

# Sepax® 2 cell processing device

Sepax 2 is the new generation of Sepax technology for automated adult stem cell processing. This mobile, closed capability system efficiently processes umbilical cord blood, bone marrow, peripheral blood or other blood-like material, as permitted by applicable regulatory requirements. The fundamental scientific technology relies on a separation chamber that provides both separation through rotation of the syringe chamber (centrifugation) and component transfer through displacement of the syringe piston. An optical sensor measures the light absorbance of the separated components and manages the flow direction of each of them in the correct output container.



	Sepax 2 features
External design	Lightweight housing, single-handed centrifuge pit closure system
Dimensions	W: 27 cm, L: 40 cm, H: 46 cm (10.6" x 15.7" x 18.1") 16.3 kg (35.9 lbs)
User interface	Color touch screen display, intuitive Graphical User Interface
Electronics and communication	Windows XP Embedded, GMAP, USB and Ethernet
Data saving capability	32 logfiles, 50 patfiles and 50 report files (.pdf)
Core technology	Electrical motor for centrifugation, pneumatic circuitry for piston drive
Optical line sensor	Red/blue transmitted LEDs Red/blue scatters LEDs
Traceability function	Barcode reader with multiple code reading capabilities and desktop printer; Full procedure data management with .pdf report and procedure graph



## **Unique features**

- Sepax 2 is equipped with a newly-developed user interface combining touch-screen technology and
  active guidance throughout the procedure. Furthermore, a help guide is integrated to provide live
  assistance to the user in case of problem.
- Thanks to the intuitive user interface, the user can easily **monitor every step** throughout the whole procedure. Hence, the user knows at any time what is happening and what the machine is doing.
- Precise tracking of procedure data including automatic printing of a procedure report with all traceability IDs is integrated, reflecting the importance of efficient and secured traceability of each processed blood unit.
- The Ethernet connection port allows connecting the Sepax 2 to the user's network in order to provide online support through secured remote access.
- USB communication is also integrated through several ports, allowing communication with peripherals such as a barcode reader, a printer and a USB key used to store and transfer procedure files to a computer.

#### Safety and performance

Sepax 2 complies with the directive 93/42/CEE for medical devices, including the electrical safety standard IEC 60601-1. The design control activities have been performed to ensure the safety and performance of the Sepax 2.

#### Storage requirements

The Sepax 2 should only be operated on a flat, stable, horizontal and clean surface and be used in an open environment to allow sufficient ventilation. The Sepax 2 must only be used/ stored under the following environmental conditions:

	Operation	Storage and Transport
Temperature	+7°C to +27°C	0°C to 50°C
Relative humidity	30% to 75%, non-condensing	20% to 75%, non-condensing

### **Order information**

Sepax 2 Product #14000 Standard Traceability kit Product #14050 Advanced Traceability kit Product #14051

Sepax 2 is CE marked and FDA 510(k) cleared.

Head Office
Biosafe Group SA

Eysins, Switzerland T: +41 22 365 27 27 E: info@biosafe.ch North America Office
Biosafe America, Inc.

Houston, USA T: +1 713 936 2615 E: info@biosafeamerica.com Latin America office
Biosafe Latin America

São Paulo, Brazil T:+55 11 2655 1731 E: info@biosafelatinamerica.com Asia office
Biosafe Asia-Pacific Ltd.

Hong Kong T: +852 2956 7500 E: info@biosafeasiapacific.com China office
Biosafe Medical Device Int. Trading
(Shanghai) Co. Ltd
Shanghai, China

Shanghai, China T: +86 21 6137 3214 E: info@biosafechina.com