

QuantiPlate™ Kit for Microcystins

Highlights:

- Quantitative laboratory detection of Microcystin toxin in surface water
- Detects from 0.2 to 2.0 ppb

Contents of Kit:

- 12 strips of 8 antibody-coated wells each, in plate frame
- 1 vial of Negative Control
- 1 vial of 0.2 ppb Microcystin LR Calibrator
- 1 vial of 0.6 ppb Microcystin LR Calibrator
- 1 vial of 2.0 ppb Microcystin LR Calibrator
- 1 bottle of Assay Diluent
- 1 bottle of Microcystin-enzyme Conjugate
- 1 packet of Wash Solution salts
- 1 bottle of Substrate
- 1 bottle of Stop Solution

Precision

	Recovery	OD
	(%CV)	(%CV)
Inti	ra-Assay	n=11
0.4 ppb	8.3%	4.1%
1.0 ppb	4.6%	5.9%
Into	er-Assay	n=11
0.4 ppb	8.7%	8.7
1.0 ppb	3.6%	11.7

Cross-Reactivity

Compound	50% B ₀	LOD 80% B ₀
Microcystin LR	0.53	0.21
Microcystin LA	0.91	0.16
Microcystin RR	0.69	0.27
Microcystin YR	0.84	0.36
Nodularin	0.30	0.12

Catalog Number EP 022

Intended Use

The EnviroLogix QuantiPlate Kit for Microcystins is designed for the quantitative laboratory detection of Microcystin toxin in surface water samples, with an assay quantitation range from 0.2 to 2 parts per billion (ppb)

How the Test Works

This QuantiPlate Kit for Microcystins is a competitive Enzyme-Linked ImmunoSorbent Assay (ELISA).

In the test, Microcystin toxin in the sample competes with enzyme (horseradish peroxidase)-labeled Microcystin for a limited number of antibody binding sites on the inside surface of the test wells.

After a simple wash step, the outcome of the competition is visualized with a color development step. As with all competitive immunoassays, sample concentration is inversely proportional to color development.

Darker color = Lower concentration Lighter color = Higher concentration

Limit of Detection

The Limit of Detection (LOD) of this Kit is 0.10 ppb. The LOD was determined by interpolation at 91.6% B_0 * from a standard curve. 91.6% B_0 was determined to be 3 standard deviations from the mean of a population of negative water samples.

*100% B_0 equals the maximum amount of Microcystin-enzyme conjugate that is bound by the antibody in the absence of any Microcystin in the sample (i.e. negative control). % B_0 = (OD of Sample or Calibrator/OD of Negative Control) x 100.

Limit of Quantification

The Limit of Quantification (LOQ) of this Kit was validated at 0.2 ppb. The LOQ was determined by fortifying a population of negative water samples at 0.2 ppb. The mean recovery was 96.4% with a coefficient of variation (CV) [(standard deviation/mean) x 100] of 8.2%.

Precision

Microcystin-fortified control solutions were repetitively analyzed both within a single assay, and in different assays on different days. The data is expressed as %CV for both the recovered concentration and for absorbance (OD).

Fortification and Recovery

Four surface water samples were fortified with Microcystin to a concentration of 1.0 ppb. The average recovery was 102%, with a CV of 4.2%.

Cross-Reactivity

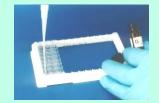
This Kit does not distinguish between the Microcystin toxin variants, but detects their presence to differing degrees. The accompanying table shows the value for 50% B_0 and the value for the 80% B_0 for four Microcystin toxin variants and nodularin toxin. Concentration is in ppb. Humic acid did not interfere in the assay up to a concentration of 100 ppm.



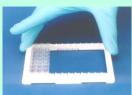
Remove unneeded strips



Select Calibrators and Control



Add controls/calibrators/sample



Mix plate



Incubate



Bottle Wash method

Materials Needed

- disposable tip adjustable air-displacement pipette which will measure 50 μL and 100 μL
- marking pen (indelible)
- tape or Parafilm®
- timer (30 minutes)
- distilled water for preparing Wash Solution
- glassware for storing Wash Solution
- wash bottle for washing strips with Wash Solution
- microtiter plate reader or strip reader
- microtiter plate washer (optional)
- twelve or 8-channel pipette that will measure 50 μL and 100 μL (optional)
- racked (glass) dilution tubes for loading samples into the plate with a 12-channel pipette (optional)
- orbital plate shaker (optional)

Preparation of Solutions

Wash Buffer:

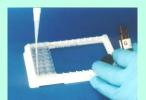
To make 1 L, add the contents of one packet of phosphate-buffered saline - Tween 20, pH 7.4 (Wash Solution salts) to 1 L of distilled water. Mix thoroughly to dissolve the salts. This can be stored at room temperature.

