FRESENIUS

Investor News

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Fresenius Kabi Receives FDA Status Upgrade for US Grand Island Facility

Fresenius Kabi has been informed by the U.S. Food and Drug Administration that its pharmaceutical manufacturing facility in Grand Island, N.Y., has achieved the upgraded status of voluntary action indicated (VAI) following an October 2014 inspection. The status change is an improvement from the “official action indicated” status the facility had been operating under.

The new VAI classification permits FDA approval of new Fresenius Kabi products at the plant.

The status change reflects the improvements that have been made at the plant since receiving a warning letter in 2012. Fresenius Kabi remains committed to continuous improvement and compliance in its operations worldwide.

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Fresenius is a global health care group, providing products and services for dialysis, hospital and outpatient medical care. In 2013, Group sales were €20.3 billion.

For more information visit the Company’s website at www.fresenius.com.

Fresenius SE & Co. KGaA

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Chairman of the Supervisory Board: Dr. Gerd Krick

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