Filtered Pipette Tips

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Filter pipette tips are used for protection against contamination.
Pipette tips with filter barrier protects pipettors and samples from contamination.
RNase, DNase and Pyrogen Free.
Filtered tips,Filter Tips Pipette, universal and fit most popular brands of pipettors.

Sterile pipette tips.
PP Material, clear color.
Volume:10ul-1250ul.
Package Style: Bulked: 1000tips/bag, 10bags/ctn
Racked: 96tips/rack, 10racks/pk, 50racks/ctn

Cat No.	Description	Qty/Case(pcs)
640401	Racked filter tips,96tips/rack, sterile, 10ul	4800
640402	Racked filter tips,96tips/rack, sterile, 10ul long	4800
640403	Racked filter tips,96tips/rack, sterile, 20ul	4800
640404	Racked filter tips,96tips/rack, sterile, 50ul	4800
640405	Racked filter tips,96tips/rack, sterile, 100ul	4800

640406	Racked filter tips,96tips/rack, sterile, 200ul	4800
640407	Racked filter tips,96tips/rack, sterile, 1000ul	4800
640408	Racked filter tips,96tips/rack, sterile, 1250ul (1000ul long)	4800
640409	Racked filter tips,96tips/rack, sterile, 200ul long	4800
640410	Racked filter tips,96tips/rack, sterile, 300ul	4800
640411	Bulked filter tips,1000tips/bag, sterile, 10ul	10000
640412	Bulked filter tips,1000tips/bag, sterile, 10ul long	10000
640413	Bulked filter tips,1000tips/bag, sterile, 20ul	10000
640414	Bulked filter tips,1000tips/bag, sterile, 50ul	10000
640415	Bulked filter tips,1000tips/bag, sterile, 100ul	10000
640416	Bulked filter tips,1000tips/bag, sterile, 200ul	10000
640417	Bulked filter tips,1000tips/bag, sterile, 1000ul	10000
640418	Bulked filter tips,1000tips/bag, sterile, 1250ul (1000ul long)	10000
640419	Bulked filter tips,1000tips/bag, sterile, 200ul long	10000
640420	Bulked filter tips,1000tips/bag, sterile, 300ul	10000

EG-KONFORMITÄTSERKLÄRUNG · EC DECLARATION OF CONFORMITY
DÉCLARATION CE DE CONFORMITÉ · DICHIARAZIONE CE DI CONFORMITÀ

Name und Adresse des Herstellers: / Name and address of the manufacturer: / Nom et adresse du fabricant: / Nome e indirizzo del fabbricante:	**BOEN HEALTHCARE CO., LTD Unit 602, International Center, No.535, Shenxu Road, Suzhou, 215021, Jiangsu, China**

Wir erklären in alleiniger Verantwortung, dass / We declare under our sole responsibility that / Nous déclarons sous notre propre responsabilité que / Dichiariamo sotto la sola responsabilità che

das Medizinprodukt: / the medical device: / le dispositif médical: / il dispositivo medico:	**Filtered Pipette Tips**
der Klasse: / of class: / de la classe: / di classe:	**Common/Others IVD (Devices of NOT Annex II and NOT self-test)**

(IVDD, Artikel 9 Absatz 1) nicht Teil der Liste A und B von Anhang II sein / (IVDD, Article9(1)) not be part of list A & B of annex II (IVDD, article 9, paragraphe 1) ne fait pas partie de la liste A et B de l'annexe II / (IVDD, articolo 9, paragrafo 1) non fanno parte dell'elenco A e B dell'allegato II

den einschlägigen Bestimmungen der Medizinprodukte-Richtlinie 98/79/EG und deren Umsetzungen in nationale Gesetze entspricht. Die Erklärung gilt in Verbindung mit dem zum Produkt gehörigen „Endprüfprotokoll“. /

meets the provisions of the directive 98/79/EC and its transpositions in national laws which apply to it. The declaration is valid in connection with the “final inspection report” of the device. /

remplit toutes les exigences de la directive sur les dispositifs médicaux 98/79/EC et de ses transpositions en droit national qui le concernent. La déclaration est valable si elle est associée au «rapport de l’inspection finale» du produit. /

soddisfa tutte le disposizioni della direttiva 98/79/EC e della loro trasposizione nel diritto nazionale che lo riguardano. Questa dichiarazione è valida in congiunzione con il “rapporto di ispezione finale” del prodotto.

Konformitätsbewertungsverfahren: / Conformity assessment procedure: / Procédure d’évaluation de la conformité: / Procedura di valutazione della conformità:	**Anhang III (voraussichtlicher Punkt 6) der IVDD 98/79 / EG Annex III (expect point 6) of IVDD 98/79/EC Annexe III (sauf le point 6) de l'IVDD 98/79 / CE Allegato III (aspettarsi il punto 6) dell'IVDD 98/79 / CE**
Registrier-Nr.: / Registration No.: / N°d’enregistrement: / Numero di registrazione:	
Benannte Stelle: / Notified Body: / Organisme notifié: / Organismo notificato:	

CE

Suzhou, 2022.01.01

General Manager

Ort, Datum / Place, date / Lieu, date / Luogo, data

Name und Funktion / Name and function / Nom et fonction / Nome e funzione