DECLARATION OF CONFORMITY for EU and UK

Radio Equipment Product Name:	Teladoc Health TV Pro 300 (a.k.a. TV Pro 300)
Model number:	20-20444
Product type/description:	Telehealth product for audio / video telecommunication
Supplied accessories, if any:	Remote control, wall mounting assembly, cables

Name and Address of the Manufacturer:
Teladoc Health, Inc.
7406 Hollister Avenue
Goleta, CA 93117

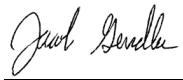
Route to Compliance
The route of compliance for the TV Pro 300 radio equipment is Annex II conformity assessment Module A – internal production control of EU Directive 2014/53/EU and Schedule 2 Module A Conformity Assessment of the UK Radio Equipment Regulations 2017. Fulfilment of the essential requirements set out in Article 3 of EU Directive 2014/53/EU and in Part 2 Chapter 1 of UK Radio Equipment Regulation 2017 have been demonstrated.

Harmonized Standards Applied
Please refer to Appendix A attached to this declaration.

I hereby declare the above product is in conformity with the EU Radio Equipment Directive 2014/53/EU, UK Radio Equipment Regulations 2017, and Council Directive 2011/65/EU (RoHS). The product has been evaluated according to the conformity assessment procedure mentioned above in section 'Route to Compliance'. This declaration of conformity is issued under the sole responsibility of the manufacturer, Teladoc Health, Inc.

Please refer to Appendix B for a history of this declaration.

Signed for and on behalf of Teladoc Health, Inc.



December 6, 2023 San Jose, CA USA
(Signature) (Date) (Place of issue)

Jacob Gendler
VP Quality Assurance & Regulatory Affairs
Teladoc Health, Inc.

Appendix A – Applied Standards

Harmonized Standards		
Number	**Edition/Year**	**Title**
2014/53/EU	2014	EU Radio Equipment Directive
2011/65/EU, EU 2015/863	2011, 2015	Restriction of the use of certain hazardous substances (RoHS) in electrical and electronic equipment
Other technical specifications		
Number	**Edition/Year**	**Title**
EN 62368-1	2014+A11:2017	Audio/Video, Information and Communication Technology Equipment – Part 1: Safety Requirements
EN 60601-1-2	2015/A1:2021	Medical electrical equipment-General requirements for basic safety and essential performance. Collateral Standard: Electromagnetic disturbances. Requirements and tests
EN 60601-1	2006+A1:2013+A12:2014+A2:2021	Medical electrical equipment-Part 1: General requirements for basic safety and essential performance