



Contacts: Melissa Neill
[Risdall McKinney Public Relations](mailto:mneill@risdall.com)
(651) 286-6736
mneill@risdall.com

Jim Bartel
[ATEK Companies](mailto:jbartel@atekcompanies.com)
(763) 392-5897
jbartel@atekcompanies.com

ATEK Medical Hires for New Regulatory and Quality Position

New position will manage increasing regulatory and compliance responsibilities necessary for international operations

GRAND RAPIDS, MICH. (July 24, 2008) – [ATEK Medical](http://www.atekmedical.com), a division of [ATEK Companies](http://www.atekcompanies.com), has hired Felix Le as director of quality and regulatory affairs. In this position, Le will be responsible for maintaining ATEK’s compliance with FDA regulations as well as international regulatory policies and overseeing the company’s continuous quality improvement processes.

“The medical industry is one of continuous improvements and consistently changing regulations,” said Chris Oleksy, president, ATEK Medical. “With the opening of our Costa Rican plant and increased production volume due to growth in our clients’ businesses, maintaining the highest levels of quality and regulatory compliance is essential. The addition of Le to the ATEK team solidifies our commitment to manufacturing the best product with the most efficient process, while still maintaining a highly controlled environment.”

Le brings to ATEK more than two decades of experience in regulated industries — nearly 17 years of which he spent at Medtronic where he held positions of increasing responsibility. Most recently Le served as senior manufacturing and pharmaceutical quality manager for Medtronic’s neuromodulation business unit. Prior to Medtronic, he held a number of positions at FMC Corporation in its naval systems division. Additionally, Le is trained in SixSigma and is an ASQ certified quality auditor, quality engineer, software quality engineer and quality manager.

For additional information about ATEK Medical, please visit www.atekmedical.com

About ATEK Medical

ATEK Medical is a total solutions partner serving the medical, disposable, electro-mechanical and implantable device markets. From product design and launch to distribution and post-production fulfillment, the ATEK Medical team is dedicated, skilled and cross-trained to serve your company’s needs. ATEK Medical capabilities include product development and launch; medical device manufacturing; packaging, labeling and sterilization services; microbiology services; supply chain management; and post-production fulfillment services. ATEK Medical is ISO 13485:2003 certified and FDA registered to design, produce and launch Class I, II and III medical devices. Since 1979, ATEK Medical has developed and launched more than 300 unique

products and specializes in product management and full-service manufacturing. For more information on ATEK Medical, visit www.atekmedical.com

About ATEK Companies

ATEK Companies is the preferred manufacturing partner serving diverse industries including medical, aerospace, defense, communications, and transportation. ATEK Companies distinguishes itself from the competition through its 60-year history of delivering unparalleled value, high quality products, and creative solutions. ATEK Companies offers total project management, including electrical and mechanical product design and launch, supply chain management, production, packaging and distribution, and post-production fulfillment services. ATEK Companies is comprised of five businesses, which include ATEK Medical, ATEK Manufacturing, ATEK Plastics, ATEK Products, and Progress Casting. Facilities are located in Minneapolis and Brainerd, Minn.; Kerrville, Texas; New Hampton, Iowa; Grand Rapids, Mich.; and Heredia, Costa Rica. For more information visit www.atekcompanies.com

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