



### Cannabis Analysis Submission Form

Company Name:		<b>Internal Use Only</b>
Address:		Date Received:
City:	Zip:	Received By:
Contact Name:		Total Submission UID # Range:
Contact Email:		
Phone:	Date:	Notes:
Billing Email:		

**RGA Submission Agreement (required for all submissions)**

**As per 16.8.2 NMAC:** By signing below, I attest I have reviewed this sheet, and I am either:

- (1) A person authorized to collect samples for a licensed cannabis establishment and have adhered to the "Self-Sampling Guidance for Testing" (<https://www.rld.nm.gov/wp-content/uploads/2022/06/RLD-Mandatory-Self-Sampling-Guidance.pdf>) issued by the CCD during the collection of these samples; or
- (2) That I am over 21 years of age, cannot use any Certificates of Analysis generated for sale or transfer, and I will receive unofficial results denoted "UNOFFICIAL RESULTS. NOT FOR TRANSFER OR SALE." for any products submitted for R&D or personal consumer purposes.

Signature: \_\_\_\_\_ Date: \_\_\_\_\_

**Please note:** We require six grams of dried flower for full panel tests. Two grams are required for flower potency and terpene profiling, residual pesticides, or microbial screening. Three grams of concentrate is required for full panel tests, and half a gram is required for individual potency and terpene profiling, microbial, or residual solvents tests, while one gram is required for residual pesticides.

**Testing Prices (per sample):**

CCD Full Panel Certification for Flower & Trim (HPLC potency, GC terpenes, moisture, microbials, residual pesticides)	\$200
CCD Full Panel Certification for Concentrates (HPLC potency, GC terpenes, microbials, and residual pesticides; <b>if needed add \$50 for residual solvents, box must be checked</b> )	\$200
Potency & Terpene Profiling Only (13 cannabinoids and 19 terpenes reported)	\$60
Residual Pesticide Analysis (21 pesticides quantified)	\$75
Residual Solvent Analysis (23 residual solvents quantified)	\$50
Microbial Panel Only (6 microbes pass/fail via qPCR)	\$60

Check box for each test needed				
CCD Full Panel via HPLC	Potency & Terpene Profiling Only	Residual Pesticide Analysis	Residual Solvent Analysis	Microbial Panel Only
<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
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UID #: <small>Internal Use Only</small>	Sample Name <small>(Print exactly as to appear on CoA)</small>	Form <small>(flower, oil, etc.)</small>	BioTrack ID#:

Rio Grande Analytics LLC will make every effort to provide accurate analysis of samples received, with liability limited to the cost of analysis. For reasonable cause, tests will be repeated. No other warranties, expressed or implied, are given. Prices subject to change without notice. Visit [www.riograndeanalytics.com](http://www.riograndeanalytics.com) for current pricing.

# From NMAC 16.8.7, Rules Currently in Effect as of 5/6/24

## Required Testing of Cannabis Products

**16.8.7.15 REQUIRED TESTING OF CANNABIS PRODUCTS:** A cannabis establishment shall segregate a batch of cannabis product and arrange for samples to be collected and tested by a cannabis testing laboratory if required by this section. The batch must pass all required tests prior to the sale or delivery to a qualified patient, primary caregiver or consumer.

**A. Required testing:** Unless an exception applies:

(1) A cannabis producer, cannabis producer microbusiness, vertically integrated cannabis establishment, or integrated cannabis microbusiness shall arrange for and pay for the testing specified in Table 1, *Required Testing of Cannabis Products*, below, of any cannabis flower and trim that it harvests prior to:

- (a) packaging for retail sale;
- (b) transfer to another cannabis establishment for the purposes of retail sale;
- (c) retail sale; or
- (d) delivery to a patient or consumer.

(2) A cannabis manufacturer, vertically integrated cannabis establishment, or integrated cannabis microbusiness shall arrange for and pay for the testing specified in Table 1 of any cannabis product, including but not limited to a concentrate or extract, that it manufactures prior to:

- (a) packaging for retail sale
- (b) transfer to another cannabis establishment for the purposes or retail sale;
- (c) retail sale; or
- (d) delivery to a qualified patient, primary caregiver or consumer.

(3) A cannabis retailer, vertically integrated cannabis establishment, or integrated cannabis microbusiness shall not sell or deliver to a patient or consumer any cannabis product unless the cannabis product has undergone all testing required by this section.

Product category	Potency	Homogeneity of Batch	Visual Inspection	Microbiological	Residual Pesticides	Residual Solvents
Flower	X		X	X	X	
Trim	X		X	X	X	
Concentrate (volatile solvent)	X			X	X	X
Kief	X		X	X	X	
Pre-rolls	X			X	X	
Concentrate (non-volatile solvent)	X		X	X	X	
Extract – alcohol	X			X	X	
Extract – other liquid	X			X	X	
Topical	X			X		
Edible	X			X	*	
Other inhalable	X				*	X
Other	X			X	*	X

\*Pesticide testing required unless exempted by Subsection E, below.

**E. Exceptions to required testing:**

(1) A cannabis establishment shall not be required to have tested for pesticide residue any cannabis product made from cannabis concentrate or cannabis extract with verified pesticide residue test results, so long as the establishment can demonstrate that the resulting product will not exceed action levels for that type of cannabis product.