How to Run the Assay

- Read all of these instructions before running the kit.
- Allow all reagents to reach room temperature before beginning (at least 30 minutes with un-boxed strips and reagents at room temperature do not remove strips from bag with desiccant until they have warmed up).
- Organize all samples, reagents and pipettes so that steps 1 and 2 can be performed in 10 minutes or less.
- If more than three strips are to be run at one time, the 10 minutes is likely to be
 exceeded, and the use of a multi-channel pipette is recommended (see "Note"
 below).
- If three or fewer strips are to be run, use a disposable-tip air-displacement pipette and a clean pipette tip to add each Calibrator and sample to the wells. Assay Diluent, Conjugate, Substrate, and Stop Solution may be added in the same manner; alternatively, use a repeating pipette with a disposable tip on the end of the Combitip for these three reagents.
- If fewer than all twelve strips are used, reseal the unneeded strips and the desiccant in the plastic bag provided.
- Use the well identification markings on the plate frame to guide you when adding the samples and reagents. Two strips may be used to run the Negative Control (NC), three Calibrators (C1-C3) and four samples, in duplicate. More samples require more strips. For an example plate layout see Figure 1.
- 1. Rapidly add $50~\mu L$ of Microcystin Assay Diluent to each well that will be used, preferably with a repeating or multi-channel pipetter.
- 2. Immediately add 50 μ L of Negative Control (NC), 50 μ L of each Calibrator (C1-C3) and 50 μ L of each sample (S1-S8) to their respective wells, as shown at left. (Follow this same order of addition for all reagents.) Do not add Microcystin-enzyme Conjugate in this step.



Strip Plate Wash option

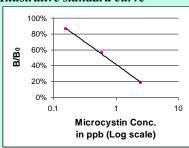


Complete protocol and add Stop Solution



Read plates in a Plate Reader within 30 minutes of the addition of Stop Solution

Illustrative standard curve



3. Thoroughly mix the contents of the wells by moving the strip holder in a rapid circular motion on the benchtop for a full 20-30 seconds. Be careful not to spill the contents!

NOTE: In order to minimize setup time it is recommended that a multi-channel pipette be used in steps 1, 2, 5, 8 and 10 when more than 3 strips are used.

- 4. Cover the wells with tape or Parafilm to prevent evaporation and incubate at ambient temperature for 30 minutes. If an orbital shaker is available shake at 200 rpm.
- 5. Add **50 μL** of **Microcystin-enzyme Conjugate** to each well. Do not empty the well contents or wash the strips at this time.
- 6. Thoroughly mix the contents of the wells as in step 3. Cover the wells with tape or Parafilm and incubate at ambient temperature for 30 minutes. Use orbital shaker if available.
- 7. After incubation, carefully remove the covering and vigorously shake the contents of the wells into a sink or other suitable container. Flood the wells completely with **Wash Solution**, then shake to empty. Repeat this wash step four times. Slap the plate on a paper towel to remove as much Wash Solution as possible. Alternatively, use a microtiter plate washer with **Wash Solution** for the wash step.
- 8. Add 100 μL of Substrate to each well.
- 9. Thoroughly mix the contents of the wells, as in step 3. Cover the wells with <u>new</u> tape or Parafilm and incubate for 30 minutes at ambient temperature. Use orbital shaker if available.

Caution: Stop Solution is 1.0 N Hydrochloric acid. Handle carefully.

10. Add **100 \muL** of **Stop Solution** to each well and mix thoroughly. This will turn the well contents yellow.

NOTE: Read the plate within 30 minutes of the addition of Stop Solution.

