

**medax**

# PERFECTUS

BONE MARROW ASPIRATION SYSTEM "PS"  
ALSO AVAILABLE IN THE EXPLANT/TRANSPLANT  
VERSION "PF"



EXPLANT/TRANSPLANT VERSION

Bone Marrow aspiration system "PS",  
also available in the explant/transplant version "PF".

- Triple sharpened tip cannula to allow a non traumatic, fast and easy penetration into the bone marrow cavity.
- Adjustable depth setter.
- Luer Lock connection for syringe.
- Sterilized by ETO, shelf life 5 years.

**medax**  
medical devices

Your partner in Biopsy and Special Needles

# PERFECTUS

BONE MARROW ASPIRATION SYSTEM "PS"

ALSO AVAILABLE IN THE EXPLANT/TRANSPLANT VERSION "PF"

GAUGE	COLOUR	LENGTH (MM)							
14	● Green	14 020	14 030	14 035	14 040	14 045	14 050	14 055	14 060
15	● Blue	15 020	15 030	15 035	15 040	15 045	15 050	15 055	15 060
16	○ White	16 020	16 030	16 035	16 040	16 045	16 050	16 055	16 060
18	● Pink	18 020	18 030	18 035	18 040	18 045	18 050	18 055	18 060

GAUGE	COLOUR	LENGTH (MM)								
14	● Green	14 065	14 070	14 075	14 080	14 085	14 090	14 095	14 100	14 150
15	● Blue	15 065	15 070	15 075	15 080	15 085	15 090	15 095	15 100	15 150
16	○ White	16 065	16 070	16 075	16 080	16 085	16 090	16 095	16 100	16 150
18	● Pink	18 065	18 070	18 075	18 080	18 085	18 090	18 095	18 100	18 150

PERFECTUS "PF", length 20 mm, 30 mm, 35 mm, 40 mm, 45 mm not available

### Ordering information:

PERFECTUS ASPIRATION TIP



PERFECTUS EXPLANT/TRANSPLANT TIP



Single box: 10 PCS.

**Medax Srl Unipersonale**

**Headquarters:** Via S. Pertini, 4 • 41039 • San Possidonio (MO) • Italy  
 Company direct No. : +39 0535 1812757 • Fax No : +39 0535 1812744 • email: [customercare@medax.it](mailto:customercare@medax.it) • PEC: [medax@legalmail.it](mailto:medax@legalmail.it) • [www.medax.it](http://www.medax.it)

**Registered Office:** Via R. Piva, 1/A • 46025 • Poggio Rusco (MN) • Italy  
 Vat N. /Fiscal Code N. Iscriz. Reg. Impr.: MN 02669860369 • N. REA: MN 233527, MO 403036 • Capitale Sociale Euro 100.011,00 i.v.



Medax's quality management system is certified to ISO 13485: 2016 standards.



In accordance with the requirements of the medical device 93/42/EEC directive and its relevant updates. All products undergo intensive clinical testing and are fully EC and FDA approved.