	Document Owner Department: QC	MBD-BT-SPEC-0511
		Page 1 of 5
		Rev 07
OXOID QUALITY ASSURANCE PRODUCT SPECIFICATION		
HALF FRASER SELECTIVE SUPPLEMENT SR0166E		

HALF FRASER SELECTIVE SUPPLEMENT

SR0166E

Formula

Vial contents (each vial is sufficient to supplement 225ml of medium)

Ammonium iron (III) citrate	112.50	mg
Nalidixic acid	2.25	mg
Acriflavine	2.8125	mg

Description

A selective supplement for the isolation of *Listeria* spp.

Directions

Aseptically add 4ml 1:1 ethanol:sterile distilled water to 1 vial and invert gently to dissolve. Aseptically add the vial contents to 225ml of sterile Fraser Broth Base (CM0895) prepared as directed and cooled to 50°C. Mix well and aseptically dispense into sterile containers.

Physical Characteristics

Orange/green pellet
Sterility - passes test

Microbiological Tests Using Optimum Inoculum Dilution


Control Media: Brilliance™ *Listeria* Agar (ISO), Tryptone Soya Agar or Columbia Blood Agar Base enriched with 5% v/v horse blood, where appropriate

Tested in Fraser Broth Base CM0895

Reactions after incubation at 30 ± 2°C for 24 ± 2 hours

Inoculate 10ml quantities of medium to achieve 1-10 colony-forming units/ml (cfu/ml) of *Listeria monocytogenes*. Incubate broths at 30 ± 2°C for 24 ± 2 hours. Subculture onto Brilliance™ *Listeria* Agar (ISO) (CM1212, SR0257 & SR0258) and incubate plates at 37 ± 2°C for 24 ± 2 hours.

<i>Listeria monocytogenes</i>	ATCC®7644
<i>Listeria monocytogenes</i>	ATCC®13932

	Document Owner Department: QC	MBD-BT-SPEC-0511
		Page 2 of 5
		Rev 07
OXOID QUALITY ASSURANCE PRODUCT SPECIFICATION		
HALF FRASER SELECTIVE SUPPLEMENT SR0166E		

A satisfactory result is represented by recovery of positive strains equal to or greater than a 4 log(10) increase.

Positive strains shall produce aesculin hydrolysis after 24 hours.

Reactions after incubation at 30 ± 2°C for 24 ± 2 hours

Inoculate 10ml quantities of medium to achieve >1E+03 cfu/ml. Incubate broths at 30 ± 2°C for 24 ± 2 hours.

<i>Bacillus cereus</i>	ATCC®10876	No aesculin hydrolysis (no blackening)
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Negative strains are inhibited or shall produce a negative diagnostic reaction.


Productivity determined by qualitative testing in accordance with the methods and criteria described in ISO 11133:2014

Inoculation with mixed cultures

Inoculate 10ml quantities of medium to achieve 1-10 colony-forming units/ml (cfu/ml) of *Listeria monocytogenes*, to each add 1E+02 to 1E+03 cfu/ml of *Escherichia coli* and 1E+02 to 1E+03 cfu/ml of *Enterococcus faecalis*. Incubate broths at 30 ± 2°C for 25 ± 1 hour. Subculture onto Brilliance™ *Listeria* Agar (ISO) (CM1212, SR0257 & SR0258) and incubate plates at 37 ± 2°C for 24 ± 2 hours

Reactions after incubation at 30 ± 2°C for 25 ± 1 hour

<i>Listeria monocytogenes</i>	ATCC®13932	WDCM00021	0.5-1.0mm blue colonies with halo
+ <i>Escherichia coli</i>	ATCC®8739	WDCM00012	No growth
+ <i>Enterococcus faecalis</i>	ATCC®19433	WDCM00009	No growth
<i>Listeria monocytogenes</i>	ATCC®13932	WDCM00021	0.5-1.0mm blue colonies with halo
+ <i>Escherichia coli</i>	ATCC®25922	WDCM00013	No growth
+ <i>Enterococcus faecalis</i>	ATCC®19433	WDCM00009	No growth
<i>Listeria monocytogenes</i>	ATCC®13932	WDCM00021	0.5-1.0mm blue colonies with halo
+ <i>Escherichia coli</i>	ATCC®8739	WDCM00012	No growth
+ <i>Enterococcus faecalis</i>	ATCC®29212	WDCM00087	No growth

	Document Owner Department: QC	MBD-BT-SPEC-0511
		Page 3 of 5
		Rev 07
OXOID QUALITY ASSURANCE PRODUCT SPECIFICATION		
HALF FRASER SELECTIVE SUPPLEMENT SR0166E		

<i>Listeria monocytogenes</i>	ATCC®13932	WDCM00021	0.5-1.0mm blue colonies with halo
+ <i>Escherichia coli</i>	ATCC®25922	WDCM00013	No growth
+ <i>Enterococcus faecalis</i>	ATCC®29212	WDCM00087	No growth

<i>Listeria monocytogenes</i>	ATCC®35152	WDCM00109	0.5-1.0mm blue colonies with halo
+ <i>Escherichia coli</i>	ATCC®8739	WDCM00012	No growth
+ <i>Enterococcus faecalis</i>	ATCC®19433	WDCM00009	No growth

<i>Listeria monocytogenes</i>	ATCC®35152	WDCM00109	0.5-1.0mm blue colonies with halo
+ <i>Escherichia coli</i>	ATCC®25922	WDCM00013	No growth
+ <i>Enterococcus faecalis</i>	ATCC®19433	WDCM00009	No growth

<i>Listeria monocytogenes</i>	ATCC®35152	WDCM00109	0.5-1.0mm blue colonies with halo
+ <i>Escherichia coli</i>	ATCC®8739	WDCM00012	No growth
+ <i>Enterococcus faecalis</i>	ATCC®29212	WDCM00087	No growth

<i>Listeria monocytogenes</i>	ATCC®35152	WDCM00109	0.5-1.0mm blue colonies with halo
+ <i>Escherichia coli</i>	ATCC®25922	WDCM00013	No growth
+ <i>Enterococcus faecalis</i>	ATCC®29212	WDCM00087	No growth

A satisfactory result is represented by recovery of >10 cfu of *Listeria monocytogenes* on Brilliance™ Listeria Agar (ISO).

Selectivity determined by qualitative testing based on the methods described in ISO 11133:2014

Inoculation with pure cultures

Inoculate 10ml quantities of medium to achieve 1E+03 to 1E+04 colony-forming units/ml (cfu/ml) of *Escherichia coli* and *Enterococcus faecalis*. Incubate broths at 30 ± 2°C for 25 ± 1 hour. Subculture onto Brilliance™ Listeria Agar (ISO) (CM1212, SR0257 & SR0258) and Tryptone Soya Agar (CM0131) and incubate plates at 37 ± 2°C for 24 ± 2 hours.


Reactions after incubation at 30 ± 2°C for 25 ± 1 hour

<i>Escherichia coli</i>	ATCC®8739	WDCM00012	No growth (CM1212, SR0257 & SR0258) No growth or cream colonies (CM0131)
<i>Escherichia coli</i>	ATCC®25922	WDCM00013	No growth (CM1212, SR0257 & SR0258) No growth or cream colonies (CM0131)

	Document Owner Department: QC	MBD-BT-SPEC-0511
		Page 5 of 5
		Rev 07
OXOID QUALITY ASSURANCE PRODUCT SPECIFICATION		
HALF FRASER SELECTIVE SUPPLEMENT SR0166E		

Revision History

Section / Step	Description of Change	Reason for Change	Reference
Microbiological characteristics	Change of Listeria plating medium	Change control	MOC-2023-0965

	Document Owner Department: QC	BT-SPEC-0047
		Page 1 of 3
OXOID QUALITY ASSURANCE PRODUCT SPECIFICATION		
NUTRIENT AGAR CM0003		

NUTRIENT AGAR

CM0003

Typical Formula*

'Lab-Lemco' powder	grams per litre	1.0
Yeast extract		2.0
Peptone		5.0
Sodium chloride		5.0
Agar		15.0

* adjusted as required to meet performance standards

Directions

Suspend 28g in 1 litre of distilled water. Bring to the boil to dissolve completely. Sterilize by autoclaving at 121°C for 15 minutes. Cool to 50°C. Mix well and pour into sterile Petri dishes.

Physical Characteristics

Straw, free-flowing powder
 Colour on reconstitution - straw 1-2
 Moisture level - less than 7%
 pH 7.4 ± 0.2 at 25°C
 Clarity - clear
 Gel strength - firm, comparable to 15.0g/litre of agar

The medium is tested for compatibility using 7% v/v oxalated horse blood, defibrinated horse blood or defibrinated sheep blood. There shall be no evidence of lysis or darkening, after incubation at 37°C, 25°C and 4°C for 72 hours.

Thermophiles shall be absent after incubation at 55°C for 3 days.

Microbiological Tests Using Optimum Inoculum Dilution

Control Medium: Nutrient Agar


Medium is challenged with 10-100 colony-forming units

Reactions after incubation at $37 \pm 2^\circ\text{C}$ for 24 ± 2 hours

Plain plates

<i>Staphylococcus aureus</i>	ATCC® 25923	1-2mm white/straw colonies
<i>Pseudomonas aeruginosa</i>	ATCC® 27853	1-3mm straw colonies

A satisfactory result is represented by recovery of positive strains equal to or greater than 70% of the control medium.

	Document Owner Department: QC	BT-SPEC-0047
		Page 2 of 3
OXOID QUALITY ASSURANCE PRODUCT SPECIFICATION		
NUTRIENT AGAR CM0003		

Enriched with 7% v/v horse blood

<i>Streptococcus pyogenes</i>	ATCC® 19615	0.25-1mm colourless colonies, β haemolysis
<i>Streptococcus pneumoniae</i>	ATCC® 6303	1-2mm grey/green colonies, α haemolysis
<i>Streptococcus pneumoniae</i>	ATCC® 6305	0.5-1mm grey/green colonies, α haemolysis

A satisfactory result is represented by recovery of positive strains equal to or greater than 70% of the control medium.

