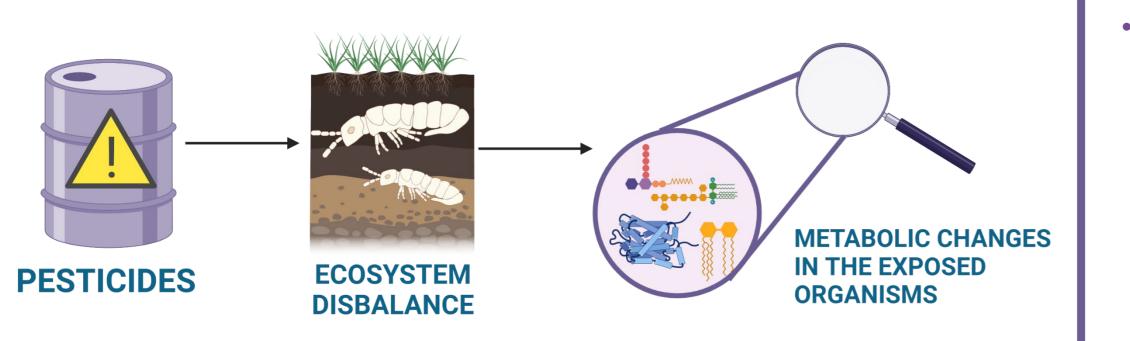
Low levels of Teflubenzuron disrupt lipid metabolism in springtails

Untargeted lipidomics to assess the response of the non-targeted species *Folsomia candida* to sub-lethal concentrations of Teflubenzuron

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INTRODUCTION

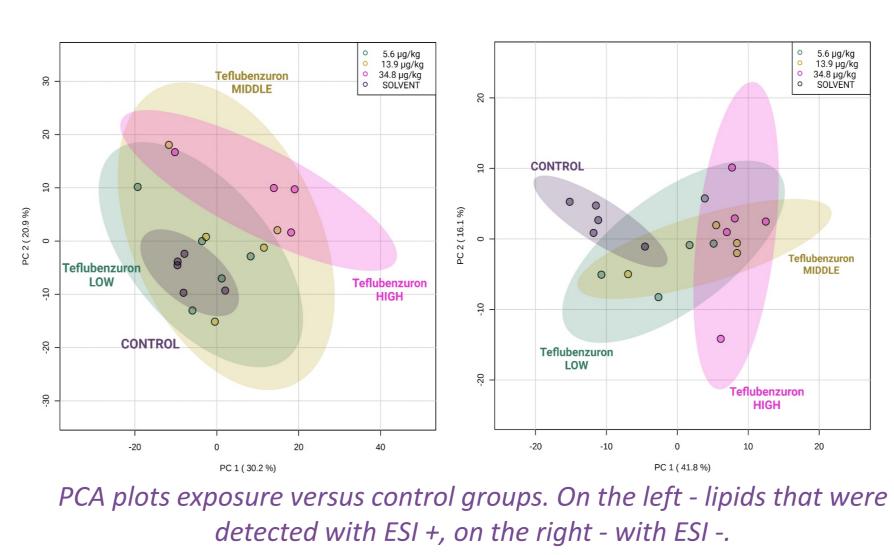


Untargeted screening of metabolic changes in the organism provides a more detailed picture of the sublethal effects of chemicals present in low concentrations in the environment.

The aim of the study was using untargeted lipidomics to investigate potential toxicity side-effects of the insecticide **Teflubenzuron on Folsomia candida exposed to** environmentally relevant concentrations.

RESULTS

- The major affected lipid classes due to exposure to TEF were:
 - Phospholipids (PC and PE) • Diacylglycerols (DG)
 - Triglycerols (TG) • Fatty acids (FA)

