

Bulletin Number: LA_IB_0013	Distribution Date: 3/24/20	Effective Date: Ongoing						
Contact Point: Metrc [®] Support	Subject: Remediation Best Pra	actices						
Reason: Metrc is providing best practices for remediating product								

Greetings Metrc® Users,

The LDAF and Metrc would like to provide further guidance on properly remediating items that has failed testing using the LDAF approved remediation methods. Those remediation methods are: High Heat and/or Further Extraction, and Repackaging/Remixing.

**Note: If a product fails testing and that product is allowed to be remediated for potency by the LDAF, it should be repackaged into a new package of a new item with the correct Unit THC and CBD content to be retested. If a package fails for residual solvents, it will require the user to be repackaged through a production batch. If a package fails for homogeneity, it will require the user to remediate without repackaging prior to retesting.

Please see the following pages for the guidance on Remediation for these remediation methods:



Step 1: Navigate to the active Packages screen

The user will select the packages area on the top navigational bar and then select the active tab on the Packages screen.

<mark>∛</mark> n	netrc	Plants 🔻	Packages	Transfe	rs∣ ▼ F	Reports -	Admin	•
Pa	acka	ges						
	Active	On Hold	Inactive					
	New F	Packages	Submit for Tes	ting Re	mediate	Create Plant	ings	New Transfer

Figure 1: Navigation to Active Packages

Step 2: Identify the package that requires Remediation

The user will identify the package with a "**TestFailed**" testing status that is going to be remediated. It is important to understand the reason the product failed testing and identify the appropriate method for remediation.

In Figure 2, two packages are shown:

- 1. A package of tincture that failed for high levels of the extraction solvent.
- 2. A package of tinctures that failed for incorrect potency.

Both packages can be remediated but it will require **different** methods to perform each task. (Figure 3).

oure	onnoid	maotivo											
Nev	w Packages	Submit for	Testing	Test Status	Record Tes	sts Remediat	e New T	ransfer	Change Items	Change	e Notes	Adjust	inish
			_										<u>ه</u> -
► a	ABCDEF01234	5670000016101		ABCDEF0123	45670000016100	M00000001601: Cherry Tincture 50 mg THC	Tinctures		Package Room	95 ea		TestFail	ed
► q	ABCDEF01234	5670000016098	Banana Hammock	ABCDEF0123	45670000015891	Jaguar CO2 Oil	Concentrate		Manufacturing Room	60 g	PB123	TestFail	ed







Figure 3: Remediation Methods

Step 3: Perform Remediation Actions

If a product fails because the levels of solvent used for extraction exceeds LDAF testing regulations, the user has the ability to remediate through further extraction.

*Please note that selecting the Remediate button for any item other than "Test Failed" will open a new Remediate Package window.

Some products in order to be remediated must undergo a process changing the product's physical/chemical form which may include adding an additive, lipid or flavoring. To record this process, the user would create a new production batch in Metrc and select the Remediate Product box and complete the required information as described and illustrated below.

This step must be done in Metrc and cannot be done via the API.

To properly repackage into a Production Batch and Remediate simultaneously, the user would first select the product to be extracted and then click on the <u>New Packages</u> button (**Figure 5**). This will trigger an action window to appear.

New Packages	Submit for Testing	Remed	iate	Create Plantir	ngs	New	Transfer			
Change Rooms	Change Items	Adjust	Fin	ish		_			ē.] (≡•]
Tag	:	Src. H's	:	Src. Pkg's	Ro	oom :	Item :	Category :	Item Strain	Quantity
Q # ABCDEFC	12345670000015181	Metrc Bliss 6/13/2019			Pro Ro	ocessing om	Shake/Trim - Metrc Bliss	Shake/Trim (by strain)	Metrc Bliss	477 g

Figure 4: Select Package for Extraction

Once the action window appears, the user should ensure that:

- "Production Batch" and "Remediate Product" boxes are checked.
- All form fields must be populated, including the **Remediation Method**, **Remediation Date**, and **Remediation Steps**.

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- Remediation Steps should be written clearly to show the steps taken in the remediation process.
- Once the information is entered, the user would select Create Packages to complete the process (Figure 6).