Reactions after incubation in 5% CO₂ at 37 ± 2°C for 24 ± 2 hours (for details, refer to Oxoid Manual - Atmosphere Generation Systems)

Enriched with 7% v/v horse blood

<i>Haemophilus influenzae</i>	ATCC® 19418	Pinpoint-0.25mm colourless colonies
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A satisfactory result is represented by recovery of positive strains equal to or greater than 70% of the control medium.

Testing performed in accordance with ISO11133:2014

Medium is challenged with 50-120 colony-forming units

Reactions after incubation at 30 ± 2°C for 24 ± 2 hours


<i>Yersinia enterocolitica</i>	ATCC® 23715	WDCM00160	1-3mm straw colonies
<i>Yersinia enterocolitica</i>	ATCC® 9610	WDCM00038	1-3mm straw colonies

A satisfactory result is represented by recovery of positive strains equal to or greater than 70% of the control medium.

Reactions after incubation at 37 ± 2°C for 24 ± 2 hours

<i>Escherichia coli</i>	ATCC® 25922	WDCM00013	1-3mm straw colonies
<i>Escherichia coli</i>	ATCC® 8739	WDCM00012	1-3mm straw colonies
<i>Salmonella typhimurium</i>	ATCC® 14028	WDCM00031	1-3mm straw colonies
<i>Salmonella enteritidis</i>	ATCC® 13076	WDCM00030	1-3mm straw colonies

A satisfactory result is represented by recovery of positive strains equal to or greater than 70% of the control medium.

	Document Owner Department: QC	BT-SPEC-0047
		Page 3 of 3
OXOID QUALITY ASSURANCE PRODUCT SPECIFICATION		
NUTRIENT AGAR CM0003		

Revision History

Section / Step	Description of Change	Reason for Change	Reference
Creation of ISO11133 section	Update to include testing of ISO11133:2014	Change control	BT-CC-1196

Distribution: Central File

Date: 27/08/14

Supersedes: 05/10/12

OXOID QUALITY ASSURANCE PRODUCT SPECIFICATION

UREA AGAR BASE

CM0053

Typical Formula*

Peptone	grams per litre	1.0
Glucose		1.0
Sodium chloride		5.0
Di-sodium phosphate		1.2
Potassium dihydrogen phosphate		0.8
Phenol red		0.012
Agar		15.0

* adjusted as required to meet performance standards

Directions

Suspend 2.4g in 95ml of distilled water. Bring to the boil to dissolve completely. Sterilize by autoclaving at 115°C for 20 minutes. Cool to 50°C and aseptically add the contents of 1 vial of Urea 40% Solution (SR0020K). Mix well, aseptically dispense 10ml amounts into sterile containers and allow to set in the slope position.

Physical Characteristics

Orange/pink, free-flowing powder
 Colour on reconstitution - orange
 Moisture level - less than 7%
 pH 6.8 ± 0.2 at 25°C
 Clarity - clear
 Gel strength - firm, comparable to 15.0g/litre of agar

Microbiological Tests Using Optimum Inoculum Dilution

Inoculate slopes of the medium with the test organisms.

Medium is challenged with 1E+04 to 1E+06 colony-forming units

Reactions after incubation at 30°C for 6 hours

<i>Morganella morganii</i>	ATCC® 25830	Weak urease +ve, pale pink slope, or negative
<i>Proteus mirabilis</i>	ATCC® 29906	Urease +ve, pink slope

Reactions after incubation at 30°C for 24 hours

<i>Morganella morganii</i>	ATCC® 25830	Urease +ve, pink slope
<i>Klebsiella pneumoniae</i>	ATCC® 13883	Weak urease +ve, pale pink slope
<i>Shigella sonnei</i>	ATCC® 25931	Urease -ve, no colour change
<i>Enterobacter aerogenes</i>	ATCC® 13048	Urease -ve, no colour change
<i>Escherichia coli</i>	ATCC® 25922	Urease -ve, no colour change
<i>Escherichia coli</i>	ATCC® 11775	Urease -ve, no colour change


Salmonella nottingham NCTC 7832 Urease -ve, no colour change

Reactions after incubation at 30°C for up to 72 hours

Candida albicans ATCC® 10231 Urease -ve, no colour change

Cryptococcus albidus ATCC® 34140 Weak urease +ve, pink slope

A satisfactory result is represented by reactions in accordance with the specification.

	Document Owner Department: QC	BT-SPEC-0118
		Page 1 of 3
OXOID QUALITY ASSURANCE PRODUCT SPECIFICATION		
BLOOD AGAR BASE NO.2 (CM0271)		

BLOOD AGAR BASE NO.2

CM0271

Typical Formula*

Proteose peptone	grams per litre	15.0
Liver digest		2.5
Yeast extract		5.0
Sodium chloride		5.0
Agar		12.0

* adjusted as required to meet performance standards

Directions

Suspend 40g in 1 litre of distilled water. Bring to the boil to dissolve completely. Sterilize by autoclaving at 121°C for 15 minutes. Cool to 50°C. Mix well and pour into sterile Petri dishes. For blood agar, enrich with 7% v/v sterile defibrinated blood.

Physical Characteristics

Straw, free-flowing powder
 Colour on reconstitution - straw 2-3
 Moisture level - less than 7%
 pH 7.4 ± 0.2 at 25°C
 Clarity - clear
 Gel strength - firm, comparable to 12.0g/litre of agar

The medium is tested for compatibility using 7% v/v oxalated horse blood, defibrinated horse blood or defibrinated sheep blood. There shall be no evidence of lysis or darkening, after incubation at 37°C, 25°C and 4°C for 72 hours.

ThermoFisher SCIENTIFIC	Document Owner Department: QC	BT-SPEC-0118 Page 2 of 3
OXOID QUALITY ASSURANCE PRODUCT SPECIFICATION		
BLOOD AGAR BASE NO.2 (CM0271)		

Microbiological Tests Using Optimum Inoculum Dilution

Control Medium: Blood Agar Base No.2

Medium is challenged with 10-100 colony-forming units

Reactions after incubation at 37°C for 18 hours

Plain plates

<i>Staphylococcus aureus</i>	ATCC® 25923	1-2mm cream colonies
<i>Staphylococcus aureus</i>	ATCC® 9144	0.5-1.5mm cream colonies
<i>Pseudomonas aeruginosa</i>	ATCC® 27853	1-4mm irregular straw or straw/green colonies
<i>Escherichia coli</i>	ATCC® 25922	1-2mm straw colonies

Enriched with 7% v/v horse blood

<i>Streptococcus pyogenes</i>	ATCC® 19615	0.25-1mm pale straw colonies, β haemolysis
<i>Streptococcus pneumoniae</i>	ATCC® 6303	1-4mm grey/green colonies, α haemolysis
<i>Streptococcus pneumoniae</i>	ATCC® 6305	pinpoint-0.25mm grey/green colonies, α haemolysis
<i>Haemophilus influenzae</i>	ATCC® 19418	pinpoint-0.25mm colourless colonies


NOTE: White centre may be present in colonies of *Streptococcus pneumoniae* when grown in aerobic conditions which is due to hydrogen peroxide production.

A satisfactory result is represented by recovery of positive strains equal to or greater than 70% of the control medium.

Zones of growth/no growth surrounding X, V and X+V factor discs (DD0003, DD0004 and DD0005) when plain plates are inoculated with the following organisms and incubated at 37°C for 18 hours:


		X	V	X+V
<i>Haemophilus influenzae</i>	ATCC® 49247	0	0	≥ 15mm
<i>Haemophilus parainfluenzae</i>	ATCC® 33392	0	≥ 20mm	≥ 20mm

Zones of inhibition with Bacitracin discs (DD0002) shall be 10-20mm when 7% v/v horse blood plates are inoculated with *Streptococcus pyogenes* ATCC® 19615 and incubated at 37°C for 18 hours.

	Document Owner Department: QC	BT-SPEC-0118
		Page 3 of 3
OXOID QUALITY ASSURANCE PRODUCT SPECIFICATION		
BLOOD AGAR BASE NO.2 (CM0271)		

Revision History

Section / Step	Description of Change	Reason for Change	Reference
Entire Document	<ul style="list-style-type: none"> Update to new document template Inclusion of statement regarding <i>S. pneumoniae</i> growth 	Change control	BT-CC-1845

	Document Owner Department: QC	BT-SPEC-0120
		Page 1 of 4
OXOID QUALITY ASSURANCE PRODUCT SPECIFICATION		
TRIPLE SUGAR IRON AGAR (ISO) (CM0277)		

TRIPLE SUGAR IRON AGAR (ISO)

CM0277

Typical Formula*

Meat extract	grams per litre	3.0
Yeast extract		3.0
Peptone		20.0
Sodium chloride		5.0
Lactose		10.0
Sucrose		10.0
Glucose		1.0
Iron (III) citrate		0.3
Sodium thiosulphate		0.3
Phenol red		0.024
Agar		12.0


* adjusted as required to meet performance standards

Directions

Suspend 65g in 1 litre of distilled water. Bring to the boil to dissolve completely. Mix well and distribute into final containers. Sterilize by autoclaving at 121°C for 15 minutes. Allow to set as slopes with 2.5cm butts.

Physical Characteristics

Straw, free-flowing powder
 Colour on reconstitution - red
 Moisture level - less than 7%
 pH 7.4 ± 0.2 at 25°C
 Clarity - clear
 Gel strength - firm, comparable to 12.0g/litre of agar

	Document Owner Department: QC	BT-SPEC-0120
		Page 2 of 4
OXOID QUALITY ASSURANCE PRODUCT SPECIFICATION		
TRIPLE SUGAR IRON AGAR (ISO) (CM0277)		

Microbiological Tests Using Optimum Inoculum Dilution


Reactions after incubation at 37°C for 18 hours

Medium is challenged with 1E+04 to 1E+06 colony-forming units

	Slope	Butt	H₂S	Gas
<i>Escherichia coli</i> ATCC® 8739	A	AG	Negative	Positive
<i>Escherichia coli</i> ATCC® 11775	A	AG	Negative	Positive
<i>Shigella sonnei</i> ATCC® 25931	NC	A	Negative	Negative
<i>Salmonella enteritidis</i> ATCC® 13076	NC	AG	Positive	Positive
	Slope	Butt	H₂S	Gas
<i>Salmonella typhimurium</i> ATCC® 14028	NC	AG	Positive	Positive
<i>Salmonella abony</i> NCTC 6017	NC	AG	Positive	Positive
<i>Salmonella nottingham</i> NCTC 7832	NC	AG	Positive	Positive
<i>Proteus hauseri</i> ATCC® 13315	A	A	Positive	Negative
<i>Enterobacter aerogenes</i> ATCC® 13048	A	AG	Negative	Positive
<i>Pseudomonas aeruginosa</i> ATCC® 9027	Alk	Alk	Negative	Negative
<i>Staphylococcus aureus</i> ATCC® 6538	A	A	Negative	Negative

A satisfactory result is represented by reactions in accordance with the specification.

