

# Build a bioanalytical strategy that will ENSURE SPEED AND AGILITY



The complexities of building a robust bioanalytical strategy for drug discovery and development can often be overwhelming and arduous. To build, execute, and optimize your strategy, it's important to work with a bioanalytical partner that has not only proven operational excellence across a range of traditional and cutting-edge technologies, but also the experience to support the unique needs of your development program.

**5M+**  
SAMPLES  
ANALYZED



**150**  
BIOLOGICAL  
FLUIDS & TISSUES



**48-HOUR**  
TURNAROUNDS  
IN DISCOVERY



## WORK WITH OUR TEAM OF DEDICATED SCIENTISTS, PROGRAM MANAGERS, QUALITY SPECIALISTS, AND INDUSTRY EXPERTS TO:

- Gain access to a robust suite of mass spectrometry platforms for small and large molecule programs
- Achieve quality data for filing your IND, NDA, and CTA with speed
- Forecast technical issues or potential roadblocks that may cause delays
- Elucidate drug efficacy in pertinent spatial context down to the cellular level, for more efficient and effective drug candidate evaluation and selection
- Maximize the growth of your drug discovery and development investments

OFFERING	PLATFORMS	COMPOUNDS	REGULATIONS	
<ul style="list-style-type: none"> <li>■ Discovery</li> <li>■ Method development</li> <li>■ Method validation</li> <li>■ Sample analysis</li> </ul>	<ul style="list-style-type: none"> <li>■ LC-MS/MS</li> <li>■ LC-HRMS</li> <li>■ LC-Fluorescence detection</li> </ul>	<ul style="list-style-type: none"> <li>■ Small molecules</li> <li>■ Biomarkers</li> <li>■ OGNs therapeutics</li> <li>■ Biologics</li> <li>■ Endogenous</li> </ul>	<ul style="list-style-type: none"> <li>■ GLP</li> <li>■ FDA</li> <li>■ CDER</li> <li>■ CBER</li> </ul>	<ul style="list-style-type: none"> <li>■ VICH</li> <li>■ CVM</li> <li>■ EMA</li> <li>■ OECD</li> </ul>

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data for **life** >

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