How to Interpret the Results

Spectrophotometric Measurement

- 1. Set the wavelength of your microtiter plate reader to 450 nanometers (nm). (If it has dual wavelength capability, use 600, 630 or 650 nm as the reference wavelength.)
- If the plate reader does not auto-zero on air, zero the instrument against 200 μL
 water in a blank well. Measure and record the optical density (OD) of each
 well's contents. Alternatively, measure and record the OD in every well, then
 subtract the OD of the water blank from each of the readings.
- 3. A semi-log curve fit should be used for the standard curve if the microtiter plate reader you are using has data reduction capabilities. If not, calculate the results manually as described in the next section.

How to Calculate the Quantitative Results

1. After reading the wells, average the OD of each set of calibrators and samples, and calculate the $\%B_0$ as follows:

%B₀=<u>average OD of Calibrator or sample</u> x 100 average OD of Negative Control

Precautions and Notes

- Store all components at 4°-8°C (39°-46°F) when not in use.
- Do not expose components to temperatures greater than 37°C (99°F) or less than 2°C (36°F).
- Allow all reagents to reach ambient temperature (18°C to 27°C or 64°F to 81°F) before use.
- Do not use kit components after the expiration date.
- Do not use reagents or test well strips from one QuantiPlate Kit with reagents or test well strips from a different QuantiPlate Kit.
- Do not expose Substrate to sunlight during pipetting or while incubating in the test wells.
- Do not dilute or adulterate test reagents or use samples not called for in the test procedure.
- As with all tests, it is recommended that results be confirmed by an alternate method when necessary.
- Observe any applicable regulations when disposing of samples and kit reagents.
- Microcystin LR in aqueous solution will stick to plastics such as polypropylene. Collect and process samples in glass containers. Clear samples free of organic material can be stored refrigerated for up to two weeks before analysis.

The $\%B_0$ calculation is used to equalize different runs of an assay. While the raw OD values of Negative Controls, Calibrators, and samples may differ from run to run, the $\%B_0$ relationship of calibrators and samples to the Negative Control should remain fairly constant.

The CV for each pair of Calibrator and sample OD values should not exceed 15%.

- 2. Graph the %B₀ of each Calibrator against its Microcystin concentration on a semi-log scale (see Illustrative Standard Curve, left).
- 3. Determine the Microcystin concentration of each sample by finding its B_0 value and the corresponding concentration level on the graph.
- 4. Interpolation of sample concentration is only possible if the $\%B_0$ of the sample falls within the range of $\%B_0$'s of the Calibrators.

If the $\%B_0$ of a sample is <u>higher</u> than that of the <u>lowest</u> Calibrator, the sample must be reported as less than 0.2 ppb.

If the $\%B_0$ of a sample is <u>lower</u> than that of the <u>highest</u> Calibrator, the sample must be reported as greater than 2.0 ppb. If a concentration must be determined for these high level samples, dilute the sample 1:8 in distilled water. Run this dilution in a repeat of the immunoassay. If the result now falls within the range of the $\%B_0$'s of the Calibrators, you must then multiply the concentration measured in the diluted sample by a factor of 8.

Figure 1a. Example of a typical plate setup. (1 x 8 strips)

	1	2	3	4	5	6	7	8	9	10	11	12
Α	NC	NC										
В	C1	C1										
С	C2	C2										
D	C3	C3										
Е	S1	S1										
F	S2	S2										
G	S 3	S3										
Н	S4	S4										

Figure 2a. Illustrative quantitative calculations

Well contents	OD	Average OD	%CV	%B ₀	Microcystin Concentration (ppb)
Negative	1.398				
Control	1.347	1.373	2.628	100	NA
0.2ppb	1.184				
Calibrator	1.177	1.181	0.419	86	NA
0.6 ppb	0.773				
Calibrator	0.776	0.775	0.274	56.4	NA
2.0 ppb	0.246				
Calibrator	0.250	0.248	1.14	18.1	NA
	0.573				
Sample	0.567	0.570	0.744	41.5	1.01

^{*}Actual values may vary; this data is for demonstration purposes only.



For Technical Support Contact Us At:

EnviroLogix

500 Riverside Industrial Parkway Portland, ME 04103-1486 USA

Tel: (207) 797-0300 Toll Free: 866-408-4597 Fax: (207) 797-7533

e-mail: info@envirologix.com

website: www.envirologix.com



LIMITED WARRANTY

EnviroLogix Inc. ("EnviroLogix") warrants the products sold hereunder ("the Products") against defects in materials and workmanship when used in accordance with the applicable instructions for a period not to extend beyond a product's printed expiration date. If the Products do not conform to this Limited Warranty and the customer notifies EnviroLogix in writing of such defects during the warranty period, including an offer by the customer to return the Products to EnviroLogix for evaluation, EnviroLogix will repair or replace, at its option, any product or part thereof that proves defective in materials or workmanship within the warranty period.