Equivalent results obtained after incubation at 30-35°C for 18-24 hours.

	Document Owner Department: QC	BT-SPEC-0120
		Page 3 of 4
OXOID QUALITY ASSURANCE PRODUCT SPECIFICATION		
TRIPLE SUGAR IRON AGAR (ISO) (CM0277)		

Key


AG	=	Acid (yellow) with gas formation
A	=	Acid (yellow)
NC	=	No change
Alk	=	Alkaline (red)

Hydrogen sulphide (H₂S)

Positive	=	Blackening
Negative	=	No blackening


Gas

Positive	=	Bubbles or splitting of agar
Negative	=	No bubbles or splitting of agar

	Document Owner Department: QC	BT-SPEC-0120
		Page 4 of 4
OXOID QUALITY ASSURANCE PRODUCT SPECIFICATION		
TRIPLE SUGAR IRON AGAR (ISO) (CM0277)		

Revision History

Section / Step	Description of Change	Reason for Change	Reference
N/A	Update to template and ISO	Change control	BT-CC-1901

	Document Owner Department: QC	BT-SPEC-0156
		Page 1 of 5
OXOID QUALITY ASSURANCE PRODUCT SPECIFICATION		
X.L.D. MEDIUM CM0469		

X.L.D. MEDIUM

CM0469

Typical Formula*

Yeast extract	grams per litre	3.0
L-Lysine HCl		5.0
Xylose		3.75
Lactose		7.5
Sucrose		7.5
Sodium desoxycholate		1.0
Sodium chloride		5.0
Sodium thiosulphate		6.8
Ammonium iron (III) citrate		0.8
Phenol red		0.08
Agar		12.5

* adjusted as required to meet performance standards

Directions

Suspend 53g in 1 litre of distilled water. With frequent agitation, bring to the boil to dissolve completely. Cool to 50°C. Mix well and pour into sterile Petri dishes. DO NOT AUTOCLAVE. DO NOT OVERHEAT.

Physical Characteristics

Straw/pink, free-flowing powder
 Colour on reconstitution - red
 Moisture level - less than or equal to 7%
 pH 7.4 ± 0.2 at 25°C
 Clarity - clear
 Gel strength - firm, comparable to 12.5g/litre of agar

Microbiological Tests Using Optimum Inoculum Dilution

Control Medium: Tryptone Soya Agar

Reactions after incubation at 37 ± 2°C for 24 ± 3 hours

Inoculation with mixed cultures using diminishing sweep technique

Medium is challenged with 1E+03 to 1E+05 colony-forming units (cfu) of *Salmonella* and *Shigella* spp. and 1E+05 to 1E+07 cfu for *Escherichia coli* ATCC® 8739.

Salmonella abony

NCTC6017

1-3mm red colonies, black centre

ThermoFisher SCIENTIFIC	Document Owner Department: QC	BT-SPEC-0156
		Page 2 of 5
OXOID QUALITY ASSURANCE PRODUCT SPECIFICATION		
X.L.D. MEDIUM CM0469		

<i>Salmonella enteritidis</i>	ATCC® 13076	1-2mm red colonies, black centre
<i>Salmonella typhimurium</i>	ATCC® 14028	1-2mm red colonies, black centre
<i>Salmonella virchow</i>	NCTC5742	1-2mm red colonies, black centre
<i>Salmonella arizonae</i>	ATCC® 13314	1-3mm red colonies, black centre
<i>Salmonella nottingham</i>	NCTC7832	1-3mm red colonies, black centre
<i>Shigella sonnei</i>	ATCC® 9290	0.5-7mm irregular/smooth red colonies
<i>Shigella flexneri</i>	ATCC® 12022	0.5-2mm irregular, red colonies

In mixed culture, using the diminishing sweep technique, a satisfactory result is represented by diagnostic reactions of *Salmonellae* and *Shigellae* strains and *Escherichia coli*. Clear differentiation must be seen and is based on the colour and morphology of the colonies.

Inoculation with pure cultures

Medium is challenged with 10-100 colony-forming units

<i>Pseudomonas aeruginosa</i>	ATCC® 9027	No growth or 0.5-2mm red colonies
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For *Pseudomonas aeruginosa* ATCC® 9027, a satisfactory result is represented by recovery equal to or less than 90% of the control medium.

<i>Proteus mirabilis</i>	ATCC® 12453	0.5-2mm orange/red colonies, with or without black centre, no swarming
<i>Proteus mirabilis</i>	ATCC® 29906	0.5-2mm orange/red colonies, with or without black centre, no swarming
<i>Serratia marcescens</i>	ATCC® 8100	1-2mm orange/yellow colonies
<i>Citrobacter freundii</i>	ATCC® 8090	0.5-2mm yellow colonies
<i>Klebsiella pneumoniae</i>	ATCC® 29665	2-4mm yellow, mucoid colonies

Other pure cultures are inhibited or shall produce colonies with a negative diagnostic reaction.

Medium is challenged with 1E+04 to 1E+06 colony-forming units

<i>Staphylococcus aureus</i>	ATCC® 6538	No growth
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Negative strains are inhibited.

Inoculation using diminishing sweep technique

Medium is challenged with 1E+04 to 1E+06 colony-forming units

<i>Escherichia coli</i>	ATCC® 11775	No growth or 0.5-4mm yellow colonies
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Escherichia coli ATCC® 11775 is inhibited or shall produce colonies with a negative diagnostic reaction.

	Document Owner Department: QC	BT-SPEC-0156
		Page 3 of 5
OXOID QUALITY ASSURANCE PRODUCT SPECIFICATION		
X.L.D. MEDIUM CM0469		

Medium is challenged with 1E+04 to 1E+06 colony-forming units

Shigella sonnei ATCC®25931 0.5-7mm irregular/smooth red colonies

Shigella sonnei ATCC®25931 shall produce colonies with a positive diagnostic reaction.

Equivalent results are obtained after incubation at 30-35°C for 24 hours.

Testing performed in accordance with ISO11133:2014

Reactions after incubation at 37 ± 2°C for 24 ± 3 hours

Medium is challenged with 50-120 colony-forming units

<i>Salmonella enteritidis</i>	ATCC®13076	WDCM00030	1-3mm red colonies, black centre
<i>Salmonella typhimurium</i>	ATCC®14028	WDCM00031	1-3mm red colonies, black centre

A satisfactory result is represented by recovery of equal to or greater than 70% of the control medium.

Medium is challenged with 1E+04 to 1E+05 colony-forming units

<i>Escherichia coli</i>	ATCC®8739	WDCM00012	No growth or 0.5-4mm yellow cols
<i>Escherichia coli</i>	ATCC®25922	WDCM00013	No growth or 0.5-4mm yellow cols


Inhibited strains shall produce no growth or at least a 1 log (10) reduction with a negative diagnostic reaction when compared to the control medium.

Inoculation using diminishing sweep technique

Medium is challenged with 1E+04 to 1E+06 colony-forming units

<i>Enterococcus faecalis</i>	ATCC®29212	WDCM00087	No growth
<i>Enterococcus faecalis</i>	ATCC®19433	WDCM00009	No growth

Negative strains are inhibited.

	Document Owner Department: QC	BT-SPEC-0156
		Page 4 of 5
OXOID QUALITY ASSURANCE PRODUCT SPECIFICATION		
X.L.D. MEDIUM CM0469		

Testing performed in accordance with current CLSI M22 A

Reactions after incubation at 35°C for 18-24 hours

Medium is challenged with 10-100 colony-forming units

<i>Shigella flexneri</i>	ATCC®12022	0.5-2mm irregular, red colonies
<i>Salmonella typhimurium</i>	ATCC®14028	1-2mm red colonies, black centre

A satisfactory result is represented by recovery of equal to or greater than 70% of the control medium.

Medium is challenged with 1E+04 to 1E+06 colony-forming units


<i>Enterococcus faecalis</i>	ATCC®29212	No growth
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Negative strains are inhibited.

Medium is challenged with 1E+04 to 1E+06 colony-forming units


<i>Escherichia coli</i>	ATCC®25922	No growth or 0.5-4mm yellow cols
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Inhibited strains shall produce no growth or at least a 1 log (10) reduction with a negative diagnostic reaction when compared to the control medium.

	Document Owner Department: QC	BT-SPEC-0156
		Page 5 of 5
OXOID QUALITY ASSURANCE PRODUCT SPECIFICATION		
X.L.D. MEDIUM CM0469		

Revision History

Section / Step	Description of Change	Reason for Change	Reference
Microbiological Tests	Update to specification for <i>Shigella sonnei</i>	Change control	BT-CC-1911
Microbiological Tests	Salmonella and Shigella mixed culture testing changed from low number quantitative to high number qualitative testing.	Change control	BT-CC-2398

	Document Owner Department: QC	BT-SPEC-0200
		Page 1 of 3
OXOID QUALITY ASSURANCE PRODUCT SPECIFICATION		
CAMPYLOBACTER BLOOD-FREE SELECTIVE AGAR BASE CM0739		

CAMPYLOBACTER BLOOD-FREE SELECTIVE AGAR BASE

CM0739

Typical Formula*

Nutrient Broth No. 2	grams per litre	25.0
Activated carbon		4.0
Casein hydrolysate		3.0
Sodium desoxycholate		1.0
Iron (II) sulphate		0.25
Sodium pyruvate		0.25
Agar		12.0

*adjusted to meet performance standards

Directions

Suspend 22.75g in 500ml of distilled water. Bring to the boil to dissolve completely. Sterilize by autoclaving at 121°C for 15 minutes. Cool to 50°C and aseptically add the contents of 1 vial of CCDA Selective Supplement (SR0155E) reconstituted as directed. Mix well and pour into sterile Petri dishes.

