

Single Cell Analysis and Imaging core facility (SCAI-Lab) at the Centre for Applied Biomedical Research (CRBA)

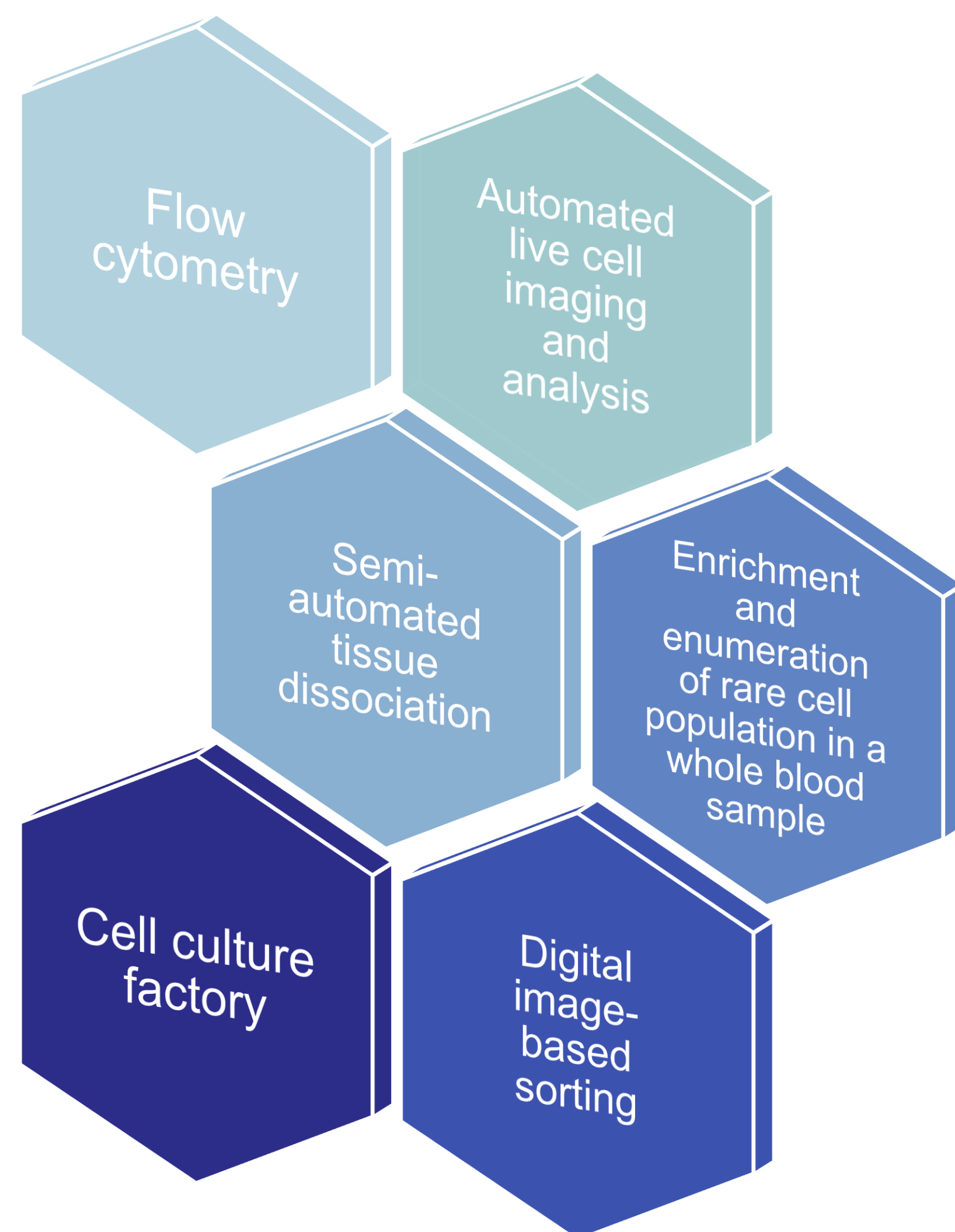


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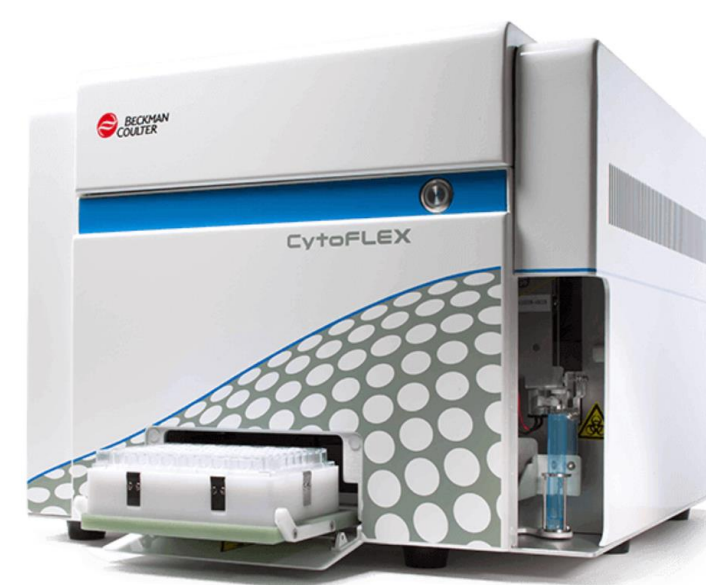
Technologies and services at SCAI-Lab

The SCAI-Lab (Single Cell Analysis and Imaging) facility was founded in 2020 and houses a range of cutting-edge instrumentation for cell study and isolation.



The **IncuCyte S3** system (Sartorius) allows to monitor and quantify the behavior of cells/spheroids/organoids directly in culture in real time.

The **CytoFLEX** Flow Cytometer (Beckman Coulter) guarantees high performances and provides superior sensitivity and resolution, that enable the detection of nanoparticles.



