



Benannt durch/Designated by
 Zentralstelle der Länder
 für Gesundheitsschutz
 bei Arzneimitteln und
 Medizinprodukten
 www.zifa.de
 BS-IVDR-099



Product Service

EU Quality Management System Certificate (IVDR)

Pursuant to Regulation (EU) 2017/746 on in Vitro Diagnostic Medical Devices,
 Annex IX Chapters I and III (Class A Devices in Sterile Condition)

No. V11 042464 0039 Rev. 00

Classification: Class A
Device Group: W050101 - BLOOD COLLECTION DEVICES
Intended Purpose: IVR 0803 - Specimen receptacles referred to in point 2.5 (rule 5),
 under c), of Annex VIII to Regulation (EU) 2017/746

The validity of this certificate depends on conditions and/or is limited to the following:

Revision History:

Rev.	Dated	Report	Description
00	2023-04-11	SH2211102	Initial issuance



