Ohio Administrative Code

Rule 901:14-2-15 Laboratory reporting.

Effective: January 31, 2020

(A) A licensed processor shall ensure that the testing laboratory selected create a unique certificate of analysis for each batch or lot tested on its behalf, which shall include, at minimum:

(1) The name and address of the testing laboratory where the analysis was performed;

(2) The ISO certification number of the testing laboratory;

(3) The name of the processor from whom the sample was received;

(4) The address of the processing facility where the sample was processed;

(5) The name of the product tested;

(6) A unique batch or lot number that will match the hemp product, in order to facilitate any warnings or recalls the department deems appropriate;

(7) The date or dates on which each test was performed;

(8) The date or dates the sample was collected;

(9) The cannabinoid profile of the sample, including the percentage content by weight for, at minimum:

(a) Delta-9-tetrahydrocannabinol (THC);

(b) Delta-9-tetrahydrocannabinolic acid (THCA);

(c) Cannabidiol (CBD);

(d) Cannabidiolic acid (CBDA); and

(e) All other cannabinoids listed on the product label.

(10) Results of analysis for microbial contamination, as applicable;

(11) Quantitative results of analysis for heavy metal contamination, as applicable;

(12) Quantitative results of analysis for mycotoxins, as applicable;

(13) Quantitative results of analysis for residual solvents, as applicable; and

(14) The signature of the laboratory manager or scientific director certifying the analysis.

(B) The certificate of analysis may contain the following:

(1) Results of quantitative analysis of additional cannabinoids for which the laboratory is able to obtain a standard for comparison; and

(2) Results of quantitative analysis of terpenes for which the laboratory is able to obtain a standard for comparison.