

## Thermo Fisher expands GlobalAccess sequencing programme

04 August 2020 | News

Company extends programme benefits to oncology laboratories and fast-tracks access to comprehensive molecular profiling on the Genexus System



Thermo Fisher Scientific has announced the expansion of its GlobalAccess Sequencing programme to include laboratories working in oncology.

Originally introduced to accelerate multi-institutional-led studies focused on SARS-CoV-2, the expanded programme now provides support to labs facing significant constraints as a result of the global pandemic by offering faster access to comprehensive, single-day molecular profiling of tumor tissue on Thermo Fisher's Ion Torrent Genexus System.

"We know that the COVID-19 pandemic has interrupted the way laboratories function on a daily basis. For some, it has deeply impacted their ability to serve customers in circumstances where time is of the essence," saidGarret Hampton, president of clinical next-generation sequencing and oncology at Thermo Fisher Scientific. "At this unprecedented time, sustaining the momentum in oncology is paramount. Extending the benefits of GlobalAccess to this community is a natural extension of our mission to enable our customers to help make the world healthier."

Under the GlobalAccess Sequencing programme, Thermo Fisher will subsidize a limited number of Genexus Systems\* for a short time in alignment with the company's commitment to serve its customers and their important work during the global pandemic. The programme, which is open to pathology laboratories that currently run or wish to begin sequencing samples with oncology assays, is available immediately and will continue through the end of this year.

The Genexus System was launched in November 2019 with the Oncomine Precision Assay\*, which most recently received Breakthrough Device Designation by the U.S. Food and Drug Administration (FDA) to identify low-grade glioma (LGG) patients with isocitrate dehydrogenase 1 and 2 (IDH1 and IDH2) mutations who may be eligible for vorasidenib (AG-881). It is a turnkey next-generation sequencing (NGS) solution featuring an automated specimen-to-report workflow that delivers results in a single day.

With only five minutes of hands-on time required, the system is designed to minimize user intervention and is conducive for maintaining social distancing requirements in a decentralized lab setting. It also requires minimal amounts of tissue sample and can run small batches cost-effectively. All together, these features position the Genexus System to enable a future in

which local hospitals can adopt NGS testing in-house and se NGS information in parallel with first-line testing modalities, suc	ets the stage for molecular pathologists to eventually analyze h as immunohistochemistry (IHC).