

Press Release

Matthias Link
Corporate Communications

Fresenius SE & Co. KGaA
Else-Kröner-Straße 1
61352 Bad Homburg
Germany
T +49 6172 608-2872
F +49 6172 608-2294
matthias.link@fresenius.com
www.fresenius.com

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Fresenius Kabi reaches a global agreement with AbbVie regarding Fresenius Kabi's adalimumab, MSB11022, a biosimilar candidate of AbbVie's Humira®

Fresenius Kabi has signed a worldwide settlement and license agreement with AbbVie, settling all pending patent litigations between the two companies. Under the terms of the agreement and subject to marketing authorization by the health authorities, Fresenius Kabi's biosimilar candidate of Humira®* (adalimumab), MSB11022**, could be commercialized in the United States from September 30, 2023.

On October 17, 2018 licenses under the agreement came into effect in certain countries in Europe in which AbbVie owns intellectual property. The application for marketing authorization for MSB11022 was submitted by Fresenius Kabi to the European Medicines Agency (EMA) at the end of last year. The dossier is currently under review. A first launch in Europe is expected in the first half of 2019.

"This agreement is a major step on our way to successfully developing and commercializing our biosimilar portfolio," said Dr. Michael Schönhofen, Member of the Fresenius Kabi Management Board and President of the Pharmaceuticals Division. "In line with our caring for life philosophy, our aim is to contribute to broader access to affordable therapies for chronic and acute diseases. Biosimilar drugs are an increasingly important component in reaching this aim – to the benefit of patients and healthcare systems. The agreement with AbbVie provides further

clarity regarding when we will be able to commercialize our biosimilar candidate of Humira®.”

*Humira® (adalimumab) is a registered trademark of AbbVie Biotechnology Ltd.

**MSB11022 is being developed as a biosimilar candidate of Humira® and is not yet approved by health authorities.

About Fresenius Kabi

Fresenius Kabi is a global health care company that specializes in lifesaving medicines and technologies for infusion, transfusion and clinical nutrition. The company's products and services are used to help care for critically and chronically ill patients. Fresenius Kabi's product portfolio comprises a comprehensive range of I.V. generic drugs, infusion therapies and clinical nutrition products as well as the medical devices for administering these products. Within transfusion technologies, Fresenius Kabi offers products for collection and processing of blood components and for therapeutic treatment of patient blood by apheresis systems.

In the field of biosimilars, Fresenius Kabi develops products with a focus on oncology and autoimmune diseases. With its corporate philosophy of "caring for life", the company is committed to putting essential medicines and technologies in the hands of people who help patients and finding the best answers to the challenges they face.

Fresenius Kabi employs over 36,000 people worldwide. In 2017 the company reported sales of more than €6 billion. Fresenius Kabi AG is a wholly owned subsidiary of the Fresenius SE & Co. KGaA healthcare group.

For more information visit the Company's website at www.fresenius-kabi.com

This release contains forward-looking statements that are subject to various risks and uncertainties. Future results could differ materially from those described in these forward-looking statements due to certain factors, e.g. changes in business, economic and competitive conditions, regulatory reforms, results of clinical trials, foreign exchange rate fluctuations, uncertainties in litigation or investigative proceedings, and the availability of financing. Fresenius Kabi does not undertake any responsibility to update the forward-looking statements in this release.

Management Board: Mats Henriksson (Chairman), Marc Crouton, John Ducker, Chris Harrison, Dr. Christian Hauer, Dr. Michael Schönhofen, Gerrit Steen
Chairman of the Supervisory Board: Stephan Sturm
Registered Office: Bad Homburg, Germany
Commercial Register: Amtsgericht Bad Homburg - HRB 11654