ENVIROLOGIX MAKES NO OTHER WARRANTIES, EXPRESS OR IMPLIED, INCLUDING BUT NOT LIMITED TO ANY IMPLIED WARRANTIES OF MERCHANTABILITY OR FITNESS FOR A PARTICULAR PURPOSE. The warranty provided herein and the data, specifications and descriptions of EnviroLogix products appearing in EnviroLogix published catalogues and product literature are EnviroLogix' sole representations concerning the Products and warranty. No other statements or representations, written or oral, by EnviroLogix' employees, agents or representatives, except written statements signed by a duly authorized officer of EnviroLogix Inc., are authorized; they should not be relied upon by the customer and are not a part of the contract of sale or of this warranty.

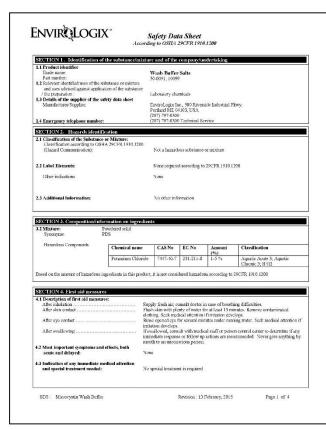
EnviroLogix does not warrant against damages or defects arising in shipping or handling, or out of accident or improper or abnormal use of the Products; against defects in products or components not manufactured by EnviroLogix, or against damages resulting from such non-EnviroLogix made products or components. EnviroLogix passes on to customer the warranty it received (if any) from the maker thereof of such non-EnviroLogix made products or components. This warranty also does not apply to Products to which changes or modifications have been made or attempted by persons other than pursuant to written authorization by EnviroLogix.

THIS WARRANTY IS EXCLUSIVE. The sole and exclusive obligation of EnviroLogix shall be to repair or replace the defective Products in the manner and for the period provided above. EnviroLogix shall not have any other obligation with respect to the Products or any part thereof, whether based on contract, tort, strict liability or otherwise. Under no circumstances, whether based on this Limited Warranty or otherwise, shall EnviroLogix be liable for incidental, special, or consequential damages.

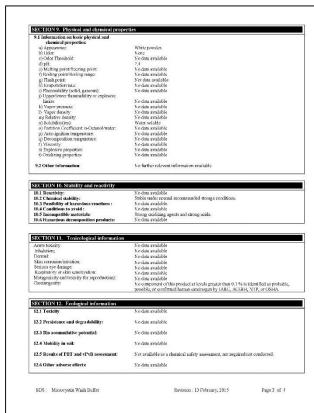
This Limited Warranty states the entire obligation of EnviroLogix with respect to the Products. If any part of this Limited Warranty is determined to be void or illegal, the remainder shall remain in full force and effect.

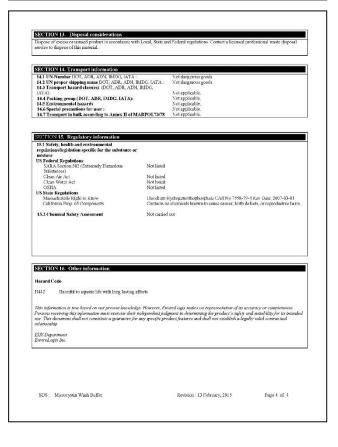
Parafilm is a registered trademark of American Can Corporation EnviroLogix, the EnviroLogix logo, and QuantiPlate are trademarks of EnviroLogix Inc.

