

# Checkliste

## How to Select the Right Serum for Your Cell Culture Application

### 1. Define Your Application

- ☐ Which cell type am I using?
- ☐ Primary cells, stem cells, viral production lines, or stable cell lines?
- ☐ Is the system immune-sensitive (PBMC, NK, T-cells)?
- ☐ Application area: diagnostics, virology, bioprocessing, basic research?

Rule of thumb: The more human-relevant the model, the stronger the case for Human Serum.

### 2. Choose Between FBS and Human Serum

- ☐ FBS for broad compatibility and routine growth
- ☐ Human Serum for immunology, PBMC, HEK293, AAV, diagnostic assays
- ☐ Do I require species matching?

Ask yourself:

- Do I need maximum reproducibility?
- Are immunoassays or diagnostic readouts involved?
- Is xenogeneic interference a risk?

### 3. Evaluate Endotoxin Requirements

- ☐ Is the application endotoxin-sensitive?  
(PBMCs, viral titers, cytokine assays, immunoassays)
- ☐ Do I require Low Endotoxin (<5 EU/mL) or Ultra-Low?
- ☐ Are complete endotoxin test results available?

If YES: choose Low or Ultra-Low Endotoxin serum.

### 4. Check Documentation & Traceability

- ☐ Country of origin clearly defined
- ☐ Viral testing panel available
- ☐ Batch CoA + QC data provided
- ☐ Full traceability ensured
- ☐ cGMP or equivalent standards met

#### **Warning:**

Missing documentation = risk for publications, audits, and QC deviations.

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## 5. Select Origin Based on Regulatory Needs

- ☐ US origin
- ☐ Australian origin
- ☐ South American origin
- ☐ EU origin (human)

Tip: Align regulatory requirements early.

## 6. Verify Processing Method

- ☐ Sterile filtered (0.1 µm / 0.2 µm)
- ☐ Gamma irradiated
- ☐ Heat inactivated
- ☐ Dialyzed
- ☐ IgG-stripped
- ☐ Charcoal-treated

Rule: More sensitive applications require higher processing control.

## 7. Perform Lot Qualification

- ☐ Lot sample requested
- ☐ Compared with current serum
- ☐ Cell proliferation tested
- ☐ Performance validated in target application
- ☐ Media and buffer compatibility confirmed

Best practice: Use only the lot that was tested and approved.

## 8. Plan for Reproducibility & Supply Stability

- ☐ Lot reservation possible?
- ☐ Required duration (3–12 months)?
- ☐ Storage conditions ensured (–20°C to RT)?
- ☐ Quantity and forecast planned

## 9. Review Critical QC Parameters

- ☐ Endotoxin
- ☐ Osmolality
- ☐ pH
- ☐ Protein concentration
- ☐ Sterility
- ☐ Cell-growth performance

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## 10. Determine Batch Configuration

- ☐ Single donor
- ☐ Pooled batch
- ☐ Human Type AB
- ☐ Human mixed donor

## 11. Verify Ethical & Regulatory Compliance

- ☐ Human donations from certified EU/US institutions
- ☐ Animal sera ethically sourced under global guidelines
- ☐ Transparent, documented origin

## 12. Confirm Logistics & Availability

- ☐ Full cold-chain compliance documented
- ☐ Realistic delivery timelines
- ☐ Lot reservation without prepayment possible
- ☐ Free testing volumes available?
- ☐ Import, export & customs clearances prepared

## Why SeamlessBio?

**SeamlessBio** provides reliable, fully documented and reproducible biological materials – from serum to specialty reagents.

We support your research with certified quality, stable supply chains and secure batch reservations.

### Advantages:

- Test samples & reservation for 6 weeks
- Fully documented (CoA, Health Certificate, EMEA available)
- Flexible time for order after fixing the contract
- Best choice for the buyer