

Item Approval Application

The Division of Cannabis Regulation (DCR) requires a full and complete Item Approval Application submission for items to be reviewed for compliance with 19 CSR 100-1.120(2). Licensees are required to submit in Metrc and follow-up with the additional required information through email at CannabisProductCompliance@health.mo.gov. If all required information is not submitted the same day, the application may be denied as an incomplete submission. An Item Approval Application is required for new Template, Approved Template-based, and Non Template-based item submissions.

Licensee Number:						
Licensee Name:						
Select one based on the type	of item(s):					
New Template	Item Name:					
(Skip to Section 2)						
Non Template-based	Item Name:					
(Skip to Section 2)						
Approved Template-base	ed					
Approved Template Nam						
Approved lemplate Nam	ie.					
Approved Template Num	nber:					
 TEMPLATE-BASED ITEM SUBMISSION (Section reserved for Approved Template-based items only) Instructions: A licensee will use this section when submitting final marijuana product(s) based on a previously approved template item. 						
		ation from the approved template item to the new template-based n, flavor and/or ingredients(s).				
Instructions: If Licensee <u>has</u> up	loaded a photo of the p	PHOTO (Section reserved for all item submissions) hysical packaging into Metrc, see 2A. If Licensee has not see 2B. Only select one option below.				
2A . Check here if the lice within Metrc.	ensee <u>has</u> uploaded a _l	photo of the physical packaging in the Packaging Photo field				
		upload in to Metrc a PDF or JPEG file of the approved final ng the approved final marijuana product to a dispensary				

3.	REQUEST Fitems only)		NATIVE LABEL PLACEMENT (Section reserved for New Template and Non Template-based			
		see may request approval for alternative placement of "Marijuana", the universal symbol, ompleting this application section.				
	For which	labeling re	quirement(s) is the Licensee requesting alternative placement? (Check all that apply):			
	"Marijı	uana"				
	Univer					
	Label r	equired by	Rule			
	Provide de	etail about	the alternative placement. Details must include:			
	•	Why the	icensee cannot meet the Rule requirements:			
	•	How the	abel placement will differ from Rule:			
	•	How the product:	proposed alternative label placement maintains and promotes safe use of the marijuana			
	•					
			censee attests the alternative label placement cannot be easily removed or separated g and will meet all other requirements as put forth in Rule.			
4.	IDENTIFIC	ATION OF I	OGOS OR SYMBOLS (Section reserved for New Template and Non Template-based items			
	only)					
	<u>Instructions</u> : A licensee will use this secon to identify all logos and/or symbols on the package's design by uploading a PDF or JPEG file for each logo and/or symbol utilized and providing the file name. Identify the logo or symbol file names uploaded with the application.					
	•	Logo 1:				
	•	Logo 2:				

5.	IDENTIFICATION OF SIDE EFFECTS OR BEHAVIORAL EFFECTS OF USAGE (Section reserved for New Template and				
	Non Template-based items only) Instructions: A licensee will use this section to identify any text indicating side effects or behavioral effects of usage.				
	• Text:				
	Attestation: The licensee attests the text indicating side effects or behavioral effect of usage will not obstruct any other required information on the packaging and labeling; will not make any unfounded nutritional, health, structure/function, and/or implied claims; is not a slogan or statement of aspirational or ambiguous meaning; and will meet all other requirements as put forth in Rule.				
6.	IDENTIFICATION OF QR CODE (Section reserved for New Template and Non Template-based items only)				
	<u>Instructions</u> : A licensee will complete this section if utilizing a QR Code.				
	Attestation: The Licensee attests the QR Code will not make any unfounded nutritional, health, Structure/function and/or implied claims, and will meet all other requirements as put forth by Rule.				
7.	REQUIRED ATTESTATIONS (Section required for ALL item submissions)				
	The Licensee attests the word "Marijuana" is at least as large as any other words used on the packaging for items referenced by this application.				
	The Licensee attests that the word "Marijuana" is at least as wide or tall as the widest part of all logos And/or symbol on the packaging for item(s) referenced by this application.				
	The Licensee attests to retain a current and valid copy of the Child Resistant Certification for the packaging submitted for item(s) referenced by this application.				
	The Licensee attests to retain a current and valid copy of the Food Grade Statement for the packaging submitted for item(s) referenced by this application.				
	The Licensee attests this final marijuana product originated from marijuana grown and cultivated in a licensed cultivation facility located in Missouri pursuant to 19 CSR 100-1.100(4)(1). For manufacturing facility licensees only: additionally, the Licensee attests any tetrahydrocannabinol (THC), such as THC-A, Delta-8, or Delta-10 in this final marijuana product, as defined in 19 CSR 100-1.170(2)(E).				
	The Licensee attests this final marijuana product, as defined in 19 CSR 100-1.010(33), is intended for human use; and the label required by 19 CSR 100-1.120(1)(C)2 includes all ingredients, whether or not the ingredients contain cannabinoids.				

The License attests the information provided for item(s) referenced by this application are true and Accurate.					
I hereby certify that the information provided is true and correct.					
Agent Printed Name (Last, First):					
Agent Business Title:					
Agent Signature:					
Date:					
Agent Email Address:					
Agent Primary Phone Number:					

The Licensee attests this final marijuana product does not contain chemical modifications, conversations, or synthetic derivation of cannabinoids to produce intoxicating cannabinoids isomers, including those created by heat or other process during use by a patient or consumer, and all cannabinoids acquired from entities other than marijuana facilities for purpose of inclusion in marijuana product were accompanied by a Certificate of Analysis at the time of acquisition that identified the testing lab that tested the product

and listed the product's ingredients, pursuant to 19 CSR 100-1.170(2)(F).