New Packages													
New Packa	ge # 1					(cle	ar)						
New Tag	ABCDEF01234567000001610	7 Q	Package # 1	ABCDEF01234	45670000016104	Q	•						
Location	Manufacturing Room	Q	Quantity	9.9999	Each	÷							
Item	M0000001601: Cherry Tinctu	re Q	_	Packaging - nev	v total will be 88.0001	ea							
Quantity	10 Each	φΣ	(contents)										
Note													
Package Date	iii 02/20/2020 today												
	Production Batch												
Prod. Batch	22												
No.													
	Remediate Product												
Method	High Heat and/or Further Extra	actic 🕈											
<u>Rem.</u> Date	🗰 02/20/2020 today												
Remediation Steps	see lab manual on how to												
+ 1													
	Create Packages	Cancel											

Figure 5: Creating a Production Batch Package to Remediate Product

Remediation Via Repackaging/Remixing

For items requiring repackaging and remixing, the steps will be the same, *except the user will instead select the "Repackaging/Remixing*" method.

• This method would be used in the case of a **tincture** that failed the acceptable potency range or homogeneity. With prior approval of the department, the product can be remediated by re-sizing or re-formulating.

Figure 7 illustrates the action window for creating the production batch and remediating tincture that failed potency testing.





New Package	S				×
New Packa	ge # 1				(clear)
New Tag	ABCDEF012345670000016108	Q	Package # 1	ABCDEF012345670000016106	a 🖸
Location	Manufacturing Room	۹	Quantity	10 Each	\$
Item	M0000001601: Cherry Tincture	Q		Packaging - new total will be 0 ea Finish Package	
Quantity	10 Each	Σ	(contents)		
Note			_		
Package Date	iii 02/20/2020 today				
	Production Batch				
Prod. Batch	23				
110.	Remediate Product				
Method	Repackaging/Remixing	\$			
Rem, Date	🗃 02/20/2020 today				
Remediation Steps	repackage per protocol				
+ 1					
	Create Packages	Cancel			

Figure 6: Verify Remediation Method

For items requiring remixing to remediate for homogeneity, the steps will be the same, *except the user will not repackage or select production batch.*

This method would be used in the case of a **tincture** that failed the acceptable range for homogeneity. With prior approval of the department, the product can be remediated by remixing. The user will first select the product to be remixed and then click on the Remediate button as shown in **Figure 7**. This will trigger an action window to appear as seen in **Figure 8** below.

New Packages Submit for Testing Test Status Record Tests Remediate						fer Change Ite	ems Adjust	Change Notes	Finish			
Tag		Src. H's	Src. Pkg's	:	Location :	Item :	Category :	Item Strain	Quantity :	P.B. No.	Lab Testing	:
🕨 Q 🎇 ABCDEFI	012345670000016106				Manufacturing Room	M00000001601: Cherry Tincture 50 mg THC	Tinctures		9.988 ea		TestFailed	

Figure 7: Select Item Package with Test Failed Status to be Remediated



Once the action window is opened, the user will choose the Remediation Method, Remediation Date and then enter the Remediation Steps clearly for how the product was remediated.

Package #	1	900				(cl	ear)
Package	AB Re	CDEF0123456	70000016104 nixing	Q.	Remediation Steps	Remixing/Repackaging for homogeniety	
Rem, Date		03/13/2020	today				

Figure 8: Remediate package with Remixing/Repackaging Method

The importance of using the remixing method for packages is to ensure the THC/CBD content is consistent throughout the product to aid in dosing recommendations. The difference between remixing and repackaging is that repackaging allows for the reduction of package size to ensure the THC/CBD content is within the LDAF acceptable range. If the product is being remixed and no changes are being made to the package, it would not be considered a production batch and would not need to be repackaged.

Step 4: Verify that the Remediated Products are in a "Remediated" Testing State

The user should now check that the package(s) on which the remediation was performed correctly show "Remediated" in the "Lab Testing" column. The user should also see the triangular symbol next to the tag number denoting that the package contains the remediated product (**Figure 9**).

▶ Q III ABCDEF012345670000016107	ABCDEF012345670000016104	M00000001601: Cherry Tincture 50 mg THC	Tinctures	Manufacturing Room	10 ea	22	Remediated
▶ Q # ABCDEF012345670000016101	ABCDEF012345670000016100	M00000001601: Cherry Tincture 50 mg THC	Tinctures	Package Room	85 ea		Remediated

Figure 9: Verify Product is in a Remediated Testing State



Step 5: Create New Test Samples

Once the product has been remediated, the user can submit the remediated product for testing. The test sample action window is illustrated in **Figure 10**. It is important that the user **ensures the product is listed as remediated before any new samples are pulled**. If the samples are pulled before the product is remediated, the item will need to be retested (as in the case of a suspected false-positive).

