

Brussels, 11 February 2020

Fraud in German laboratory casts additional doubts on the 2017 re-approval of glyphosate and on the entire EU pesticide safety evaluation procedure

The Laboratory of Pharmacology and Toxicology (LPT) Hamburg that was found recently to commit fraud in a series of regulatory tests had also carried out many of the tests in the glyphosate re-approval dossier in 2017, new study reveals. At least one in 7 glyphosate regulatory studies, with the certificate "Good Laboratory Practice" (GLP), come from LPT Hamburg, the same laboratory that was caught manipulating GLP toxicity studies by replacing dead animals will living ones, changing tumour data to "inflammations" and generally distorting the data to please its clients. It is highly concerning that GLP studies are still considered the golden scientific standard by regulatory authorities who seem to believe that cheating under GLP is impossible.

PAN Europe asks to the European Commission to discard the studies carried out by LPT laboratory from the glyphosate dossier currently undergoing a re-evaluation at EU-level, and from any other dossier.

Based on testimonies from LPT's employees and evidence of fraud carried out in LPT Hamburg, a major GLP laboratory in Germany, the survey¹ carried out by the organisations PAN Germany, Global2000 and Corporate Europe Observatory reveals that at least 14% of the new regulatory studies submitted for the re-approval of glyphosate in 2017 were conducted by LPT Hamburg. The number could be higher, as this information in the dossiers often remains undisclosed to the public. The laboratory is currently facing criminal charges, and although it is impossible to know whether the fraud occurred only in the glyphosate-related studies, any tests delivered by LPT Hamburg must be considered unreliable and thus discarded from the re-assessment procedure.

Good Laboratory Practice is a mandatory standard in regulatory studies obliging laboratories to write down a series of endpoints in a specific format. This system allows for a higher level of standardisation of the reporting and easier control by regulatory authorities.

Angeliki Lyssimachou, environmental toxicologist at PAN Europe, said: "The vast majority of studies leading to the approval of a pesticide are carried out by the pesticide industry itself, either directly or via contract laboratories such as LPT Hamburg. We have criticized this conflict of interest for many years. Our 140+ NGO coalition "Citizens for Science in Pesticide Regulation" regularly calls on the Commission to quit this scandalous process: tests must be carried out by independent laboratories under public scrutiny, while the financing of studies should be supported by industry".

Hans Muilerman, chemical policy officer at PAN Europe, added: "For years, European Member States, EFSA and the Commission have been defending the belief that an industry-funded GLP study is more reliable than a non-industry, non-GLP study. Thousands of relevant independent pesticide studies showing harm to humans

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 $^{^1\} https://www.global2000.at/sites/global/files/2020-GoodLaboratoryPractice-en.pdf$

² https://citizens4pesticidereform.eu/

or nature have been discarded by regulators because they are not GLP, under the Klimisch scoring principle³".

"By including non-GLP studies, the International Agency for Research on Cancer (IARC) has classified glyphosate as a probable carcinogen. It is by giving so little weight to all non-GLP studies that the European Commission and Member States came to a different conclusion. It is time for citizens' health to take precedence over companies' profit!" Lyssimachou concluded.

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³ <u>Klimisch score</u> is a creation of BASF employees, a company producing pesticides. It is supposed to assess the reliability of ecotoxicology studies and uses GLP certification as criteria for discarding non-industry studies. Indeed, the vast majority of university studies are non-GLP as the certification is very expensive, little flexible and not suited for academic research.