1	A	dverse Effects of Reproductive Toxicity and Metabolism in Mice by
2		Polystyrene Microplastics
3		許瑜真(5123)
4		2023/3/22
5		Outline
6	1.	Introduction
7	2.	Reproductive toxicity of polystyrene microplastics: In vivo experimental study on
8		testicular toxicity in mice
9	3.	Adverse effects of pristine and aged polystyrene microplastics in mice and their
10		Nrf2-mediated defense mechanisms with tissue specificity
11	4.	Conclusion
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13		Abstract
14		Microplastics (MPs) are plastic particles with a size of less than 5 mm and are

15 derived from any kind of plastic. Humans may be exposed to MPs by ingesting 16 contaminated food and water, and MPs may infiltrate and accumulate in human tissues, 17 posing a health risk. However, research on the toxic effects of MPs on the reproductive system on mammals is relatively limited as well as health hazards associated with pristine 18 19 and ultraviolet-aged MPs remain largely unknown. Therefore, this study evaluated the 20 reproductive toxicity and changes in toxicity of pristine and aged MPs to different organs 21 in mice. Both MPs and aged MPs caused structural damage to the mouse gut, liver, spleen, 22 and testis. MPs may cause damage to mouse sperm production through testicular injury, 23 which in turn leads to a decrease in sperm quality. Increases in serum alanine 24 aminotransferase and immunoglobulin A levels in 1 mg/day aged MPs group 25 demonstrated that aged MPs exposure could induce greater liver and spleen functional 26 damage than PS. Nuclear factor erythroid 2-related factor 2 (Nrf2) and heme oxygenase-27 1 (HO-1) levels in the MPs and aged MPs groups showed significant increases in the 28 liver and testis, and a significant decrease in the spleen, which were analyzed to get a 29 first survey for Nrf2/HO-1-mediated tissue-specific defense mechanisms. In conclusion, 30 exposure to MP and aged MPs induced potential metabolic disorders, and aged MPs 31 could produce more serious immune damage and reproductive toxicity. 32

1	Reference
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