## **SUPPLEMENTARY MATERIALS**

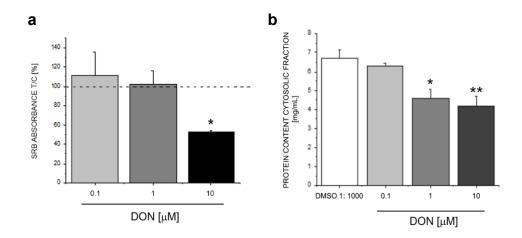
Deoxynivalenol induces structural alterations in epidermoid carcinoma cells A431 and impairs the response to biomechanical stimulation

Giorgia Del Favero<sup>1\*</sup>, Lydia Woelflingseder<sup>1</sup>, Lukas Janker<sup>2</sup>, Benjamin Neuditschko<sup>2</sup>, Stefano Seriani<sup>3,4</sup>, Paolo Gallina<sup>3</sup>, Orfeo Sbaizero<sup>3</sup>, Christopher Gerner<sup>2</sup>, Doris Marko<sup>1</sup>

- 1 Department of Food Chemistry and Toxicology, Faculty of Chemistry, University of Vienna. Währingerstr. 38-40, 1090 Vienna, Austria
- 2 2 Department of Analytical Chemistry, Faculty of Chemistry, University of Vienna. Währingerstr. 38-40, 1090 Vienna, Austria
- 3 Department of Engineering and Architecture, University of Trieste Via A. Valerio 10, 34127 Trieste, Italy
- 4 Robotik und Mechatronik Zentrum, Deutsches Zentrum für Luft- und Raumfahrt e.V. (DLR), Oberpfaffenhofen, Germany
- \* Corresponding author. Dr. Giorgia Del Favero (giorgia.del.favero@univie.ac.at)

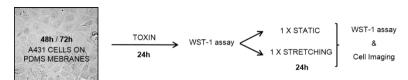
Department of Food Chemistry and Toxicology, Faculty of Chemistry, University of Vienna. Währingerstr. 38-40, 1090 Vienna, Austria.

Fig. S1



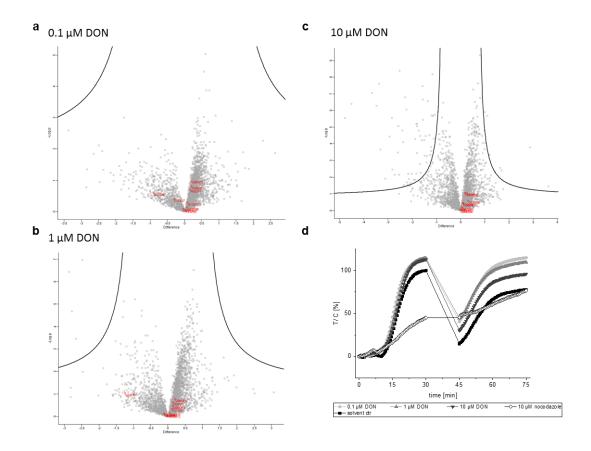
**Supplementary Materials Fig. S1.** Protein quantification in A431 cells after 24h incubation with DON. (a) SRB assay. Data are expressed as test / control [%] and depicted as mean  $\pm$  S.E. n=3-4 independent experiments performed in triplicate. (b) Bradford assay quantification of protein content in the cytosolic fraction. \*/\*\* indicates at 0.05 and 0.01 level different population means in comparison to incubation with 0.1  $\mu$ M DON (a) or solvent control (DMSO 1:1000; b) ANOVA.

Fig. S2



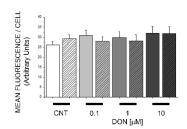
**Supplementary Materials Fig. S2.** Schematic representation of the experimental workflow of the results presented in Figures 4; 5; 6.

Fig. S3



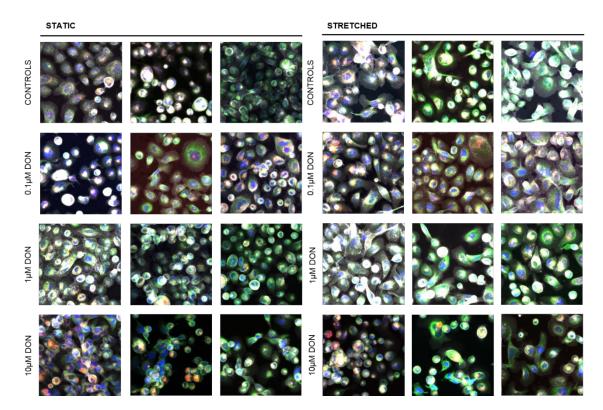
**Supplementary Materials Fig. S3.** Volcano plots describing DON-induced proteome alterations in A431 cells. In red tubulin isoforms (Q9BQE3; P68366; P07437; Q13885; Q13509; P68371; Q9BUF5; Q3ZCM7; P23258) expressed by A431 cells and relative abundance after incubation with DON 0.1 μM (a), 1 μM (b) and 10 μM (c). Effect of DON (0.1-10 μM) on tubulin polymerization (d) in comparison to solvent control (black) and tubulin polymerization inhibitor Nocodazole (10 μM, white). Tubulin polymerization assay was performed with a commercial kit (17-10194 *In vitro* Tubulin Polymerization Assay Kit ( $\geq$  99% Pure Bovine Tubulin) from Chemicon, Merck Millipore according to the specification of the supplier.

Fig. S4



**Supplementary Materials Fig. S4**. Mean fluorescence signal of cellular nuclei stained with Hoechst 33258. Solid bars refer to static incubation and striped bars refer to stretched incubation of control cells (white background) and pre-incubation with DON (grey background). Data are expressed as mean  $\pm$  S.E. and quantification is the result of n  $\geq$  45 cells randomly from 3 independent experiments.

Fig. S5



**Supplementary Materials Fig. S5.** Additional representative images acquired during the performance of the experiments. Static and Stretched membranes were always incubated in parallel and each column has been obtained with a different cell preparation.