

MEDIA RELEASE



The news follows the recent US Food and Drug Administration (FDA) approval of two of the company's hematology analyzers, the [Abacus 3CP](#), 3-part differential analyzer and [Abacus 5](#), 5-part differential analyzer.

With more than 30,000 hematology analyzers already sold in over 100 international markets, the creation of the new production facilities, together with the US FDA approvals, are important milestones that have strengthened Diatron's global position, opening up many additional opportunities in the US market.

According to Ronald Backes, Vice President of International Business Development at Diatron, "One of the outcomes of the new production facility will be to concentrate our research and design efforts. This will allow us to ensure we are producing analyzers that match our customer's needs and provide an optimum service in terms of reliability, efficiency and cost-effectiveness".

For more information please visit the Diatron Group website: <http://www.diatron.com>

Editors' Notes

The Diatron Group

Diatron's vision since the company's foundation in 1989, is to become a worldwide leader *in-vitro* diagnostics, providing customers with a broad portfolio of high-quality tailored solutions. In its new production facility in Budapest, Hungary, the extensive R&D/Engineering team develops and manufactures compact hematology analyzers and reagents for the human medical and veterinary markets.

Diatron is a portfolio company of, The Riverside Company, a global private equity firm focused on acquiring growing businesses valued at up to \$200 million (€200 million in Europe). Founded in 1988, Riverside's international portfolio now includes 80 companies, and it has more than \$3 billion/€2.5 billion in assets under management.

See www.riversidecompany.com for more information