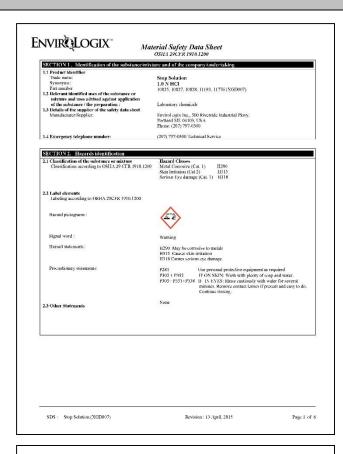
© EnviroLogix 2016



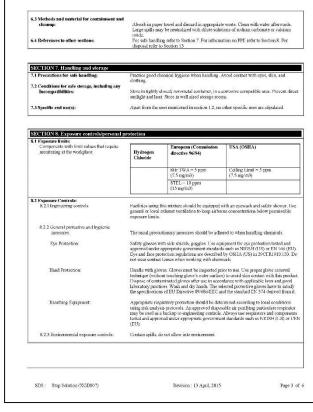
5.1 Extinguishing media:	
Suitable extinguishing agents:	CO2, extinguishing powder or water spray. Fight larger fires with water spray or alcohol resistant fram.
5.2 Special hazards arising from the substance or mixture:	Carbon oxides, Oxides of Phosphorous, Potassium, Sodium, Hydrogen Chloride gas
5.2 Advice for firefighters:	Wear protective equipment appropriate for fire conditions including respiratory protective gear
SECTION 6. Accidental release measures	
6.1 Personal precautions, protective equipment and emorgency procedures:	Use PPE, avoid dust formation, ensure adequate ventilation, avoid breathing dust
6.2 Environmental precautions:	Prevent further leakage or spillage if safe to do so. Do not let product enter drains. Discharge to the environment must be avoided.
6.3 Methods and material for containment and clean up:	Pick up and arrange disposal without creating dust. Sweep up and shovel. Keep in suitable closed containers for disposal
6.4 Reference to other sections:	For safe handling refer to Section 7, For information on PPE refer to Section 8. For disposal, refer to Section 13.
SECTION 7. Handling and storage	
7.1 Procautions for safe handling:	Practice good chemical hygiene when handling. Avoid contact with eyes, skin and clothing. Prevent formation of dust.
7.2 Conditions for safe storage, including any Incompatibilities:	Keep containers closed, store in a dry, well ventilated space.
менирацияния	
7.3 Specific end use(s): SECTION 8. Exposure controls/personal pro	Apart from the uses mentioned in section 1.2, no other end uses are stipulated.
7.3 Specific end use(s):	
7.3 Specific end use(s): SECTION 8. Expassure controls/personal pro 8.1 Control parameters: Components with workplace centrol	dection
7.3 Specific end use(s): SECTION 8. Exposure controls/personal pro SECTION 9. Exposure controls/personal pro Components with workplace control Parameters. 8.2 Exposure controls	dection Contains no substances with occupational exposure limit values
7.3 Specific end use 6: SECTION 8. Expansive controls/personal pre SECTION 9. Expansive controls/personal pre SECTION 9. Expansive controls 8.1 Exposure controls 8.2 I Appropriate engineering controls 8.2 Personal Protective Faujament: 8.2 Personal Protective Faujament:	Contains no solutaness with occupational exposure limit values Ensure operwish and safety shower are marby, provide ventilation if moreovery Safety glances with nide shoulds, gapples. Use confirment for eye, protessive saded and segrowed under appropriate powerment shadatals such as NESH (USE) or PA. 166 (PA). For and five proceeding or sufficient scale and segrowed under appropriate powerment shadatals such as NESH (USE) or PA. 166 (PA). For and five proceeding or sufficient scale and solutions are described by OSIA (PA) in 20-CER (PA) (PA).
7.3 Specific end use 6: SECTION 8. Exposure controls/personal processor and control parameters. Componers with workplace control parameters. 8.2 Exposure controls. 8.2.1 Appropriate augmenting controls. 8.2.2 Pensonal Protective Equipment. Eyes.	Centains no substances with occupational exposure limit values Braure grounds and safety shower are marby, provide ventilation if moreosary Sufray planes with side shields, googles. Use outprenent for eye protection tested and agreeousl under agreeoping government shouldness such as NUSH (US) or 8th 166 (03). Eye and fine proceeding regulations are described by OSHA (US) in 29CER1910.133 Dr. and were contactly leases with reduting by the theretools and were contactly because with neutron of which theretools record to the contact of the co
7.3 Specific end use q: SECTIONS. Exposure controls/personal pre SI Control parameters: Componers with workplace control Parameters: 8.2 Exposure controls 8.2 Lapprograms augmenting controls 8.2 Lapprograms augmenting controls 8.2 Personal Probedive Equipment Flyes Hands	Contains no solutaness with occupational exposure limit values. Ensure operand and safety shower are nearby, provide ventilation if moreovary. Safety glasses with side sholds, gogeles. Use outpressed for eye protection tested and exproved under apprepriate government shoulded, such as NOSH (US) or NO 166 (93). Do not were contact lenses when working with chemical Do not were contact lenses when working with chemical Do not were contact lenses when working with chemical residence of contaminated places after not use. Use proper place removal technique (without teaching glow's cours surface) to avoid dain contact with this contact of contaminated places after not moved men with appreciable lower and good absoratory practices. With mind day hands. The selected proceeding with appreciable lower and good absoratory practices. With mind day hands. The selected procedure given the control of the contact of the seasofth IN 5-742 derived. Exposure places of the University 800000000 and the seasofth IN 5-742 derived Exposure. Appropriate respiratory protection should be determined according to local conditions using rid analysis protecols. An approved disposable are purifying particulate requires to ested and suppress under superprint government standards each and NISM (IS) or
7.3 Specific end use 6: SECTIONS. Expanser controls personal pre SECTIONS. Expanser controls Component with reveloplese centrel Patameters. SE Exposure controls SE.1 Appropriate aggineering controls 8.2.1 Personal Protective Equipment: Syss. Hands Respiratory protection	Centains ne-solvances with occupational exposure limit values Ensure operwish and safety shower are marby, provide ventilation if moreovary. Suffey planes with nide should, grapher. Use outpressed for grey, pro-solves tested and sepressed used as approach government for grey, pro-solves tested and sepressed used as approach as positions and as NESH (GIS) or No. 166 (202). Eva and few pro-cention regulations are described by CoEIA (US) in 20CET 191 (13.3). Do not were contact treation working with chemical processes of the safety of the processes of the safety of the safety of the processes of the safety of th







