



LABORATORY PROTOCOL

PCR for plasmid-mediated colistin resistance genes, *mcr-1* and *mcr-2* (multiplex)

(protocol optimized at National Food Institute, Denmark)

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Version 2

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HISTORY OF CHANGES				
Version	Sections changed	Description of change	Date	Approval
1	New document	-	December 2015	Authors
2	Throughout document	Addition of <i>mcr-2</i> and adjustments to use Dream Taq Green Mastermix in the reaction	October 2016	Authors

PROTOCOL

PCR protocol for the *mcr-1* and *mcr-2* genes (multiplex)

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Materials

Equipment

- PCR thermocycler
- Pipettes for 1 μ L to 1000 μ L
- Electrophoresis unit
- Microwave or autoclave
- Eppendorf centrifuge
- Photo camera
- UV-transilluminator
- Water bath 50°C

Materials

- Molecular marker (Ladder 100bp)
- Ethidium bromide solution (1%)
- Staining bath
- Electrophoresis buffer TAE or TBE (see composition of media and reagents)
- Eppendorf tubes
- Tips (filter) for pipettes 1 μ L to 1000 μ L
- Agarose
- Primers
- Dream Taq Green PCR Master Mix (2X) (Thermo Fisher)
- DNA template (boiling lysates)
- TE buffer (Tris:EDTA 10:1)
- TrisHCl buffer
- Crushed ice
- Mineral oil (if necessary)

Safety

Carry out all procedures in accordance with the local codes of safe practice.

For staining the DNA, ethidium bromide is used. This dye is carcinogenic. Therefore gloves and proper clothes should be worn (appendix 1: laboratory safety).

Visualisation of the stained DNA is done by use of UV transilluminator. UV light is harmful for skin and eyes. Therefore proper protection (facemask, glasses) should be worn (Appendix 1: laboratory safety).

PCR detection of *mcr-1* and *mcr-2* (multiplex)

The *mcr-1* gene was described by Liu *et al.* who published their findings in a Lancet article in the end of 2015. This gene encodes phosphoethanolamine transferase enzyme family, with expression in *E. coli* resulting in the addition of phosphoethanolamine to lipid A and in this way confers resistance to colistin. A similar gene, *mcr-2* was described in Belgian isolates by Xavier *et al.*, 2016 and published in Eurosurveillance.

This protocol has been modified by adding the *mcr-2* reaction to the previous protocol optimized at Statens Serum Institute (SSI, 2015) and by using a different Taq polymerase containing mastermix.

Control strains

E. coli carrying *mcr-1* (ID: 2012-60-1176-27)

E. coli containing *mcr-2* (ID: KP37) obtained from Xavier *et al.*

DNA extraction

The template DNA used consisted of boiling lysates prepared from the strains. A brief description: a loopful of culture was suspended in 100 µl of sterile TE buffer, boiled 10 min at 100°C, centrifuged 5 min at 6000 G. For use as template in the PCR, the DNA supernatant was further diluted at 1:10 in TrisHCl buffer.

Primers

Target gene	Primer sequences	Reference
<i>mcr-1</i> (35-343)	CLR F 5'-CGGTCAGTCCGTTTGTTC-3'	Liu <i>et al.</i> , 2015
	CLR R 5'-CTTGGTCGGTCTGTAGGG-3'	
<i>mcr-2</i> (494-1060)	MCR2 IF 5'- TGTTGCTTGTGCCGATTGGA-3'	Xavier <i>et al.</i> , 2016
	MCR2 IR 5'-AGATGGTATTGTTGGTTGCTG-3'	

Reaction mix

Prepare the following mix in a microcentrifuge tube (for a 25µL reaction; see the example of PCR set up). Additionally, prepare a blank reaction without template DNA as negative control. This version of the PCR protocol is optimized for the Dream Taq Green PCR Master Mix (2X) (https://tools.thermofisher.com/content/sfs/manuals/MAN0012704_DreamTaq_Green_PCR_MasterMix_K1081_UG.pdf) which can be replaced by another polymerase, although the protocol might need some optimization to adjust for the particular conditions at your laboratory in which we can give you some assistance.

