Exposure to micro- and nanoplastics in Schmidtea mediterranea causes impaired (neuro)regeneration

Citation for published version (APA):

Tytgat, J., Leynen, N., Van Belleghem, F. G. A. J., Bijnens, K., Saenen, N., & Smeets, K. (2022). Exposure to micro- and nanoplastics in Schmidtea mediterranea causes impaired (neuro)regeneration. Poster session presented at 5th European Meeting on Planarian Biology, Sant Feliu de Guixols, Spain.

Document status and date: Published: 03/10/2022

Document Version:

Publisher's PDF, also known as Version of record

Please check the document version of this publication:

• A submitted manuscript is the version of the article upon submission and before peer-review. There can be important differences between the submitted version and the official published version of record. People interested in the research are advised to contact the author for the final version of the publication, or visit the DOI to the publisher's website.

• The final author version and the galley proof are versions of the publication after peer review.

• The final published version features the final layout of the paper including the volume, issue and page numbers.

Link to publication

General rights

Copyright and moral rights for the publications made accessible in the public portal are retained by the authors and/or other copyright owners and it is a condition of accessing publications that users recognise and abide by the legal requirements associated with these rights.

• Users may download and print one copy of any publication from the public portal for the purpose of private study or research.

- You may not further distribute the material or use it for any profit-making activity or commercial gain
 You may freely distribute the URL identifying the publication in the public portal.

If the publication is distributed under the terms of Article 25fa of the Dutch Copyright Act, indicated by the "Taverne" license above, please follow below link for the End User Agreement:

https://www.ou.nl/taverne-agreement

Take down policy

If you believe that this document breaches copyright please contact us at:

pure-support@ou.nl

providing details and we will investigate your claim.

Downloaded from https://research.ou.nl/ on date: 19 Nov. 2022



Exposure to micro- and nanoplastics in *Schmidtea mediterranea* causes

impaired (neuro)regeneration

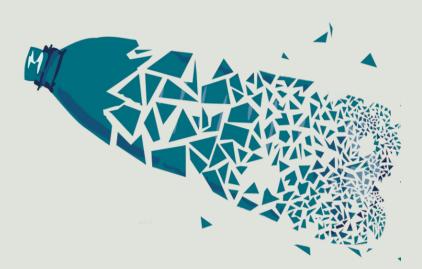
Tytgat Julie¹; Leynen Nathalie¹; Van Belleghem Frank^{1, 2}; Saenen Nelly¹; Bijnens Karolien¹; Smeets Karen¹

1: Centre for Environmental Sciences, Zoology, Biodiversity and Toxicology; Hasselt University; Hasselt; Belgium.

2: Department of Environmental Sciences; Faculty of Science; Open Universiteit; Heerlen; the Netherlands.

Background

- Due to the lack of proper waste management and disposal, plastic degrades to micro- and nanoplastics (MNPs) and ends up in the aquatic environment.
- MNPs are plastic fragments smaller than **5 mm** and **100 nm** respectively.
- Studies showed that MNPs are potentially harmful but detailed knowledge on developing organisms and the link between the physicochemical properties is missing, which is needed for sufficient risk assessment strategies.
- We focused on particle characterization since knowledge of the physicochemical characteristics is necessary to gain insights into the mechanisms of particle toxicity
- Schmidtea mediterranea is used as a proxy to study developmental toxicity and is exposed to carboxylated polystyrene MNPs of 49 nm (PS49), 215 nm (PS215), 1 μm (PS01) and 2 μm (PS02) in diameter. This creates an *in vivo* model system in line with the 3R guidelines to study particle toxicity.



Aim

To link physicochemical properties with induced effects to properly define MNP-specific adverse outcome pathways in the function of **new risk assessment strategies** by focusing on developing tissues in the planarian model system.