Physical Characteristics

Black, free-flowing powder
 Colour on reconstitution - black
 pH 7.4 ± 0.2 at 25°C
 Clarity - opaque
 Gel strength – firm, comparable to 12g/litre of agar

Microbiological Tests Using Optimum Inoculum Dilution


Control Medium : Columbia Blood Agar Base enriched with 7% v/v laked horse blood and Campylobacter Growth Supplement SR0232

Reactions after incubation at 37 ± 2°C for 48 hours under microaerophilic conditions

Tested with the addition of CCDA Selective Supplement SR0155

Medium is challenged with 10-100 colony-forming units

<i>Campylobacter jejuni</i>	ATCC®33560	0.5-2mm grey colonies
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	Document Owner Department: QC	BT-SPEC-0200
		Page 2 of 3
OXOID QUALITY ASSURANCE PRODUCT SPECIFICATION		
CAMPYLOBACTER BLOOD-FREE SELECTIVE AGAR BASE CM0739		

A satisfactory result is represented by recovery of positive strains equal to or greater than 50% of the control medium.

Medium is challenged with 1E+04 to 1E+06 colony-forming units

<i>Campylobacter lari</i>	ATCC®35221	0.5-2mm grey colonies
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For *Campylobacter lari* ATCC®35221, a satisfactory result is represented by growth and a positive diagnostic reaction in accordance with the specification.

Testing performed in accordance with ISO11133:2014

Reactions after incubation at 41.5 ± 2°C for 44 ± 4 hours

Medium is challenged with 50-120 colony-forming units


<i>Campylobacter jejuni</i>	ATCC®29428	WDCM00156	0.5-2mm grey colonies
<i>Campylobacter jejuni</i>	ATCC®33291	WDCM00005	0.5-2mm grey colonies
<i>Campylobacter coli</i>	ATCC®43478	WDCM00004	0.5-2mm grey colonies

A satisfactory result is represented by recovery of positive strains equal to or greater than 50% of the control medium.

Medium is challenged with 1E+04 to 1E+06 colony-forming units


<i>Escherichia coli</i>	ATCC®25922	WDCM00013	No growth
<i>Escherichia coli</i>	ATCC®8739	WDCM00012	No growth
<i>Staphylococcus aureus</i>	ATCC®25923	WDCM00034	No growth

Negative strains are inhibited.

	Document Owner Department: QC	BT-SPEC-0200
		Page 3 of 3
OXOID QUALITY ASSURANCE PRODUCT SPECIFICATION		
CAMPYLOBACTER BLOOD-FREE SELECTIVE AGAR BASE CM0739		

Revision History

Section / Step	Description of Change	Reason for Change	Reference
Physical Characteristics	Removal of moisture value	Change control	BT-CC-1617
Microbiological Characteristics	Change of testing for <i>Campylobacter lari</i> ATCC®35221 changed from low number quantitative to high number qualitative testing.	Change control	BT-CC-2939

	Document Owner Department: QC	BT-SPEC-0211
		Page 1 of 3
OXOID QUALITY ASSURANCE PRODUCT SPECIFICATION		
LISTERIA SELECTIVE AGAR BASE (OXFORD FORMULATION) CM0856		

LISTERIA SELECTIVE AGAR BASE (OXFORD FORMULATION)

CM0856

Typical Formula*

	grams per litre
Columbia Blood Agar Base	39.0
Aesculin	1.0
Ferric ammonium citrate	0.5
Lithium chloride	15.0

* adjusted as required to meet performance standards

Directions

Suspend 27.75g in 500ml of distilled water. Bring to the boil to dissolve completely. Sterilize by autoclaving at 121°C for 15 minutes. Cool to 50°C and aseptically add the contents of 1 vial of Listeria Selective Supplement (SR0206E or SR0140E) reconstituted as directed. Mix well and pour into sterile Petri dishes.

Physical Characteristics

Straw, free-flowing powder
 Colour on reconstitution - pale green
 Moisture level - less than or equal to 7%
 pH - 7.0 ± 0.2 at 25°C
 Clarity - clear
 Gel strength - firm, comparable to 10.0g/litre of agar

Microbiological Tests Using Optimum Inoculum Dilution

Control Medium: Columbia Blood Agar Base enriched with 5% v/v horse blood


Reactions after incubation at 37°C for 48 hours

Tested with the addition of Listeria Selective Supplement (Oxford Formulation) SR0140

Medium is challenged with 10-100 colony-forming units

<i>Listeria monocytogenes</i>	ATCC® 7644	0.25-1.0mm brown/black dimpled colonies and halo
<i>Listeria monocytogenes</i>	ATCC® 13932	0.25-1.0mm brown/black dimpled colonies and halo

A satisfactory result is represented by recovery of positive strains equal to or greater than 50% of the control medium.

	Document Owner Department: QC	BT-SPEC-0211
		Page 2 of 3
OXOID QUALITY ASSURANCE PRODUCT SPECIFICATION		
LISTERIA SELECTIVE AGAR BASE (OXFORD FORMULATION) CM0856		

Medium is challenged with 10-100 colony-forming units

Staphylococcus aureus ATCC®25923 No growth or pinpoint-1.5mm yellow colonies

Staphylococcus aureus ATCC®25923 is inhibited or shall produce a negative diagnostic reaction from an inoculum of 10-100 cfu

Medium is challenged with 1E+04 to 1E+06 colony-forming units

Enterococcus faecalis ATCC®29212 No growth


Enterococcus faecalis ATCC®19433 No growth

Escherichia coli ATCC®25922 No growth

Escherichia coli ATCC®8739 No growth


Candida albicans ATCC®10231 No growth or minimal growth

Negative strains are inhibited. *Candida albicans* ATCC®10231 shall be inhibited or produce pinpoint colourless colonies with no blackening of the media.

	Document Owner Department: QC	BT-SPEC-0211
		Page 3 of 3
OXOID QUALITY ASSURANCE PRODUCT SPECIFICATION		
LISTERIA SELECTIVE AGAR BASE (OXFORD FORMULATION) CM0856		

Revision History

Section / Step	Description of Change	Reason for Change	Reference
Microbiological characteristics	Change to <i>Staphylococcus aureus</i> growth characteristics	Change control	MOC-2022-0180

	Document Owner Department: QC	MBD-BT-SPEC-0220
		Page 1 of 5
		Rev 06
OXOID QUALITY ASSURANCE PRODUCT SPECIFICATION		
FRASER BROTH BASE (ISO) CM0895		

FRASER BROTH BASE (ISO)

CM0895

Typical Formula*

Proteose peptone	grams per litre	5.0
Tryptone		5.0
Meat extract		5.0
Yeast extract		5.0
Sodium chloride		20.0
Di-sodium hydrogen phosphate		12.0
Potassium dihydrogen phosphate		1.35
Aesculin		1.0
Lithium chloride		3.0

* adjusted as required to meet performance standards

Directions

To make Half Fraser Broth

Dissolve 12.9g in 225ml of distilled water. Sterilize by autoclaving at 121°C for 15 minutes. Cool to 50°C and aseptically add the contents of 1 vial of Half Fraser Selective Supplement (SR0166E) reconstituted as directed. Mix well and dispense into sterile containers.

Alternatively, dissolve 129.2g in 2.25 litres of distilled water. Sterilize by autoclaving at 121°C for 15 minutes. Cool to 50°C and aseptically add the contents of 1 vial of Half Fraser Selective Supplement (SR0166G) reconstituted as directed. Mix well and dispense into sterile containers.

To make Fraser Broth

Dissolve 28.7g in 500ml of distilled water. Sterilize by autoclaving at 121°C for 15 minutes. Cool to 50°C and aseptically add the contents of 1 vial of Fraser Selective Supplement (SR0156E) reconstituted as directed. Mix well and dispense into sterile containers.

Physical Characteristics


Straw, free-flowing powder

Colour on reconstitution - straw 2-3

Moisture level - less than or equal to 7%

pH 7.2 ± 0.2 at 25°C

Clarity - clear

	Document Owner Department: QC	MBD-BT-SPEC-0220
		Page 2 of 5
		Rev 06
OXOID QUALITY ASSURANCE PRODUCT SPECIFICATION		
FRASER BROTH BASE (ISO) CM0895		

Microbiological Tests Using Optimum Inoculum Dilution

Control Media: Brilliance™ Listeria Agar (ISO) or Columbia Blood Agar Base enriched with 5% v/v horse blood, where appropriate.

Tested with the addition of Fraser Selective Supplement SR0156

Reactions after incubation at 37 ± 2°C for 24 ± 2 hours

Inoculate 10ml quantities of medium to achieve 1-10 colony-forming units/ml (cfu/ml) of *Listeria* spp. Incubate broths at 37 ± 2°C for 24 ± 2 hours. Subculture onto Brilliance™ Listeria Agar (ISO) (CM1212, SR0257 & SR0258) and incubate plates at 37 ± 2°C for 24 - 48 hours.

Listeria monocytogenes ATCC®7644

Listeria monocytogenes ATCC®13932

A satisfactory result is represented by recovery of positive strains equal to or greater than a 3 log(10) increase.

Positive strains shall produce aesculin hydrolysis after 48 hours.

Reactions after incubation at 37 ± 2°C for 48 ± 2 hours

Inoculate 10ml quantities of medium to achieve 1E+03 to 1E+04 cfu/ml. Incubate broths at 37°C for 48 hours.

Bacillus cereus ATCC®10876 No aesculin hydrolysis (no blackening)


Negative strains shall produce no aesculin hydrolysis after 48 hours.

Testing performed in accordance with ISO11133:2014

Tested with the addition of Fraser Selective Supplement SR0156. For testing with the addition of Half Fraser Selective Supplement SR0166 refer to Half Fraser Selective Supplement SR0166E or SR0166G product specifications.

