Public declaration regarding the manufacture and use of in-house devices by health institutions

Name of health institution: Suomen Terveystalo Oy

Address: Jaakonkatu 3 A, 00100 Helsinki

Suomen Terveystalo Oy declares that the devices described in the accompanying table are only manufactured and used in Suomen Terveystalo Oy and do meet the applicable general safety and performance requirements (GSPR) of the medical devices Regulation (EU 2017/745). A reasoned justification is provided in case applicable general safety and performance requirements are not fully met.

Date and location: 20.10.2026 Tampere

Name, function and signature of responsible person(s):



Annika Rökman, SaMD Quality Manager

Table of in-house devices:

Device identification (e.g. name, description, reference number)	**Device type (IVD/ MD)**	**Risk class of the device**	**Intended purpose**	**Applicable GSPR fully met? (Y/N)**	**Information on and justification for applicable GSPR that are not fully met (using the numbering as in Annex I of the IVDR/MDR)**
Survey Result Calculator (SRC) version 1.X.Y	MD	I	The Survey Result Calculator (SRC) is a software application intended to process, calculate, and present responses from	Y	

			questionnaires and other structured data collection instruments. The software generates predefined qualitative and quantitative outputs for information and documentation purposes. Thereby SRC facilitates documentation, communication, information management, and administrative patient management within healthcare processes. It is intended for use by patients or persons acting on their behalf, and by healthcare professionals, through Terveystalo user interfaces. The software does not perform diagnostic, prognostic, therapeutic, monitoring, or clinical decision-support functions.		