



Published on 30 January 2019 by **Thomas Caveng**, Legal Translator / Marketing Director t.caveng@soulier-avocats.com

Tel.: + 33 (0)4 72 82 20 80

Read this post online

US Regulation of Food Products: What to Expect in 2019



Title: US Regulation of Food Products: What to Expect in 2019

Jurisdiction: USA

Authors: Raqiyyah Pippins, Dana Weekes, Pari R. Mody and Elizabeth Trentacost

Law firm: Arnold & Porter Kaye Scholer LLP

Subject:

The US Food and Drug Administration (FDA or the Agency) and the US Department of Agriculture's (USDA's) 2018 policy and regulatory activities have set the foundation for a robust 2019 policy agenda — once the partial shutdown of the US government ends. While we await the resolution of the shutdown — which has suspended all but the most urgent food regulatory tasks — and issuance of FDA's strategic plan for 2019, there are key FDA, USDA and legislative developments that occurred towards the end of 2018 that should be on the radar of every company that manufactures, sells or markets conventional food or dietary supplement products in the United States.

Read the contribution



<u>Soulier Avocats</u> is an independent full-service law firm that offers key players in the economic, industrial and financial world comprehensive legal services.

We advise and defend our French and foreign clients on any and all legal and tax issues that may arise in connection with their day-to-day operations, specific transactions and strategic decisions.

Our clients, whatever their size, nationality and business sector, benefit from customized services that are tailored to their specific needs.

For more information, please visit us at www.soulier-avocats.com.

This material has been prepared for informational purposes only and is not intended to be, and should not be construed as, legal advice. The addressee is solely liable for any use of the information contained herein.