CONGEN

Flow Chart for SureFood® PREP Basic

Art. No. S1052

November 2017

Information

(1) Preparation of the basic material

(2) Lysis of basic material

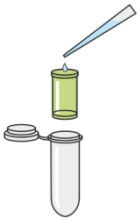
Add **400 µl Lysis Buffer** (Code L)
and **20 µl Proteinase K** (Code K)

Mixing

Incubation **30 min 65°C** by shaking

Centrifugation **1 min 12000 rpm**

(3) Pre - filtration and setting of optimal binding conditions



Place the **green Spin Filter** (Code F)
into a **clear Receiver Tube** (Code R)

Add liquid supernatant

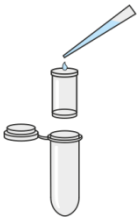
Centrifugation **1 min 12000 rpm**

Discard Spin Filter

(4) Binding of the nucleic acids

Add **200 µl Binding Buffer** (Code B)

Mixing



Place a **clear Spin Filter** (Code S)
into a new **clear Receiver Tube** (Code R)

Transfer the complete solution
onto the Spin Filter (Code S)

Incubation **1 min at room temperature**

Centrifugation **1 min 12000 rpm**

Discard the filtrate

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(5) **Purification of bound nucleic acids &**

(6) **Drying of the Spin Filter**

Add **550 µL Pre-Wash Buffer** (Code P)

Centrifugation **1 min 12000 rpm**

Discard filtrate
Place Spin Filter back into Receiver Tube (Code R)

Add **550 µL Wash Buffer** (Code W)

Centrifugation **1 min 12000 rpm**

Discard filtrate
Place Spin Filter back into Receiver Tube (Code R)

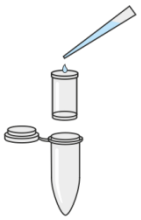
Add **550 µL Wash Buffer** (Code W)

Centrifugation **1 min 12000 rpm**

Discard filtrate
Place Spin Filter back into Receiver Tube (Code R)

Centrifugation **2 min 12000 rpm**

(7) **Elution of nucleic acids**



Place **Spin Filter** into a
clear 1.5 ml Receiver Tube (Code T)

Add **100 µL preheated Elution Buffer** (Code E)

Incubation **3 min 65°C**

Centrifugation **1 min 10000 rpm**

Discard Spin Filter

The eluted DNA is ready-to-use for the PCR