

## **DNA** isolation

Using ZymoBIOMICS™ 96 MagBead DNA Kit for isolating DNA from swabs

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	01 <b>Vortex</b> the collection tube with the swab for several seconds, follow by <b>quick spin.</b>
	02 Transfer <b>250 μL</b> of the supernatant from the collection tube to the ZR BashingBead™ Lysis Tubes.
	03 Add <b>750</b> µL of ZymoBIOMICS™ Lysis Solution the Lysis Tubes. <b>Vortex at maximum</b> speed for <b>10 minutes</b> at the <b>RT</b> .
	04 Centrifuge the tubes at <b>10 000 g for 1 minute</b> .
	05 Transfer <b>200 μL</b> of the supernatant into a new 1.5 mL tube and add <b>200 μL</b> of ZymoBIOMICS™ MagBinding Buffer.
	06 Dispense <b>25 μl</b> of ZymoBIOMICS™ MagBinding Beads to each well
	07 Transfer the tubes to a magnetic stand until the beads pellet, then aspirate and discard the supernatant and remove the tube from the stand.
	08 Dispense <b>500 μl</b> of ZymoBIOMICS™ MagBinding Buffer and mix well by pipette.
	09 Transfer the tubes to a magnetic stand until the beads pellet, then aspirate and discard the supernatant and remove the tube from the stand.
	10 Dispense <b>500 μl</b> of ZymoBIOMICS™ MagWash 1 and mix well by pipette.
	11 Transfer the tubes to a magnetic stand until the beads pellet, then aspirate, discard the supernatant, and remove the tube from the stand.
	12 Dispense <b>900 μl</b> of ZymoBIOMICS™ MagWash 2 and mix well by pipette.
	13 Transfer the tubes to a magnetic stand until the beads pellet, then aspirate, discard the supernatant, and remove the tube from the stand.
	14 Repeat the wash (Steps 12-13)
	15 Dispense <b>50 μl</b> of ZymoBIOMICS™ DNase/RNase Free Water and mix well by pipette.
	16 Transfer the tubes to a magnetic stand until the beads pellet.
	17 Transfer the supernatant (containing the eluted DNA) to a clean tube.