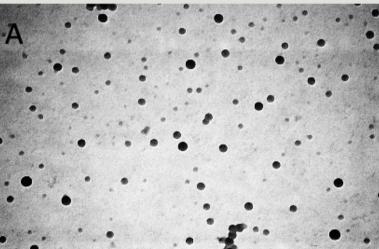


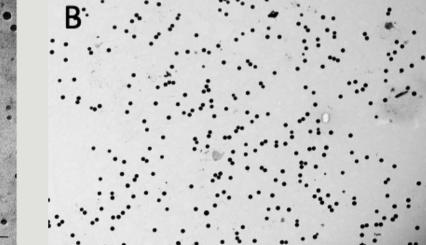
Impaired (neuro)regeneration

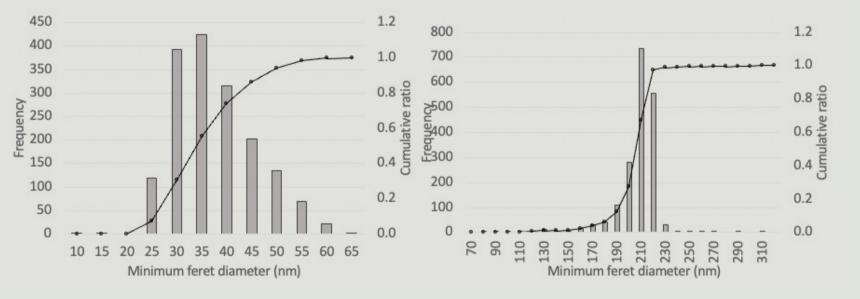
Delayed eye development

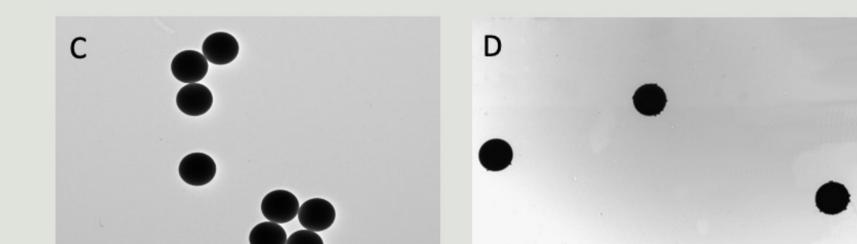
Eye development after exposure to MNPs. Depiction of the eye development in tail fragments. Eye development is delayed after MNPs and is exposure to concentration size and dependent. For PS01 no effects At least 7 were observed. replicates were used per condition and experiments were repeated. 7 dpa.

