Test of Larval Growth and Survival Using Inland Silverside

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Parameter Sample Preparation	Specification	Met s	fics?			
Filtering	If indigenous organisms, filter through a sieve (60 µm) (Must)					
D.O. Measurement	In each sample prior to filtering and after T° adjustment					
Pre-aeration	None unless D.O. is < 4 mg/L, then aerate all test solutions (Must) for a few					
	minutes at a rate not exceeding 100 bubbles/min, until the D.O. is ≥ 4mg/L					
pH Adjustment	pH measured in each sample each day before new test solutions are made					
	A second (pH adjusted) test might be run if pH is outside 6.0 to 9.0					
T° Adjustment	T° to be measured in sample on arrival at lab					
	Sample adjusted to 25 ± 1°C prior to test initiation (approximately 1h)					
Salinity Adjustment	Salinity of each sample measured before starting the test					
	Sample adjusted to 28 - 32 g/kg using hypersaline brine (HSB) (as per EC					
	guidance on salinity adjustment) (Must)					
Test Conditions						
Test Facility	Isolated from general laboratory disturbances					
,	Instruments available to measure basic water quality variables (T°, D.O., pH,					
	salinity) and lab prepared for other analyses					
Test Type	Static renewal					
Test Duration	7 days (Must)					
Test T°	25 ± 1°C (Must)					
Light Quality	Ambient laboratory illumination.					
Light Intensity	10 - 20 μE/m ² /s					
Photoperiod	16 ± 1h light; 8 ± 1h dark					
Salinity	28 - 32 g/kg; preferably 30 g/kg; each test solution within 1 g/kg of the control;	•••		•••		
Samily	adjust using HSB (with a salinity of 90 ± 1g/kg) or deionized water					
	Nominal test conc. adjusted and reported in consideration of any salinity					
	adjustments (Must).					
D.O. Range	D.O. in test solutions not be permitted to fall below 4 mg/L (Must)					
Aeration	None, unless D.O. < 4 mg/L, then aerate all chambers at a rate not exceeding					
	100 bubbles/min					
Vessel Size & Type	Glass chamber with sump area; borosilicate glass or non toxic disposable					
	plastic labware					
	600 mL - 1L containers; covered during test with safety glass plates or sheet					
	plastic (6mm thick)					
Test Volume	500 - 750 mL/replicate; water depth 5 cm					
	Minimum of 50 mL of solution per larvae (Must)					
Renewal of Solution	≤ 24 h for test duration (Must)					
	80 - 95% of solution replaced; dead brine shrimp and detritus removed; new					
	test solution added slowly and cautiously to avoid injury to the fish					
Dilution/Control Water.	Filtered (60 µm) uncontaminated lab seawater, reconstituted seawater, or					
	filtered (60 µm) upstream receiving water					
	Salinity: 28 - 32 g/kg (Must); recommend 30 g/kg; salinity adjusted using					
	aged HSB with a salinity of 90 ± 1g/kg or deionized water, distilled water or					
	uncontaminated freshwater			l		
	Any HSB used, be from the same source as that used to adjust the salinity of					
	the sample or test solutions (Must)					
	Adjusted to 25 ± 1°C before use					
	If the test organisms have been cultured in water which is different from the					
	test control/dilution water, a second set of controls, using culture water, is to					
	be included in the test					
	If any HSB is added to sample or test solutions to adjust salinity, the toxicity		l	l		
	test include a set of controls prepared using only this HSB and deionized					
	water, adjusted to the test salinity 30 ± 2 g/kg (Must)	•••				
	If uncontaminated receiving water used as control/dilution water, an additional					
	lab seawater control is to be run (Must)		•••			
	Any test using dilution water (eg: natural seawater) which differs from this					
	HSB control include a separate set of controls prepared using this same					
	dilution water (Must).	<u> </u>				

