NOBOMED

QUALITY POLICY STATEMENT

NoBoMed Zrt. provides its conformity assessment activities in compliance with the requirements of the MSZ EN ISO/IEC 17021-1:2016 standard and Regulation (EU) 2017/746 (hereinafter: IVDR) referred to as governing the placing on the market of in vitro diagnostic medical devices.

We ensure that our employees understand and accept the provisions of the quality policy.

Our goal is to serve our customers reliably in the long term, while complying with standards, and legal requirements, ,and taking into account common specifications, professional guidelines containing, best practices recommendations. To this end, our management ensures:

- the continuous improvement of the quality management system and conformity assessment activities;
- the necessary resources;
- adequate staffing levels;
- appropriate environmental conditions;
- regular monitoring of our processes; and the provision of services that meet our customers' expectations.

The primary goal of our efforts—to which we subordinate all our activities—is to achieve and maintain customer satisfaction through the continuous improvement of the quality of our services and the quality management system of NoBoMed Zrt.

We support and require the acquisition and application of up-to-date knowledge. Through regular training, we ensure our employees are prepared to perform the tasks ahead of us.

For the management of NoBoMed Zrt., ensuring impartiality and independence is of special importance in connection with its conformity assessment activities; the principles thereof are set forth in the Statement on Impartiality, Independence, and Confidential Information Management.

Our quality objectives have been defined based on the quality policy , and all our employees are responsible for their implementation.

Budapest, June 08, 2026.

Dr. Csilla Pozsgay m.p.
Chair of the Board of Directors