

**Materials List for:****Toxin Induction and Protein Extraction from *Fusarium spp.* Cultures for Proteomic Studies**Matias Pasquali<sup>1</sup>, Frédéric Giraud<sup>1</sup>, Jean Paul Lasserre<sup>1</sup>, Sébastien Planchon<sup>1</sup>, Lucien Hoffmann<sup>1</sup>, Torsten Bohn<sup>1</sup>, Jenny Renaud<sup>1</sup><sup>1</sup>Department of Environment and Agro-Biotechnologies (EVA), Nutrition and Toxicology Unit (NuTox), Centre de Recherche Public-Gabriel LippmannCorrespondence to: Matias Pasquali at [pasquali@lippmann.lu](mailto:pasquali@lippmann.lu)URL: <https://www.jove.com/video/1690>DOI: [doi:10.3791/1690](https://doi.org/10.3791/1690)**Materials****Media and solutions**PDA: 39 g Potato Dextrose Agar (Difco); 1 L distilled water.V8 agar: 200 ml V8 (Campbell's, USA), 2g CaCO<sub>3</sub> (Sigma), 16g Agar (DIFCO), 800 ml distilled waterToxin inducing media: 1g K<sub>2</sub>HPO<sub>4</sub>, 0.5g KCl, 0.5g MgSO<sub>4</sub>·7H<sub>2</sub>O, 10 mg FeEDTA, 2g L glutamic acid, 10g sucrose in 1L of distilled water.Lysis Buffer:

	100mL	Final concentration	
Tris-HCl pH8	5mL	50mM	1M
SDS	2g or 10mL	2%	20%
DTT	500mL	10mM	2M
EDTA	20mL	0.1mM	0.5M
PMSF	200mL		
Complete mini protease inhibitor	1 tablet		
Water	Qsp		

Precipitation Buffer:

	100mL	Final concentration
TCA	20mL	20%
DTT	0.1mL	0.1%
Aceton	Qsp	

Washing Buffer:

	100mL	Final concentration
DTT	0.1mL	0.1%
Aceton	100mL	

Labelling Buffer (store at -18°C in small aliquots):

	1mL	Final concentration
Urea	140 mg	7M
Thiourea	50 mg	2M
CHAPS	40 mg	4%
Tris	30 µL	30 mM
Water	1 mL	