



Sample Evaluation Checklist

Systematic protocol — 20 criteria across 4 evaluation dimensions

Complete within 7 days of receipt · Score 1 (Fail) to 5 (Excellent) per criterion

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HOW TO USE: Click any score box (1–5) per criterion. Add notes in gray fields. Save and email back.

Score Guide: 1 = Fail (does not meet spec) 2 = Poor (significant concern) 3 = Pass (meets minimum)

4 = Good (meets expectation) 5 = Excellent (exceeds expectation)

Total Score: 80%+ = Approve 60–79% = Conditional <60% = Reject (Max 95 points, 19 scored criteria)

– SECTION 1 / 4

Sample Identification & Receipt Information

NuCoreBio Order / Sample Reference

Date Sample Received

e.g. 2026-03-15

Product Name / Formula

Batch / Lot Number (from COA)

Must match label

Dosage Form

Capsule / Powder / Tablet / Gummy

Net / Fill Weight per Unit

e.g. 500mg/capsule

Your Company / Brand Name

Evaluator Full Name

Person completing this form

Packaging / Shipping Condition on Arrival:

Excellent — intact, undamaged

Good — minor cosmetic issues

Concern — describe in notes

Damaged — describe in notes

— SECTION 2 / 4

Physical & Organoleptic Evaluation

Evaluation Criterion	1-Fail	2-Poor	3-Pass	4-Good	5-Excel	Notes / Specific Obs
2.01 Appearance — Color & Surface Quality <i>Color uniformity, no mottling, correct size/shape/color for dosage form</i>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
2.02 Fill Weight Consistency <i>Weigh 10 units. Each within ±5% of stated fill weight. CV <3% ideal.</i>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
2.03 Odor / Aroma Profile <i>Characteristic of active ingredients; absent: rancidity, off-notes, chemical odors</i>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
2.04 Capsule / Package Seal Integrity <i>No cracks, leaking, splitting; band intact; blister seal unbroken</i>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
2.05 Texture / Dissolution (Powders/Gummies) <i>Particle size uniform; powder dissolves easily; gummy texture appropriate</i>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
2.06 Disintegration Time <i>Capsule disintegrates <30 min in 37°C water per USP <701>. Record actual time</i>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
2.07 Supplement Facts Panel Accuracy <i>Every ingredient, dose, and unit matches COA. % Daily Values correct.</i>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
2.08 Packaging Information Completeness <i>Lot number, mfg date, expiry, directions, warnings, contact info all present</i>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	

— SECTION 3 / 4

Analytical Documentation Review (COA Verification)

Review the Certificate of Analysis and all supporting documents provided with this sample.

Evaluation Criterion	1-Fail	2-Poor	3-Pass	4-Good	5-Excel	Notes / Specific Obs
3.01 COA from ISO/IEC 17025 Accredited Lab <i>Lab accreditation status. Supplier-only COA = lower confidence. 3rd-party preferred</i>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
3.02 Active Marker Potency — within ±5% of label claim <i>e.g. "Withanolides 5.2%" vs stated "≥5%". HPLC method stated.</i>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
3.03 Heavy Metals — all 4 within USP <232> limits <i>Pb ≤1ppm, Cd ≤0.3ppm, As ≤1ppm, Hg ≤0.1ppm. ICP-MS preferred.</i>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
3.04 Microbial Panel — complete and passing <i>TPC, TYMC, E. coli (absence/g), Salmonella (absence/25g), S. aureus (absence/25g)</i>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
3.05 Pesticide Residues — screened and within limits <i>EU MRL or USP <561>. GC-MS or LC-MS multi-residue. Method and limits stated</i>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
3.06 Batch / Lot Number Cross-Reference Consistent <i>COA lot = label lot = shipping document. Any mismatch = document integrity issue</i>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
3.07 Dates Consistent (mfg date + shelf life = expiry) <i>Mfg date + shelf life claimed = expiry date. Logical and consistent.</i>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	