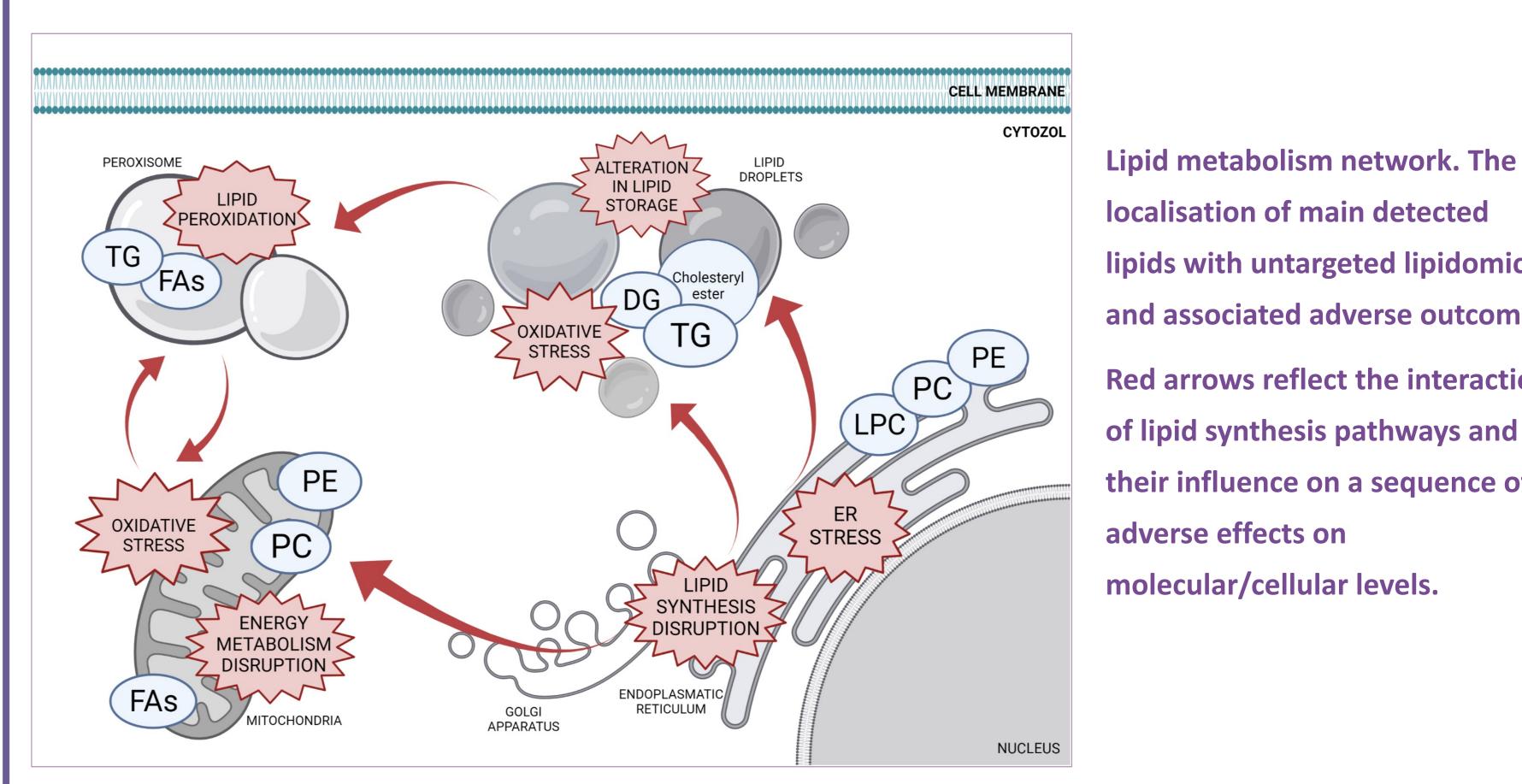


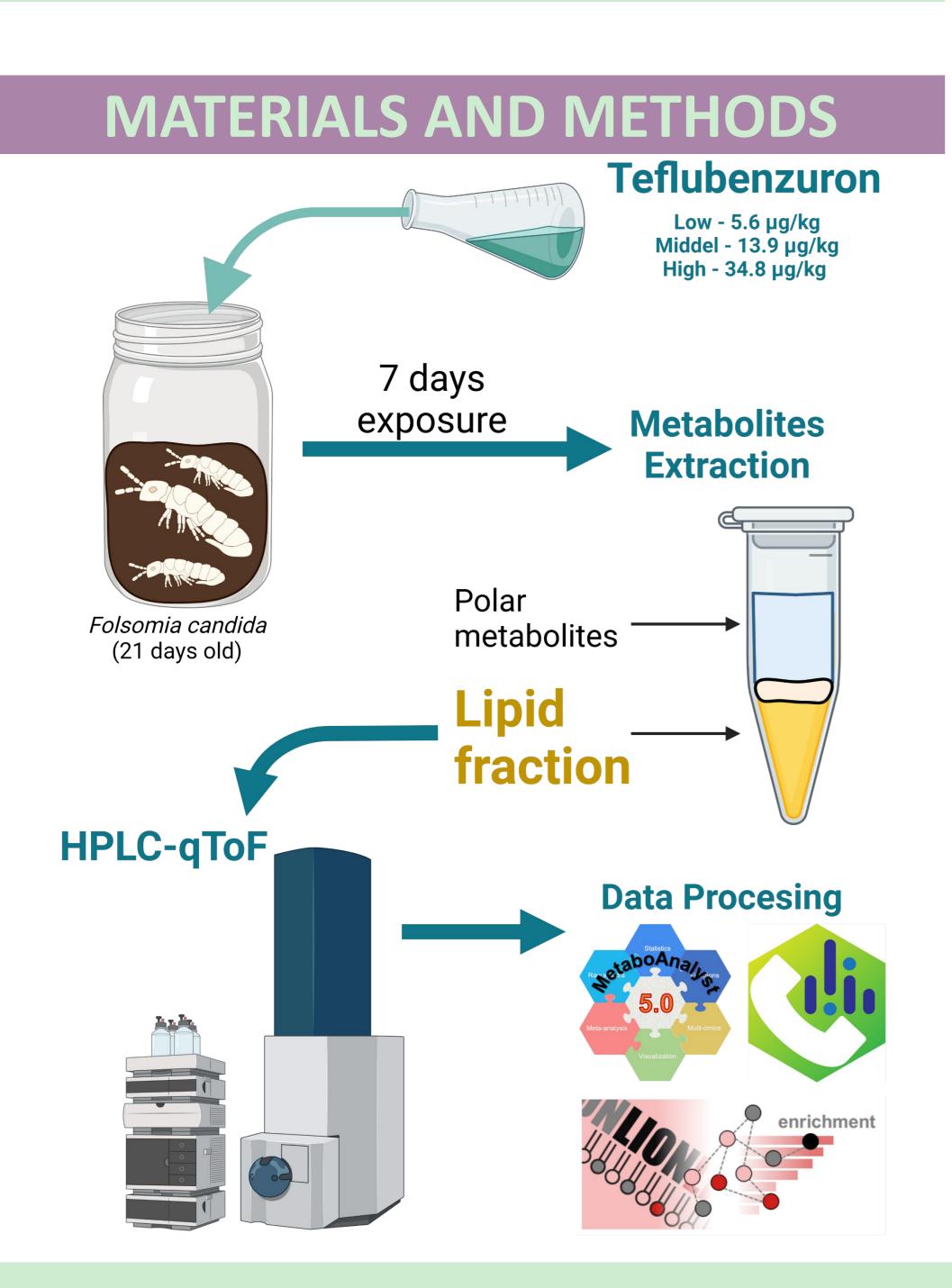
• PCA demonstrated the separation of control replicates with exposed groups.

