kikkoman®

ATP+AMP surface hygiene monitoring ~Application manual for food and beverage industries~

A majority of food poisoning incidents are caused by cross-contamination due to ineffective or improper cleaning. ATP+AMP hygiene monitoring is a convenient test method for quickly measuring the degree of cleanliness on site. This provides an excellent tool for hygiene training and cleanliness control to prevent food poisoning.

Reagents and instrument required



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Principle of ATP plus AMP detection

What is ATP? What is AMP?

ATP (adenosine triphosphate) is the primary molecule involved in metabolism in all living organisms. AMP (adenosine monophosphate) is derived from ATP during the processing, such as heat treatment and fermentation.

ATP cycling method

Kikkoman has created a method using the ATP regeneration enzyme PPDK to measure both ATP and AMP as part of the ATP cycle. This method provides enhanced sensitivity.





* PPDK · · · Pyruvate orthophosphate dikinase

Achievement of super-high sensitivity with both ATP and AMP detection



Measurable even with food residues containing little ATP, such as beer and sausage.

measured using as total ATP+AMP, which microorganisms and

Object to be measured

Cutting board



In ATP+AMP hygiene monitoring, the degree of contaminant is

What is the clean state?

Difference in risk perception

ATP+AMP Conventional method Cutting board method CONCLUSION (culture procedure) CONCLUSION contamination condition microorganism food residu Contaminated Contaminated Cause of microorganism proliferation No Contaminated contamination food residues Potential risk of Possibility of overlooking potential microorganism proliferation is able to risk of microorganism Defective cleaning condition proliferation be detected. No No contamination Cutting board contamination Both cleaning and disinfection are OK.

When food residue is present, there is a possibility for microorganisms to proliferate quickly. The conventional culture procedure can detect microorganisms but the ATP+AMP method also detects food residues. Consequently, ineffective cleaning can be accurately determined.





Let's achive a clean condition (lower luminescence intensity) free of microorganisms and food residue.

Operation example of ATP+AMP hygiene monitoring

	pore		inpre					9			
Es	stablis	h test loc	ations	Test lo	Test locations should be established at the following points:						
				•Areas	 Areas difficult to wash and easy for contaminants to remain. 						
				• Areas	•Areas where not only cleaning but also disinfection and sterilization are conducted.						
					•Areas in contact with ready to eat foods.						
					•Areas at risk of cross-contamination, such as hands and fingers of employees. etc.						
_	$\mathbf{\Lambda}$										
Establish benchmark values				etc.) a	Set the benchmark value as 200 RLU for flat and smooth surfaces (metal, glass, etc.) and 500 RLU for things with surface irregularities and susceptible to scratches (plastic products, etc.).						
				•Thes set r •Deci	<note> Note> These recommended values are not always applicable to any locations. The goal is to set reasonable targets that can be met with rigorous testing and proper cleaning. Decide a swabbing method in accordance with the material and shape of the location to be examined and implement. </note>						
	Example										
	Example (Hands and fingers)Swab every direction of the palm, between fingers, fingertips, etc. The pass and fail le									inderting etc. The pass and fail levels are	
(Hands and fingers) Swab every direction of the paim, between fingers, fingertips, etc. 1500 RLU and 3000 RLU, respectively.								ingerups, etc. The pass and fail levels are			
			L	D							
	- Contraction			- AG	- man 23						
				<u>+ +</u>			·				
	<pre>Kitchen> Portions for</pre>			for the tes	r the test Pass (RLU) Fail (RLU) Swabbing method						
	-		Cutting board 500 1000 10 cm square around the center								
	-		Colander and bowl 200 400 10 cm square of the center bottom portion and top end portion of the inside								
	к		Kitchen	counter	20	0 4	400 10 cm squares at five points on the surface				
			Knife200400Both overall blade surfaces, joint between handle and blade							•	
	Stain		Stainles	s vat	20	0 4	400 Corners where contaminants are likely to remain.				
	Round			20		00 Three inside areas (bottom, middle stand, upper stand)					
	Refrigera		ator (hand	or (handle) 200 4			00 Inside and outside of the overall handle				
	Refrigerato			ator (insid	or (inside) 500 1000 All directions of 10 cm square at the shelf center						
	(Manufactur	ing line)	Valve po	ortions a	und joints	where con	taminan	ts are li	kelv to remain.	
	(Manufacturing line) Valve portions and joints where contaminants are likely to remain.										
	(Environmental inspection) High-frequency contact locations such as telephone sets, door knobs, keyboards and mic of personal computers, etc.										
Ψ											
Es	stablis	h analysi	s scheo	dule C	Ile Conduct the test after cleaning and before disinfection and sterilization.						
(Conducting the test while foods are being handled will not re											
Operation exmple The table below provides an example of location									nple of	operational for hygiene control at each	
location.											
	Tested places Pass/Fail (RLU Lavel 1							Second		Setting of Pass/Fail criteria	
					U) not Lavel 2 measurement		Improvement measures		rement	Not more than level 1 - Pass	
			1,500	3,000	2,412	Caution	Re-cleaning	1,323	Pass	More than fail level 2 - Fail	
	Cutting board 500		500	1,000	760	Caution	Re-cleaning	349	Pass	Between level 1 and level 2 - Caution	
	Bowl		200	400		Passed					

Pass

101

Re-cleaning

130 Passed

44 Passed

820 Failed

200

200 200

Kitchen counter

Refrigerator handle

Vat

400

400

400

Procedures for swab test

Preparation

LuciPac Pen

Remove LuciPac Pen from a refrigerator (2-8C) and allow it to reach room temperature.

*Measuring with the LuciPac Pen while it is still cold does not produce correct measurement results.

Lumitester PD-30

Press "POWER **1**" key to turn ON. After counting down 8 seconds, Lumitester PD-30 is ready to measure.

Use two AA alkaline batteries or two AA nickel metal-hydride batteries. When battery indicator displayed exhaustion sign, turn off the power and replace batteries.



Swabbing

1 Pull out a cotton swab stick (orange) and moisten a swab with tap water.

In the event that the surface to be swabbed is wet with water, the swab does not need to be pre-moistened.





3 Put the swab stick back into the main body and push it through all the way.





When pushing it to the main body, take care not to get your finger caught in. Pressing the bottom by hand or against a table helps you push it in more easily.

2 Swab the object to be tested. See page 3 for the swabbing method.



*In the event that alcohol or other disinfectants remain on the surface to be swabbed, results may not be accurate.It is recommended to carry out the test after washing and before using disinfectants. If the test is conducted after using disinfectants, rinse the surface with water or wipe with paper towel, and conduct the test.

4 Shake off LuciPac Pen to drop the releasing reagent to the bottom of the reaction tube and dissolve the luminescent reagent.

*Make sure all liquid in the capsule falls into the reaction tube.





Measurement

- **5** Insert LuciPac Pen to the measurement chamber of Lumitester PD-30.
- 6 Press the "ENTER ITTER " key of Lumitester PD-30 to start measurement. In 10 seconds, the measurement result is displayed.







*Use the case to stand it if possible. If the stand is not used, do not incline the device more than 60 degrees.

Upon completion of measurement, be sure to remove LuciPac Pen from Lumitester PD-30. Leaving LuciPac Pen inserted in Lumitester may cause problems resulting from liquid leakage, etc.

After measurement

Dispose of used LuciPac Pen in conformity to regulations of local government.

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