

## Set high-quality mass spectrometry to warp speed: accelerating drug discovery



On average, it takes 10-15 years and \$1-2 billion to approve a new pharmaceutical for clinical use. Since approximately 90% of new drug candidates fail in clinical development, the ability to make early, informed and accurate decisions on the safety and efficacy of new leads is key to maximizing success.<sup>2</sup> This infographic explores the benefits of Acoustic Ejection Mass Spectrometry (AEMS)-

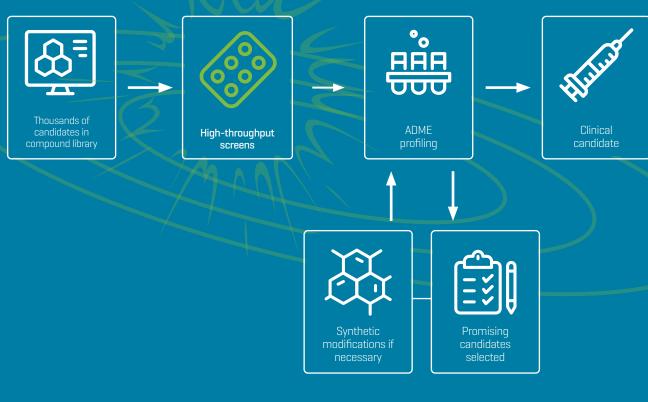
based workflows for efficient and rapid lead optimization of drug candidates.

# Drug discovery demands quality data and high throughput at the lowest possible cost. By driving

Why is high-throughput screening needed?

clinical candidates.

key decisions earlier in the process, these attributes increase the likelihood of developing successful

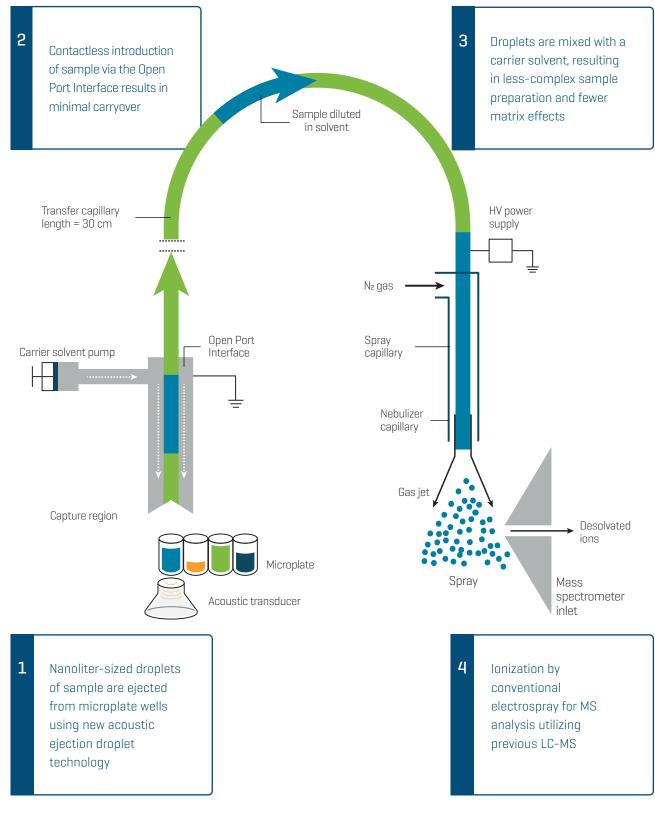


Key decisions about drug candidates depend on the quality of the data analysis. Liquid

Addressing the challenges of traditional methods

assays. However, more high-throughput assays that are just as selective and sensitive are now needed to meet the demands of drug discovery. The Echo® MS system produces results with the precision and accuracy of traditional LC-MS techniques in a far shorter time frame.

