



# **SBT RESOLVER<sup>TM</sup> DRB1**

**PCR Amplification and Sequencing of HLA DRB1 Full Exon 2**

## **Instructions for Use**

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For Research Use Only. Not for use in diagnostic procedures. No claim or representation is intended to provide information for the diagnosis, prevention, or treatment of a disease.



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# Contents

<b>PRINCIPLE .....</b>	<b>3</b>
<b>KIT COMPOSITION .....</b>	<b>4</b>
<b>STORAGE REQUIREMENTS.....</b>	<b>5</b>
<b>MATERIALS, REAGENTS AND EQUIPMENT NOT SUPPLIED.....</b>	<b>5</b>
<b>SAMPLE REQUIREMENTS.....</b>	<b>6</b>
<b>WARNINGS AND SAFETY PRECAUTIONS .....</b>	<b>6</b>
<b>PROCEDURE.....</b>	<b>7</b>
1. PCR .....	7
2. AGAROSE GEL ELECTROPHORESIS .....	8
3. PURIFICATION OF PCR PRODUCT .....	8
4. SEQUENCING REACTION .....	9
5. PURIFICATION OF SEQUENCING REACTION PRODUCTS.....	10
6. DENATURATION & ELECTROPHORESIS OF SEQUENCING REACTION PRODUCTS .....	11
7. EDITING AND ANALYSIS OF ELECTROPHEROGRAMS .....	11
<b>PERFORMANCE CHARACTERISTICS .....</b>	<b>12</b>
<b>LIMITATIONS AND CAUTIONS .....</b>	<b>12</b>
<b>LICENSE .....</b>	<b>12</b>
<b>TROUBLESHOOTING.....</b>	<b>13</b>
<b>RELATED PRODUCTS.....</b>	<b>15</b>
<b>SUPPORT AND CONTACT DETAILS.....</b>	<b>16</b>

## **Principle**

The HLA Sequence Based Typing (SBT) procedure described here, involves the initial amplification of the target sequences followed by treatment with ExoSAP-IT<sup>®</sup> to remove unincorporated primers and dNTPs. The amplicon is then used as a template for direct automated fluorescent DNA sequencing using customized sequencing primers and the Big Dye<sup>®</sup> Terminator sequencing chemistry available from Applied Biosystems<sup>™</sup> by Life Technologies<sup>™</sup>. The extension products are purified according to the ethanol precipitation method and denatured using Hi-Di<sup>™</sup> formamide available from Applied Biosystems<sup>™</sup> by Life Technologies<sup>™</sup>, before separation and detection on an automated fluorescent DNA sequencer. It is recommended that the resulting data is then analysed with Assign<sup>™</sup> sequence analysis software from Conexio Genomics Pty Ltd.

## Kit Composition

Kit	REF	$\Sigma$	PRE-PCR Contents <sup>†</sup> (No of vials)		POST-PCR Contents (No of vials)		
HLA-DRB1	HH-PD5.2-5(20)	20 tests	<b>DNA POL – DRB1</b>	1 x 9 $\mu$ L	<b>DRB1EX2F</b>	<b>DRB1EX2R-2</b>	1 x 44 $\mu$ L each
			<b>HLA-DRB1 MIX</b>	1 x 370 $\mu$ L	<b>DRB1EX3R-2</b>	<b>RB-TG344-R</b>	
	HH-PD5.2-5(50)	50 tests	<b>DNA POL – DRB1</b>	1 x 18 $\mu$ L	<b>DRB1EX2F</b>	<b>DRB1EX2R-2</b>	1 x 110 $\mu$ L each
			<b>HLA-DRB1 MIX</b>	1 x 920 $\mu$ L	<b>DRB1EX3R-2</b>	<b>RB-TG344-R</b>	

<sup>†</sup>The PRE-PCR kit contains a vial of a locus-specific PCR mix (i.e. **HLA-DRB1 MIX**) consisting of PCR buffer, dNTPs, MgCl<sub>2</sub>, and locus specific PCR primers, along with a single vial of DNA polymerase (**DNA POL – DRB1**).

The POST-PCR kit contains sequencing primers (e.g. **DRB1EX2F**).

