



BD FACSymphony™ A1 Cell Analyzer

Premium performance in a benchtop footprint





BD FACSymphony™ A1 Cell Analyzer features:

- Premium high-end BD FACSymphony™ instrument technology scaled to fit on your benchtop
- Flexibility to meet a broad spectrum of research needs from **small particle research** to **16-color immunophenotyping**
- Industry standard **BD FACSDiva™ Software** for streamlined workflow from system setup to data acquisition and analysis

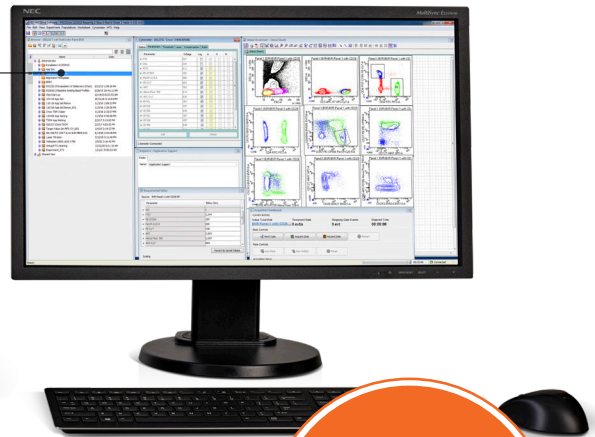
BD® Small Particle Detector

Optional detector for small particle side scatter (SP SSC) for resolving particles as small as 90 nm



Familiar Workflow

Industry standard BD FACSDiva™ and FlowJo™ Software enable easy system QC, data acquisition and analysis



Trusted Partner
Supported by
over 45 years of
flow cytometry
expertise

BD FACSymphony™ Instrument Technology

Reduce background noise and increase sensitivity with low-noise electronics, high-powered lasers and tight beam spot, shared across the premium BD FACSymphony™ platforms. The BD FACSymphony™ A1 Analyzer is compatible with BD Horizon™ Dyes and supports up to 16 colors or 19 parameters simultaneously.



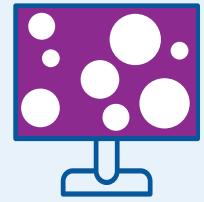
Premium BD FACSymphony™ instrument technology

delivered in a compact size

Up to

16 fluorochromes and 19 parameters

to conduct deep and broad phenotyping

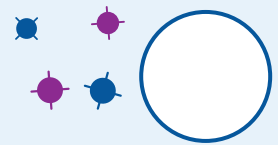


Enhance detection sensitivity

with four high-powered 100 mW lasers: Violet (405 nm), Blue (488 nm), Yellow-Green (561 nm) and Red (637 nm)

BD® Small Particle Detector Option for analysis of small particles

such as extracellular vesicles including exosomes



Identify and analyze rare cell types and events

with our redesigned optics including small beam spots combined with low-noise electronics



Gain rich scientific insights

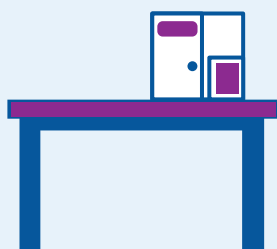
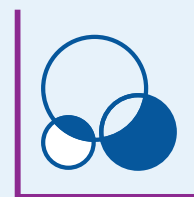
by leveraging BD Horizon Brilliant™ Reagents

Enable easy system QC

using industry-standard BD FACSDiva™ Software and BD® CS&T Beads



Utilizes FlowJo™ Software, the leading bioinformatics platform* for flow cytometry analysis



Ideal for labs with limited space

Small footprint (58 x 61 x 59 cm)

Automated sample processing in high-throughput mode

using the BD® High-Throughput Sampler Option



*In 2020, FlowJo™ Software was cited in leading immunology peer-reviewed journals more often than any other flow cytometry analysis software.

