

Johnson & Johnson Shareholder Meeting Question
Scott Shepard, Free Enterprise Project Coordinator
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I'm Scott Shepard with the National Center for Public Policy Research.

As Johnson & Johnson revealed in December, FDA testing errors resulted in the company fearing that its baby powder had been contaminated with asbestos – an extremely serious charge, and error.¹ More recently, the FDA's long and cumbersome regulatory process has proven a hindrance to speedy production of protective and medical equipment, and to the development of tests, treatments and cures for COVID-19, which has resulted in some waivers and exceptions to normal processes.² Given these experiences, will Johnson & Johnson call for review and streamlining of the FDA approval and regulatory processes, and for a halt to deeper intrusions of government bureaucracy into the U.S. healthcare system?

*Contact: Judy Kent at (703) 759-0269 or cell (703) 477-7476 or JKent@nationalcenter.org
and David W. Almasi at (703) 568-4727 or DAlmasi@nationalcenter.org
National Center for Public Policy Research
20 F Street, NW, Suite 700, Washington, DC 20001
www.nationalcenter.org • [@NationalCenter](https://twitter.com/NationalCenter) • [@FreeEntProject](https://twitter.com/FreeEntProject)*

¹ See, e.g., Noah Manskar, *Johnson & Johnson Blames FDA for Baby Powder Asbestos Scare*, NEW YORK POST (Dec. 4, 2019), available at <https://nypost.com/2019/12/04/johnson-johnson-blames-fda-for-baby-powder-asbestos-scare/> (last accessed April 15, 2020).

² See, e.g., Lisa Winter, *First Saliva Test for COVID-19 Approved for Emergency Use by FDA*, THE SCIENTIST (April 14, 2020), available at <https://www.the-scientist.com/news-opinion/first-saliva-test-for-covid-19-approved-for-emergency-use-by-fda-67416/> (last accessed April 15, 2020); Madlin Mekelburg, *Did FDA Regulations Slow Testing for the Coronavirus? Fact-Checking Rep. Dan Crenshaw's Claim*, SAN ANTONIO EXPRESS-NEWS (April 13, 2020), available at <https://www.expressnews.com/politics/texas/article/Did-FDA-regulations-slow-testing-for-the-15190265.php> (last accessed April 15, 2020); Robin Patel & Stefano Bertuzzi, *Expediting COVID-19 Testing and Expanding Access to Clinical Labs*, AMERICAN SOCIETY FOR MICROBIOLOGY (March 9, 2020), available at <https://asm.org/Articles/2020/January/Expediting-COVID-19-Testing-and-Expanding-Access-t> (last accessed April 15, 2020).