SEC	TION 3. Compositi	ion/inform	ation on ingr	edients				
3.2	Mixture Aqueous solution			200 200 C 20				
	Chemical name	Amount (%)	CAS No	Classification According to OSHA 29CFR 1910.1200				
	Hydrochloric acid	1-4%	EC No	Hazard Classification	Hazard Code			
	nydrodiinie aut	104.29	7647-01-0	May be Corrosive to Metals	H290			
			231-595-7	Causes Skin Irritation	H315			
				and September 1982 and	H318			
				Causes Serious Eye Damage	011318			
	ETION 4. First aid r							
	After inhalation :	measures		In case of inhalation. Remove to fresh air. It				
A	After skin contact :			respiration. Get medical attention immediate In case of skin contact. Remove contaminates	d clothing and shoes immediately.			
				Wash affected area with mild scap or deterge evidence of chemical remains.	mt for al least 10 minutes or until no			
A	After eye contact :			In case of eye contact, immediately flush eye minutes. Lifting evelids occasionally, until n				
				medical attention immediately.				
A	After swallowing			In case of ingestion. DO NOT Induce vomiti medical personnel. Never give anything by a physician immediately.	ng unless directed to do so by mouth to an unconscious person. Call			
		ame and off	ects, both acut					
	dost important sympt And delayed:	oms and en		May cause skin irritation and eye damage				
4.3 It	and delayed: ndication of any imme	ediate medi		ıd	2 a 2 52 111			
4.3 It	And delayed:	ediate medi			opt to neutralize the acid.			
4.3 In	and delayed: ndication of any imms pecial treatment need	ediate medi led:	cal aftention an	ıd	upt to neutralize the acid.			
4.3 II 8 SEC	and delayed: Indication of any imms Indication of an	ediate medi led:	cal attention an	ıd				
4.3 It s	and delayed: Indication of any imms Special treatment need "TION 5. Firefightic axtinguishing media:	ediate medi led: ng measur	cal aftention an	id IX) NOT use sodium bioarborate in an attem				
4.3 In 8	and delayed: Indication of any imms Indication of an	ediate medi led: ng measur	cal attention an	td IX) NOT use sodium bleathorate in an attem CO2, orthographing powder or water spray. Fight I				
5.1 E	and delayed: Indication of any imms Special treatment need THON 5. Firefighti Extinguishing media; Special hazards arising	ediate medicled:	es ubstance or	to IX) NOT use seekum bloochorate in an aftern IX (NOT) and IX (NOT) and IX (NOT) are seekum bloochorate in an aftern IX (NOT) are seekum bloochorate in a seekum bloochorate in	arger fires with water spray or alcohol			
5.1 E	and delayed: ndication of any imma pecial treatment need TION 5. Firefightif Extinguishing media: special hazards arising nlyture:	ediate medicled:	es ubstance or	td 150 NOT me wedum bloorbornts in an aftern CO22, actinguishing powder or water spray. Fight I field that flows West protective gest appropriate for fire condition.	arger fires with water spray or alcohol			
8120 5.1 E 5.2 S	and delayed: ndication of any imma pecial treatment need TION 5. Firefightif Extinguishing media: special hazards arising nlyture:	ediate medicled: ng measur g from the s	cal aftention an	td 150 NOT me wedum bloorbornts in an aftern CO22, actinguishing powder or water spray. Fight I field that flows West protective gest appropriate for fire condition.	arger fires with water spray or alcohol			
A.3 In SECOND SE	and delayed: inflication of any immunopecial treatment need TION 5. Fireflightif axtinguishing media: special barards arising ulxture: A.dvice for fireflighters	ediate medicled: ng measur g from the s	ubstance or neasures quipment	td 150 NOT me wedum bloorbornts in an aftern CO22, actinguishing powder or water spray. Fight I field that flows West protective gest appropriate for fire condition.	arger fires with vorter spray or alcohol- as including respiratory protective			