## Storage Requirements

The PRE- and POST-PCR boxes may be separated and stored in designated PRE- and POST-PCR freezers. When stored at -20°C (temperature range of -15°C to -25°C is acceptable), the kit components can be used until the expiry date indicated on the outer labels and can tolerate up to 25 freeze-thaw cycles.

To maintain optimal kit performance, the kit components should be removed from the -20°C storage location and thawed rapidly at room temperature before use. The kit components, with the exception of the polymerase, should then be gently vortexed to ensure that the components of each tube are appropriately mixed after thawing. After use, the kits/components should be returned immediately to -20°C.

## Materials, Reagents and Equipment Not Supplied

### PCR

1. Sterile water
2. Electronic or mechanical pipettes and aerosol-resistant tips
3. Thermal cycler with heated lid  
These kits have been validated using the following thermal cyclers:  
MJ Research PTC 225 DNA Engine DYAD™, Applied Biosystems™ by Life Technologies™ Gene Amp® PCR System 9700, and Eppendorf Mastercycler® Pro.  
**Use of other thermal cyclers with these kits requires validation by the user.**
4. 0.2mL thin-walled thermal cycling reaction tubes (8 well strips or 96 well plates).  
Use those recommended for use with your thermal cycler.
5. Sterile 1.5mL tubes
6. Sterile work area such as biological safety cabinet or hood.
7. Table top centrifuge with plate adapters and capacity to reach 2500 x g
8. Vortex

### Agarose Gel Electrophoresis

9. Agarose gel electrophoresis apparatus
10. 1% agarose (molecular biology grade) TBE gel containing 0.1µg/mL ethidium bromide.
11. Loading buffer
12. PCR Marker suitable to cover range of 300 – 1300 bp
13. UV transilluminator

### Purification of PCR Product

14. ExoSAP-IT® (USB® Products Cat No 78200 for 100 reactions)
15. 2mM MgCl<sub>2</sub> (available for purchase from Conexio Genomics, product code MgCl2-1.0(50) or MgCl2-1.0(3000))
16. Shaker

**The use of alternative PCR purification techniques requires validation by the user prior to use.**

## Sequencing Reaction

17. BigDye® Terminator Cycle Sequencing Kit v3.1 or v1.1, Applied Biosystems™ by Life Technologies™.
18. 5x Sequencing Reaction Buffer (Conexio Genomics, product code SEQ BUF-2.0(400) or SEQ BUF-2.0(5000)) or BigDye® Terminator v3.1 or v1.1 5X Sequencing Buffer, Applied Biosystems™ by Life Technologies™.

## Purification of Sequencing Reaction Products

19. 125mM EDTA, pH8.0 (available for purchase from Conexio Genomics, product code EDTA-3.0(200) or EDTA-3.0(5000))
20. Absolute and 80% Ethanol. Each run requires freshly prepared 80% ethanol consisting of absolute ethanol and sterile water. DO NOT USE DENATURED ETHANOL.

**The use of alternative sequencing purification techniques requires validation by the user prior to use.**

## Denaturation and Electrophoresis of Sequencing Reaction Products

21. Hi-Di™ Formamide, Applied Biosystems™ by Life Technologies™, product code 4311320
22. Automated DNA Sequencer and accessories (e.g. Applied Biosystems™ by Life Technologies™ ABI Prism® 3730), including data collection and software.  
  
These kits have been tested and validated on the Applied Biosystems™ by Life Technologies™ 3100, 3730 and 3730xl capillary sequencers and software.

**The use of other denaturation techniques and sequencing platforms requires validation by the user prior to use.**

23. HLA Sequencing Analysis Software (e.g. Assign™ SBT, version 3.5 or higher or Assign™ ATF, Conexio Genomics Pty Ltd).

## Sample Requirements

1. Sterile water (negative/ no template control)
2. High molecular weight human genomic DNA (concentration range of 20-100ng/μL in Tris/EDTA buffer and  $OD_{260/280} > 1.8$ ) extracted from ACD or EDTA anticoagulated whole blood specimens. Do NOT use whole blood specimens containing heparin.



## Warnings and Safety Precautions

- This kit must be used by trained and authorized laboratory personnel.
- All samples, equipment and reagents must be handled in accordance with good laboratory practice. In particular, all patient material should be considered as potentially infectious. The use of gloves and laboratory coats is strongly

recommended. Handle and dispose of all sample material according to local and national regulatory guidelines.