Able to detect up to 16 colors and resolve rare cell subsets

Table 1. Instrument configuration and reagents in the cytotoxic immune cells panel

Laser	Filter	Fluorochrome	Specificity
Violet 405 nm	450/50	BV421	Perforin
	525/50	BV480	CD159a (NKG2A)
	610/20	BV605	CD19
			CD14
			CD123
			CD141
		FV5575V	-
670/30	BV650	CD3	
710/50	BV711	CD314 (NKG2D)	
780/60	BV786	HLA-DR	
Blue 488 nm	530/30	FITC	CD57
	710/50	PerCP-Cy5.5	CD8
Yellow-Green 561 nm	586/15	PE	CD158 (KIRs)
	610/20	PE-CF594	CD56
	670/30	PE-Cy5	CD95 (Fas)
	710/50	PE-Cy5.5	CD127 (IL7R- α)
780/60	PE-Cy7	CD38	
Red 637 nm	670/30	AF647	Granzyme K
	710/50	R718	Granzyme B
	780/60	APC-H7	CD16 (Fc γ RIII)

BV, BD Horizon Brilliant Violet™; FVS, BD Horizon™ Fixable Viability Stain; AF, Alexa Fluor™

Figure 1

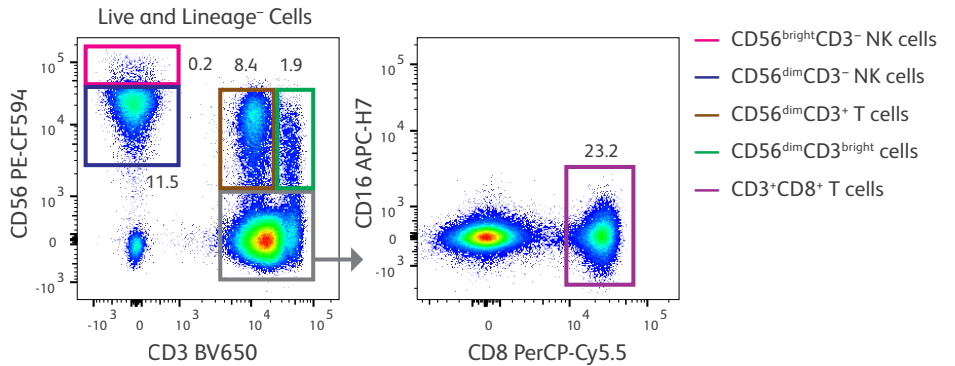


Figure 1. Identification of cytotoxic immune cell populations in healthy human peripheral blood
Within live and lineage negative cells, analysis of CD56 versus CD3 revealed various cell populations that were color coded as cytokine-producing NK cells (pink), cytotoxic NK cells (blue), CD56⁺ T cells containing NKT cells (brown), CD56⁺ T cells containing $\gamma\delta$ T cells (green) and cytotoxic CD8⁺ T cells (purple).



To learn more, download the panel sheet *Characterization of Cytotoxic Immune Cells in Human Peripheral Whole Blood* from bdbiosciences.com