1 Information on basic physical and				
chemical properties: a) Appearance:	Clear figurd, colorless to slight	entlane.		
a) Appearance: b) Other	Present (slight)	yenow.		
c) Odor Threshold:	No Data Available			
d) pH:	pH 1			
c) Melting point/freezing point	No Data Available			
f) Boiling point/Boiling range:	No Data Available.			
e) Flash point:	Not applicable.			
h) Evaporation rate:	0.36 (Water) compared with n-	Butyl Acetate = 1		
i) Flammability (solid, gaseous): j) Upper/lower flammability or explosive	No Data Available			
limits:	No Data Available			
k) Vapor pressure:	No Data Available			
Vapor density mt Relative density.	No Data Available No Data Available			
n) Keianve censity; n) Solubilityfies);	Fully miscible, water.			
o) Partition Coefficient: n+Octanol/water:	No Data Available			
p) Auto-ignition temperature:	No Data Available			
g) Decomposition temperature:	No Deta Available			
r) Viscosity:	No Data Available but should	e similar to that of y	soler.	
s) Explosive properties:	No Data Available.			
t) Oxidizing properties:	No Data Available			
.2 Other information:	No further relevant information	t available.		
CTION 10. Stability and reactivity				
.1 Reactivity:	No data available			
0.2 Chemical Stability:	Stable under normal tempera	tures and pressures.		
3.3 Possibility of hazardous reactions:	Under normal conditions of s	torage and use, haza	rdous reactions	will not occur.
4 Conditions to avoid:	No specific data			
0.5 Incompatible materials:	Metals, Alkali metals, bases,	Armines.		
0.6 Hazardous decomposition products:	Under normal conditions of s not be produced.	kerage and use, haza	rdous decemper	sitions products should
CTION 11. Toxicological information formation on Toxicological Effects		Effect Dose 1.050=900mg/s;	s	stions products should
CTION 11. Toxicological information formation of Toxicological Effects	not be produced. 7647-01-0 HCI Acute onal toxocity Acute demail toxocity	Effect Dose 1.050=900mg/kg No data	S	pecies
CTION 11. Toxicological information formation on Toxicological Effects	not be produced. 7647-01-0 HC1 Acute oral teodoity	Effect Dose 1.D30=900mg/kg	S	pecies Elbiri
CTION 11. Toxicological information information on Toxicological Hiers care effices (routiny rosts).	7647-01-0 HCI Acute oral lexicity Acute inhalative toxicity No sensitizing effects known	Effect Dose 1.050=900mg/sq No dala LCSG = 3124 mg	S	pecies Elbiri
CCTION 11. Toxicological information formation on Toxicological Effects curie effects (rocicity rosts): emissionic missionic MM (carenogenisty, mutagenesty and rescis)	7647-01-0 HCI Acute oral lexicity Acute inhalative toxicity No sensitizing effects known	Effect Dose 1.050=900mg/sq No dala LCSG = 3124 mg	S	pecies Elbiri
CTION 11. Toxicological information formation on Toxicological Iffects care effects (rotatily rosts); retailmation: M.S. (narron-pariety, mutagenesty and toxicily or equivalence effects.	not be produced 7647-01-0 HCI Acute and tocomy Acute dermal tocomy Acute inhalaire tocomy No sensitizing effects known	Effect Dose 1.050=900mg/sq No dala LCSG = 3124 mg	S	pecies Elbiri
CTION 11. Toxicological information formation on Toxicological Iffects care effects (rotatily rosts); retailmation: M.S. (narron-pariety, mutagenesty and toxicily or equivalence effects.	not be produced 7647-01-0 HCI Acute onli tenedry Acute demail investry Acut a reliable ve toolvily No sensitizing effects known No CNR effects.	Effect Dose 1.050=900mg/sq No dala LCSG = 3124 mg	S	pecies Elbiri
