sec.gov.

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