January 1, 2025

Subject: **European Union Medical Device Regulation**

Dear Valued Business Partner,

Arrow Electronics, Inc. or one of its affiliates or wholly owned subsidiaries (collectively, "Arrow") is in receipt of your inquiry regarding the European Union Medical Device Regulation ("EU MDR"). It is Arrow's practice to respond to such matters through its corporate offices.

EU MDR "*lays down rules concerning the placing on the market, making available on the market or putting into service of medical devices for human use and accessories for such devices*." As Arrow is not a manufacturer, importer, nor distributor of "medical devices," the obligations of EU MDR do not apply to our business. Moreover, the electronic component parts distributed by Arrow are not intended for direct contact with the human body in medical applications and are therefore beyond the scope of both the letter and spirit of EU MDR.

Medical device manufacturers are the best source of accurate information regarding the composition, safety, and performance of their products.

For any additional inquiries regarding the application of EU MDR to Arrow's operations, please reach out to compliance@arrow.com.

Thank you for your continued confidence in Arrow.

Sincerely,

Arrow Electronics, Inc.

9151 E Panorama Cir
Centennial, CO 80112
P 303 824 4000
arrow.com