Figure 2A

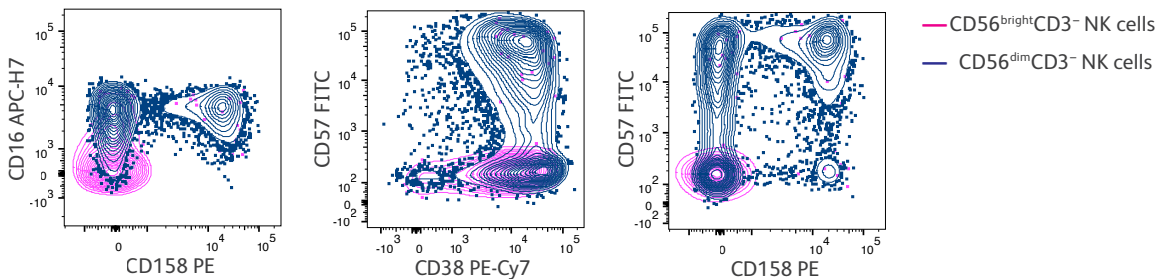


Figure 2B

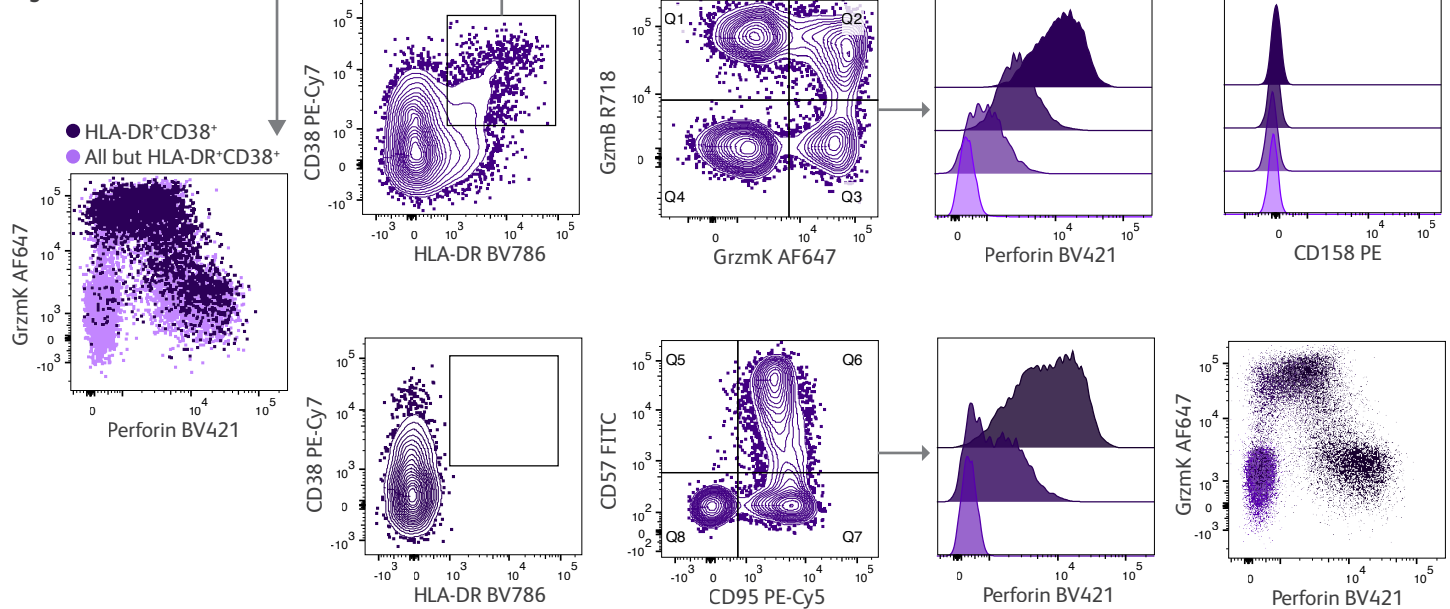
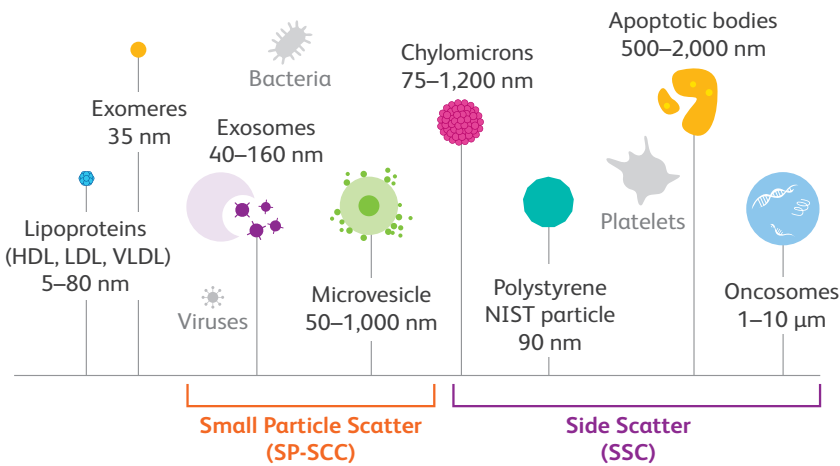
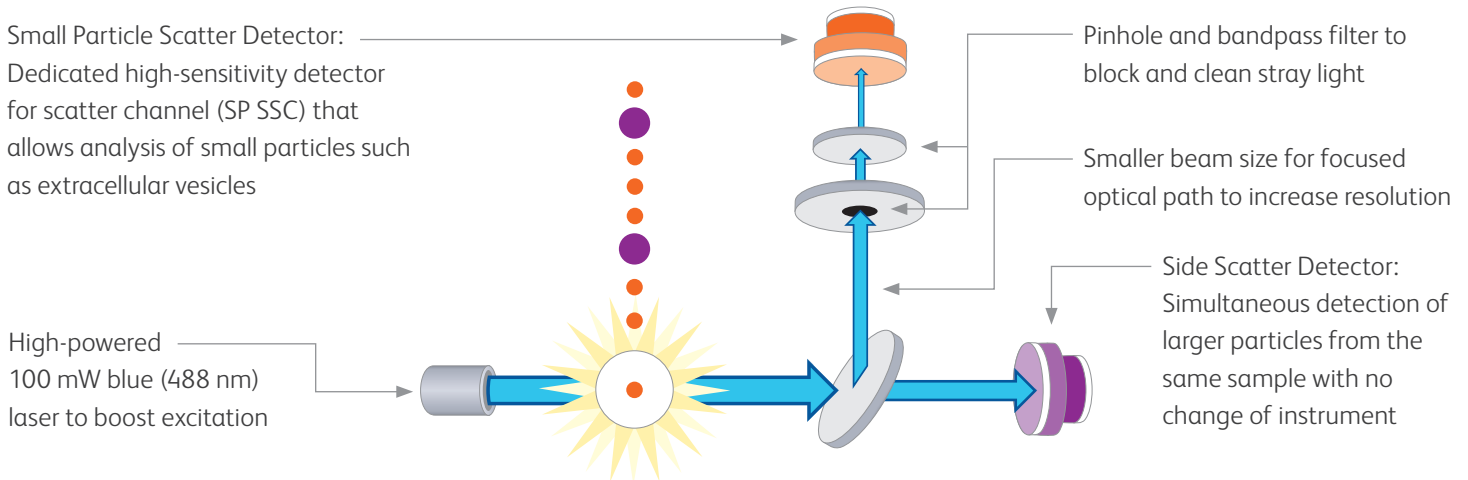