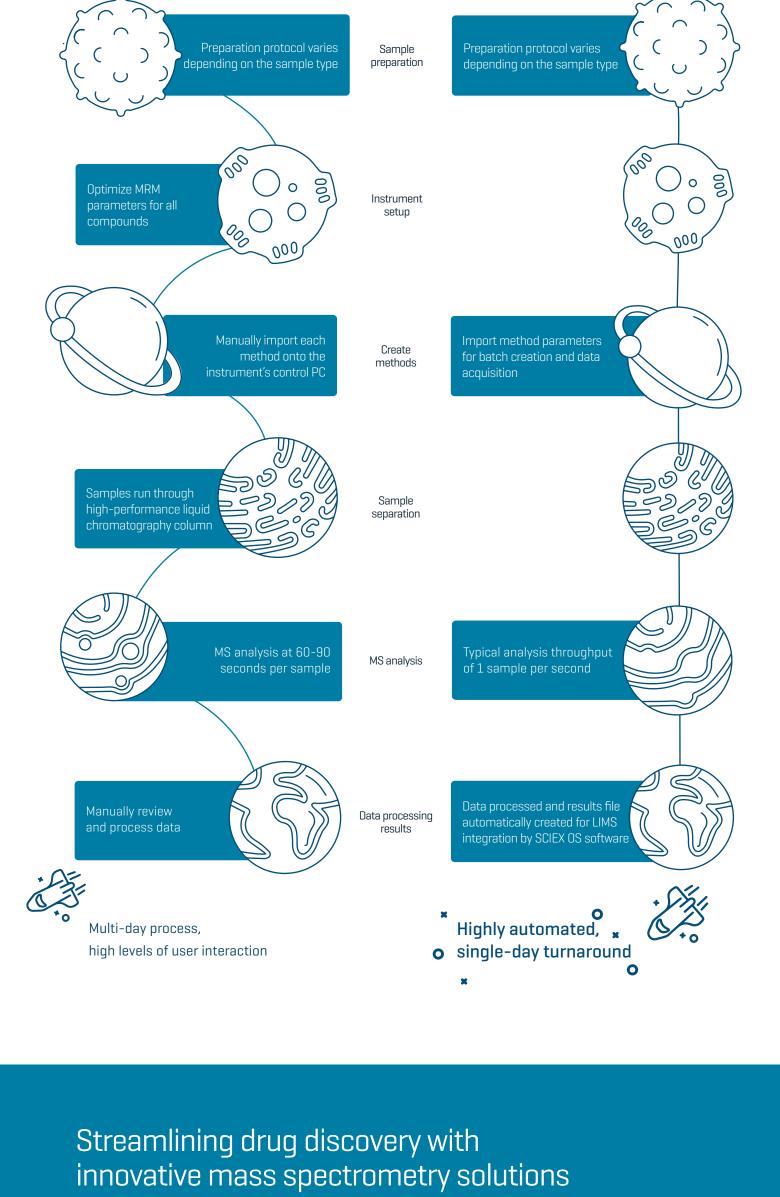
chromatography-mass spectrometry (LC-MS) has been the gold standard for lead drug identification



### Traditional LC-MS Echo® MS system

Shoot for the moon: optimizing high-throughput

mass spectrometry screening



### Acoustic ejection Nanoliter sample reduces the risk of consumption allows repeated chromatography



analysis of the same sample

issues and sample

carryover



system to accelerate drug discovery.

Easy integration into existing workflows

- References Sun D, Gao W, Hu H, Zhou S. Why 90% of clinical drug development fails and how to improve it? Acta Pharmaceutica Sinica B. 2022;12[7]:3049-3062. doi:10.1016/j.apsb.2022.02.002 Lowe D. The latest on drug failure and approval rates. Science. https://www.science.org/content/blog-post/latest-drug-
- failure-and-approval-rates. Published May 9 2019. Accessed October 28, 2022. The SCIEX clinical diagnostic portfolio is For In Vitro Diagnostic Use. Rx Only. Product(s) not available in all countries. For information on availability, please contact your local sales representative or refer to www.sciex.com/diagnostics. All other products are For Research Use Only. Not for use in Diagnostic Procedures. Trademarks and/or registered trademarks mentioned herein, including associated logos, are the property of AB Sciex Pte. Ltd. or their respective owners in the United States and/or certain other countries (see www.sciex.com/trademarks). Echo® and Echo® MS are

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