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**PUROLITE RECEIVES FOOD AND DRUG CERTIFICATION AND GMP
COMPLIANCE CERTIFICATION FOR PUROLITE S.R.L.**

BALA CYNWYD, PA – September 10, 2009 – Purolite Corporation, the preferred global supplier of ion exchange resins, has recently received FDA and GMP compliance registration for the manufacturing of bulk pharmaceutical/biotech products for the medical related industries.

“As a leading global manufacturer of traditional and specialty ion exchange resins for industrial, commercial, and municipal water needs, Purolite has been also expanding into the medical, pharmaceutical and biotech industries for a number of years. These regulated industries require strict government audits of manufacturing processes and facilities to ensure quality and safety metrics are consistently achieved.

“Purolite is fully committed to meeting these stringent medical regulations”, says Don Brodie, Vice President of Operations at Purolite. “We are dedicated to anticipating the needs of our global customers and exceeding their expectations with new technology, products, and applications. Purolite continues to grow through our commitment to providing the highest quality product and technical support to every customer”

Purolite Corporation was founded in 1981 and is a leading manufacturer of ion exchange, catalyst, adsorbent and specialty resins and the only company to focus exclusively on this market. Headquartered in Bala Cynwyd, PA, the company has ISO-9000:2001 certified sales offices in more than 30 countries, as well as manufacturing and R&D facilities in the USA, China and Romania. Purolite also has a dedicated central research and development facility in the United Kingdom.