February 2019

# QlAsymphony® SP Protocol Sheet

Casework\_1000\_V8 and Casework\_1000\_H2O\_V8 protocol

This document is the Casework\_1000\_V8 and Casework\_1000\_H2O\_V8 *QlAsymphony SP Protocol Sheet*, R1, for QlAsymphony DNA Investigator® Kit.



#### General information

The QIAsymphony DNA Investigator Kit is intended for molecular biology applications. This product is not intended for the diagnosis, prevention, or treatment of a disease.

These protocols are for purification of total DNA from samples encountered in forensic, human identity, and biosecurity applications using the QIAsymphony SP and the QIAsymphony DNA Investigator Kit.

Since the type of samples that can be processed using the QIAsymphony DNA Investigator Kit can vary greatly, there is also a variety of different pretreatments, optimized for specific sample types. For the CW\_1000\_V8 and CW\_1000\_H2O\_V8 protocol, samples are lysed under denaturing conditions in the presence of proteinase K and Buffer ATL in a total volume of 1000 µl.

The Elution can be performed either with the buffer ATE ( $CW_1000_V8$ ) or with the buffer AVE ( $CW_1000_V8$ ).

**Note**: It is the user's responsibility to validate performance using this combination for any procedures used in their laboratory.

#### CW\_1000\_H2O\_V8

Kit	QIAsymphony DNA Investigator Kit
Sample material	Large-volume forensic samples
Protocol name	CW_1000_H2O_V8
Default Assay Control Set	ACS_CW_1000_H2O_V8
Elution solution	Buffer AVE
Elution volume	ابر 150 بار 150 بار 100 با
Required software version	Version 5.0 or higher

## CW\_1000\_V8

Kit	QIAsymphony DNA Investigator Kit
Sample material	Large-volume forensic samples
Protocol name	CW_1000_V8
Default Assay Control Set	ACS_CW_1000_V8
Elution solution	Buffer ATE
Elution volume	100 µl, 150 µl, 200 µl
Required software version	Version 5.0 or higher

# Materials required but not provided

## For all sample types

- Vortexer
- Thermomixer or shaker-incubator

# "Sample" drawer

Sample type	Large-volume forensic samples
Sample volume	ابر 1000
Primary sample tubes	See www.qiagen.com/goto/qsdnainvestigator for more information
Secondary sample tubes	See www.qiagen.com/goto/qsdnainvestigator for more information
Inserts	See www.qiagen.com/goto/qsdnainvestigator for more information
Other	n/a

n/a = not applicable.

# "Reagents and Consumables" drawer

Position A1 and/or A2	Reagent cartridge (RC)
Position B1	n/a
Tip rack holder 1–17	Disposable filter-tips, 200 µl
Tip rack holder 1–17	Disposable filter-tips, 1500 µl
Unit box holder 1–4	Unit boxes containing sample prep cartridges
Unit box holder 1-4	Unit boxes containing 8-Rod Covers

n/a = not applicable.

### "Waste" drawer

Unit box holder 1–4	Empty unit boxes
Waste bag holder	Waste bag
Liquid waste bottle holder	Liquid waste bottle

#### "Eluate" drawer

Elution rack (we recommend using slot 1, cooling position)  See www.qiagen.com/goto/qsdnainvestigator for more information	Elution rack (we recommend using slot 1, cooling position)	
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## Required plasticware

	One batch, 24 samples*	Four batches, 96 samples*
Disposable filter-tips, 200 µl†‡	28	112
Disposable filter-tips, 1500 µl†‡	56	224
Sample prep cartridges§	15	60
8-Rod Covers¶	3	12

<sup>\*</sup> Use of less than 24 samples per batch decreases the number of disposable filter-tips required per run.

**Note**: Numbers of filter-tips given may differ from the numbers displayed in the touchscreen depending on settings, for example, number of internal controls used per batch.

# Preparation of sample material

When working with chemicals, always wear a suitable lab coat, disposable gloves, and protective goggles. For more information, consult the appropriate material safety data sheets (MSDSs), available from the product supplier.

#### Important points before starting

- QlAsymphony magnetic particles copurify RNA and DNA if both are present in the sample. If RNA-free DNA is required, add RNase A to the sample in the step indicated in the respective pretreatment protocol.
- Before beginning the procedure, read "Important Notes", page 12 of the QIAsymphony DNA Investigator Handbook.

<sup>&</sup>lt;sup>†</sup> There are 32 filter-tips/tip rack.

<sup>&</sup>lt;sup>‡</sup> Number of required filter-tips includes filter-tips for 1 inventory scan per reagent cartridge.

<sup>§</sup> There are 28 sample prep cartridges/unit box.

<sup>1</sup> There are twelve 8-Rod Covers/unit box.

#### Large-volume samples

This protocol is for isolation of total (genomic and mitochondrial) DNA from forensic samples that require increased volumes for thorough lysis, e.g., diffuse stains on fabric or paper. The pretreatment includes lysis of samples using proteinase K.

#### Things to do before starting

- Before using Buffer ATL, check that it does not contain a white precipitate. If necessary, incubate for 30 minutes at 70°C with gentle agitation.
- Set a thermomixer or shaker-incubator to 56°C for use in step 4.
- If processing semen samples, prepare an aqueous 1 M DTT\* stock solution. Store aliquots at -20°C. Thaw immediately before use.

#### Pretreatment protocol for large-volume samples

- 1. Transfer samples to a 2 ml microcentrifuge tube (not provided). Large samples can be cut into smaller pieces to fit more conveniently.
- 2. Add 960 µl Buffer ATL.
- 3. Add 40  $\mu$ l proteinase K, and mix by vortexing. If processing semen samples, add 40  $\mu$ l 1 M DTT
- 4. Place the tube in a thermomixer or heated orbital incubator, and incubate with shaking at 900 rpm at 56°C for at least 15 min.
- 5. Carefully transfer the lysate to sample tubes or plates that are compatible with the sample rack of the QIAsymphony SP.

See www.qiagen.com/QlAsymphony/Resources for a full list of compatible vessels. We recommend using 2 ml tubes (e.g., Sarstedt, cat. no. 72.693 or 72.608) or S-Blocks (cat. no. 19585).

**Note**: Do not transfer any solid material as this may clog the tips during automated DNA purification.

Lysate remaining in solid sample material (e.g., denim) can be harvested by transferring the material to a QIAshredder spin column (not supplied) and centrifuging at full speed for 2 min in a microcentrifuge. Transfer the flow-through to the sample tube or plate.

- 6. Continue with the protocol "DNA Purification from Casework and Reference Samples" (page 19 in the *QlAsymphony DNA Investigator Handbook*).
- \* When working with chemicals, always wear a suitable lab coat, disposable gloves, and protective goggles. For more information, consult the appropriate material safety data sheets (MSDSs), available from the product supplier.

# Revision history

Document revis	ion history
R1 02/2019	Update for QIAsymphony Software version 5.0

For up-to-date licensing information and product-specific disclaimers, see the respective QIAGEN® kit handbook or user manual. QIAGEN kit handbooks and user manuals are available at **www.qiagen.com** or can be requested from QIAGEN Technical Services or your local distributor.

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