

Bacterial Ribosome sequencing

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Overview:

Sequencing 16S bacterial ribosome DNA is useful for species identification. Attostar provides a sequencing PCR master mix with primers, dNTPs, buffer and the sequencing primer. The PCR master mix amplifies a conserved region found in many bacteria. This region in the *E. coli* ribosome spans the positions 8 to 926. The exact numbering for the span in other bacteria may vary. This PCR master mix amplifies a hypervariable region, useful for species identification.

After the PCR reaction, the sample is treated to remove primers and nucleotides, then, sent to a sequencing facility of your choice.

The sequence information may then be used at NCBI Blast for identification.

For technical support, contact Attostar@Attostar.com

Products:

Products	Catalog No.	Quantity
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Attostar 16S ribosome sequencing kit	SEQ16S-KIT	1 kit
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(Store at –20C)

Includes: 4 vials master mix, SEQ16S-MM, 1 vials of Sequencing Primer, SEQ16S-P.

Typical use: Prepare bacterial DNA, add DNA and Taq to master mix, do PCR, treat PCR product and sequence.

16S ribosome master mix	SEQ16S-MM	0.350 ml
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(Store at –20C)

Typical use: Add DNA and Taq master mix, do PCR, treat PCR product and sequence.

Sequencing Primer	SEQ16S-P	0.5 ml
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(Store at –20C)

Typical use: Send 10 ul to sequencing facility with PCR product.

BSA 1 mg/ml	BSA100	0.1 ml
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(Store at –20C)

Typical use: Add to master mix if PCR is done in LightCycler glass capillaries.

Reagents to be obtained by the user:

AmpliTaq Gold LD. Obtain from Applied Biosystems.

Catalog number 4338857(1000 units) or 438856 (250 units)

<https://www2.appliedbiosystems.com/>

PrepMan Ultra Sample Preparation Reagent

Catalog number 4318930

<https://www2.Appliedbiosystems.com/>

ExoSAP-IT

Catalog number 78200 100 reactions

<http://www.usbweb.com>

Sybr GreenI

Catalog number S9430 0.5 ml

<http://www.sigmaaldrich.com>

Typical use: Dilute a small aliquot 1:1000 in water and store at –20C.

PROCEDURE:

Prepare bacterial DNA:

- With a 1ul loop, add bacteria from an isolated colony to a 0.2 ml tube containing 50 ul PrepMan Reagent. This should be a visibly cloudy suspension.
- Vortex 10-30 seconds.
- Heat 100C for 10 min.
- Place 2 ul of this heated solution into 0.1 ml DNase RNase free water.
- Do PCR with one of the 3 methods described below.

ExoSAP-IT treatment of PCR product:

- Mix 5 ul PCR product with 2 ul of ExoSAP-IT.
- Incubate at 37C for 15 minutes.
- Incubate at 80C for 15 minutes to inactivate ExoSAP-IT.
- Store treated PCR products at -20C if it is not shipped the same day.

Ship PCR product to sequencing facility:

- Include your PCR product and the sequencing primer SEQ16S-P. A single primer is usually sufficient for an excellent result. Other sequencing primers may be used and provided on request.
- Request a FASTA report via email.

Enter the PCR results at rdna.ridom.de :

- In the “**Diagnostics**” text box, choose “Identify a microorganism by ribosomal DNA sequence”.

Or

Enter the PCR results into Ribosomal database project

<http://rdp.cme.msu.edu/index.jsp>

- Go to Seq Match: http://rdp.cme.msu.edu/seqmatch/seqmatch_intro.jsp

Or

Enter the PCR results into European Ribosomal RNA database

<http://bioinformatics.psb.ugent.be/webtools/rRNA/>

- Go to rRNA
BLAST:<http://bioinformatics.psb.ugent.be/webtools/rRNA/blastrrna.html>

Enter the PCR results into NCBI BLAST:

- Use this URL:

http://www.ncbi.nlm.nih.gov/BLAST/Blast.cgi?CMD=Web&LAYOUT=TwoWindows&AUTO_FORMAT=Semiauto&ALIGNMENTS=50&ALIGNMENT_VIEW=Pairwise&CLIENT=web&DATABASE=nr&DESCRIPTIONS=100&ENTREZ_QUERY=%28none%29&EXPECT=10&FILTER=L&FORMAT_OBJECT=Alignment&FORMAT_TYPE=HTML&NCBI_GI=on&PAGE=Nucleotides&PROGRAM=blastn&SERVICE=plain&SET_DEFAULTS.x=34&SET_DEFAULTS.y=8&SHOW_OVERVIEW=on&END_OF_HTTPGET=Yes&SHOW_LINKOUT=yes&GET_SEQUENCE=yes

- Cut and paste the FASTA sequence results into the “enter Query sequence text box”.
- Click BLAST.
- The results will be returned very shortly.