Dream Taq PCR Master Mix*	12.5 µL
Primer Mix**	2 µL
DNA template	2 µL
Water up to	25 µL

*This PCR at the EURL-AR laboratory is optimized for the Dream Taq Green PCR Master Mix (Thermo Fisher) which already contains MgCl₂.

**The primer mix contains 2µM of each primer in water or TE buffer

Conditions for the PCR

94°C 15 min + 25X (94°C 30 sec + 58°C 90 sec + 72°C 60 sec) + 72°C 10min.

References

Liu YY, Wang Y, Walsh TR, Yi LX, Zhang R, Spencer J, Doi Y, Tian G, Dong B, Huang X, Yu LF, Gu D, Ren H, Chen X, Lv L, He D, Zhou H, Liang Z, Liu JH, Shen J. Emergence of plasmid-mediated colistin resistance mechanism MCR-1 in animals and human beings in China: a microbiological and molecular biological study. *Lancet Infect Dis.* 2015 Nov 18. pii: S1473-3099(15)00424-7. doi: 10.1016/S1473-3099(15)00424-7. [Statens Serum Institute, working protocol (Personal communication Frank Hansen and Henrik Hasman)]

Xavier BB, Lammens C, Ruhel R, Kumar-Singh S, Butaye P, Goossens H, Malhotra-Kumar S. Identification of a novel plasmid-mediated colistin-resistance gene, *mcr-2*, in *Escherichia coli*, Belgium, June 2016. *Euro Surveill.* 2016;21(27):pii=30280. DOI: <http://dx.doi.org/10.2807/1560-7917.ES.2016.21.27.30280>

Detailed Procedure

Preparation of the samples

1. Transfer 100 μL of TE buffer to a 1.5 mL Eppendorf tube. Using a disposable inoculation loop (white; 1 μL), pick a loop full of bacteria from a plate and transfer to the Eppendorf tube.
2. Boil the suspension (or heat at 95°C) for 5-10 minutes.
3. Centrifuge at 6000 G for 5 min.
4. Dilute the supernatant lysed DNA 10-fold in TrisHCl

Preparation of the mix

1. Check the number of samples and calculate the amount of PCR master mix needed.
2. Prepare the PCR master mix (you may do it in a tray of crushed ice as mentioned in appendix 2).
3. Aliquot the PCR master mix to the required number of PCR tubes (23 μL per tube).
4. Depending on the PCR machine that is used, one drop of mineral oil could be added.

Theory / comments

Make a homogeneous cell suspension in an Eppendorf tube, using a loop or cotton swab. Shake or vortex suspension just before use. Only very little cell mass is needed. If too many bacteria are used, it might cause inhibition of the PCR.

TE is used because it contains EDTA that binds divalent ions needed by enzymes that would be able to degrade DNA. Since EDTA can inhibit the PCR reaction, dilute in TrisHCl

Boiling breaks down the bacterial cell wall and allows release of DNA.

Make a ventilation hole in the lid of the Eppendorf tube using a needle, alternatively the eppendorf tubes can be capped with a lidlock.

Always prepare mix for at least 1-2 additional samples ($n \text{ mix} = n \text{ samples} + 1 \text{ or } 2$). Use crushed ice if the PCR master mix is prepared at temperatures above 25°C as this may affect the result.

If the PCR machine has no heat in the lid use oil as a lid to avoid the mixture to vaporise and condensate in the lid of the tubes

Running the PCR

1. Add 2 μL of sample to the sample tube and close the lid.
2. Add 2 μL of water to the negative control tube and close the lid
3. Finish the procedure by adding the positive control DNA and close the lid.
4. Place the tubes into the PCR thermocycler.
5. Program the PCR thermocycler (or select the requested program) as mentioned in appendix 2 (Example of PCR set-up).
6. Run the program.

Preparation of the agarose gel

1. Assemble the gel tray and make a proper set-up.
2. Prepare a 1.5% agarose solution in TBE buffer 1X by boiling the solution a few minutes until completely dissolved by the use of a water bath or microwave oven.
3. Cool the agarose to 40-50°C in a water bath and pour the agarose into the gel tray.
4. Let the gel solidify for 15-30 minutes.
5. Prepare a staining bath containing a final concentration of 5 $\mu\text{g}/\text{mL}$ ethidium bromide.