Inoculation with mixed cultures

Inoculate 10ml quantities of medium to achieve 1-10 colony-forming units/ml (cfu/ml) of *Listeria* spp., to each add 1E+02 to 1E+03 cfu/ml of *Escherichia coli* and 1E+02 to 1E+03 cfu/ml of


	Document Owner Department: QC	MBD-BT-SPEC-0220
		Page 3 of 5
		Rev 06
OXOID QUALITY ASSURANCE PRODUCT SPECIFICATION		
FRASER BROTH BASE (ISO) CM0895		

Enterococcus faecalis. Incubate broths at $37 \pm 2^{\circ}\text{C}$ for 24 ± 2 hours. Subculture onto Brilliance™ Listeria Agar (ISO) (CM1212, SR0257 & SR0258) and incubate plates at $37 \pm 2^{\circ}\text{C}$ for 24 ± 2 hours.

Reactions after incubation at $37 \pm 2^{\circ}\text{C}$ for 24 ± 2 hours

<i>Listeria monocytogenes</i>	ATCC®13932	WDCM00021	0.5-1.0mm blue colonies with halo
+ <i>Escherichia coli</i>	ATCC®8739	WDCM00012	No growth
+ <i>Enterococcus faecalis</i>	ATCC®19433	WDCM00009	No growth
<i>Listeria monocytogenes</i>	ATCC®13932	WDCM00021	0.5-1.0mm blue colonies with halo
+ <i>Escherichia coli</i>	ATCC®25922	WDCM00013	No growth
+ <i>Enterococcus faecalis</i>	ATCC®19433	WDCM00009	No growth
<i>Listeria monocytogenes</i>	ATCC®13932	WDCM00021	0.5-1.0mm blue colonies with halo
+ <i>Escherichia coli</i>	ATCC®8739	WDCM00012	No growth
+ <i>Enterococcus faecalis</i>	ATCC®29212	WDCM00087	No growth
<i>Listeria monocytogenes</i>	ATCC®13932	WDCM00021	0.5-1.0mm blue colonies with halo
+ <i>Escherichia coli</i>	ATCC®25922	WDCM00013	No growth
+ <i>Enterococcus faecalis</i>	ATCC®29212	WDCM00087	No growth
<i>Listeria monocytogenes</i>	ATCC®35152	WDCM00109	0.5-1.0mm blue colonies with halo
+ <i>Escherichia coli</i>	ATCC®8739	WDCM00012	No growth
+ <i>Enterococcus faecalis</i>	ATCC®19433	WDCM00009	No growth
<i>Listeria monocytogenes</i>	ATCC®35152	WDCM00109	0.5-1.0mm blue colonies with halo
+ <i>Escherichia coli</i>	ATCC®25922	WDCM00013	No growth
+ <i>Enterococcus faecalis</i>	ATCC®19433	WDCM00009	No growth
<i>Listeria monocytogenes</i>	ATCC®35152	WDCM00109	0.5-1.0mm blue colonies with halo
+ <i>Escherichia coli</i>	ATCC®8739	WDCM00012	No growth
+ <i>Enterococcus faecalis</i>	ATCC®29212	WDCM00087	No growth
<i>Listeria monocytogenes</i>	ATCC®35152	WDCM00109	0.5-1.0mm blue colonies with halo
+ <i>Escherichia coli</i>	ATCC®25922	WDCM00013	No growth
+ <i>Enterococcus faecalis</i>	ATCC®29212	WDCM00087	No growth

A satisfactory result is represented by recovery of >10 cfu of *Listeria monocytogenes* on Brilliance™ Listeria Agar (ISO).

	Document Owner Department: QC	MBD-BT-SPEC-0220
		Page 4 of 5
		Rev 06
OXOID QUALITY ASSURANCE PRODUCT SPECIFICATION		
FRASER BROTH BASE (ISO) CM0895		


Inoculation with pure cultures

Inoculate 10ml quantities of medium to achieve 1E+03 to 1E+04 colony-forming units/ml (cfu/ml) of *Escherichia coli* and *Enterococcus faecalis*. Incubate broths at 37 ± 2°C for 24 ± 2 hours. Subculture onto Brilliance™ Listeria Agar (ISO) (CM1212, SR0257 & SR0258) and Tryptone Soya Agar (CM0131) then incubate plates at 37 ± 2°C for 24 ± 2 hours.

Reactions after incubation at 37 ± 2°C for 24 ± 2 hours


<i>Escherichia coli</i>	ATCC® 8739	WDCM00012	No growth (CM1212, SR0257 & SR0258)
<i>Escherichia coli</i>	ATCC® 8739	WDCM00012	No growth or cream colonies (CM0131)
<i>Escherichia coli</i>	ATCC® 25922	WDCM00013	No growth (CM1212, SR0257 & SR0258)
<i>Escherichia coli</i>	ATCC® 25922	WDCM00013	No growth or cream colonies (CM0131)
<i>Enterococcus faecalis</i>	ATCC® 19433	WDCM00009	No growth (CM1212, SR0257 & SR0258)
<i>Enterococcus faecalis</i>	ATCC® 19433	WDCM00009	No growth or straw colonies (CM0131)
<i>Enterococcus faecalis</i>	ATCC® 29212	WDCM00087	No growth (CM1212, SR0257 & SR0258)
<i>Enterococcus faecalis</i>	ATCC® 29212	WDCM00087	No growth or straw colonies (CM0131)

A satisfactory result is represented by no growth of *Escherichia coli* and *Enterococcus faecalis* on Brilliance™ Listeria Agar (ISO) and <100 cfu on Tryptone Soya Agar.

	Document Owner Department: QC	MBD-BT-SPEC-0220
		Page 5 of 5
		Rev 06
OXOID QUALITY ASSURANCE PRODUCT SPECIFICATION		
FRASER BROTH BASE (ISO) CM0895		

Revision History

Section / Step	Description of Change	Reason for Change	Reference
Microbiological Characteristics	Change of Listeria plating medium	Change control	MOC-2023-0965

	Document Owner Department: QC	BT-SPEC-0228
		Page 1 of 4
OXOID QUALITY ASSURANCE PRODUCT SPECIFICATION		
TRYPTONE BILE X-GLUCURONIDE MEDIUM (TBX) CM0945		

TRYPTONE BILE X-GLUCURONIDE MEDIUM (TBX)

CM0945

Typical Formula*

Tryptone	grams per litre	20.0
Bile salts No.3		1.5
X-glucuronide		0.075
Agar		15.0

* adjusted as required to meet performance standards

Directions

Suspend 36.6g in 1 litre of distilled water. Bring gently to the boil to dissolve completely. Sterilize by autoclaving at 121°C for 15 minutes. Cool to 50°C and pour 15ml of the medium into sterile Petri dishes or hold at 45°C when using the pour plate technique.

Physical Characteristics

Straw, free-flowing powder
 Colour on reconstitution - straw 2-3
 Moisture level - less than 7%
 pH 7.2 ± 0.2 at 25°C
 Clarity - clear
 Gel strength - firm, comparable to 15.0g/litre of agar

ThermoFisher SCIENTIFIC	Document Owner Department: QC	BT-SPEC-0228 Page 2 of 4
OXOID QUALITY ASSURANCE PRODUCT SPECIFICATION		
TRYPTONE BILE X-GLUCURONIDE MEDIUM (TBX) CM0945		

Microbiological Tests Using Optimum Inoculum Dilution

Control Medium: Tryptone Soya Agar

Reactions after incubation at $44 \pm 2^\circ\text{C}$ for 21 ± 3 hours

Stack all plates not more than 3 high in plastic bags containing damp cotton wool, seal bags with tape.

Inoculation using pour plate technique

Medium is challenged with 30-100 colony-forming units

<i>Escherichia coli</i>	ATCC®11775	1-2mm blue/green colonies
<i>Klebsiella pneumoniae</i>	ATCC®29665	1-2mm straw colonies

A satisfactory result for *Escherichia coli* strains is represented by recovery equal to or greater than 70% of the control medium.

For *Klebsiella pneumoniae* ATCC®29665, a satisfactory result is represented by recovery of positive strains equal to or greater than 50% of the control medium.

Medium is challenged with $1\text{E}+04$ to $1\text{E}+06$ colony-forming units

<i>Klebsiella aerogenes</i>	NCTC9528	No growth
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Negative strains are inhibited.

Testing performed in accordance with ISO11133: 2014


Reactions after incubation at $44 \pm 2^\circ\text{C}$ for 21 ± 3 hours

Inoculation using pour plate technique

Medium is challenged with 50-120 colony-forming units

<i>Escherichia coli</i>	ATCC®25922	WDCM00013	1-2mm blue/green colonies
<i>Escherichia coli</i>	ATCC®8739	WDCM00012	1-2mm blue/green colonies
<i>Escherichia coli</i>	NCTC13216	WDCM00202	1-2mm blue/green colonies

A satisfactory result for *Escherichia coli* strains is represented by recovery equal to or greater than 70% of the control medium.

	Document Owner Department: QC	BT-SPEC-0228
		Page 3 of 4
OXOID QUALITY ASSURANCE PRODUCT SPECIFICATION		
TRYPTONE BILE X-GLUCURONIDE MEDIUM (TBX) CM0945		

Inoculation using surface plate technique

Medium is challenged with 50-120 colony-forming units

<i>Escherichia coli</i>	ATCC®25922	WDCM00013	1-2mm blue/green colonies
<i>Escherichia coli</i>	ATCC®8739	WDCM00012	1-2mm blue/green colonies
<i>Escherichia coli</i>	NCTC13216	WDCM00202	1-2mm blue/green colonies

A satisfactory result for *Escherichia coli* strains is represented by recovery equal to or greater than 70% of the control medium.

Medium is challenged with 1E+04 to 1E+06 colony-forming units

<i>Enterococcus faecalis</i>	ATCC®19433	WDCM00009	No growth
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Negative strains are inhibited.

Medium is challenged with 1E+04 to 1E+05 colony-forming units

<i>Enterococcus faecalis</i>	ATCC®29212	WDCM00087	No growth
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Negative strains are inhibited.