Distance tree of results:

- Scroll down and click the “distance tree of results” for a nice display.

Distance tree of results **NEW**

Sequences producing significant alignments:	Score (Bits)	E Value
gi 18644240 gb AF371531.1 Uncultured bacterium clone p-2526-...	1633	0.0
gi 46560660 gb AY587799.1 Bacterium Te91A 16S ribosomal RNA gen	1633	0.0
gi 84778493 dbj AB246407.1 Enterococcus faecium gene for 16S...	1633	0.0
gi 27368101 gb AY172570.1 Enterococcus faecium 16S ribosomal RN	1633	0.0
gi 17977836 emb AJ420800.1 EFA420800 Enterococcus faecium 16S rR	1633	0.0
gi 89357459 gb DQ411813.1 Enterococcus faecium strain ATCC 1...	1629	0.0
gi 2828134 gb AF039901.1 AF039901 Enterococcus faecium 16S ribos	1629	0.0
gi 55975493 gb AY653231.1 Enterococcus faecium 16S ribosomal RN	1626	0.0
gi 67511364 emb AJ874342.1 Enterococcus faecium 16S rRNA gene,	1626	0.0
gi 5732228 gb AF070223.1 AF070223 Enterococcus faecium 16S ribos	1626	0.0
gi 5578754 emb Y18294.1 EFE18294 Enterococcus faecium 16S rRNA g	1626	0.0

- You may need to elect Sequence title.
- You may need to scroll down to find your unknown

Blast Tree View Widget - Microsoft Internet Explorer

File Edit View Favorites Tools Help

Address http://www.ncbi.nlm.nih.gov/blast/treeweb/blast_tree_view.cgi?request=page&rid=1152248784-25350-56013527972.BLASTQ4&dbnarr

Tree view for rid: 1152248784-25350-56013527972.BLASTQ4, query ID: lcl|1_25350, database: nr

This tree was produced using BLAST pairwise alignments. [more...](#)

Tree method: Fast Minimum Evolution
 Sequence Label: Sequence Title (if available)
 Max Seq Difference: 0.75

rectangle slanted radial force Show distance

Mouse over an internal node for a subtree or alignment

Blast names color map

unknown
eubacteria

- Enterococcus sp. A20 16S ribosomal RNA gene, partial sequence
- Enterococcus sp. 9c 16S ribosomal RNA gene, partial sequence
- Enterococcus sp. Pd-2006 16S ribosomal RNA gene, partial sequenc
- Enterococcus sp. C1-2006 16S ribosomal RNA gene, partial sequenc
- Enterococcus mundtii 16S ribosomal RNA gene, partial sequence
- Enterococcus sp. CF-2005 16S ribosomal RNA gene, partial sequenc
- Enterococcus mundtii 16S rRNA gene
- Enterococcus mundtii gene for 16S rRNA, partial sequence
- Enterococcus sp. T4-2006 16S ribosomal RNA gene, partial sequence
- Enterococcus pseudoavium 16S rRNA gene, strain LMG 11426
- Enterococcus hirae 16S rRNA gene, strain LMG 6399
- Enterococcus hirae 16S rRNA gene, strain DSM20160
- Enterococcus azikeevi partial 16S rRNA gene, strain IB-A35
- Enterococcus hirae 16S ribosomal RNA gene, partial sequence
- Enterococcus hirae 16S rRNA gene
- Enterococcus hirae isolate C17456 16S ribosomal RNA gene, partial sequence
- Enterococcus hirae partial 16S rRNA gene, strain CECT 4081
- Uncultured bacterium clone OTU7 16S ribosomal RNA gene, partial sequence
- Enterococcus hirae gene for 16S rRNA, partial sequence, strain: DST 2010
- Uncultured bacterium clone OTU9 16S ribosomal RNA gene, partial sequence
- Bacterium mpn-isolate group 13 16S ribosomal RNA gene, partial sequence
- Enterococcus hirae 16S rRNA gene, strain CECT279T
- Enterococcus hirae 16S rRNA gene
- uncultured bacterium partial 16S rRNA gene, clone PEH55
- Enterococcus sanguinicola strain ss1743 16S ribosomal RNA gene, partial sequence
- Enterococcus sanguinicola strain BAA-781 16S ribosomal RNA gene, partial sequence
- Enterococcus sp. A3 16S ribosomal RNA gene, partial sequence