- There are NO dangerous substances contained in any of the kit components.
- Do NOT use reagents beyond their expiration date.
- The use of kit components from different kit batches is NOT recommended. Such use may affect the assay's performance.
- Use of reagents not included in this kit or not listed under "Materials, Reagents and Equipment Not Supplied" (e.g. alternative *Taq* DNA polymerases) is NOT recommended. Such use may affect the performance of the assay.
- Care should be taken to prevent cross-contamination of DNA specimens. Change tips between DNA specimens wherever possible.
- Pre- and Post-PCR activities must be strictly physically separated. Use specifically designated equipment, reagents and laboratory coats.
- Ethidium bromide is a potential carcinogen. Protective gloves must always be used when preparing and handling gels. Dispose of ethidium-bromide gels and buffers according to local and national guidelines.
- While viewing and photographing agarose gels under UV light, always avoid direct exposure and use appropriate UV-blocking face protection, disposable gloves and laboratory coats.

## Procedure

### 1. PCR

- 1.1. A separate PCR reaction will need to be set up for each sample to be tested. Each run should include appropriate positive control/s of known genotype, and at least one negative control.
- 1.2. Prepare a fresh solution of PCR master mix each time a PCR is performed. Quickly thaw the locus-specific PCR mix at room temperature. Once thawed, vortex briefly.
- 1.3. Dispense the required volume of PCR mix and DNA polymerase into a sterile tube for the number of samples to be tested (refer to Table 1 below for the volume per reaction). Pulse vortex the solution 3-4 times.

Component	Volume
HLA-DRB1 MIX	16.7 $\mu$ L
DNA POL – DRB1	0.3 $\mu$ L

**Table 1: Composition of the master mix required per sample.**

- 1.4. Dispense 17 $\mu$ L of the master mix into each reaction well.
- 1.5. Add 3 $\mu$ L of sample DNA or appropriate positive control/s to each reaction well. Add 3 $\mu$ L of sterile water to the negative control reaction well.

- 1.6. Seal the reaction wells. Mix gently by vortexing and centrifuge briefly.
- 1.7. Place the reaction wells into a thermal cycler and run according to the thermal cycling conditions below.

95°C - 10 mins  
96°C - 20 secs } 33 cycles  
60°C - 30 secs }  
72°C - 3 mins }  
15°C - hold

- 1.8. Amplification takes approximately 2.5 hours to complete.
- 1.9. When the PCR is completed, remove the reaction wells/plate from the thermal cycler and either proceed directly to gel electrophoresis or store at 4°C until required.

**NOTE:** Purification of amplicons by ExoSAP-IT<sup>®</sup> treatment should occur within 24 hours of completion of PCR.

## 2. Agarose Gel Electrophoresis

- 2.1. Confirm successful amplification by agarose gel electrophoresis using 2µL of each PCR product combined with 5µL loading buffer (alternative volumes of loading buffer should be validated prior to use). The use of 1% agarose gels is recommended.

The number and expected sizes of the resultant amplicons will vary according to the sample genotype. Expected PCR amplicon sizes range between ≈ 450 bp - 850bp.

## 3. Purification of PCR Product

**NOTE:** Purification systems other than EXOSAP-IT<sup>®</sup> (e.g. Agencourt<sup>®</sup> AMPure<sup>®</sup> XP or column-based systems) can be used to purify these PCR products. It is strongly recommended that users validate these procedures before proceeding. If EXOSAP-IT<sup>®</sup> is to be used it is recommended that users follow the procedure described below.

- 3.1. Prepare a mastermix consisting of 4µL of ExoSAP-IT<sup>®</sup> and 8µL of 2mM MgCl<sub>2</sub> per sample to be purified. Dispense 12µL of the mastermix into the reaction well of each reactive sample. Seal the wells and vortex to mix. Centrifuge briefly before placing into the thermal cycler. Run the thermal cycler according to the following profile:

37°C - 30mins  
80°C - 15mins  
4°C - hold

- 3.2. Upon completion, dilute the purified product 1:4 with sterile water. This dilution step will ensure that there is sufficient template to perform the sequencing reactions and ensure that the concentration of the template is sufficient to produce good quality sequence data.

**NOTE:** A higher dilution factor (e.g. 1:8) may be required if consistently high signals and associated noise and artefacts are observed. Weaker PCR products may require a lower dilution factor.



