

# Verici Receives Medicare Coverage for Tutivia

FRANKLIN, TN, UNITED STATES, April 11, 2025 /EINPresswire.com/ -- Verici Dx plc (AIM: VRCI), a developer of advanced clinical diagnostics for organ transplant is pleased to announce a positive Technical Assessment by Palmetto GBA, a Medicare Administrative Contractor (MAC) that assesses diagnostic technologies through its Molecular Diagnostic Services Program (MoIDX) allowing for Medicare coverage of its Tutivia assay, a diagnostic test for acute rejection. Patients with a high-risk Tutivia score are nearly six times as likely to experience acute rejection over those with low-risk results.

Tutivia is the Company's post kidney transplant blood test that reports the patient's risk of all forms of acute rejection, including borderline, T cell-mediated, and antibody-mediated rejections. The test therefore supports clinical care for the estimated c.100,000 global patients who undergo kidney transplant procedures annually. Tutivia delivers a significant improvement in biomarker offerings, particularly early post-transplant when patient management can be complex and other biomarkers are contraindicated or less informative. Tutivia's has an important role in helping improve patient outcomes with significant implications for clinical care, providing relevant information for immunosuppression management and supporting clinical decision making with regard to kidney biopsy.

Following a review of clinical validity, analytical validity, and clinical utility, MoIDX determined that Tutivia has met the criteria for coverage under LCD L38568, MoIDX: Molecular Testing for Solid Organ Allograft Rejection for renal transplant patients. The MoIDX coverage decision signifies that Tutivia meets requirements for Medicare coverage and will support broader access for renal transplant patients.

Commenting on the approval, Sara Barrington, CEO of Verici Dx, said:

"The achievement of this key milestone reflects the exciting work that is ongoing at Verici and the quality of its science. Securing a positive MoIDX coverage decision will improve patient access to the Tutivia assay, advancing both availability and adoption. With this decision, we hope that more patients can benefit from a significant step forward in enhancing diagnostic accuracy and post-transplant patient care. With Tutivia already seeing strong positive feedback from early adopters and the accelerated sales already coming through, we look forward to continued success in a commercial setting."

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