Medium is challenged with 1E+03 to 1E+04 colony-forming units

<i>Citrobacter freundii</i>	ATCC®43864	WDCM00006	No growth or 1-2mm white to green/beige colonies
<i>Pseudomonas aeruginosa</i>	ATCC®27853	WDCM00025	No growth or 1-2mm white to green/beige colonies

Inoculation using membrane filtration technique

Medium is challenged with 50-120 colony-forming units


<i>Escherichia coli</i>	ATCC®25922	WDCM00013	1-2mm blue/green colonies
<i>Escherichia coli</i>	ATCC®8739	WDCM00012	1-2mm blue/green colonies
<i>Escherichia coli</i>	NCTC13216	WDCM00202	1-2mm blue/green colonies

	Document Owner Department: QC	BT-SPEC-0228
		Page 4 of 4
OXOID QUALITY ASSURANCE PRODUCT SPECIFICATION		
TRYPTONE BILE X-GLUCURONIDE MEDIUM (TBX) CM0945		

A satisfactory result for *Escherichia coli* strains is represented by recovery equal to or greater than 70% of the control medium.

Revision History

Section / Step	Description of Change	Reason for Change	Reference
Entire Document	Reformatting to new template Update to test specification	Change control	BT-CC-1561
Entire document	Change title typographical error. Addition of <i>Klebsiella aerogenes</i> NCTC9528	Change control	BT-CC-2204

	Document Owner Department: QC	BT-SPEC-0287
		Page 1 of 5
OXOID QUALITY ASSURANCE PRODUCT SPECIFICATION		
BRAIN HEART INFUSION CM1135		

BRAIN HEART INFUSION

CM1135

Typical Formula*

Brain infusion solids	grams per litre	12.5
Beef heart infusion solids		5.0
Proteose peptone		10.0
Glucose		2.0
Sodium chloride		5.0
Di-sodium phosphate		2.5

* adjusted as required to meet performance standards

Directions

Dissolve 37g in 1 litre of distilled water. Mix well and distribute into final containers. Sterilize by autoclaving at 121°C for 15 minutes.

Physical Characteristics

Straw, free-flowing powder
 Colour on reconstitution - straw 3-4
 Moisture level - less than 7%
 pH 7.4 ± 0.2 at 25°C
 Clarity - clear

ThermoFisher SCIENTIFIC	Document Owner Department: QC	BT-SPEC-0287 Page 2 of 5
OXOID QUALITY ASSURANCE PRODUCT SPECIFICATION		
BRAIN HEART INFUSION CM1135		

Microbiological Tests Using Optimum Inoculum Dilution

Control Media: Tryptone Soya Agar, Columbia Blood Agar Base enriched with 5% v/v horse blood, Columbia Blood Agar Base enriched with 5% v/v chocolate horse blood or Sabouraud Dextrose Agar, where appropriate

Reactions after incubation at 37 ± 2°C for 18 hours

Medium is challenged with 10-100 colony-forming units

<i>Streptococcus pyogenes</i>	ATCC® 19615	Turbid growth
<i>Streptococcus pneumoniae</i>	ATCC® 6303	Turbid growth
<i>Streptococcus pneumoniae</i>	ATCC® 6305	Turbid growth
<i>Enterococcus faecalis</i>	ATCC® 19433	Turbid growth
<i>Pseudomonas aeruginosa</i>	ATCC® 27853	Turbid growth

A satisfactory result is represented by visible growth.

Reactions after incubation at 37 ± 2°C for 18 hours under anaerobic conditions (for details, refer to Oxoid Manual - Atmosphere Generation Systems)

Medium is challenged with 10-100 colony-forming units

<i>Streptococcus pneumoniae</i>	ATCC® 6305	Turbid growth
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A satisfactory result is represented by visible growth.

Reactions after incubation at 37 ± 2°C for 48 hours

Medium is challenged with 10-100 colony-forming units

<i>Candida albicans</i>	ATCC® 10231	Turbid growth
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A satisfactory result is represented by visible growth.

ThermoFisher SCIENTIFIC	Document Owner Department: QC	BT-SPEC-0287 Page 3 of 5
OXOID QUALITY ASSURANCE PRODUCT SPECIFICATION		
BRAIN HEART INFUSION CM1135		

Reactions after incubation at 37 ± 2°C for 48-72 hours

Tested with the addition of 0.1% w/v bacteriological agar (1 Agar Tablet (CM0049) in 100ml medium)

Medium is challenged with 10-100 colony-forming units

Bacteroides fragilis ATCC® 25285 Turbid growth

A satisfactory result is represented by visible growth.

Reactions after incubation at 37 ± 2°C for 24 hours

Enriched 1 x 10ml volume with two X factor discs (DD0003) and two V factor discs (DD0004)

Medium is challenged with 10-100 colony-forming units

Haemophilus influenzae ATCC® 33391 Turbid growth

A satisfactory result is represented by visible growth.

Enriched with 10% v/v horse blood

Reactions after incubation at 37 ± 2°C for 48 hours and subculture onto chocolate agar plates

Incubate plates at 37 ± 2°C for 24-48 hours in CO₂ atmosphere

(for details, refer to Oxoid Manual - Atmosphere Generation Systems)

Neisseria meningitidis ATCC® 13077 1-2mm grey/brown colonies

Neisseria gonorrhoeae NCTC 11148 1-2mm grey/brown colonies

Neisseria gonorrhoeae ATCC® 19424 1-2mm grey/brown colonies

A satisfactory result is represented by a positive diagnostic reaction, on subculture.

ThermoFisher S C I E N T I F I C	Document Owner Department: QC	BT-SPEC-0287 Page 4 of 5
OXOID QUALITY ASSURANCE PRODUCT SPECIFICATION		
BRAIN HEART INFUSION CM1135		

Reactions after incubation at 37 ± 2°C for 4 hours

Tube Coagulase Test

Medium is challenged with 1E+04 to 1E+06 colony-forming units

<i>Staphylococcus aureus</i>	ATCC® 9144	Coagulase positive
<i>Staphylococcus epidermidis</i>	ATCC® 14990	Coagulase negative

A satisfactory result is represented by the appropriate coagulase reaction.

Testing performed in accordance with ISO11133:2014

Reactions after incubation at 37 ± 2°C for 24 ± 2 hours

Medium is challenged with 10-100 colony-forming units

<i>Staphylococcus aureus</i>	ATCC® 25923	WDCM00034	Turbid growth
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A satisfactory result is represented by visible growth.


Tested in accordance with current CLSI M22 A

Reactions after incubation at 35 ± 2°C for 18 hours

Medium is challenged with 10-100 colony-forming units


<i>Escherichia coli</i>	ATCC® 25922	Turbid growth
<i>Staphylococcus aureus</i>	ATCC® 25923	Turbid growth

A satisfactory result is represented by visible growth.

	Document Owner Department: QC	BT-SPEC-0287
		Page 5 of 5
OXOID QUALITY ASSURANCE PRODUCT SPECIFICATION		
BRAIN HEART INFUSION CM1135		

Revision History

Section / Step	Description of Change	Reason for Change	Reference
Update to 10% Horse Blood Reactions	Change of incubation parameters	Change control	BT-CC-1736

	Document Owner Department: QC	BT-SPEC-0777
		Page 1 of 3
OXOID QUALITY ASSURANCE PRODUCT SPECIFICATION		
M.R.S. (ISO) AGAR CM1153		

M.R.S. (ISO) AGAR

CM1153

Typical Formula*

Enzymatic digest of casein	grams per litre	10.0
Meat extract		10.0
Yeast extract		4.0
Tri-ammonium citrate		2.0
Sodium acetate		5.0
Magnesium sulphate heptahydrate		0.2
Manganese sulphate tetrahydrate		0.05
Di-potassium hydrogen phosphate		2.0
Sorbitan mono-oleate		1.08
Glucose		20.0
Agar		12.37

*adjusted to meet performance standards

Directions

Suspend 66.7g in 1 litre of distilled water. Bring to the boil to dissolve completely. Sterilize by autoclaving at 121°C for 15 minutes. Mix well and pour into sterile Petri dishes.

Physical Characteristics

Dark straw, free-flowing powder
 Colour on reconstitution – brown/orange
 Moisture level- less than or equal to 7%
 pH 5.7 ± 0.1 at 25°C
 Clarity - clear
 Gel strength - firm comparable to 12.37g/litre of agar

Microbiological Tests Using Optimum Inoculum Dilution

Control Medium: M.R.S. Agar or Tryptone Soya Agar

Reactions after incubation at 30 ± 2°C for 72 ± 3 hours under microaerophilic conditions

Medium is challenged with 10-100 colony-forming units

<i>Lactobacillus gasseri</i>	ATCC®19992	0.5-2mm pale straw colonies
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ThermoFisher SCIENTIFIC	Document Owner Department: QC	BT-SPEC-0777
OXOID QUALITY ASSURANCE PRODUCT SPECIFICATION		Page 2 of 3
M.R.S. (ISO) AGAR CM1153		

A satisfactory result is represented by recovery of positive strains equal to or greater than 70% of the control medium.

Testing performed in accordance with ISO11133:2014

Reactions after incubation at $30 \pm 2^{\circ}\text{C}$ for 72 ± 3 hours under microaerophilic conditions

Medium is challenged with 50-120 colony-forming units


<i>Lactobacillus sakei</i>	ATCC®15521	WDCM00015	0.5-2mm pale straw colonies
<i>Lactococcus lactis</i>	ATCC®19435	WDCM00016	0.5-2mm cream colonies
<i>Pediococcus pentosaceus</i>	ATCC®33316	WDCM00158	ppt-3mm cream colonies

A satisfactory result is represented by recovery of positive strains equal to or greater than 70% of the control medium.

Medium is challenged with $1\text{E}+04$ to $1\text{E}+06$ colony-forming units


<i>Escherichia coli</i>	ATCC®25922	WDCM00013	No growth
<i>Escherichia coli</i>	ATCC®8739	WDCM00012	No growth
<i>Bacillus cereus</i>	ATCC®11778	WDCM00001	No growth

Negative strains are inhibited.

