



PRESS RELEASE  
ImmunoSafe Platform® - Non-clinical ISR assessment

testament to our commitment in revolutionizing drug development. By offering a comprehensive solution for local toxicity assessment, we aim to provide our biotech and pharma partners with actionable first-in-human data, ahead of clinical trials, ensuring safer and more effective therapeutic solutions."

Genoskin recently showcased its innovative approach at Eurotox 23 in September, presenting a poster titled "[Automated prediction of drug reactogenicity at the site of injection, by combining natural human skin & computational inference of biological pathway](#)." This presentation further emphasizes the company's dedication to advancing the field of non-clinical local toxicity assessment and its potential impact on the future of drug development.

For more information about Genoskin and its pioneering service offering, ImmunoSafe: ISR Platform® with AUDACY, please visit <https://www.genoskin.com>.

About Genoskin:

Genoskin provides transformative non-clinical platforms leveraging human skin to test therapeutic and non-therapeutic products. By generating actionable human data, Genoskin provides a reliable alternative to animal experiments, ensuring safer and more effective drug development. The company uses real human tissues, prepared from donated human skin leftovers ethically sourced from plastic surgeries, and innovative technologies to maintain viability, immunocompetency and functionality of the human skin samples in a ready-to-use *ex vivo* culture system. Genoskin was founded in 2011, as a spin-off of the French National Center for Scientific Research (CNRS) and The Paul Sabatier University in Toulouse.



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