

MEDTRONIC PLC

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Manufacturing Site: Parkmore Business Park West, Galway, Ireland

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Ref No: 0086869

Date: 20/07/2026

To: Procurement Committee / Tender Authorities of the Republic of Armenia

MANUFACTURER'S AUTHORIZATION LETTER

We, Medtronic plc, as an official manufacturer of medical devices having our registered operational headquarters at 710 Medtronic Parkway, Minneapolis, MN 55432, USA and manufacturing facilities at Galway, Ireland, do hereby authorize "NORD" LLC (TIN: 09433452, Address: 12 M. Harutyunyan str., apt. 63, Kapan, Syunik, Republic of Armenia) to submit a bid, negotiate and sign the contract concluded as a result of the procurement procedures for the supply of the following goods manufactured by us:

Product Name: Zotarolimus-Eluting Coronary Stent System (Resolute Onyx™)

We hereby certify and guarantee that the offered product fully complies with all technical specifications requested for the tender:

1. Material & Structure: Platinum-iridium (Pt-Ir) core with cobalt alloy casing shell, designed with a unique single-wire continuous sinusoid technology.
2. Coating & Polymer: Biocompatible polymer (BioLinx™) with an outer hydrophilic coating.
3. Drug formulation: Zotarolimus, with a precise drug density of 1.6 µg/mm² (±1%).

4. Crossing Profile: Stent crossing profile does not exceed 0.041 inches.
5. Size Matrix: Diameters of 2.25, 2.50, 2.75, 3.00, 3.50, and 4.00 mm. Each diameter option provides at least 9 distinct lengths, where the shortest available length is no more than 8 mm, and the longest length is not less than 38 mm.
6. Quality Certification: CE MARK and/or FDA quality certificates are fully valid and guaranteed for any supplied batch.
7. Condition: Brand new, unused, in factory sterile packaging. Includes the original factory manual containing technical indicators, features, and instructions for use.

Furthermore, in compliance with Article 2 of the RA Law HO-325-N "On Medical Care and Services to the Population" and specific tender logistics, we guarantee and undertake the following:

- A Certificate of Conformity and a Manufacturer's Authorization Letter are provided directly with the Bid.
- A Certificate of Origin for the specific delivered products shall be officially presented during the contract performance stage.
- At the time of delivery, the remaining shelf life shall strictly meet the following thresholds:
 - * For products with a total shelf life of up to 1 year: at least 75% remaining.
 - * For products with a total shelf life of 1 to 2 years: at least 2/3 remaining.
 - * For products with a total shelf life of more than 2 years: at least 15 months remaining.
- The delivery shall be fully executed by the supplier to the destination address: Gyumri, Garegin Nzhdeh str. 3/3, strictly before 16:00 on the day of delivery.
- The initial delivery phase shall be executed within a minimum of 20 calendar days from the effective date of the contract, and any subsequent stage-by-stage deliveries shall be carried out within 3 working days upon the customer's request. The final delivery deadline shall not exceed December 25, 2026.
- All product transportation and unloading processes shall be executed entirely at the supplier's expense and resources.

This authorization letter is valid until December 31, 2026.

For and on behalf of Medtronic plc:

 / 销售经理 /
(Signature of Authorized Signatory)

[Official Corporate Stamp / Seal]