Newmarket Biomedical Ltd.
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DATE: 15 January 2025

Quality Policy Statement

At Newmarket Biomedical Ltd, we are dedicated to delivering in-vitro diagnostic medical devices and exceptional product support that consistently meet or exceed customer expectations.

Our commitment to quality is unwavering and grounded in compliance with ISO 13485:2016, IVDR (EU) 2017/746, Medical Device Single Audit Program (MDSAP), and specific business requirements across all operational territories.

To achieve this, we prioritize:

- Continual Improvement: Regularly refining and enhancing the effectiveness of our quality management systems to adapt to emerging needs.
- Quality Objectives: Establishing and reviewing measurable objectives to ensure alignment with our mission and stakeholder expectations.
- Infrastructure and Environment: Maintaining robust infrastructure and a safe, supportive work environment that fosters staff development and ensures the sustainability of our business operations.

By embedding these principles in our culture and operations, Newmarket Biomedical Ltd strives to be a trusted partner, delivering products and services that stand as their own best endorsement of quality and reliability.

Colin A Knox
Chief Executive Officer

Mark R Bates
Chief Operations Officer

Helen Masters
Technical & Regulatory Director