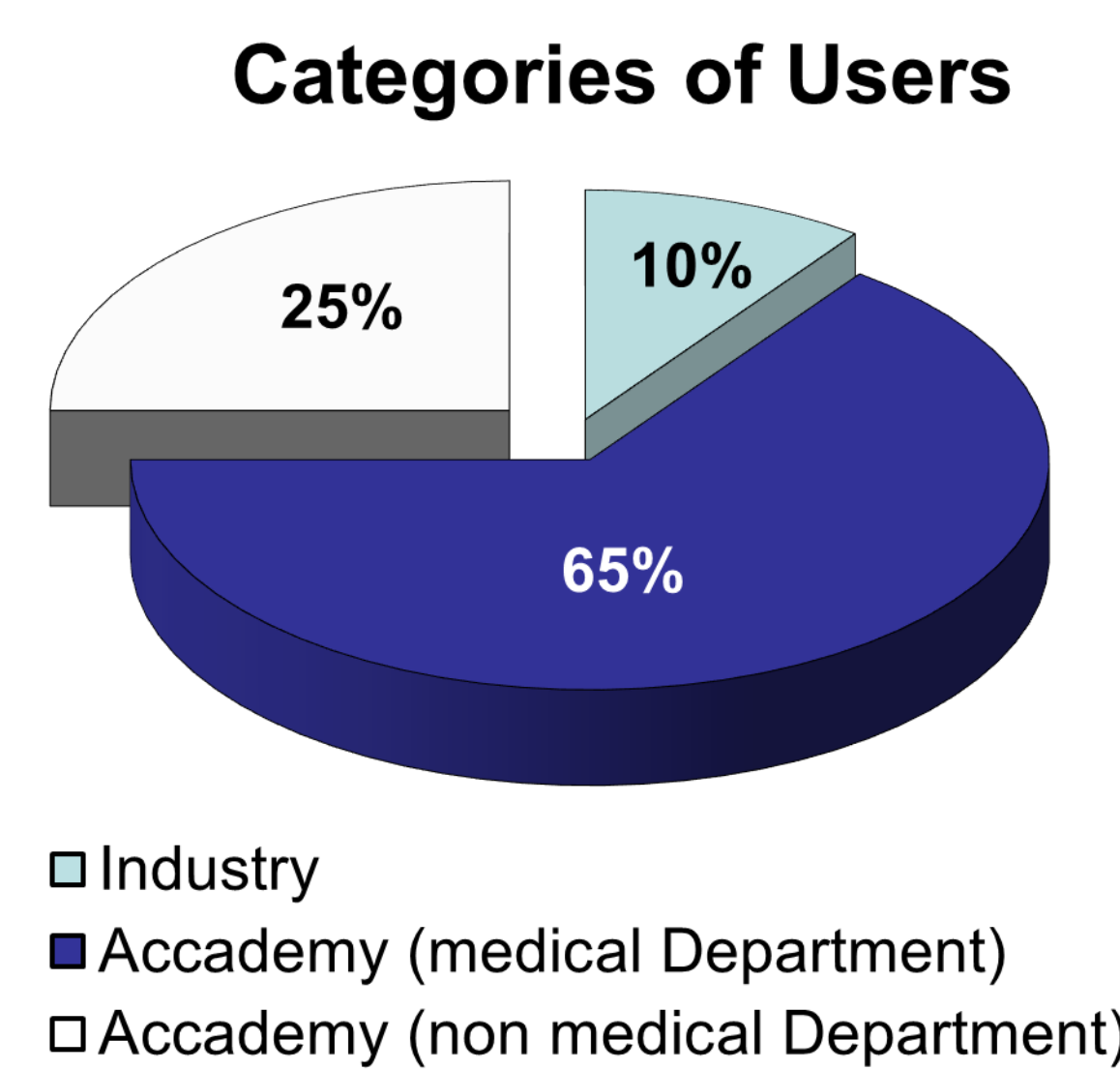
The **CellSearch** platform (Menarini Silicon Biosystems) allows the enrichment and enumeration of circulating tumor cells (CTCs) and circulating endothelial cells (CEC) from whole blood.

The **DEPArray NxT** (Menarini Silicon Biosystems) digital image-based sorter, guarantees, with a high degree of purity, the recovery of the single cell of interest.



The **gentleMACS Octo Dissociator** with Heaters (Miltenyi) offers a fully automated workflow for tissue dissociation.