• The decrease of TGs in the organisms can be connected to the usage of stored lipids as an additional energy source to cope with toxic effects.

ESI +				
Lipid classes	Annotated lipids	FDRq-value	ES	Regulation
Glycerolipids [TG,DG]	235	1.70E-30	-0.71	DOWN
Triacylglycerols[TG]	166	1.70E-30	-0.97	DOWN
Glycerophospholipids [PC,PE]	161	1.70E-30	0.642	UP
Steryl esters [CE]	3	5.87E-02	0.70	UP
Cellular compartment				
Lipid droplets [TG]	169	3.00E-30	-0.98	DOWN
Endoplasmatic reticulum (ER) [PC,PE] 161	3.00E-30	0.64	UP
Endosome/Lysosome [CE]	22	1.36E-09	0.68	UP
ESI -				
Lipid classes	Annotated lipids	FDRq-value	ES	Regulation
Glycerophospholipids [PC,PE,PG]	126	5.50E-30	-0.68	DOWN
Fatty acids [FA]	58	1.47E-15	0.71	UP
Cellular compartment				
Mitochondrion [PE]	66	2.80E-15	-0.62	DOWN
Endoplasmatic reticulum (ER) [PC,PE] 112	4.80E-14	-0.57	UP

Enrichment analysis of lipid classes and their regulation in cellular compartments, for all exposure concentrations versus control.





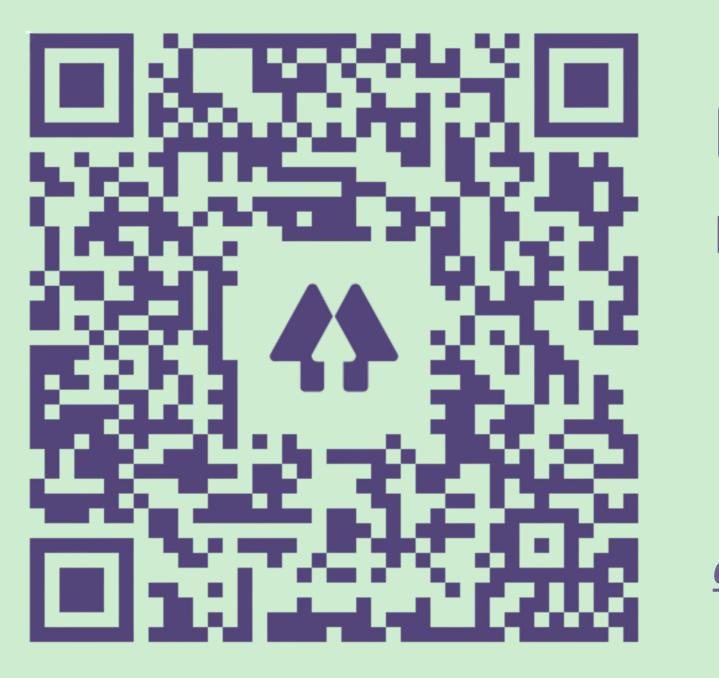
localisation of main detected lipids with untargeted lipidomics and associated adverse outcomes. **Red arrows reflect the interaction** of lipid synthesis pathways and their influence on a sequence of

adverse effects on molecular/cellular levels.

CONCLUSION

- > Teflubenzuron present in sub-lethal concentrations can cause significant effects on lipid metabolism.
 - > The main lipids dysregulated by Teflubenzuron were phospholipids, triglycerols, and fatty acids.
- > The dysregulation of the lipids can cause malfunctions in lipid membrane

synthesis, energy storage, oxidative stress (lipid peroxidation) and lipid metabolism.



Find out more about me and my project

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