Theory / comments

Always end the set up with the positive control DNA to avoid contaminations of the test tubes, causing false positive results.

For PCR a positive control and a negative control (sterile water) should be taken along.

If oil is used, make sure DNA is dispensed below the oil phase.

Depending of electrophoresis system use TBE or TAE buffer

Wear suitable protection for working with ethidium bromide (gloves, clothes). Appendix 1: laboratory safety.

Assembling the results

1. Put the gel into the electrophoresis unit and if necessary refill with buffer.
2. Load 8 μL of each PCR sample into the wells of the gel. Finish off by loading at least one molecular marker.
3. Replace the lid of the unit and run the gel by starting the electrophoresis process.
4. After a complete run of 30-45 minutes, remove the lid of the unit and place the gel in a staining-bath for about 30 minutes. Rinse shortly in water before visualising the gel / bands.
5. Place the gel / tray on top of the UV-transilluminator.
6. Visualise the results by switching on the UV-lamp.
7. Look for the presence of specific bands (Appendix 2: Example of PCR set up).

Theory / comments

TBE contains boric acid. See Appendix 1 for safety sheet.

The Dream Taq Green Master Mix contains two tracking dyes and a density reagent that allows the direct loading of PCR product

It is important to use a proper marker in order to notice whether the PCR product has the right size.

Electrophoresis can be done at different voltages/amperages. Normally, 70mA seems to be fine. Running time depends on several parameters like buffer composition, resistance, current.

Be extremely careful and wear the correct protective gloves when dealing with the ethidium bromide staining bath, please read the safety precautions in Appendix 1)

UV light is harmful for skin and eyes. Wear proper protection (facemask) see Appendix 1, laboratory safety

The whole PCR process is very sensitive towards contaminations that can affect the result as false positive results. It is therefore recommended to perform the different steps, if possible, in different rooms e.g.:

Room 1: Preparing the PCR master mix into tubes.

Room 2: Adding the samples to the tubes and running the samples in the PCR thermocycler.

Room 3: Running the electrophoresis and visualising the DNA.

Composition and preparation of media and reagents

Reagents can be made as described below and/or are commercially available from companies like Invitrogen Life Technologies and Roche Applied Science.

TAE (Tris-Acetate EDTA) buffer

Working solution

- 0.04 M Tris Acetate
- 0.001 M EDTA

Concentrated stock solution (50X)

Per liter

- Tris base 242 g
- Glacial acetic acid 57.1 mL
- 0.5 M EDTA (pH 8.0) 100 mL

TBE (Tris-Borate EDTA) buffer

Working solution

- 0.089 M Tris borate
- 0.089 M boric-acid
- 0.002 M EDTA

Concentrated stock solution (5X)

Per liter

- Tris base 54 g
- Boric acid 27.5 g
- 0.5 M EDTA (pH 8.0) 20 mL

Tris-EDTA buffer (TE 10:1) 1 L (pH 8)

- 1 M Tris-HCl (pH 8) 10 mL
- 0.5 M EDTA (pH 8.0) 2 mL
- Water 988 mL

Tris-HCl buffer 1 L (pH 8)

1 M Tris-HCl (pH 8) 10 ml
Water 990 ml

Ethidium bromide (10mg/mL)

- Add 1 g of ethidium bromide to 100 mL of H₂O. Stir on a magnetic stirrer for several hours to ensure that the dye has dissolved. Wrap the container in aluminum foil or transfer to a dark bottle and store at 4°C.
- Ethidium bromide can be bought ready-made as a 10 mg/ml solution

APPENDIX 1 (laboratory safety)

Safe Work Procedure ETHIDIUM BROMIDE

Use

Ethidium bromide is added to electrophoresis gels for visualisation of nucleic acids

Hazards

Class 6 – Toxic. Potent mutagen

Risk control measures

Only use ethidium bromide (EtBr) after receiving safety training (laboratory induction / authorisation). Wear safety glasses when using ethidium bromide. Avoid skin contact; ethidium bromide may be absorbed through the skin. Wear latex gloves, laboratory coat. Always dispose of gloves after use. Do not touch equipment, door handles, phone, keyboard, etc.