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	Specification	Met : Y	Speci N	fics'
Vessel Identification	Test chambers labeled with test conc. and replicate number			
# Test Conc	≥ 5 plus control to calculate ICp and LC50 (Must); dilution factor ≥ 0.5			
	1 plus control for single conc. test			
# Replicates/Conc	≥ 3 per test conc. and controls (Must); 4 replicates recommended			
	Test start with equal number of replicates for each test conc. and controls			
# Organisms/Vessel	≥ 10 larvae randomly distributed to each test chamber on the day preceding			
	the test (Must); 15 larvae per chamber recommended			
	The test is started by removing ~90% of the clean seawater from each test			
	chamber and replacing with the appropriate test solution			
Vessel Randomization.	Test chambers placed in a randomized position in water bath			
	Before beginning the test, remove and replace any dead larvae from each test chamber			
Removal of Dead	Dead organisms discarded daily during the test.			•••
Feeding Regime	Feed daily during test with newly hatched (< 24h old) brine shrimp nauplii			•••
recally regime	(0.10 g wet weight per replicate on days 0-2; 0.15 g wet weight per replicate			
	on days 3-6) from days 0-6; larvae not fed on day 7			
	Equal amounts of <i>Artemia</i> be fed to each replicate test chambers (Must)			
Cleaning	All non-disposable test vessels and equipment to be thoroughly cleaned and		'''	
	rinsed in accordance with section 5.3 (Must).			
	Siphon bottom of test chamber daily immediately before test solution renewal			
	and feeding			
Endpoints	Mortality and growth: if multi conc. test, LC50 for mortality and ICp for mean			
	dry weight for surviving fish (both with their 95% confidence limits) (Must)			
salinity	At least at start and end (just before or immediately after renewal) of each 24-hour exposure in representative concentrations (high, medium, low, and controls) in both the fresh and used solution (Must)			
	intervals of 24 h from the start until the end of test at 7 d of exposure # of fish showing loss of equilibrium or abnormal swimming behaviour			
Growth	determined for each test vessel		•••	
Growth	Fish dried immediately at 105 °C for 6 h or at 60 °C for 24 h			
	Upon removal from oven, boats moved immediately to dessicator			
	Thereafter, the boats be individually and randomly removed from the			
	dessicator and weighed on a balance the measures consistently to 10 μ g			
	Rapid weighing and standard timing among weigh boats is necessary			
Test Organism	Manidia hamilia			
Species	Menidia beryllina			
Source	From in-house cultures or commercial suppliers			
Age	7 to 11-days post hatch; 24h range in age			
Age	In a given test, all organisms be approximately the same age and be taken			•••
	from the same source			
Health Criteria	A group of organisms not be used for a test if they appear to be unhealthy, discolored, or otherwise stressed, or if mortality exceeds 10 % preceding the		•••	
	test; upon failure of these criteria, the entire group is to be discarded and a			
	new group obtained (Must)			
Culture/Holding Conditions				
	25 ± 1°C (Must) ; rate of change ≤ 2°C/day for new adult fish batches 6.0 - 9.0			

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Parameter	Specification	Met Y	ifics?	
Salinity	28 - 32 g/kg (Must) ; ideally 30 g/kg; ≤ 3 g/kg change over 12 h			
Light Quality	Ambient laboratory illumination			
Light Intensity	10 - 20 μE/m²/s			
Photoperiod	16 ± 1h light; 8 ± 1h dark			
Feeding	Adults: flake food or frozen brine shrimp twice daily and <i>Artemia</i> nauplii (< 24h old) once daily			
	Newly hatched fish:with about 500 rotifers <i>Brachionus plicatilis</i> per larvae per day from hatch through 4d post-hatch; on days 5 and 6, newly hatched (<12h			
Cleaning	old) <i>Artemia</i> mixed with the rotifers; after 7d, <i>Artemia</i> (< 24h old) only Siphoning of debris daily or as required			
Culture Water	Filtered (≤ 60 µm) uncontaminated natural seawater, or reconstituted seawater; or filtered (≤ 60 µm) receiving water			
	T°, D.O., pH and salinity monitored in culture tanks daily.			
Morbidity/Mortality	Adult and pre-adult fish being cultured inspected daily for signs of disease			
morbialty/mortality	Mortality rates and any evidence of disease recorded at least 5 d/w			
	Dead and moribund individuals removed immediately			
Obtaining Eggs	If embryos, tile should be removed and placed in hatching tray			
Hatching Eggs	Aerate tile or remove eggs from tiles and aerate in separatory funnel			
	Inspect incubating embryos daily			
Facility & Apparatus	Remove and discard dead embryos or those with fungus daily		•••	
Facility & Apparatus	Vessels and accessories contacting organisms and culture media made of non-toxic material (Must)			
	Culture facility located away from physical disturbances and preferably	•••		
	separate from test containers			
QA/QC	200/ summingling controls (Marca)			
Test Validity Criteria	≥ 80% survival in controls (Must)			
	0.50 mg average dry weight of control larvae where test starts with 7-days old larvae and dried immediately after test termination; or 0.43 mg if fish are first			
	preserved (not more than 7 days) in 4% formalin or 70% ethanol (Must)			
Reference Toxicant	Monthly and following the same procedure as the definitive test (Must);			
	ideally with larvae from culture that are used in toxicity test			
	Standard test of 7 d with LC50 and ICp endpoints (Must)			
	Sodium chloride, potassium chloride, cadmium chloride, copper sulfate,			
	sodium dodecyl sulfate and potassium dichromate are suitable			
	Using same water as culture dilution/water			
Warning Chart	Prepared for each reference toxicant and continually updated			
	Within acceptable warning limits (± 2 SD on log scale)			
Sample Handling Sample Collection	For off-site effluent tests, either 3 subsamples from a single sampling or ≥ 3 separate samples are collected (Must); for on-site tests, samples are			
	collected daily and used within 24 h			
Volumes	Volumes of 6L per day recommended			
Containers	Non-toxic materials for sample and transport containers, new containers or			
Labeling	thoroughly rinsed used containers			
Labeling	Upon collection, sample containers filled, sealed and labeled/coded Include at least sample type, source, date and time of collection and name of		•••	
	sample collectors			
Holding Time	Test to be initiated within 3 d after sampling (Must); recommend within 1d			
Holding Conditions	Keep samples cool throughout their period of transport at 4 °C using regular			l
g conditionon	ice or frozen gel packs			
	Upon collection, if sample > 4 °C, cool to 4 °C with regular ice or frozen gel			
	packs (not dry ice)			
	The portion(s) of sample or subsamples required for solution renewals be			
	stored in darkness in sealed containers without air headspace at 4 °C			
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TEST SPECIFIC CHECKLIST