CCTION 11. Toxicological information formation or Toxicological Effects care effects (rocicity tests): custination: M4 (carenogenicity, mutageneity and rocicity effects) effects (rocicity effects) effects (defining locicological information):	not be produced 7647-01-0 HCI Acute onli tenedry Acute demail investry Acut a reliable ve toolvily No sensitizing effects known No CNR effects.	Effect Dose 1.050=900mg/sq No dala LCSG = 3124 mg	S	pecies Elbiri
CTION 11. Toxicological information formation at Toxicological Thects care offices (roughly rosts): Toxicological Thects care offices (roughly rosts): Toxicological Thects of toxicological toxicological toxicological information: CTION 12. Ecological Information	2015 be produced. 2017-81-81-101 Actual and tesceity. Actual darmal tesceity. Actual charmal tesceity. No sensitizing effects learners. No CANR effects. No Additional Information	Effect Dose 1,D36=900mp3c; No dala 1,CS9 = 3124 mg	S 5 15 15 11 11 11 11 11 11 11 11 11 11 1	pocies Histi
CCTION 11. Toxicological information formation or Toxicological Effects care effects (rocicity tests): custination: M4 (carenogenicity, mutageneity and rocicity effects) effects (rocicity effects) effects (defining locicological information):	7647-01-0 HCI Actus oral kossety Actus oral kossety Actus imbalative toxicity No sensitizing effects knewn No CMM effects No Additional Information Aquatic toxicity (18 HC3)	Effect Dose 1.1356-90tmp3c; No data 1.CS0 = 3124 mg	S S III III III III III III III III III	pecies hibbit
CTION 11. Toxicological information formation in Toxicological Tifects care effects (roughly rosts): Also feare-specially, mutagereity and rosicity are production of the production of the second of the control of th	7617-01-0 HCI Acute onl toucotry Acute inhalidry toucity Acute inhalidry toucity Acute inhalidry toucity Acute inhalidry toucity Acute finding officels known Acute find toucity Aquatic touci	Effect Dose 1.D30=900mp3c; No dala 1.CS0 = 3124 mg Effect dose 1.CS0=220 mg/L	S 5 15 15 11 11 11 11 11 11 11 11 11 11 1	pocies Histi
CTION 11. Toxicological information formation in Toxicological Tifects care effects (roughly rosts): Also feare-specially, mutagereity and rosicity are production of the production of the second of the control of th	7647-01-0 HCI Actus oral kossety Actus oral kossety Actus imbalative toxicity No sensitizing effects knewn No CMM effects No Additional Information Aquatic toxicity (18 HC3)	Effect Dose 1.1356-90tmp3c; No data 1.CS0 = 3124 mg	S S III III III III III III III III III	pecies hibbit

QuantiPlate Kit for Microcystins Page 8 of 8

12.2 Persistence and degradability: No Data Available

12.4 Mobility in soil: No Data Available

12.5 Results of PBT and vPvB assessment: No Data Available

12.6 Other adverse effects

No Data Available

12.6 Other adverse effects

No Data Available

No Data A

Not carried out

Revision: 13 April, 2015 Page 5 of 6

15.2 Chemical Safety assessment

SDS: Stop Solution (XGD007)

SECTION 16. Other information

This information is true based on our present bouledge. However, Emvedages nodes no expression of its accuracy or completeness, true and true true to the true present of the accuracy or completeness. The information of the accuracy or completeness are related to the control of the present of the accuracy or completeness are related to the control of the accuracy of

Revision: 13 April, 2015

Page 6 of 6

SDS: Stop Solution (XGD007)