Quick-Start Protocol

February 2018

# MagAttract<sup>®</sup> 96 cador<sup>®</sup> Pathogen Kit

The MagAttract 96 *cador* Pathogen Kit (384) (cat. no. 947457) can be stored at room temperature (15–25°C). For expiry date information, consult the label on the kit box.

### Further information

- MagAttract 96 cador Pathogen Handbook: <u>www.qiagen.com/handbooks</u>
- Technical assistance: toll-free 00800-22-44-6000, or www.qiagen.com/contact

### Important notes before starting

- Read the safety information in the BioSprint® 96 User Manual before use.
- Dissolve carrier RNA in Buffer AVE as indicated on the tube.
- Add isopropanol (100%) to Buffer ACB and ethanol (96–100%) to Buffers AW1 and AW2 before use. See the respective bottle labels for volumes.
- Equilibrate buffers to room temperature (15–25°C).
- If using a sample volume less than 200  $\mu l,$  add PBS to bring total sample volume to 200  $\mu l.$
- 96-Rod Covers are supplied as packs of 1 or 2, inserted into an S-Block. If using a pack of 2 covers, store the second cover in another S-Block or plate. Take care to not bend the covers.

#### Procedure

- 1. Label 5 x S-Blocks and one 96-Well Microplate MP.
- Pipet 20 µl Proteinase K into the bottom of an S-Block well and add 200 µl sample. If your sample volume is less than 200 µl, bring it to 200 µl by adding PBS.
- 3. Prepare Buffer VXL mixture (see Table 1) and mix thoroughly for 30 s.



## Sample to Insight

Note: Do not add proteinase k	( directly	to this	Buffer	VXL ı	nixture!
Table 1. Buffer VXL mixture preparation					

	I	Number of samples*			
Reagent	1	48	96		
Buffer VXL	100 µl	4.8 ml	9.6 ml		
Buffer ACB	400 µl	19.2 ml	38.4 ml		
MagAttract Suspension G	25 µl	1.2 ml	2.4 ml		
Carrier RNA (1 µg/µl)	1 µl	48 µl	96 µl		

\* Prepared volumes are 105% of required volumes to compensate for pipetting errors and possible evaporation. Excess buffer should be discarded.

- 4. Add 500 µl Buffer VXL mixture to each sample in the S-Block.
- 5. Prepare 4 additional S-Blocks (slots 2–6) and one 96-Well Microplate MP according to Table 2.

Table 2. BioSprint 96 worktable setup and reagent volumes

	volume per well (µl)
Large 96-Rod Cove	r _
croplate MP Buffer AVE	100
Ethanol (96–100%)	750
Buffer AW2	700
Buffer AW1	700
Lysate*	720
	Large 96-Rod Cover croplate MP Buffer AVE Ethanol (96–100%) Buffer AW2 Buffer AW1 Lysate*

\* Includes 20 µl Proteinase K, 200 µl sample and 500 µl Buffer VXL mixture.

- 6. Switch the BioSprint 96 on at the power switch.
- 7. Slide the front door of the protective cover open.
- 8. Select the protocol "BS96 cador v2" using the  $\wedge$  and  $\vee$  keys.
- 9. Press "Start" and follow the messages for loading the worktable as shown in Table 2.

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