Figure 2. Phenotyping of circulating cytotoxic cells using a 16-color panel

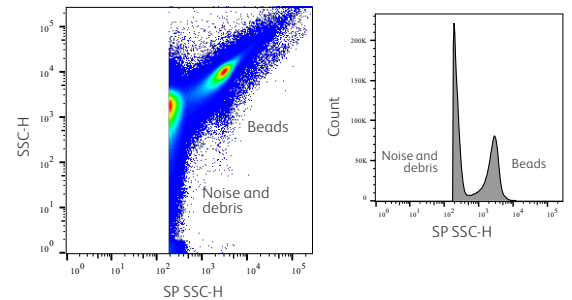
The plots represent the analysis of cytolytic proteins in combination with various cell differentiation markers, enabling a deeper characterization of the cell populations gated in Figure 1. **A.** Overlay of NK cell subsets. **B.** Identification of activated CD8 T cells based on the expression of CD38 and HLA-DR. The HLA-DR FMO staining helped to determine the gating boundaries for proper detection of the double positive cells.

Simultaneous detection of large (SSC) and small (SP SSC) particles

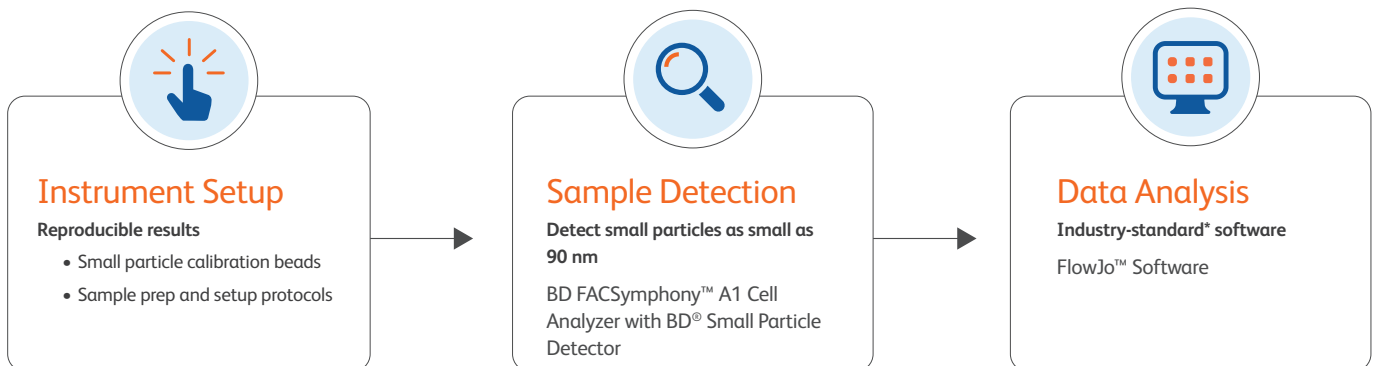
The BD FACSymphony™ A1 Cell Analyzer with optional BD® Small Particle Detector is able to resolve scatter of small particles such as extracellular vesicles, viral particle, exosomes and more.