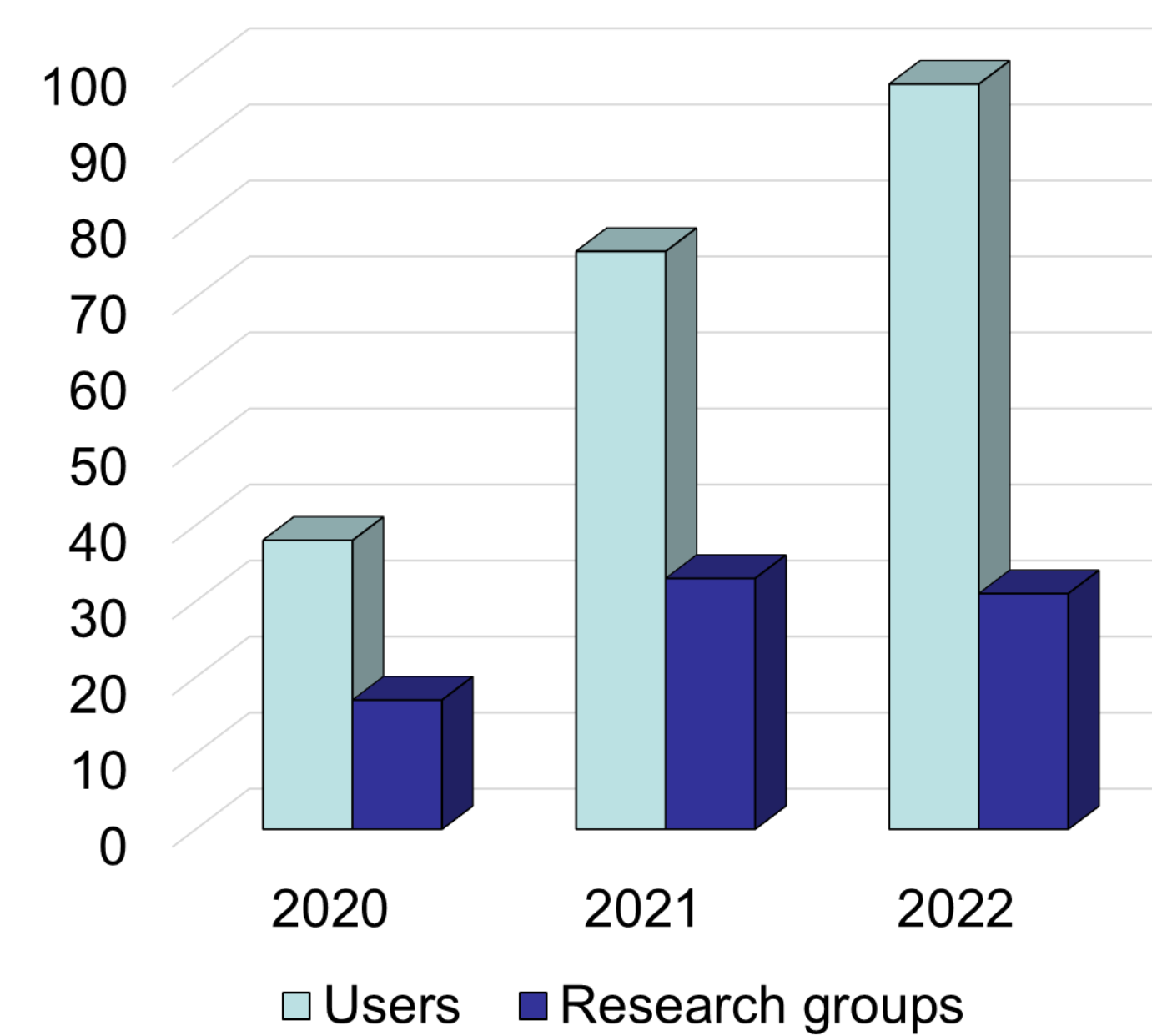
Performance analysis and assessment



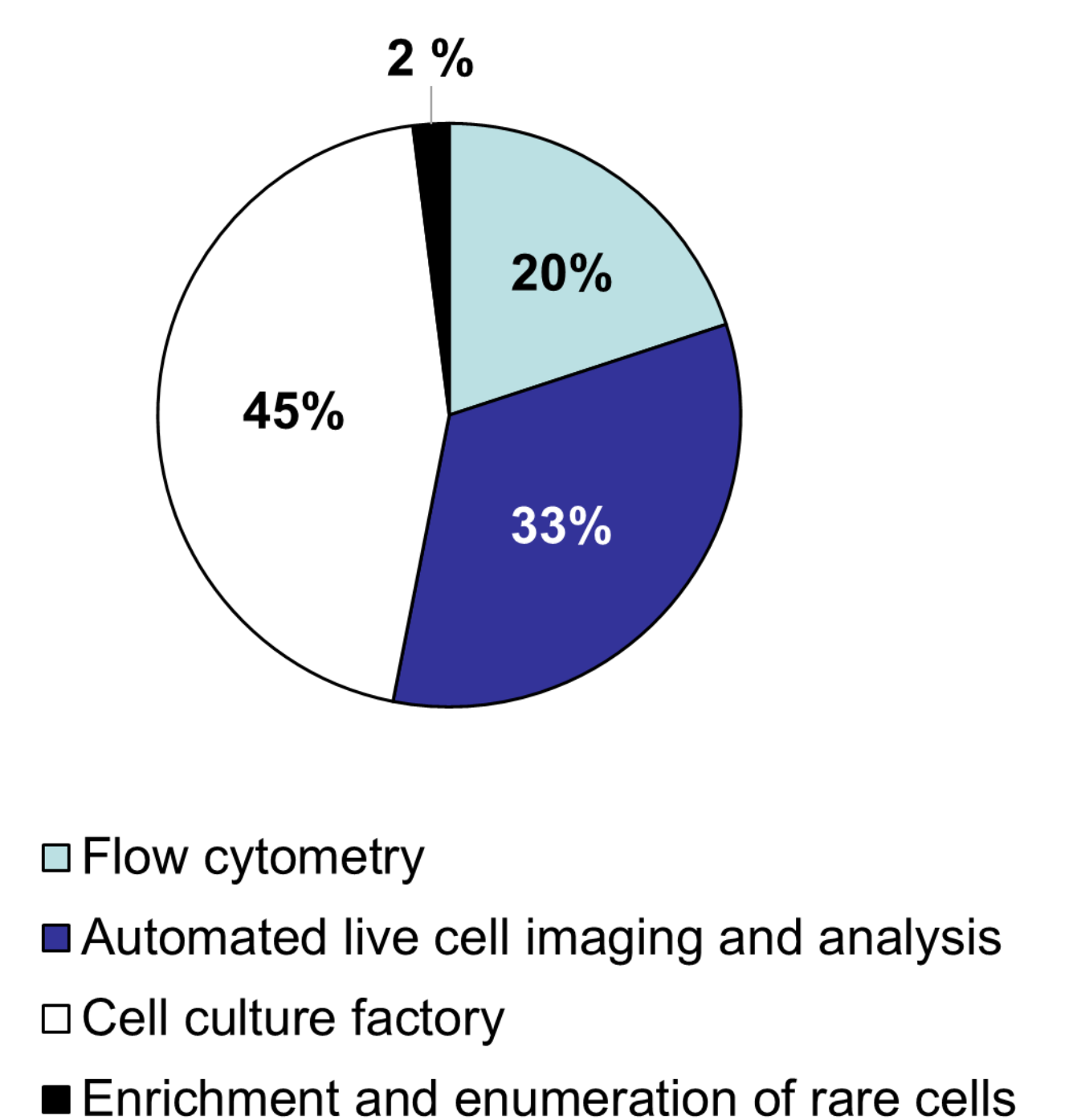
The SCAI-Lab three-year assessment shows the Medical Department as the predominant user affiliation. Among non-medical Departments, the facility welcomes a range of different disciplines.

The increasing trend shown by the number of users and research groups over time suggests an expansion of SCAI-Lab.

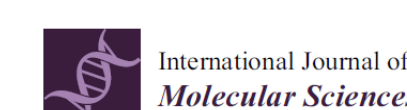
User trend with time



User distribution among technologies



Key publications



Circulating Endothelial Cells: A New Possible Marker of Endothelial Damage in Kawasaki Disease, Multisystem Inflammatory Syndrome in Children and Acute SARS-CoV-2 Infection

Marianna Fabi¹, Biljana Petrovic^{2,3}, Laura Andreozzi^{1,*}, Elena Corinaldesi⁴, Emanuele Filice⁵, Carlotta Biagi¹, Alessia Rizzello⁶, Bianca Elisa Mattesini⁶, Simone Bugani^{2,3} and Marcello Lanari¹



Table 4. CEC values during acute and subsacute stages, in the three groups of patients.