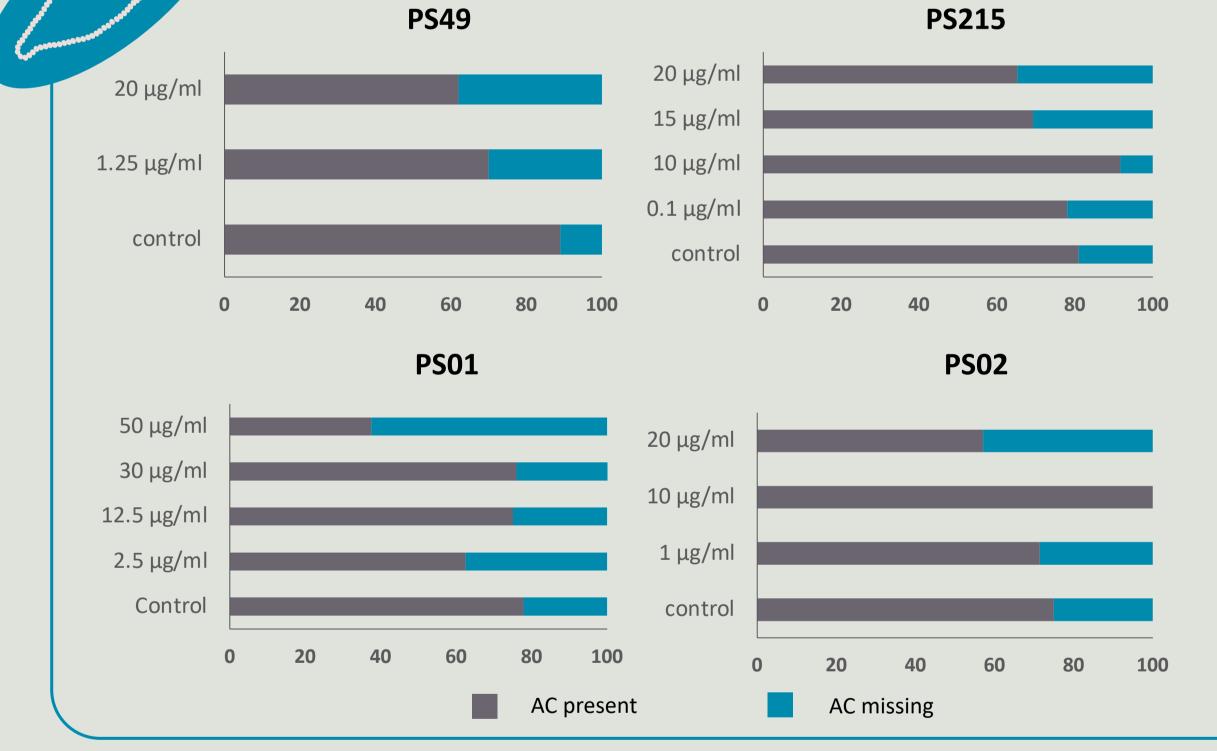
Particle characterization







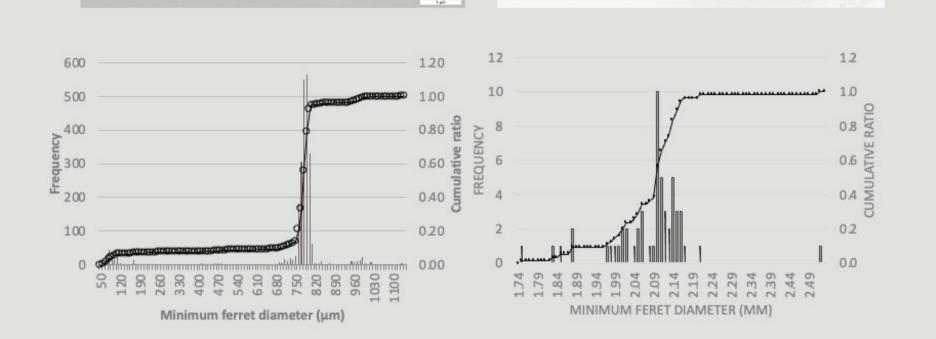




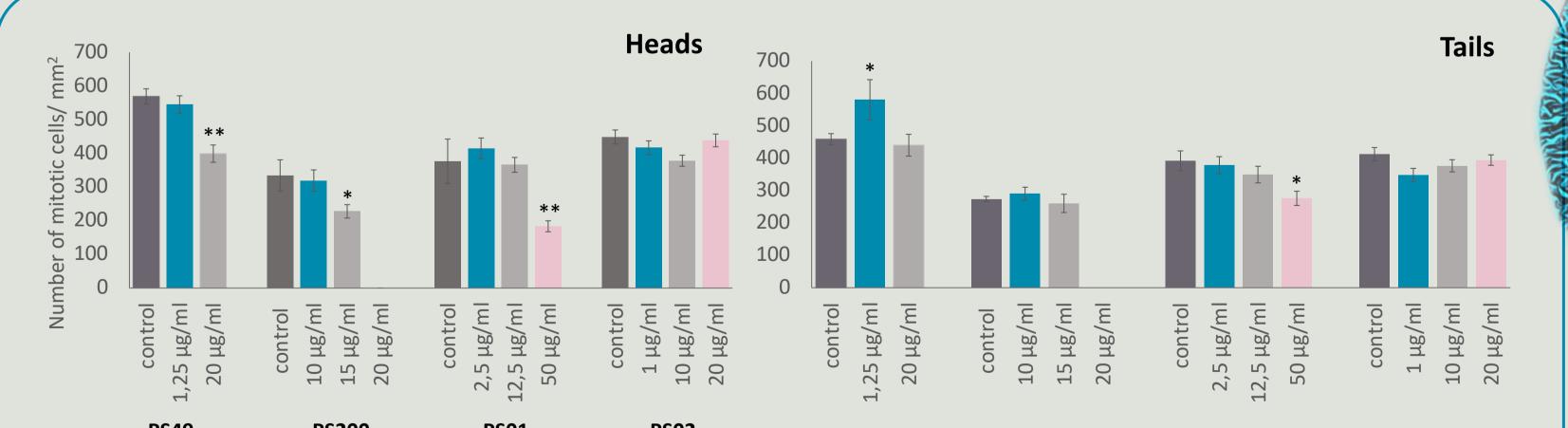
Delayed formation of the anterior commissure

The formation of the anterior commissure (AC) after exposure to MNPs. Depiction of the AC formation in tail fragments visualized by Synorf staining. The formation of the AC is delayed in all particles and is concentration-dependent. At least 7 replicates were used per condition. 7 dpa.

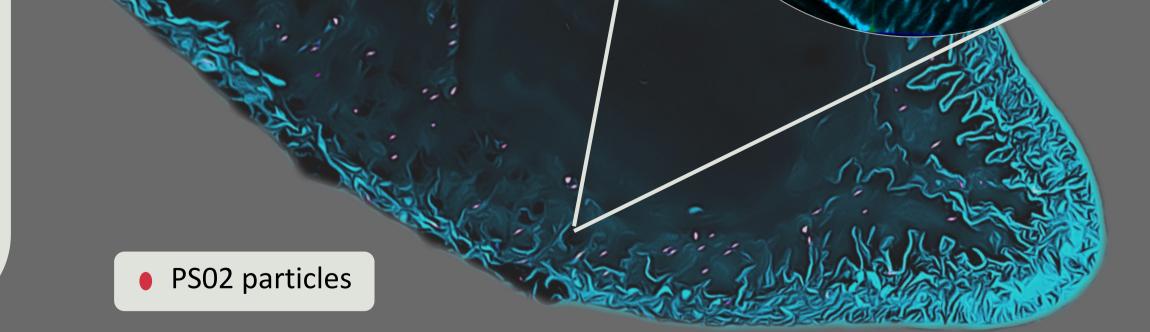
PS02



Size distribution of polystyrene MNPs. TEM Images of PS49 (A), PS215 (B), PS01 (C) and PS02 (D) were recorded with TEM with the corresponding size distributions assessed by the NanoDefine ParticleSizer software plugin in ImageJ.



PS49 PS200 PS01 Number of mitotic stem cells after exposure to MNPs. Stem cells were visualized with **Affected mitotic** H3P staining. The number of mitotic cells is affected after exposure to MNPs and is concentration and size dependent. Most effects are seen in head fragments, while tale division fragments are less affected. At least 7 replicates were used per condition and experiments were repeated. 7 dpa. * p<0.05; ** p<0.01; *** p<0.001



Conclusions and future perspectives

- All particles caused impaired (neuro) regeneration and delayed eye and anterior commissure development in a **concentration and size-dependent** way.
- An affected stem cell division underlies these effects
- Because previous studies showed that nanoparticles are located nearby mitochondria, I
- hypothesize that disturbed mitochondrial redox dynamics can explain the toxic effects.
 - By linking particle property changes within cells with the mechanistic effects, more complete risk assessment strategies can be established in the future.

