



F. BOSCH
M E D I Z I N T E C H N I K
G E R M A N Y

Friedrich Bosch GmbH & Co. KG · Postfach 5 · D-72402 Bisingen

DECLARATION OF CONFORMITY

We company **FRIEDRICH BOSCH GMBH & CO. KG, Hohenlaimstr. 30, 72406 Bisingen, Germany**

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declare under our sole responsibility, that the product:

aneroid sphygmomanometer Prakticus I

Article number:

9021.20.112 Prakticus I with velcro cuff black
9021.20.105 Prakticus I with velcro cuff burgundy
9021.20.116 Prakticus I with velcro cuff red
9021.20.117 Prakticus I with velcro cuff blue
9021.20.108 Prakticus I with velcro cuff green
9021.20.109 Prakticus I with velcro cuff yellow
9021.20.110 Prakticus I with velcro cuff orange

9022.20.112 Prakticus II with velcro cuff black
9022.20.105 Prakticus II with velcro cuff burgundy
9022.20.116 Prakticus II with velcro cuff red
9022.20.117 Prakticus II with velcro cuff blue
9022.20.108 Prakticus II with velcro cuff green
9022.20.109 Prakticus II with velcro cuff yellow
9022.20.110 Prakticus II with velcro cuff orange

class of product: class **I M**

to which this declaration relates is in conformity with the following standards
Guideline 93/42 EWG appendix VI.

The quality system is attested by the accredited body TÜV Süd PRODUCT SERVICE GmbH, Ridlerstr. 65, 80339 München with the licence number **0123**.
The mentioned product is manufactured in compliance with EN 1060-1 and 2: 1994.
The certificate has the number GM3 20267 0020 Rev. 00 and is valid until 26.05.2024.

date: 06.02.2020

Friedrich Bosch GmbH & Co. KG
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Josef Bieger
Director of quality-management

Amtsgericht Stuttgart HRA 420 693
Verwaltungsgesellschaft mbH,
Sitz Bisingen · HRB 420 287
Geschäftsführer: Josef Bieger
Dipl.-Kaufmann