3.3. ExoSAP-IT<sup>®</sup> treated samples may be stored at 4°C until ready for use. ExoSAP-IT<sup>®</sup> treated samples can be stored at 4°C for up to a week before use, but should be stored at -20°C for long term storage.

## 4. Sequencing Reaction

**NOTE:** In instances where heterozygous ambiguities are to be resolved with hemizygous sequencing primers such as HARPS<sup>®</sup>, please refer to the SBT Resolver™ HARPS<sup>®</sup> Instructions for Use.

4.1. Table 2 lists the sequencing primers that are to be used for this kit.

HLA-DRB1 <sup>†</sup>	
DRB1EX2F	DRB1EX2R-2
DRB1EX3R-2	RB-TG344-R <sup>†</sup>

**Table 2: Sequencing primers provided for use with the HH-PD5.2-5 kits.**

<sup>†</sup>RB-TG344-R is a HARP<sup>®</sup> directed to the codon 86 dimorphism. Its use is optional.

4.2. Prepare a fresh solution of sequencing primer mix on ice each time a sequence reaction is performed. The composition and volumes for the mix indicated below are **per sample**.

Component	Volume
Sequencing primer	2 µL
Sterile water	11.5 µL
BigDye <sup>®</sup> Terminators	1 µL
5x Seq Rxn Buffer	3.5 µL

4.3. Mix each sequencing reaction mixture gently by pulse vortexing.

4.4. Dispense 18µL of the sequencing reaction mix into each appropriate reaction well.

**NOTE:** For runs which involve few samples with many sequencing primers, it is acceptable to dispense the sequencing primer (2µL) directly into the individual reaction wells. A master mix may then be created composing of sterile water, BigDye<sup>®</sup> Terminators and 5x Seq Rxn Buffer, of which 16µL is to be dispensed into each reaction well. It is strongly recommended that use of this alternative procedure is validated by the user prior to implementation.

4.5. Add 2µL of purified PCR product to each appropriate well.

**NOTE:** Care must be taken to prevent cross-contamination of sequence reactions.

4.6. Seal the reaction wells, mix gently and centrifuge briefly to ensure that the contents are located at the base of each reaction well.

4.7. Place the reaction wells into a thermal cycler and run according to the following profile:

Number of cycles	Temperature and time
25	96°C – 10sec 50°C – 5sec 60°C – 2min
1	4°C - hold

- 4.8. Once the program is complete, remove the reaction wells/plate from the thermal cycler and either proceed directly to purification of the reaction products or store in the dark at 4°C until required. It is recommended that samples are purified and run on the DNA sequencer within 24 hours.

## 5. Purification of Sequencing Reaction Products

**NOTE:** Purification of the reaction products may be carried out by procedures other than the ethanol precipitation method described here. It is strongly recommended that users validate these procedures before proceeding.

- 5.1. Briefly centrifuge the reaction wells/plates before proceeding. If reusable lids/caps have been used during thermal cycling, label the lids/caps to avoid cross-contamination.
- 5.2. Carefully remove the seals.
- 5.3. To each reaction well add 5µL of 125mM EDTA, pH8.0. Ensure that the EDTA reaches the base of the reaction well.
- 5.4. Add 60 µL of 100% ethanol to each reaction well. Seal the wells/plate and vortex briefly but thoroughly to ensure thorough mixing.
- 5.5. Pellet the extension products by centrifuging at 2000g for 45 minutes. **IMMEDIATELY PROCEED TO THE NEXT STEP.** If this is not possible, re-centrifuge for an additional 10 minutes before proceeding.
- 5.6. Remove the seals to the reaction wells and discard the supernatant by inverting the reaction wells onto paper towel or tissues.
- 5.7. Place the inverted reaction wells and paper towel or tissue into the centrifuge. Centrifuge at 350g for 1 minute to remove any residual supernatant.
- 5.8. Remove the reaction wells from the centrifuge and place in an upright position on the work bench. Discard the paper towel or tissues.
- 5.9. Prepare fresh solution of 80% ethanol with absolute ethanol and sterile water.
- 5.10. Add 60µL of 80% ethanol to each well. Reseal the wells and vortex briefly.
- 5.11. Spin at 2000g for 5 minutes.
- 5.12. Repeat steps 5.6 and 5.7.
- 5.13. Remove the reaction wells from the centrifuge and discard the paper towel. Reseal the reaction wells and proceed to the denaturation step. Otherwise store at -20°C in the dark. It is recommended that the extension products are run on the DNA sequencer within 24 hours of setting up the sequencing reactions.

