Privacy protection in the reporting of adverse events

Please be informed that the administrator of your personal data provided in the application form is MagnaPharm Poland Sp. z o.o. with its registered office in Warsaw at Al. Jerozolimskie 146D, 02-305 Warsaw.

Your personal data will be processed on the basis of art. 6 par. 1 point c) and art. 9 par. 2 point i) of the General Data Protection Regulation (GDPR, EU 2016/679 dated 27 April 2016) solely for the purpose of implementing obligations in the field of monitoring the safety of medicinal products in accordance with Regulation (EU) No. 520/2012 of 19 June 2012.

Providing the reporter's data is voluntary but is a prerequisite for the acceptance of the application. Providing personal data by persons obliged to submit adverse reactions of medicinal products is mandatory and results from art. 36e of the Act of September 6, 2001, Pharmaceutical Law (Journal of Laws of 2008 No. 45, item 271, as amended).

The recipients of your personal data will be only entities authorized to obtain personal data on the basis of the law.

You have the right to access your personal data and the right to correction, delete, limit processing, the right to transfer data and the right to object to the processing after the expiration of their storage period, as well as the right to lodge a complaint with the personal data protection supervisory authority if you consider that the processing of your personal data violates the provisions of the GDPR.

Your personal data will be kept for the entire duration of the Marketing Authorization and for a period of 10 years after the Marketing Authorization expires.

For further information concerning data processing please contact us via e-mail: dane.osobowe@magnapharm.com or daneosobowe@magnapharm.eu or at the postal address of the administrator's registered office.