MACHEREY-NAGEL

NucleoMag[®] DNA Food– Support Protocol Honey

Protocol details

Application	DNA from honey
Kit	NucleoMag [®] DNA Food
REF	744945/.1/.4
Protocol name	NucleoMag DNA Food –Support Protocol Honey



Protocol steps

Steps	Procedure
Homogenize sample	Transfer 1 mL honey into 2 mL tube. Add 900 μL PCR grade water.
	Vortex sample to homogenize water with honey sample. Approximately 30 sec.
	Incubate at 55 °C for 15 Min 12000 rpm.
Precipitate	Centrifuge at 14.000 x g for 10 min.
	Discard the supernatant, avoid touching the sides of the tube.
	Note: Pellet might not be visible but forms at the bottom and on the sides of the tube.
Wash pellet	Add 400 µL PCR grade water to the pellet
	and vortex until the pellet is homogenized
	Centrifuge 14.000 x g 5 Min. Discard supernatant
Lyse sample	Add 20 μ L Proteinase K and 380 μ L Lysis Buffer CF to the pellet. Resuspend the pellet by pipetting up and down and transfer the sample to a MN Bead Tube Type B.
	Homogenize the sample in a Bead or Retsch mill (30 Hz. 3 Min).
	Centrifuge 11.000xg 1 min.
	Transfer supernatant to a new 1,5 mL Tube.
	Incubate 55 °C 10 min.
	For manual procedure transfer ~400 µL of the supernatant to a new tube or 96-well Plate and continue with step 3 of the NucleoMag DNA Food user manual.

For **automation on KingFisher**[®] **Flex** transfer ~400 μ L of the supernatant to a King-Fisher[®] Deep-well Block and follow the instructions of the Protocol Information Nucleo-Mag_DNA_Food_Flex

Additional consumables

Product	Content	REF
96-well Accessory Kit A for KingFisher®	for 4 x 96 samples including Deep-well Blocks, Elution Plate, Deep-well Tip Comb	744950
MN Bead Tube Type B	50x MN Bead Tube Type B containing 40–400 μm glass beads for homogenization	740812.50
MN Bead Plate Type B	Rack of prefilled tube strips (8 x 12) containing 40–400 µm glass beads for homogenization	740851.4



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