Internet

PCR methods

PCR with standard thermal cycler:

This is a robust reaction and almost always produces a good PCR product. However to be sure, gel electrophoresis of the product may be done.

Reagents needed for 24 ul final volume reactions:									
Reaction tube number	1	2	3	4	5	6	7	8	
Master mix SEQ16S-MM	19.6	39.2	58.8	78.4	98	117.6	137.2	156.8	μl
AmpliTaq Gold polymerase	0.4	0.8	1.2	1.6	2	2.4	2.8	3.2	μl
Dispense 20 ul /tube									
Add 4 ul DNA/tube									

- Prepare master mix using the table.
- Dispense 20 microliters into 0.2 ml PCR tube.
- Add 4 ul of the bacterial DNA to the PCR tube.
- Place into thermal cycler.

Sample program using a Perkin Elmer Gene Amp 9600 thermal cycler. (or equivalent):

Thermal cycling program:

94C 10 minutes hold to activate the AmpliTaq Gold LD polymerase.

40 cycles:

53C for 60 sec

72C for 90 sec

94 for 45 sec

Final 5 min 72C hold.

- (Optional) Gel electrophoresis of PCR product.

PCR with Real-Time thermal cycler:

Adding Sybr Green to the PCR reaction permits detection of the PCR product with several real time PCR thermal cyclers. Real time detection provides evidence for production of a PCR product suitable for sequencing.

Reagents needed for 24 ul final volume reactions								
Reaction tube number	1	2	3	4	5	6	7	8
Master mix: SEQ 16S-MM	19.6	39.2	58.8	78.4	98	117.6	137.2	156.8 μ l
AmpliTaq Gold polymerase	0.4	0.8	1.2	1.6	2	2.4	2.8	3.2 μ l
SYBR Green 1:1000	1	2	3	4	5	6	7	8 μ l
Dispense 21 ul / tube								
Add 3 ul DNA / tube								

- Prepare master mix using the table.
- Dispense 20 microliters into 0.2 ml PCR tube.
- Add 3 ul of the bacterial DNA to the PCR tube.
- Place into thermal cycler such as RotorGene.

Thermal cycling program:

94C 10 minutes hold to activate the AmpliTaq Gold LD polymerase.

40 cycles:

53C for 60 sec

72C for 90 sec Acquire on FAM/Sybr Green channel.

94 for 45 sec

Final 5 min 72C hold.

PCR with LightCycler using glass capillaries:

Adding Sybr Green to the PCR reaction permits detection of the PCR product with the LightCycler. LightCycler real time detection provides evidence for production of a PCR product suitable for sequencing.

LightCycler	Reagents needed for 24 ul final volume reactions							
Reaction tube number	1	2	3	4	5	6	7	8
Master mix: SEQ 16S-MM	19.6	39.2	58.8	78.4	98	117.6	137.2	156.8 μ l
AmpliTaq Gold polymerase	0.4	0.8	1.2	1.6	2	2.4	2.8	3.2 μ l
BSA 1 mg/ml	1	2	3	4	5	6	7	8 μ l
SYBR Green 1:1000	1	2	3	4	5	6	7	8 μ l
Dispense 22 ul / glass capillary								
Add 2 ul DNA / glass capillary								

- Prepare master mix using the table.
- Dispense 22 microliters into glass capillary.
- Add 2 ul of the bacterial DNA to the glass capillary.
- Place into LightCycler.

Thermal cycling program:

94C 10 minutes hold to activate the AmpliTaq Gold LD polymerase.

40 cycles:

53C for 60 sec

72C for 90 sec Acquire on F1 gain=1.

94 for 45 sec

72C for 5 min hold.

40C for 30 seconds cool.

- To remove from glass capillary, invert the capillary and gently centrifuge for 1 second and pipet the PCR product after removing cap.

For technical support, contact Attostar@Attostar.com

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