Weighing solid - Powder may cause irritation when inhaled - wear dust mask and use in ventilated area. Use designated micropipette, only, when dispensing the liquid.

Engineering / Ventilation controls

Ensure access to a safety shower and eye wash in areas where ethidium bromide is used. Preferably weigh the solid in a fume hood.

Storage requirements

Store in a cool, dry place away from strong oxidizing agents. Keep containers tightly closed. Use with adequate ventilation.

First aid / Spill control procedures

Wash off immediately with copious amounts of cold water (at least 10 minutes). Ethidium bromide is absorbed through the skin so follow the cold water washing with a thorough washing with warm water and soap. Contaminated clothing should be removed as soon as possible and thoroughly washed.

In case of contact with eyes, immediately flush eyes with copious amounts of water for at least 15 minutes (eye wash).

Seek medical attention.

If the spill is on equipment, use ultraviolet light (wear appropriate eye protection) to locate spill, then use the decontamination procedure outlined below.

Wear protective clothing.

Small spill: If in solution, absorb freestanding liquid using vermiculite or Polyzorb from Spill Kit. Use ultraviolet light to locate spill. Follow instructions on Spill Kit.

Large Spill: Notify others in the area of spill. Evacuate area. Barricade area with tape (in Spill Kits) to prevent entry until arrival of response personnel. Provide assistance and information to spill clean up crew.

Waste

Store waste; liquid: In proper waste container

Store waste; Solid: In proper waste container

Ethidium bromide liquid disposal

1. Add 10g activated charcoal per 2.5L waste
2. Leave for 1 hour, with occasional shaking
3. Filter contents through Whatman Number 1 filter paper.
4. Filtrate may be disposed of down the sink.
5. Charcoal & paper is treated as solid hazardous waste and disposed in the EtBr Solid Waste Bucket.

If Using 'Green Bag' (Bio-101 Cat. No. 2350-200):

- 1 For 10mg Ethidium Bromide (max) add 1 'Green Bag' to the waste bottle with a magnetic flea.
2. Place waste bottle onto a magnetic stirrer and mix the solution for 24hours.
3. Dispose of the 'Green Bag' in the dry Ethidium Bromide waste. The remaining solution may be disposed of in the sink.

Staining gels

During electrophoresis, add EtBr after boiling up the agarose - let it cool down before adding EtBr

Afterwards, soak gel in a well-marked plastic container - put name and date on container as it is possible to re-use the staining solution.

Safe Work Procedure **ULTRA VIOLET SOURCE**

Ultraviolet light

Ultraviolet radiation is the portion of the electromagnetic spectrum that falls in the region of 100 to 400nm. This spectrum has been divided into three regions:

A: 400nm to 315nm known as Near-UV or UV-A

B: 315nm to 280nm known as Mid-UV or UV-B

C: 280nm to 100nm known as Far-UV or UV-C

Hazards

Two categories of hazard are involved in the use of high intensity UV lamps: those inherent in the radiation itself and those associated with operation of the lamps. All radiation of wavelength shorter than 250 nm should be considered dangerous.

- Damage to eyes and skin caused by exposure to UV radiation. Repeated overexposure of skin to UV has been linked with premature aging, wrinkles and most seriously, skin cancer. Eye damage can result in corneal scarring and cataract formation.
- Burns caused by contact with a hot UV lamp.
- Fire ignited by hot UV lamp.
- Interaction of other nearby chemicals with UV radiation.

Damage caused to apparatus placed close to UV lamp

Risks

Damage to vision is likely following exposure to high intensity UV radiation.

Who is likely to be injured?

The user or anyone exposed to the UV light as a result of faulty procedure. Injuries may be slight to severe.

Control measures operating precautions

Lab-coats, gloves and safety glasses or other appropriate eye/skin protection such as UV protective glasses or a UV protective face shield must be worn.

Reactions using UV lamps: external irradiation sources

- These operations must never be attempted by an untrained person.
- These operations must never be attempted by a single person.
- These operations must never be attempted out of normal working hours.
- Use of UV lamps must be carried out in a fume hood with boarded up windows.
- As far as possible, the UV source should be contained in a closed radiation box.
- The fume hood sash must remain closed while the UV lamp is switched on.

