OUR BIOANALYTICAL SCIENTISTS WORK SEAMLESSLY WITH YOU, FROM DEVELOPING ASSAYS TO REPORTING DATA.

For more than 20 years, our scientists have been partnering with clients to develop challenging assays. ABS Laboratories has vast experience in conducting small-molecule and metabolite pharmacokinetic studies—both preclinical and clinical—for submission to regulatory bodies. At ABS, clients have a direct line of communication with the lead scientist involved, who provides a project overview and regular progress updates. Our analysts will follow a sample from receipt through sample preparation and analysis to results—and can assist on data interpretation—assuming full responsibility throughout the process.

We partner with you, offering a depth of integration and support.

We develop and validate assays to meet your goals and objectives. The partnership we form with you, coupled with our expertise, accelerates the process. We offer a range of validated methods for pharmaceuticals that are already on the market. We regularly validate assays for multiple analytes to meet regulatory requirements. At ABS, we've also developed stabilization procedures for hard-to-assay drugs.

Our state-of-the-art systems deliver accuracy and reliability.

We understand the need for high-quality data and the importance of finding new ways to look at clinical research. For that reason, we use only LC-MS/MS platforms to quantify and investigate the absorption and excretion of drugs and their metabolites during the drug development process. Our smarter testing methods range from simple sample dilution to complicated solid-phase extraction and derivatization methods. ABS Laboratories also owns a large library of validated bioanalytical methods for a multitude of drugs, for rapid initialization and analysis of study samples.

We understand the global regulatory landscape.

Our methods meet EMA and FDA guidelines, and our studies are conducted to GLP and GCP standards. We have a broad understanding of global regulatory processes in all areas of drug development from conducting bioanalysis studies that are ready for regulatory submission for a new drug to supporting the reformulation of existing drugs or medical devices. We are your experienced partner in navigating what can be a complicated landscape.

ABS
AN ACM GLOBAL LABORATORY

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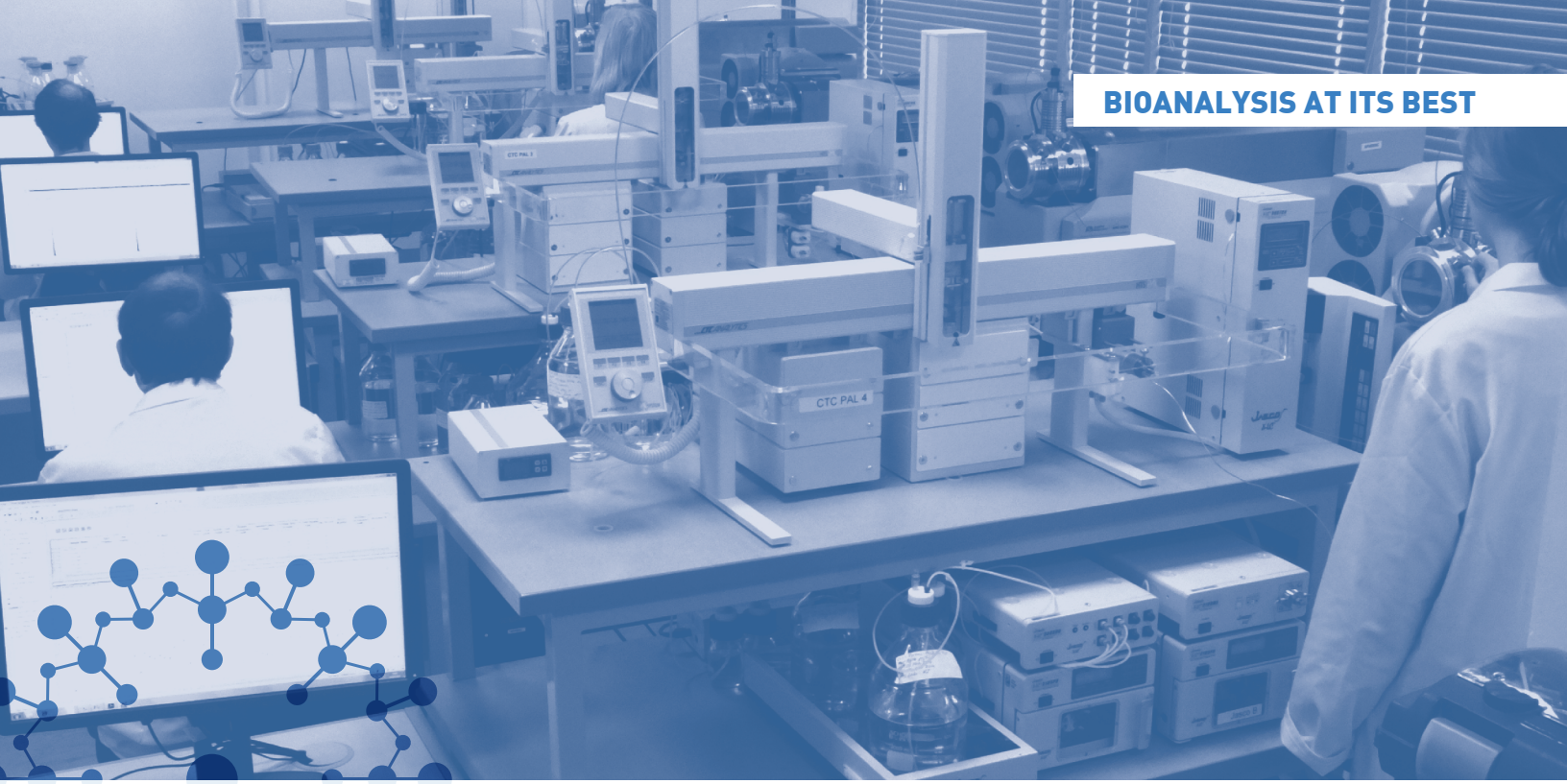

BIOANALYSIS AT ITS BEST

Our solutions and approach focus on your needs.

Our approach to development is based on your molecule and tailored to meet your specific individual requirements and goals, ensuring full satisfaction. We are proactive and take pride in the honesty and transparency we bring to the table. From initial method evaluation to study development improvements, we stay on track to meet your goals and keep you informed and involved every step of the way.

As part of ACM Global Laboratories, we offer unparalleled reach and support.

ABS is part of ACM Global Laboratories, offering an integrated, one-stop approach for all your clinical requirements. ABS Laboratories has been providing expertise and support to pharmaceutical companies, universities, healthcare providers and charitable organizations for more than four decades. ACM Global Laboratories provides general clinical laboratory services, performing more than 30 million tests annually, with operations that extend across more than 65 countries through facilities in the United States, Europe, Singapore, China and India. ACM Global can also supply sample collection kits and logistics support as required.

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