## 6. Denaturation & Electrophoresis of Sequencing Reaction Products

**NOTE:** The procedure for the denaturation of extension products in Hi-Di™ Formamide described here may not be necessary if purification procedures other than the ethanol precipitation have been used. It is strongly recommended that users validate alternative procedures before proceeding.

- 6.1. Add 12µL of Hi-Di™ Formamide to each reaction well. Vortex and centrifuge the wells/plate briefly.
- 6.2. Incubate the reaction wells at 98°C for 5 minutes. Following incubation, ensure that the reaction wells are cooled quickly to room temperature (e.g. place on ice or use the thermal cycler to perform the denaturation and cooling steps) before being placed on the sequencer. If it is not possible to run the plates immediately, store at 4°C until required.

**NOTE:** Ensure that there are no air bubbles in the reaction wells. These can enter and damage the capillary.

- 6.3. Load the reaction wells/plate onto the automated sequencer and prepare the data collection file according to the sequencer manufacturer specifications.
- 6.4. The following instrument parameters have been validated by the manufacturer using Big Dye® Terminator Sequencing Kit v3.1 and POP-7™. These parameters may require user validation for other polymers, sequencing chemistries and instruments. Please refer to the appropriate instrument user's manual for detailed instructions and guidance (e.g. ensure that the dye set setting is appropriate for the chemistry used, for example v1.1 Big Dye® Terminator sequencing chemistry will require a different dye set).

<b>Parameter</b>	<b>Setting</b>
Dye set	Z_BigDyeV3
Mobility file	KB_3730_POP7_BDTV3
Basecaller	KB.bcp
Run Module	Regular FastSeq50_POP7
Injection time	15 sec
Run time	3000 sec

- 6.5. Use the instrument's data collection software to process the raw collected data and create the sequence files. Please refer to the appropriate instrument user's manual for detailed instructions and guidance.

## 7. Editing and analysis of electropherograms

The SBT Resolver™ kits were developed and validated using the Assign™ SBT and Assign™ ATF software developed by Conexio Genomics Pty Ltd. For more details please refer to the Conexio Genomics website (<http://www.conexio-genomics.com>).

## Performance Characteristics

Conexio Genomics Pty Ltd's SBT Resolver™ kits are locus specific assays. Sequence analysis of PCR and sequencing primer sites and performance evaluation has not identified alleles that are not amplified through the recommended use of these kits.

In most instances the use of the sequencing primers incorporated in each kit will produce a HLA typing for most samples without the need for further resolution. In those instances where heterozygous ambiguities remain, the use of resolving sequencing primers (such as SBT Resolver™ HARPS®) is recommended.

It should be noted that mutations at amplification or sequencing primer sites are possible and may result in allele drop-out. Samples that suggest a homozygous typing result must be confirmed by alternative procedures.

## Limitations and Cautions

- It is strongly recommended that these kits are validated by the user prior to implementation in the laboratory using samples whose HLA type has been determined by other molecular based procedures. In particular, any deviations from this procedure (e.g. the use of alternative PCR or DNA sequencing purification procedures) must be validated by the user prior to implementation.
- These kits have been validated using panels of samples whose genotypes cover a broad range of alleles. However it should be noted that rare alleles and alleles with polymorphisms in amplification and sequencing primer sites may be encountered and these may not be amplified or sequenced.
- The nature of HLA sequence based typing is such that factors other than the PCR mix may result in preferential amplification or allele drop out. As a consequence, apparent homozygous typing results should be confirmed using alternative methods and/or family genotyping.
- A positive control (human DNA) and negative control (sterile water) must be included on every PCR run. The positive control must produce a PCR product of the appropriate size depending on the locus amplified and the resultant sequence must be in concordance with the sample's genotype. There must be no PCR products in the negative template control for each experiment. If a band is evident contamination may have occurred at some level and the run must be repeated.
- Occasionally there may be additional, fainter PCR products evident. These additional bands do not interfere with sequence results or quality.

