

## Aquadien DNA Extraction Kit, 3578121 Short Protocol Quick Guide



- Pipet 1 ml of R1 solution into a microtube
- Place a polycarbonate membrane filter on a sterilized filtration apparatus mounted on an air pump or vacuum flask
- Filter 100 ml-1 L of water



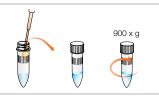
- Carefully fold the membrane in half 3 times to obtain a cone
- Using tweezers, place the membrane in the microtube containing 1 ml of R1 solution







Incubate at 95°C for 15 min at 1,300 rpm in a heating thermoshaker



3 min



- Carefully take out the membrane, pressing it to the walls of the tube to recover all the solution
- Centrifuge at 900 x g for 3 min

The DNA is contained in the 800 µl of supernatant.









- Place a purification column in a collector vial
- Transfer 500 μl of the supernatant to the purification column
- Centrifuge at 6,000 x g for 10 min



- Add 100 μl of R2 solution to the purification column; throw away the collector vial
- Cover the purification column with a clean collector vial and turn the whole unit upside down
- Centrifuge at 1,000 x g for 3 min
- Throw away the purification column



100  $\mu$ l of purified DNA solution is obtained.

Use 5 μl of the extracted DNA solution for real-time PCR analysis

For detailed instructions, review the kit user guide.



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