- The fume hood may contain only the UV lamp and associated apparatus and chemicals. No other chemicals are to be stored in the fume hood and no other reactions are to be performed in the fume hood.
- Reaction vessels containing flammable solvents must be at least 20 cm away from the lamp to avoid excessive heating.
- Flammable equipment (e.g. rubber/plastic tubing) must be positioned at least 10 cm away from the lamp.
- After the UV lamp is switched off, unless the reaction mixture requires immediate attention, the fume hood sash should remain closed for 30 minutes to allow the UV lamp to cool.

Reactions using UV lamps: low/medium pressure Hg lamps in an immersion well

- These operations must never be attempted by an untrained person.
- These operations must never be attempted by a single person.
- Low/ Medium pressure lamps are to be used ONLY in approved, water-cooled immersion well apparatus.
- The UV lamp power supplies must incorporate an electrical cutout that activates in the event of disruption to cooling water.
- The UV lamp must not be switched on until:
 - The glassware is shrouded in Al foil
 - The immersion well set-up is shielded by the appropriate metal case
 - The boarded up fume-hood doors are closed
- The UV lamp must NEVER be switched on/connected outside of the shrouded immersion well apparatus.

Training

For the use of high intensity UV sources, new users must be trained by another member of the laboratory who, in the opinion of the member of staff in charge of the laboratory, is sufficiently competent to give instruction on the correct procedure. Newly trained users should be overseen for some time by a competent person.

Emergency Procedures

UV exposure: Act according to local procedures and provide first aid to the injured. If necessary prepare a report for working accidents.

Burns: Act according to local procedures and provide first aid to the injured. If necessary prepare a report for working accidents.

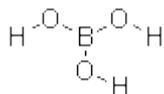
Hazard sheet **BORIC ACID**

Synonyms: Boric Acid, MB Grade (1.12015); Boron; Boricacidhighpurity; Boricacidwhitextl

Molecular Formula: **H₃BO₃**

Formula Weight: 61.83

Registry number: 10043-35-3



Registry number: 10043-35-3

Density: 1.43

Melting point: 169 °C

Hazard Symbol



Toxic

Risk Description

R60 May impair fertility.

Safety Description

S45 In case of accident or if you feel unwell, seek medical advice immediately (show the label where possible).

S53 Avoid exposure - obtain special instructions before use.

IR

Analysis

Result

Miscellaneous

545 (29.51); 641 (30.64); 813 (26.79); 1193 (21.05); 1456 (17.55);
3215 (15.75)

APPENDIX 2 (Example of PCR set up)

PCR		Date:	Initials:
Primers Forward:	CLR F 5'-CGGTCAGTCCGTTTGTTC-'3 MCR2-IF 5'-TGTTGCTTGTGCCGATTGGA-'3		
Primers Reverse:	CLR R 5'-CTTGGTCCGGTCTGTAGGG-'3 MCR2-IR 5'-AGATGGTATTGTTGGTTGCTG-'3		
DNA polymerase:	DreamTaq™ Green PCR Master Mix		
PCR product:	<i>mcr-1</i> : 309 bp <i>mcr-2</i> : 567 bp		
Remarks:	Plasmid-mediated colistin resistance Use 2µl boiling lysate as template		
Positive controls:	<i>mcr-1</i> : <i>E. coli</i> 2012-60-1176-27 <i>mcr-2</i> : <i>E. coli</i> KP37		

Number of samples	1	8	1. 15 min at 94 °C
PCR H ₂ O	8,5	68	2. 25 cycles <u>30</u> sec at <u>94</u> °C <u>90</u> sec at <u>58</u> °C <u>60</u> sec at <u>72</u> °C
2xGreen PCR Master Mix	12,5	100	
Primers F - 0,5 µl of each	1	8	3. <u>10</u> min at <u>72</u> °C
Primers R - 0,5 µl of each	1	8	
			4. _____ hold at _____ 4 °C
Total volume	23	184	

M:	100 bp Ladder plus
1	Sample 1
2	Sample 2
3	<i>mcr-1</i> positive control
4	<i>mcr-2</i> positive control
5	Sample 3
6	<i>mcr-1</i> and 2 positive control mix
7	mastermix
8	
9	
10	
11	
12	