## License

The SBT Resolver™ kits contain GoTaq® Hot Start Polymerase (DNA POL) which is manufactured by Promega Corporation for distribution by Conexio Genomics Pty Ltd. Licensed to Promega under U.S. Patent Nos. 5,338,671 and 5,587,287 and their corresponding foreign patents.

## Troubleshooting

<b>Problem</b>	<b>Possible cause(s)</b>	<b>Solution</b>
No or weak PCR product	Poor quality DNA	Assess DNA quality by gel electrophoresis. Intact DNA should be approx 3kb with little or no evidence of smearing on gel. Re-extract DNA and repeat PCR where possible.
	Insufficient quantity of DNA added to PCR.	Check concentration of DNA is between 20-100ng/ $\mu$ L. Re-extract DNA and repeat PCR where possible.
	Presence of PCR inhibitors in genomic DNA	Avoid the use of whole blood specimens containing heparin. Re-extract DNA and repeat PCR where possible.
	DNA polymerase not added to the mastermix or insufficient mixing of mastermix prior to addition to samples.	Repeat PCR. Ensure mastermix components are added and mixed sufficiently by vortexing.
	Thermal cycling problems	Check the thermal cycling run parameters. Check the run history to ensure that the run was not terminated prematurely. Ensure that the thermal cycler is operating according to manufacturer's specifications and is regularly maintained.
	No ethidium bromide added to the gel.	Submerge the gel in a staining bath containing 1X TBE with 0.5mg/mL ethidium bromide. Destain in 1X TBE before taking gel image. Ensure ethidium bromide is added to gel prior to pouring.
Incorrect band sizes	Incorrect kit used	Check that the appropriate kit is used.
	Incorrect thermal cycling program used.	Check the thermal cycle parameters.
	PCR contamination	Check the negative control for evidence of contamination. Decontaminate work area and repeat PCR. Repeat PCR to identify source of contamination. Consider using a fresh kit. If the genomic DNA of a sample appears to be contaminated, re-extract or

		obtain an alternative source of DNA.
Weak signal intensity of electropherograms	Weak PCR product	Check gel image. Sequencing weak PCR bands is NOT recommended as the sequence quality may be insufficient for SBT. Consider using a lower dilution factor (eg 1:2, 1:3) after PCR purification.
	Insufficient reaction products applied to sequencer	Check sequencer parameters. Injection time and voltage may need to be increased.
	Problems during purification of sequencer products	Use extreme care when discarding the supernatant as it may dislodge the pellet.
Signal intensity is too high (Presence of high fluorescent peaks – artefacts)	Too much PCR product	Check the gel image. Consider using a higher dilution factor following PCR purification. Check the amount of DNA polymerase used in the PCR.
	Too much reaction products applied to sequencer.	Check instrument parameters. Consider reducing the injection time and voltage.
Noisy baseline (high background)	Contaminated PCR product	Refer to corrective actions listed above.
	Amplification of closely related HLA genes	Check thermal cycling parameters. Consider adjusting the parameters if an alternative thermal cycler is used.
	Poor PCR purification	Ensure ExoSAP-IT <sup>®</sup> treatment is undertaken according to kit's user instructions. Ensure that the PCR mixture is mixed thoroughly with ExoSAP-IT <sup>®</sup> .
	Contaminated sequencing reactions	Ensure that all steps are taken to prevent cross contamination. Change pipette tips wherever possible. Add liquids at the top of the reaction wells. Prevent aerosols.
	Contaminated sequencing primer	Check sequence quality of the other sequencing primers and other samples using the same primer.
	Contaminated dye terminator mix or sequencing buffer	Repeat sequencing with fresh aliquot of reagents.
	Poor purification of sequencing products.	Repeat sequencing and ensure that purification is undertaken according to manufacturer's instructions.