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March 1998

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Parameter	Specification	Met Specific		
Minimum Level of	Do typical test reports reflect the minimum level of reporting outlined			
Reporting	below? (Must)			
Sample Data	Brief description of sample type if and as provided to the lab			l
Campio Bata	Information on labeling or coding, for each sample			
	Date of sample/subsample collection; date and time sample(s)/subsample(s)			
	received at test facility.			
	For effluent or leachate, T° of sample upon receipt at lab			
	D.O. and pH of each sample just before its preparation and use.			
	Dates or days during test when individual samples or subsamples used		l	
Test Organism	Species and source of organisms		l	
	Age at start of test.			
	Any unusual appearance, behaviour, or treatment of test organisms, before			
	their use in the test			
	Data showing health of organisms, including mean % mortality preceding test.	l		
Test Facilities	Name and address of test laboratory			
Toot i dominoo	Name of person(s) performing the test.			
	Brief description of test vessels (size, shape, type of material)			
Control/Dilution Water.	Type and source of water used as control and dilution water			
Control/Blidtion Water.	Type and quantity of any chemical(s) added to control or dilution water			
Test Method	Statement that the Environment Canada guidance document on salinity			
rest Method	adjustment has been followed			
	Citation of method used and type of test.			
	In those instances where any sample or test solutions has/have been pH			
	adjusted, and/or is/are filtered, brief description of procedure(s)			
	Description of procedure(s) for salinity adjustment of sample and dilution			
	water	•••		
	Description of procedure for preparation of hypersaline brine			
	Frequency and type of all observations and measurements made during test.			
	Name and citation of program(s) and methods used for calculating statistical			
-	endpoints			
Test Conditions	Design and description if any deviation from or exclusion of any of the			
	procedures and conditions specified in test method document			
	Manner and rate of exchange of test solutions			
	Number, concentration, volume, and depth of solutions in test vessels,			
	including controls			
	# of individuals per test vessel, and # of replicates per treatment			
	Brief statement (including procedure, rate, and duration) if any pre-aeration or			
	aeration of sample or test solutions			
	Dates when test was started and ended			
Test Results	All required measurements of T°, pH, D.O. and salinity in sample and test			
	solutions (including HSB controls and, if natural seawater has been used as			
	dilution water, natural seawater controls), before and made during the test			
	# and % of mortality of the organisms in each test chamber, as recorded daily.			
	Average dry weight per original fish in each test chamber			
	LC50 (including the associated 95% confidence limits) for survival data and			
	indication of quantal statistic method used; details regarding any			
	transformation of data that was required			l
	ICp (including the associated 95% confidence limits) for growth data and			
	indication of quantitative statistic method used; details regarding any		1	
	transformation of data that was required		1	
	Results and duration of any toxicity tests with the reference toxicant(s)			1
			1	
	performed within 30 days of the test, together with the geometric mean value		1	
	(± 2 SD) for the same reference toxicant(s) as derived at the test facility in		1	
	previous tests		1	
	Anything unusual about the test, any problems encountered, any remedial measures taken.		1	