

Short instructions

Load samples

Chill 5 Rack

A Ori

B Neg

•00000 C Pos

Chill 15 Rack

A Ori

B Neg

C Pos

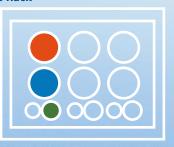


Chill 50 Rack

A Ori

B Neg

C Pos





Touch to start up

Use the touch screen to program your cell separation procedures.

Scan reagents with the 2D barcode reader





Short instructions – sample labeling and separation

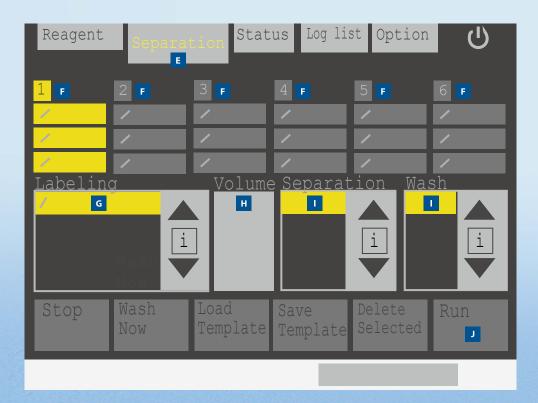
Enter reagents

- **A** Go to the **Reagent** menu and highlight a reagent rack position.
- B Press the Read Reagent button.Present a reagent vial in front of the blinking 2D code reader.
- **C** Enter up to four reagents.
- **D** Save as a template if desired.

Separation Status Log list Option Info С c С Delete Read Load Enter Stop Selected Reagent Template Template Reagent

Define the separation procedure

- **E** Go to the **Separation** menu.
- **F** Highlight one or more samples.
- **G** Select the desired **Labeling** reagent.
- **H** Touch the **Volume** submenu to enter the sample volume.
- I Select a **Separation** and a **Wash** program.
- J Place reagent vials and sample tubes on the respective racks and press **Run**.





Short instructions – maintenance

Priming

Prime the instrument after it is switched on:

- Go to the **Separation** menu and press Wash Now.
- 2 Select Rinse and press Run.

Cleaning

Before shutting down, clean the instrument:

- 1 Press the shutdown button at the upper right hand corner of the screen.
- 2 Select Yes.
- **3** Upon completion of the **Sleep** program, switch off the Instrument using the main power switch on the lower right side of the instrument.

Replace Fluid bottles

- 1 Take out an empty bottle and unscrew bottle closure counter-clockwise but do not remove it. Do not disconnect the color-coded tubing.
- 2 Place a fresh bottle into the holder, open it and fasten the bottle closure to the new bottle. Note the color-coding.

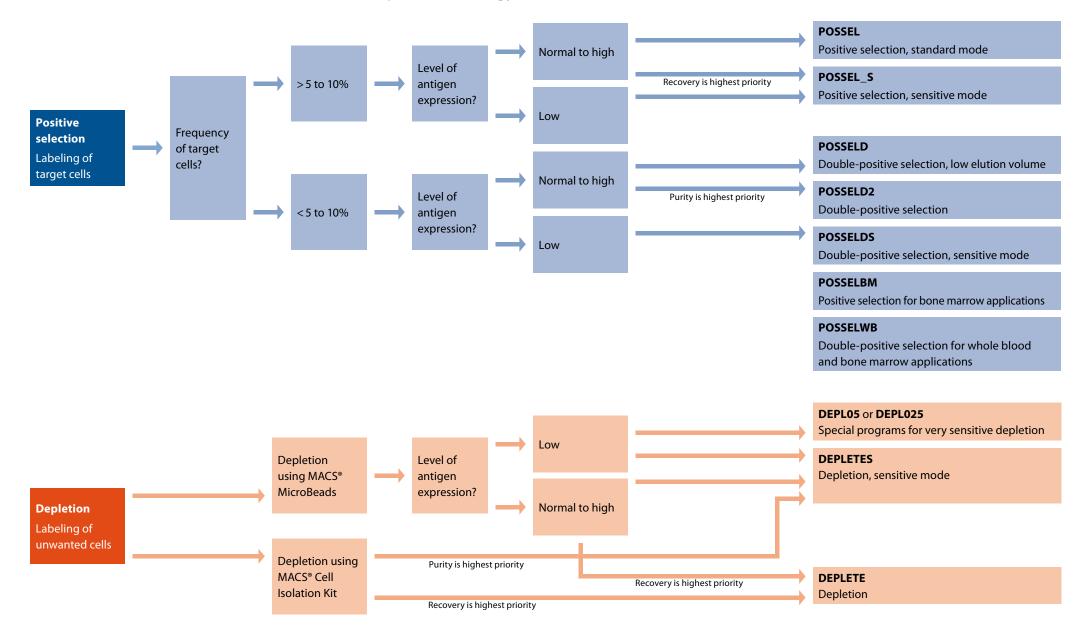


Column exchange

- 1 Open the front door.
- 2 Ensure that the fluid bottles are filled with solutions.
- 3 Go to Option > Special > Col_ex.
- **4** Press **Run**. Wait until the instrument prompts you to exchange columns.
- **5** Pull out the column using both hands.
- **6** Unscrew first bottom and then top column connector counter-clockwise.
- 7 Insert a fresh column and fasten it to the column connectors.
- **8** Press the column back into its slot until you hear a click. Repeat the whole process with column 2.
- 9 Press Done.