	Document Owner Department: QC	BT-SPEC-0777
		Page 3 of 3
OXOID QUALITY ASSURANCE PRODUCT SPECIFICATION		
M.R.S. (ISO) AGAR CM1153		

Revision History

Section / Step	Description of Change	Reason for Change	Reference
Entire document	Creation of new document	Change control	BT-CC-1368
Typical formula/ Physical characteristics	Correction of typographical errors. pH range changed to the correct limits.	Change control	BT-CC-2783

	Document Owner Department: QC	MBD-BT-SPEC-0838
		Page 1 of 4
OXOID QUALITY ASSURANCE PRODUCT SPECIFICATION		
BRILLIANCE™ LISTERIA AGAR BASE (ISO) CM1212		

BRILLIANCE™ LISTERIA AGAR BASE (ISO)

CM1212

Typical Formula*

Enzymatic digest of animal tissues	grams per litre	18.0
Enzymatic digest of casein		6.0
Yeast extract		10.0
Sodium pyruvate		2.0
Glucose		2.0
Magnesium glycerophosphate		1.0
Magnesium sulphate (anhydrous)		0.5
Sodium chloride		5.0
Lithium chloride		10.0
Di-sodium hydrogen phosphate (anhydrous)		2.5
5-Bromo-4-chloro-3-indolyl-β-D-glucopyranoside		0.05
Agar		12.0


* adjusted as required to meet performance standards

Directions

Suspend 34.5g in 480ml of distilled water. Mix well and sterilize by autoclaving at 121°C for 15 minutes. Cool to 48°C. Aseptically add the contents of 1 vial of Brilliance™ Listeria Selective Supplement (ISO) (SR0257E) reconstituted as directed, and 1 vial of Brilliance™ Listeria Differential Supplement (ISO) (SR0258E) warmed to 48°C. Mix well and pour into sterile Petri dishes.

Physical Characteristics

Straw, free-flowing powder
 Colour on reconstitution - orange/brown
 Moisture level - less than or equal to 7%
 pH 7.2 ± 0.2 at 25°C
 Clarity - clear
 Gel strength - firm, comparable to 12.0g/litre of agar

	Document Owner Department: QC	MBD-BT-SPEC-0838
		Page 2 of 4
OXOID QUALITY ASSURANCE PRODUCT SPECIFICATION		
BRILLIANCE™ LISTERIA AGAR BASE (ISO) CM1212		

Microbiological Tests Using Optimum Inoculum Dilution

Control Media: Tryptone Soya Agar, Columbia Blood Agar Base enriched with 5% v/v horse blood or Sabouraud Dextrose Agar, where appropriate

Tested with the addition of Brilliance™ Listeria Selective Supplement (ISO) SR0257 and Brilliance™ Listeria Differential Supplement (ISO) SR0258

Reactions after incubation at 37 ± 2°C for 24 ± 2 hours

Medium is challenged with 30-120 colony-forming units

<i>Listeria monocytogenes</i>	NCTC11994	0.5-2mm blue-green colonies with halo
<i>Listeria monocytogenes</i>	ATCC®7644	0.5-2mm blue-green colonies with halo

A satisfactory result is represented by recovery of positive strains equal to or greater than 50% of the control medium.

Reactions after incubation at 37 ± 2°C for 48 ± 4 hours

Medium is challenged with 30-120 colony-forming units

<i>Listeria monocytogenes</i>	NCTC11994	1-3mm blue-green colonies with halo
<i>Listeria monocytogenes</i>	ATCC®7644	1-3mm blue-green colonies with halo
<i>Listeria ivanovii</i>	NCTC12701	0.5-3mm blue-green colonies with or without halo

A satisfactory result is represented by recovery of positive strains equal to or greater than 70% of the control medium. For *Listeria ivanovii* NCTC12701, a satisfactory result is represented by recovery equal to or greater than 50% of the control medium.

Medium is challenged with 1E+04 to 1E+05 colony-forming units


<i>Bacillus cereus</i>	ATCC®10876	No growth or 1-2mm cream/blue colonies
<i>Staphylococcus aureus</i>	ATCC®25923	No growth or 0.5-1mm yellow colonies
<i>Saccharomyces cerevisiae</i>	ATCC®9763	No growth or 1-2mm cream/blue colonies

Negative strains are inhibited or shall produce at least a 2 log(10) reduction when compared to the control medium.

Medium is challenged with 1E+04 to 1E+06 colony-forming units

<i>Proteus mirabilis</i>	NCTC10975	No growth
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Negative strains are inhibited.

	Document Owner Department: QC	MBD-BT-SPEC-0838
		Page 3 of 4
OXOID QUALITY ASSURANCE PRODUCT SPECIFICATION		
BRILLIANCE™ LISTERIA AGAR BASE (ISO) CM1212		

Testing performed in accordance with ISO11133:2014

Table B.1

ISO Standard 11290-1:2017 tested with the addition of Brilliance™ Listeria Selective Supplement (ISO) SR0257 and Brilliance™ Listeria Differential Supplement (ISO) SR0258

Reactions after incubation at 37 ± 2°C for 24 ± 2 hours

Medium is challenged with 50-120 colony-forming units

<i>Listeria monocytogenes</i>	ATCC®13932	WDCM00021	0.5-2mm blue-green colonies with halo
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A satisfactory result is represented by recovery of positive strains equal to or greater than 50% of the control medium.

Reactions after incubation at 37 ± 2°C for 48 ± 4 hours

Medium is challenged with 50-120 colony-forming units

<i>Listeria monocytogenes</i>	ATCC®13932	WDCM00021	1-3mm blue-green colonies with halo
<i>Listeria monocytogenes</i>	ATCC®35152	WDCM00109	1-3mm blue-green colonies with halo

A satisfactory result is represented by recovery of positive strains equal to or greater than 70% of the control medium.

Medium is challenged with 1E+03 to 1E+04 colony-forming units


<i>Listeria innocua</i>	ATCC®33090	WDCM00017	0.5-3mm blue-green colonies without halo
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A satisfactory result is represented by good growth with a negative diagnostic reaction.

Medium is challenged with 1E+04 to 1E+06 colony-forming units


<i>Escherichia coli</i>	ATCC®25922	WDCM00013	No growth
<i>Escherichia coli</i>	ATCC®8739	WDCM00012	No growth
<i>Enterococcus faecalis</i>	ATCC®29212	WDCM00087	No growth
<i>Enterococcus faecalis</i>	ATCC®19433	WDCM00009	No growth

Negative strains are inhibited.

	Document Owner Department: QC	MBD-BT-SPEC-0838
		Page 4 of 4
OXOID QUALITY ASSURANCE PRODUCT SPECIFICATION		
BRILLIANCE™ LISTERIA AGAR BASE (ISO) CM1212		

Revision History

Section / Step	Description of Change	Reason for Change	Reference
Physical Characteristics	Clarity change from opaque to clear	Change control	MOC-2023-0118

	Document Owner Department: QC	BT-SPEC-0491
		Page 1 of 3
OXOID QUALITY ASSURANCE PRODUCT SPECIFICATION		
LISTERIA SELECTIVE SUPPLEMENT (OXFORD FORMULATION) SR0140E		

LISTERIA SELECTIVE SUPPLEMENT (OXFORD FORMULATION)

SR0140E

Formula

Vial contents (each vial is sufficient to supplement 500ml of medium)

Cycloheximide	200.0 mg
Colistin sulphate	10.0 mg
Acriflavine	2.5 mg
Cefotetan	1.0 mg
Fosfomycin	5.0 mg

Description

A selective supplement for the isolation of *Listeria monocytogenes*.

Directions

Aseptically add 5ml of 70% ethanol to 1 vial and mix gently to dissolve. Avoid frothing. Aseptically add the vial contents to 500ml of sterile Listeria Selective Agar Base (CM0856) prepared as directed and cooled to 50°C. Mix well and pour into sterile Petri dishes.

Physical Characteristics

Yellow powder/pellet
Sterility - passes test

Microbiological Tests Using Optimum Inoculum Dilution

Control Medium: Columbia Blood Agar Base enriched with 5% v/v horse blood

Reactions after incubation at 37°C for 48 hours

Tested in Listeria Selective Agar Base CM0856

Medium is challenged with 10-100 colony-forming units

<i>Listeria monocytogenes</i>	ATCC® 7644	0.25-1.0mm brown/black dimpled colonies and halo
<i>Listeria monocytogenes</i>	ATCC® 13932	0.25-1.0mm brown/black dimpled colonies and halo

A satisfactory result is represented by recovery of positive strains equal to or greater than 50% of the control medium.

ThermoFisher SCIENTIFIC	Document Owner Department: QC	BT-SPEC-0491 Page 2 of 3
OXOID QUALITY ASSURANCE PRODUCT SPECIFICATION		
LISTERIA SELECTIVE SUPPLEMENT (OXFORD FORMULATION) SR0140E		

Medium is challenged with 10-100 colony-forming units


Staphylococcus aureus ATCC®25923 No growth or pinpoint-1.5mm yellow colonies

Staphylococcus aureus ATCC®25923 is inhibited or shall produce a negative diagnostic reaction from an inoculum of 10-100 cfu.

Medium is challenged with 1E+04 to 1E+06 colony-forming units


<i>Enterococcus faecalis</i>	ATCC®29212	No growth
<i>Enterococcus faecalis</i>	ATCC®19433	No growth
<i>Escherichia coli</i>	ATCC®25922	No growth
<i>Escherichia coli</i>	ATCC®8739	No growth
<i>Candida albicans</i>	ATCC®10231	No growth or minimal growth

Negative strains are inhibited. *Candida albicans* ATCC®10231 shall be inhibited or produce pinpoint colourless colonies with no blackening of the media.