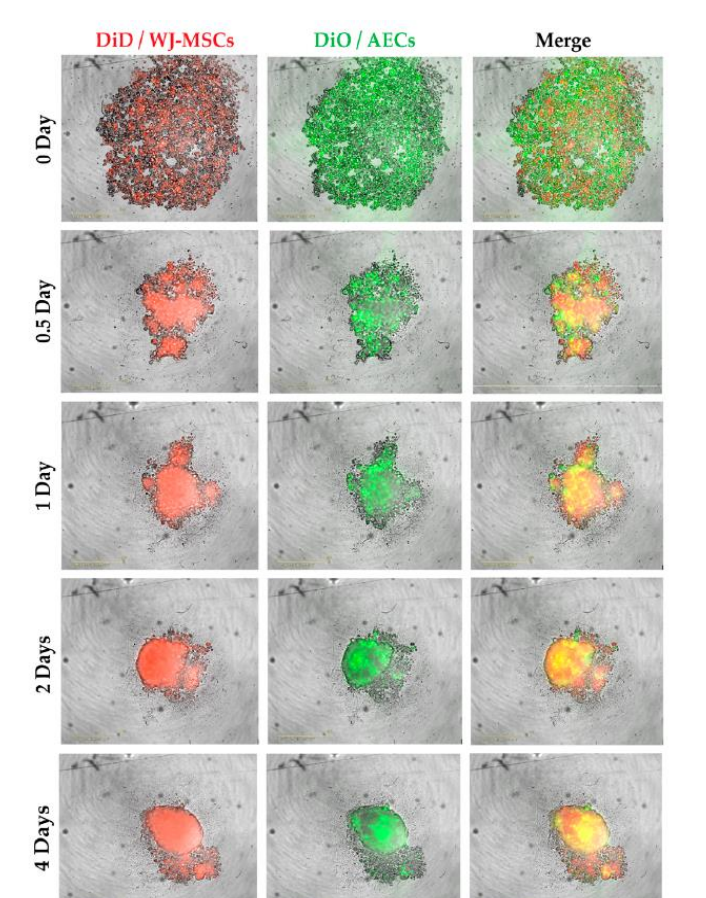
Acute Phase	KD (n = 9)	MIS-C (n = 20)	AC19 (n = 10)	p
CECs num./mL, median (IQR)	16.3 (13.6–48.8) *	5 (4–15.5) *	27.1 (9.3–101.7)	0.042
CECs > mv, n (%)	6/6 (100%) *	5/19 (26.3%) †‡	7/10 (70%) †	0.003
Synocytia, n (%)	3/6 (50%) *	3/19 (15.8%)	3/10 (30%)	n.s.
Subacute phase				
CECs num./mL, median (IQR)	45.8 (18.5–131.0) *	3.6 (1.8–21.6) *	-	0.01
Synocytia, n (%)	4/9 (44.4%)	3/19 (15.7%)	-	n.s.
CECs > mv, n (%)	7/9 (77.8%) *	6/19 (30%) *	-	0.046

KD stands for Kawasaki Disease; MIS-C, stands for Multisystem Inflammatory Syndrome in Children; AC19 stands for acute COVID-19; CECs stands for circulating endothelial cells; * stands for statistically significant difference between KD and MIS-C; † stands for statistically significant difference between MIS-C and AC19; n.s. stands for not significant.



Characterization of Perinatal Stem Cell Spheroids for the Development of Cell Therapy Strategy

Francesca Paris¹, Pasquale Marrazzo^{1,*}, Valeria Pizzuti¹, Cosetta Marchionni¹, Maura Rossi², Martina Michelotti¹, Biljana Petrovic^{2,3}, Elisabetta Ciani⁴, Giuliana Simonazzi^{2,5}, Andrea Pession⁶, Laura Bonsi^{1,*} and Francesco Alviano¹



1. Circulating Endothelial Cells: A New Possible Marker of Endothelial Damage in Kawasaki Disease, Multisystem Inflammatory Syndrome in Children and Acute SARS-CoV-2 Infection. Fabi M, Petrovic B, Andreozzi L, Corinaldesi E, Filice E, Biagi C, Rizzello A, Mattesini BE, Bugani S, Lanari M. International Journal of Molecular Science, 2022 September.
2. Characterization of Perinatal Stem Cells Spheroids for the Development of Cell Therapy Strategy against Diabetes. Paris F, Marrazzo P, Pizzuti V, Marchionni C, Rossi M, Michelotti M, Petrovic B, Ciani E, Simonazzi G, Pession A, Bonsi L and Alviano F. Bioengineering, 2023, February.

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