Resolution of 90-nm polystyrene particles with the BD® Small Particle Detector option



Seamless small particle detection workflow



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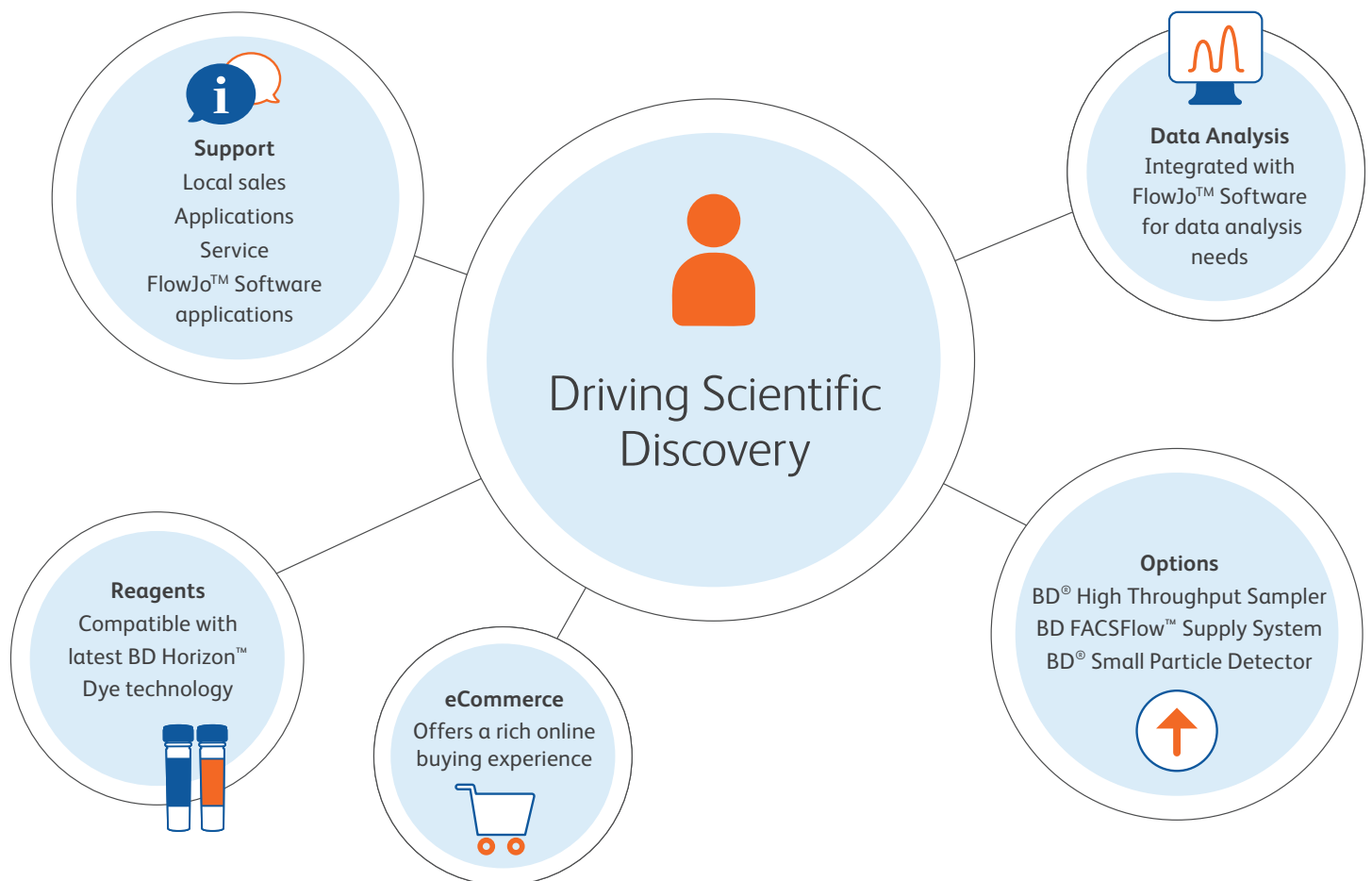
BD FACSymphony™ Systems



	BD FACSymphony™ A1	BD FACSymphony™ A3	BD FACSymphony™ A5	BD FACSymphony™ S6
Number of lasers	4	5	5–9	5–9
Fluorescent detectors	16	Up to 28	Up to 48	Up to 58
Instrument type	Analyzer	Analyzer	Analyzer	Sorter
Software	BD FACSDiva™	BD FACSDiva™	BD FACSDiva™	BD FACSDiva™
Footprint	58 x 61 cm	83.8 x 76.2 cm	101.6 x 78.7 cm	101.6 x 78.7 cm
Small particle detector	Yes	No	No	No

Backed and Supported by BD

We're committed to partnering with you to provide the mission-critical tools and support you need to advance your research.



Class 1 Laser Product.
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