	Document Owner Department: QC	BT-SPEC-0491
		Page 3 of 3
OXOID QUALITY ASSURANCE PRODUCT SPECIFICATION		
LISTERIA SELECTIVE SUPPLEMENT (OXFORD FORMULATION) SR0140E		

Revision History

Section / Step	Description of Change	Reason for Change	Reference
Microbiological characteristics	Change to <i>Staphylococcus aureus</i> growth characteristics	Change control	MOC-2022-0180

	Document Owner Department: QC	MBD-BT-SPEC-0505
		Page 1 of 5
		Rev 04
OXOID QUALITY ASSURANCE PRODUCT SPECIFICATION		
FRASER SELECTIVE SUPPLEMENT SR0156E		

FRASER SELECTIVE SUPPLEMENT

SR0156E

Formula

Vial contents (each vial is sufficient to supplement 500ml of medium)

Ammonium iron (III) citrate	250.0 mg
Nalidixic acid	10.0 mg
Acriflavine hydrochloride	12.5 mg

Description

A selective supplement for the detection of *Listeria monocytogenes*.

Directions

Aseptically add 5ml of 1:1 ethanol:sterile distilled water to 1 vial and mix gently to dissolve. Aseptically add the vial contents to 500ml of sterile Fraser Broth Base (CM0895) prepared as directed and cooled to 50°C. Mix well and aseptically dispense into sterile containers.

Physical Characteristics

Orange/green pellet
Sterility - passes test

Microbiological Tests Using Optimum Inoculum Dilution

Control Media: Brilliance™ Listeria Agar (ISO) or Columbia Blood Agar Base enriched with 5% v/v horse blood, where appropriate.

Tested in Fraser Broth Base CM0895

Reactions after incubation at 37 ± 2°C for 24 ± 2 hours

Inoculate 10ml quantities of medium to achieve 1-10 colony-forming units/ml (cfu/ml) of *Listeria* spp. Incubate broths at 37 ± 2°C for 24 ± 2 hours. Subculture onto Brilliance™ Listeria Agar (ISO) (CM1212, SR0257 & SR0258) and incubate plates at 37 ± 2°C for 24 to 48 hours.

<i>Listeria monocytogenes</i>	ATCC®7644
<i>Listeria monocytogenes</i>	ATCC®13932

	Document Owner Department: QC	MBD-BT-SPEC-0505
		Page 2 of 5
		Rev 04
OXOID QUALITY ASSURANCE PRODUCT SPECIFICATION		
FRASER SELECTIVE SUPPLEMENT SR0156E		

A satisfactory result is represented by recovery of positive strains equal to or greater than a 3 log(10) increase.

Positive strains shall produce aesculin hydrolysis after 48 hours.

Reactions after incubation at 37 ± 2°C for 48 ± 2 hours

Inoculate 10ml quantities of medium to achieve 1E+03 to 1E+04 cfu/ml. Incubate broths at 37°C for 48 hours.

<i>Bacillus cereus</i>	ATCC®10876	No aesculin hydrolysis (no blackening)
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Negative strains shall produce no aesculin hydrolysis after 48 hours.

Testing performed in accordance with ISO11133:2014

Inoculation with mixed cultures

Inoculate 10ml quantities of medium to achieve 1-10 colony-forming units/ml (cfu/ml) of *Listeria* spp., to each add 1E+02 to 1E+03 cfu/ml of *Escherichia coli* and 1E+02 to 1E+03 cfu/ml of *Enterococcus faecalis*. Incubate broths at 37 ± 2°C for 24 ± 2 hours. Subculture onto Brilliance™ *Listeria* Agar (ISO) (CM1212, SR0257 & SR0258) and incubate plates at 37 ± 2°C for 24 ± 2 hours

Reactions after incubation at 37 ± 2°C for 24 ± 2 hours

<i>Listeria monocytogenes</i>	ATCC®13932	WDCM00021	0.5-1.0mm blue colonies with halo
+ <i>Escherichia coli</i>	ATCC®8739	WDCM00012	No growth
+ <i>Enterococcus faecalis</i>	ATCC®19433	WDCM00009	No growth
<i>Listeria monocytogenes</i>	ATCC®13932	WDCM00021	0.5-1.0mm blue colonies with halo
+ <i>Escherichia coli</i>	ATCC®25922	WDCM00013	No growth
+ <i>Enterococcus faecalis</i>	ATCC®19433	WDCM00009	No growth
<i>Listeria monocytogenes</i>	ATCC®13932	WDCM00021	0.5-1.0mm blue colonies with halo
+ <i>Escherichia coli</i>	ATCC®8739	WDCM00012	No growth
+ <i>Enterococcus faecalis</i>	ATCC®29212	WDCM00087	No growth

	Document Owner Department: QC	MBD-BT-SPEC-0505
		Page 3 of 5
		Rev 04
OXOID QUALITY ASSURANCE PRODUCT SPECIFICATION		
FRASER SELECTIVE SUPPLEMENT SR0156E		

<i>Listeria monocytogenes</i>	ATCC®13932	WDCM00021	0.5-1.0mm blue colonies with halo
+ <i>Escherichia coli</i>	ATCC®25922	WDCM00013	No growth
+ <i>Enterococcus faecalis</i>	ATCC®29212	WDCM00087	No growth

<i>Listeria monocytogenes</i>	ATCC®35152	WDCM00109	0.5-1.0mm blue colonies with halo
+ <i>Escherichia coli</i>	ATCC®8739	WDCM00012	No growth
+ <i>Enterococcus faecalis</i>	ATCC®19433	WDCM00009	No growth

<i>Listeria monocytogenes</i>	ATCC®35152	WDCM00109	0.5-1.0mm blue colonies with halo
+ <i>Escherichia coli</i>	ATCC®25922	WDCM00013	No growth
+ <i>Enterococcus faecalis</i>	ATCC®19433	WDCM00009	No growth

<i>Listeria monocytogenes</i>	ATCC®35152	WDCM00109	0.5-1.0mm blue colonies with halo
+ <i>Escherichia coli</i>	ATCC®8739	WDCM00012	No growth
+ <i>Enterococcus faecalis</i>	ATCC®29212	WDCM00087	No growth

<i>Listeria monocytogenes</i>	ATCC®35152	WDCM00109	0.5-1.0mm blue colonies with halo
+ <i>Escherichia coli</i>	ATCC®25922	WDCM00013	No growth
+ <i>Enterococcus faecalis</i>	ATCC®29212	WDCM00087	No growth

A satisfactory result is represented by recovery of >10 cfu of *Listeria monocytogenes* on Brilliance™ Listeria Agar (ISO).

Inoculation with pure cultures

Inoculate 10ml quantities of medium to achieve 1E+03 to 1E+04 colony-forming units/ml (cfu/ml) of *Escherichia coli* and *Enterococcus faecalis*. Incubate broths at 37 ± 2°C for 24 ± 2 hours. Subculture onto Brilliance™ Listeria Agar (ISO) (CM1212, SR0257 & SR0258) and Tryptone Soya Agar (CM0131) then incubate plates at 37 ± 2°C for 24 ± 2 hours.

Reactions after incubation at 37 ± 2°C for 24 ± 2 hours

<i>Escherichia coli</i>	ATCC®8739	WDCM00012	No growth (CM1212, SR0257 & SR0258)
<i>Escherichia coli</i>	ATCC®8739	WDCM00012	Cream colonies (CM0131)
<i>Escherichia coli</i>	ATCC®25922	WDCM00013	No growth (CM1212, SR0257 & SR0258)
<i>Escherichia coli</i>	ATCC®25922	WDCM00013	Cream colonies (CM0131)
<i>Enterococcus faecalis</i>	ATCC®19433	WDCM00009	No growth (CM1212, SR0257 & SR0258)
<i>Enterococcus faecalis</i>	ATCC®19433	WDCM00009	Cream colonies (CM0131)

	Document Owner Department: QC	MBD-BT-SPEC-0505
		Page 4 of 5
		Rev 04
OXOID QUALITY ASSURANCE PRODUCT SPECIFICATION		
FRASER SELECTIVE SUPPLEMENT SR0156E		

Enterococcus faecalis ATCC®29212 WDCM00087 No growth (CM1212, SR0257 & SR0258)
Enterococcus faecalis ATCC®29212 WDCM00087 Cream colonies (CM0131)

A satisfactory result is represented by no growth of *Escherichia coli* and *Enterococcus faecalis* on Brilliance™ Listeria Agar (ISO) and <100 cfu on Tryptone Soya Agar.

	Document Owner Department: QC	MBD-BT-SPEC-0505
		Page 5 of 5
		Rev 04
OXOID QUALITY ASSURANCE PRODUCT SPECIFICATION		
FRASER SELECTIVE SUPPLEMENT SR0156E		

Revision History

Section / Step	Description of Change	Reason for Change	Reference
Microbiological Characteristics	Change of Listeria plating medium	Change control	MOC-2023-0965

Distribution: Central File

Date: 04/08/11

Supersedes: 12/11/10

OXOID QUALITY ASSURANCE PRODUCT SPECIFICATION

NOVOBIOCIN SUPPLEMENT

SR0181E

Formula

Vial contents (each vial is sufficient to supplement 500ml of medium at 20mg/l)

Novobiocin

10.0 mg

Directions

Aseptically add 2ml of sterile distilled water to 1 vial and invert gently to dissolve. Aseptically add the vial contents to 500ml of sterile Modified Tryptone Soya Broth (CM0989) prepared as directed and cooled to 50°C. Mix well and aseptically dispense into sterile containers.

Physical Characteristics

White pellet

Sterility - passes test

Microbiological Tests Using Optimum Inoculum Dilution

Control Medium: Tryptone Soya Agar

Medium is challenged with 10-100 colony-forming units

Reactions after incubation at 41.5°C for 18-24 hours

Tested in Modified Tryptone Soya Broth CM0989

<i>Escherichia coli</i> (O157:H7)	NCTC 12079	Turbid growth
<i>Escherichia coli</i> (O157:H7) (Verotoxin negative)	NCTC 12900	Turbid growth
<i>Escherichia coli</i>	ATCC® 25922	Turbid growth
<i>Escherichia coli</i>	ATCC® 8739	Turbid growth
<i>Klebsiella pneumoniae</i>	ATCC® 29665	Turbid growth

Medium is challenged with 1E+04 to 1E+06 colony-forming units

<i>Bacillus subtilis</i>	ATCC® 6633	No growth to minimal growth
<i>Staphylococcus aureus</i>	ATCC® 25923	No growth to minimal growth

Negative strains are inhibited or shall produce minimal growth