Short instructions – separation strategy





Short instructions – sample dilution

Cell Separation Reagent	Strategy	No. of reagents	Dilution volume	Autolabeling			
				Minimal volume*	Minimal total cell number	Maximal volume	Maximal total cell number
Chill 5 Rack ¹							
Direct MicroBeads human, rat, non-human primate	Positive selection or depletion	1	10 ⁷ cells per 80 μL	160 μL	2×10 ⁷	1600 μL	2×10 ⁸
Direct MicroBeads, mouse	Positive selection or depletion	1	10 ⁷ cells per 90 μL	180 μL	2×10 ⁷	1800 μL	2×10 ⁸
Whole Blood MicroBeads	Whole blood or bone marrow	1	Original volume	0.25 mL		1 mL	
Cell Isolation Kits	Untouched isolation	2	10 ⁷ cells per 40 μL	160 μL	4×10 ⁷	800 μL	2×10 ⁸
Cell Isolation Kits	Untouched isolation	3	10 ⁷ cells per 30 μL	120 μL	4×10 ⁷	600 μL	2×10 ⁸
Chill 15 Rack ²							
Direct MicroBeads human, rat, non-human primate	Positive selection or depletion	1	10 ⁷ cells per 80 μL	160 μL	2×10 ⁷	5200 μL	6.5×10 ⁸
Direct MicroBeads, mouse	Positive selection or depletion	1	10 ⁷ cells per 90 μL	180 μL	2×10 ⁷	5850 μL	6.5×10 ⁸
Whole Blood MicroBeads	Whole blood or bone marrow	1	Original volume	1 mL		4 mL	
Cell Isolation Kits	Untouched isolation	2	10 ⁷ cells per 40 μL	160 μL	4×10 ⁷	2600 μL	6.5×10 ⁸
Cell Isolation Kits	Untouched isolation	3	10 ⁷ cells per 30 μL	120 μL	4×10 ⁷	1950 μL	6.5×10 ⁸
Chill 50 Rack ³							
Whole Blood MicroBeads	Whole blood or bone marrow	1	Original volume	4 mL		8 mL	

 $^{^1}$ Max. number of samples: 6; min. first incubation volume: 0,2 mL; max. final labeling volume: 2 mL 2 Max. number of samples: 5; min. first incubation volume: 0,2 mL; max. final labeling volume: 6,5 mL

³ Max. number of samples: 3; min. first incubation volume: 4 mL; max. final labeling volume: 8 mL.

^{*}When working with fewer cells than the necessary minimal volume, resuspend cells in the stipulated minimal volume.



Short instructions

Chill rack specifications

Rack type and symbol	Slots	Maximal number of samples	Manual labeling	Autolabeling		
			Maximal sample volume	Minimal first incubation volume	Maximal final labeling volume	
Chill 5	24×5 mL	6 (5 mL	2.5 mL	0.2 mL	2.0 mL	
000000		tubes)		0.25 mL*	1 mL*	
Chill 15	15×15 mL 5×5 mL	5 (15 mL	12.5 mL	0.2 mL	6.5 mL	
0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0	3×3 IIIL	tubes)		1 mL*	4 mL*	
Chill 50	6×50 mL 3×15 mL	3 (50 mL	50 mL	4 mL*	8 mL*	
000	3×5 mL	tubes)				

Daily maintenance and rinsing programs

Program	Description	Recommended usage	Duration
Qrinse	Standard short rinse of separation columns and tubing system with Running Buffer	Between separations of frequent cells (>5 %)	1.5 min
Rinse	Rinse of separation columns and tubing system with Washing Solution and Running Buffer	Prior to first separation	4 min
Clean	Rinse of separation columns and tubing system with storage solution, Washing Solution, and Running Buffer	After whole blood and bone marrow applications.	7 min
Sleep	Rinse with Washing Solution followed by filling with storage solution	Before switching off the autoMACS Pro Separator	5 min

Buffer consumption

Program	Washing Solution	Running Buffer	Storage solution	MACS Bleach Solution	Time
Qrinse	-	48 mL	-	-	1.5 min
Rinse	96 mL	48 mL	-	-	4 min
Clean	96 mL	48 mL	48 mL	-	7 min
Sleep	96 mL	-	48 mL	-	5 min
Safe	96 mL	96 mL		40	21 min
Store	96 mL	-	96 mL	-	8 min
Col_ex	96 mL	96 mL	_	_	6 min

Periodic maintenance

Action	Description	Recommended usage	Duration
Column exchange using (Col_ex program)	Replacement of separation columns	Every two weeks OR after 100 separations, whichever comes first	6 min
Running the Safe program	Decontamination procedure with bleach solution (1% sodium hypochlorite)	Every 3–6 months	21 min
Cleaning the pump syringe	Cleaning of pump syringe (refer to user manual)	Every 1–3 months	
Running the Store program	Rinse with Washing Solution, followed by storage solution; replacement of columns with substitutes	Before storing the instrument for a period longer than two weeks	

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^{*}Volumes refer to whole blood samples only.