



## Safety Reporting Privacy Notice

Date: 1 February 2019

Medicines regulations require **Biogen International GmbH** ("**Biogen**") to take detailed records of every side effect, also known as an "**adverse event**" (meaning an unwanted, untoward, unintended or harmful event associated with the use of a Biogen medicinal product) reported to Biogen, in order to monitor product safety appropriately. This Privacy Notice describes how Biogen collects and processes your "**personal data**" (meaning any information relating to an identified or identifiable natural person) to help us fulfil our responsibility to monitor the safety of all medicines that we market or study in clinical trials and to comply with our legal obligations (also known as our safety reporting or pharmacovigilance obligations).

### Information we collect and how we use it

#### (a) Patients (subject of the report)

We collect personal data about a patient in relation to a reported adverse event associated with that patient. We may receive the data from the patient directly or from a third party reporting the adverse event. Reporters may include health care professionals, relatives or other members of the public. The personal data that we collect may be limited by national laws, but generally includes:

- initials;
- gender;
- weight and height;
- age or age category/group (such as adult, elderly or child) and date/year of birth;
- details of the product associated with the adverse event, including the dosage taken or prescribed, the reason for being prescribed the product and any subsequent change to the patient's usual regimen;
- details of other medicines or remedies the patient is taking or was taking at the time of the adverse event, including the dosage, the period of time the patient was taking the other medicine, the reason the patient was taking that other medicine and any subsequent change to the patient's regimen;
- details of the adverse event experienced such as the outcome, causality, etiology and diagnosis, the treatment the patient received for that adverse event, and any long-term effects the event has had on the patient's health;
- other medical information considered relevant, including risk factors and pregnancy; and
- other medical history or familial history considered relevant, including documents such as laboratory reports, medication histories and patient histories.

#### (b) Reporters

Biogen is required by law to make follow-up enquiries on reported adverse events. We must therefore keep sufficient information about reporters to allow us to contact them for additional information once we have received the adverse event report. The personal data that we may collect about reporters includes:

- name;
- contact details (address, e-mail address, phone number or fax number);
- profession/specialism (this information may determine the questions a reporter is asked about an adverse event, depending on his/her assumed level of medical knowledge); and
- relationship with the subject of the report.



Where the reporter is also the patient who is the subject of a report, this information may be combined with the information the patient provides in relation to the adverse event.

As part of our safety reporting obligations, we may use patient and reporter personal data to:

- investigate the adverse event;
- contact reporters for further information about the adverse event reported; and
- collate information about the adverse event with information about other adverse events received by Biogen to support safety monitoring of the product; and
- report to applicable competent regulatory authorities.

### **How we share personal data with others and international transfers**

As our safety reporting obligations require us to review patterns across reports received from every country where we market our products, the analysis is performed by an international group of highly-qualified safety physicians. Information provided as part of an adverse event report is shared within Biogen on a worldwide basis through Biogen's Global Safety Database. This database is hosted at Biogen's headquarters in the United States (Biogen Inc). Biogen's international headquarters, Biogen International GmbH in Switzerland, may also require access. Biogen also engages service providers to assist it in the administration of its safety reporting activities (such as IT service providers). Biogen is also obliged to transfer adverse event data to national regulatory authorities for their databases and to the European Medicine Agency's EudraVigilance database.

Such transfers may include transfers outside of your country to countries which do not implement an adequate level of protection for your personal data under your national law or European Union data protection law. Personal data collected for safety reporting may be transferred to a third party in the event that one of our products is sold, assigned or transferred. We may also share personal data with other pharmaceutical companies who are our co-marketing, co-distribution or other licence partners, where safety reporting obligations for a product require such exchange of safety information. Biogen takes appropriate steps to ensure personal data is adequately protected if transferred across national boundaries. Switzerland is a country deemed to provide an adequate level of data protection under its data protection laws by the European Commission. Biogen otherwise has EU-approved Standard Contractual Clauses in place where necessary to provide an adequate level of data protection. Upon your request, Biogen will provide you with further information on recipients of your personal data and any data transfer agreements with recipients outside the European Economic Area.

### **How we store personal data**

Because patient safety is paramount, we retain all the information we collect a result of an adverse event report indefinitely to ensure that we can continuously monitor the safety of our products over time.

### **Your rights**

You may contact Biogen at any time if you would like to access your personal data or require information about the personal data that we hold about you. You may object to the processing of your personal data for legitimate reasons, request restriction of the processing of it and you may also request the correction or erasure of it. Please note that some of these rights are limited by applicable data protection law and we have the right to collect, process and store personal data to perform our legal obligations under safety reporting laws.



### **Contact information**

Under European data protection laws, a "data controller" is the legal entity that is responsible for protecting your personal data and helping you to exercise your data protection rights. Biogen is the data controller of your personal data. If, at any time, you have questions or concerns about this Privacy Notice or the processing of your personal data, or would like to exercise your rights as outlined above, you can contact Biogen's EU Data Protection Officer at [privacy@biogen.com](mailto:privacy@biogen.com). You may, should you feel it necessary, lodge a complaint with your local data protection authority if you feel your privacy rights have been infringed.