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Frontage Labs strengthens early phase clinical capabilities across US and China

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To support pharmaceutical and biotech partners with accelerated, high?quality clinical development solutions

Frontage Laboratories, Inc., a global Contract Research, Development and Manufacturing Organization (CRDMO), has announced the expansion of its early phase clinical research capabilities across the United States (US) and China. This strategic growth enhances the company's ability to support pharmaceutical and biotech partners with accelerated, high?quality clinical development solutions.

Frontage's US operations are anchored by a modern 160?bed, 36,000?sq?ft Phase I clinical unit in Secaucus, New Jersey, designed to conduct large?scale Phase I and Bioequivalence (BE) studies as well as radiolabeled human AME research.

The company has further strengthened its C14?hAME capabilities through a dedicated unit staffed by licensed radiation experts and supported by an onsite nuclear pharmacy for sterile and non?sterile radiolabeled compounding. Frontage has successfully executed combined hAME and Absolute Bioavailability studies aligned with the FDA's 2024 guidance.

In response to growing industry demand, Frontage is also expanding into Oncology Phase I trials through partnerships with regional hospitals and clinical networks, enabling patient?based research across key cancer indications.

As the industry approaches a significant wave of patent expirations, Frontage's multiple US clinical units are positioned to manage complex, high?volume BE programmes. Collaboration with Frontage China enables flexible Multi?Regional Clinical Trial (MRCT) execution to support global programme timelines.