

## ADSORPTION OF CRYSTAL VIOLET FROM SOLUTIONS

**Diachenko Anastasiia**

10th grade student of the Scientific Lyceum

**Kaminskyi Oleksandr**

Ph.D., Associate Professor

**Titov Yurii**

Doctor of Chemical Sciences, Professor

**Panasiuk Dmytro**

Assistant

Department of Chemistry

Zhytomyr Ivan Franko State University

Under conditions of ongoing military actions, the issue of drinking water quality in Ukraine has intensified and reached a global scale, remaining highly relevant. It should also be noted that water in the countries of the European Union is officially recognized as the primary food product [1]. Moreover, one of the key objectives of sustainable development (Sustainable Development Goal 6) to be achieved by 2030 is to ensure the availability and sustainable management of water supply and sanitation for all [2].

Among the disadvantages of using crystal violet are its carcinogenic properties and the difficulty of controlling its presence in the environment. These issues are associated with its high resistance to microbial degradation and its ability to persist in various environmental media. In addition, crystal violet may cause moderate eye irritation and can even lead to prolonged damage to the cornea and conjunctiva. Inhalation of this dye may result in irritation of the respiratory tract as well as symptoms such as nausea, diarrhea, headache, dizziness, and pain. Prolonged exposure to crystal violet can also negatively affect mucous membranes and the organs of the gastrointestinal tract [3].

In this study, zinc ferrite was synthesized using a sol–gel method combined with an autocombustion process, and the adsorption of crystal violet dye from aqueous solutions onto the surface of  $\text{ZnFe}_2\text{O}_4$  was investigated.

Figure 1 illustrates the adsorption isotherm of the dye on the  $\text{ZnFe}_2\text{O}_4$  surface. The concentration of crystal violet solutions before and after adsorption was determined photocolometrically at a wavelength of 540 nm.

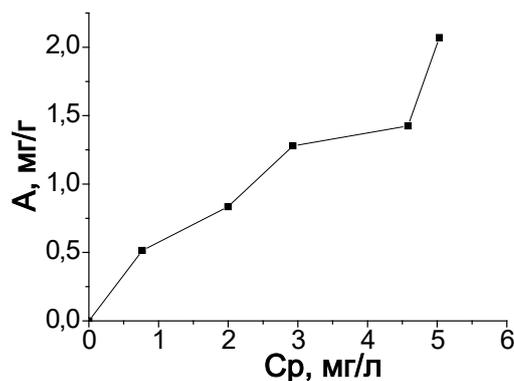


Fig. 1. Adsorption isotherm of crystal violet on the  $ZnFe_2O_4$  surface

It was established that the adsorption capacity ( $A$ ) is 2.07 mg/g, while the removal efficiency ( $R$ , %) reaches 49.69 % at an initial dye concentration ( $C_0$ ) of 10 mg/L. The shape of the isotherm curve resembles Langmuir-type isotherms (L3-type) according to the Giles classification [4]. This type of isotherm indicates weak interactions at the adsorbate–adsorbent interface, which is characteristic of nonporous adsorbents with relatively homogeneous surfaces.

It was also determined that the adsorption isotherm of crystal violet is better described by the Freundlich model than by the Langmuir model, as evidenced by the correlation coefficient ( $R^2 = 0.922$ ). This suggests that the adsorption of the dye occurs on heterogeneous surface sites with a non-uniform distribution of adsorption energies.

### References

1. European Parliament and Council of the European Union. (2020). Directive (EU) 2020/2184 of the European Parliament and of the Council of 16 December 2020 on the quality of water intended for human consumption. <https://www.legislation.gov.uk/eudr/2020/2184>
2. Ahmad, N., Kameda, T., Rahman, M. T., Rahman, F., & Lesbani, A. (2025). Preparation of a new hybrid  $MgAl-LDH@magnetite$  activated charcoal by hydrothermal method for stability and adsorption mechanism of Congo red. *Results in Surfaces and Interfaces*, 18, 100440. <https://doi.org/10.1016/j.rsurfi.2025.100440>
3. Sadoq, M., Atlas, H., Imame, S., et al. (2024). Elimination of crystal violet from aqueous solution by adsorption on natural polysaccharide: Kinetic, isotherm, thermodynamic studies and mechanism analysis. *Arabian Journal of Chemistry*, 17(1), 105453. <https://doi.org/10.1016/j.arabjc.2023.105453>
4. Giles, C. H., MacEwan, T. H., Nakhwa, S. N., & Smith, D. (1960). Studies in adsorption: Part XI. A system of classification of solution adsorption isotherms and its use in diagnosis of adsorption mechanisms and in measurement of specific surface area of solids. *Journal of the Chemical Society*, 3973-3993. <https://doi.org/10.1039/jr9600003973>