Submit for Tes	sting ×
New Packa	ige Sample # 1 (clear)
New Tag Location Item	ABCDEF012345670000016110 Q Package Room Q M00000001601: Cherry Tincture : Q Go (contents)
Quantity Note Package Date	Same item 2 Each Remediated Test Sample iiii 02/27/2020 today
Creating Sample More info	Packages immediately locks out the source Package(s) by setting the Testing State to SubmittedForTesting.
	Submit for Testing Cancel

Figure 10: Submit New Package Sample for Testing

Once the test sample has been created, then it can be transferred to the testing facility using the "Lab Transfer" Transfer type. Once the transfer is completed, the user must wait for the testing facility to complete additional testing.

Verify Remediated Product is Test Passed:

Once the remediated test sample is sent to the testing facility and the product passes, the parent packages at the manufacturing facility will reflect a test result of "TestPassed" as shown in **Figure 11**. This indicates the product can be processed into a final product or released for sale to a Pharmacy.



	Tag :	Location :	Item :	Category :	Quantity :	<u>P.B.</u> No.	Lab Testing	Date :
•	Q ■ ABCDEF012345670000016108 ▲	Manufacturing Room	Jaguar CO2 Oil	Concentrate	10 g	23	TestPassed	02/20/2020
•	Q # ABCDEF012345670000016101 🛦	Package Room	M00000001601: Cherry Tincture 50 mg THC	Tinctures	85 ea		TestPassed	10/08/2019

Figure 11: Verify TestPassed Remediated Product

Pharmacies Can Verify Remediated Products:

Once an item has been received, pharmacies can identify a package that contains remediated product the triangle symbol as seen in **Figure 12** indicates a package contains remediated product.

For the package to be transferred to a Pharmacy, the product must have a TestPassed, or RetestPassed testing state. This can be verified in the "Lab Testing" column (**Figure 12**).



Figure 12: Pharmacy Verification of Test Status

To obtain information on the lab results for the package and obtain additional history, select the arrow to the left of the magnifying glass for the package in question and select the corresponding Lab Results tab. The lab results will display with date, pass/fail, result and the test name. This is illustrated in **Figure 13**.



	Та	g			÷	Location :	tem					:	Category	:	Quantity	:	Lab Testing	÷	Date	÷	
4	٩	AE	BCDEF012345670	0000016101 🖌	8	Package Room	/00000001	601: Cher	ry T	Tincture 50	mg	THC	Tinctures		84 ea		TestPassed		10/08/20	19	X
	Lab Results History																				
																·					
	Test Date : Overall :			Test	Name	: Test F	assed	:	Result	:	Sample Package			Item	Item			: Categ	lory	÷	
		۲	03/19/2020	Passed	тно	C_%_(CONCENTRATE	TRATE) Passed			0.3		ABCDEF0	123456700000	16113	M000000010 THC	601:	Cherry Tincture 5	i0 mg	Tinctu	ires	
		×	03/19/2020	Passed	тно	CA_%_(CONCENTRAT	E) Passe	d		0.1983		ABCDEF012345670000016113		M000000010 THC	601:	Cherry Tincture 5	i0 mg	Tinctu	ires		
	► 03/19/2020 Passed			CBE	D_%_(CONCENTRATE) Passe	d		0.2	ABCDEF012345670		123456700000	16113	M000000010 THC	501:	Cherry Tincture 5	i0 mg	Tinctu	ires		
	▶ 03/19/2020 Passed			CBE	DA_%_(CONCENTRAT	E) Passe	d		0.2		ABCDEF0	123456700000	16113	M000000016 THC	601:	Cherry Tincture 5	i0 mg	Tinctu	ires		

Figure 13: Lab Test Results Tab

Support:

On the far right of the navigation toolbar, a user will see that there is a Support dropdown. Please utilize this dropdown to reference guides, Metrc Support, or training sign up. These tools can be used at any time for no additional charges as long as you are an owner, manager, administrator, or employee in an active, licensed business.

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