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**GRS Genomic DNA Kit – Blood & Cultured Cells –
#GK02.0100
(FOR RESEARCH ONLY)**

SUGGESTED PROTOCOL FOR DNA PURIFICATION FROM FUNGUS

- 1) Harvest up to 5×10^7 fungus cells by centrifugation for **10 minutes** at **5.000g**.
- 2) Discard the supernatant, and resuspend the pellet in **600µl** of **Sorbitol Buffer**¹
- 3) Add 200U of **Lyticase** or **Zymolyase**. Incubate for **30 minutes** at **30°C**.
- 4) Harvest spheroplast by centrifugation for 10 minutes at 2.000g. Remove the supernatant and resuspend pellet in **200µl** of **Buffer BC2**. Incubate at room temperature for 5 minutes. Proceed with the lysis step of the cultured cell protocol (page 6).

¹Sorbitol Buffer = 1,2M sorbitol; 10mM CaCl₂; 0.1M Tris-HCl pH 7,5; 35mM β-mercaptoethanol