Presence of Dye blobs	Poor purification of sequencing products	Purify products according to kit instructions. Ensure products are washed sufficiently with 80% ethanol.
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## Related Products

**ASSIGN<sup>™</sup> SBT 3.8+**

Product code: CGX0036+

## SBT RESOLVER<sup>™</sup>

XH-PD1.1-2(20) XH-PD1.1-2(50)	SBT Resolver <sup>™</sup> HLA-A kit (20 and 50 tests)
BS-PD2.1-2(20) BS-PD2.1-2(50)	SBT Resolver <sup>™</sup> HLA-B kit (20 and 50 tests)
HH-PD3.2-2(20) HH-PD3.2-2(50)	SBT Resolver <sup>™</sup> HLA-C kit (20 and 50 tests)
PQ-PD6.2-2(20) PQ-PD6.2-2(50)	SBT Resolver <sup>™</sup> HLA-DQB1 kit (20 and 50 tests)
HH-PD10.1(20) HH-PD10.1(50)	SBT Resolver <sup>™</sup> HLA-DPB1 kit (20 and 50 tests)
AN-PD11.0-0(20) AN-PD11.0-0(50)	SBT Resolver <sup>™</sup> HLA-DRB3 kit (20 and 50 tests)
AN-PD12.0-0(20) AN-PD12.0-0(50)	SBT Resolver <sup>™</sup> HLA-DRB4 kit (20 and 50 tests)
AN-PD13.0-0(20) AN-PD13.0-0(50)	SBT Resolver <sup>™</sup> HLA-DRB5 kit (20 and 50 tests)
LC-PD2.9(20) LC-PD2.9(50)	SBT Resolver <sup>™</sup> HLA-B57 kit (20 and 50 tests)

## SBT RESOLVER<sup>™</sup> HARPS

Product codes:

C1-TT98-F(20)	C1-AC98-F(20)	C1-TC98-F(20)	C1-TA98-F(20)	C1-CA102-F(20)
C1-CT102-F(20)	C1-CC102-F(20)	C1-AG203-F(20)	C1-GT240-F(20)	C1-TT368-F(20)
C1-GG307-R(20)	C1-GG363-AF(20)	C1-TA363-F(20)	C1-AT362-F(20)	C1-AC497-F(20)
C1-TA368-F(20)	C1-GT355-R(20)	C1-GG362-R(20)	C1-CT423-F(20)	C1-CG570-R(20)
C1-BTA-F(20)	C1-BCG-F(20)	C1-CC144-F(20)	C1-AC206-F(20)	C1-GC209-F(20)
C1-GA206-F(20)	C1-CG319-F(20)	C1-CA309-R(20)	C1-GAT309-R(20)	C1-GAA309-R(20)
C1-AG360-F(20)	C1-GC363-F(20)	C1-GG363-BF(20)	C1-TA420-F(20)	C1-AC362-F(20)
C1-CC486-F(20)	C1-CT559-R(20)	C1-GA559-R(20)	C1-AC559-R(20)	C1-GG572-R(20)
C1-CG572-R(20)	C1-GAG601-R(20)			
C1-CT97-F(20)	C1-CT112-F(20)	C1-CG134-F(20)	C1-AG270-F(20)	C1-AC302-R(20)
C1-GC302-R(20)	C1-CG343-F(20)	C1-CA343-F(20)	C1-GA361-F(20)	C1-TG539-R(20)
C1-GG539-R(20)	C1-AA601-R(20)			
RB-01-F(20)	RB-04-F(20)	RB-09-F(20)	RB-15-F(20)	RB-52-F(20)

RB-GG125-F(20)	RB-AA197-F(20)	RB-TT197-F(20)	RB-GT196-F(20)	RB-GA196-F(20)
RB-TA164-F(20)	RB-TT227-F(20)	RB-AT258-F(20)	RB-GC258-F(20)	RB-CT257-R(20)
RB-AT257-R(20)	RB-TT321-R(20)	RB-GT344-R(20)	RB-TG344-R(20)	
QB-TA173-F(20)	QB-CT173-F(20)	QB-TA185-F(20)	QB-CG353-R(20)	QB-GG353-R(20)
PB-AT251-R(20)	PB-GT313-R(20)	PB-TAC121-F(20)	PB-GG341-R(20)	PB-GC194-F(20)
PB-AG341-R(20)				

## General Purpose Laboratory Reagents

MgCl <sub>2</sub> – 1.0(50)	2mM MgCl <sub>2</sub>
MgCl <sub>2</sub> - 1.0(3000)	
SEQ BUF – 2.0(400)	5x Seq Rxn Buffer
SEQ BUF – 2.0(5000)	
EDTA – 3.0(200)	125mM EDTA, pH8.0
EDTA – 3.0(5000)	

Please contact your local distributor for further details.

## Support and Contact Details

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For ordering details, please refer to the Olerup website (<http://www.olerup.com>).

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