Company Statement of May 13, 2022

In the United States, any drug under development must be tested in a controlled environment to properly assess its safety before being tested on humans. This federally required testing is part of scientific research mandated by government agencies in the United States and around the world.

Inotiv provides biomedical research services to companies developing life-saving treatments that span a range of human diseases. In our Indiana facility, the drug being tested, if successful, will address a specific rare disease affecting newborns, infants and children. This drug must be tested and confirmed to be safe both in toxicity and in dosage before it advances to clinical trials in humans. For more information about the federally-mandated drug development process, please find several links to government information and other publicly available resources below.

Inotiv complies with all applicable federal, state and local regulations, as well as the Animal Welfare Act, and, at our Indiana and Maryland facilities, is accredited by AAALAC, the Association for the Accreditation and Assessment of Laboratory Animal Care International.

FDA

FBR 1

FBR 2