Lionbridge LIFE SCIENCES

Expert Language Translation. Global Success.

Adverse Events Translation

How to Ensure a Successful Strategy

Thomson Reuters estimates that more than 60% of today's clinical studies are conducted in multiple geographies and that number is increasing.

Lionbridge Life Sciences has an established track record of reducing translation turnaround times for clients by more than 30%.

Adverse event reporting is tedious and laborious, particularly when you factor in translation, but help is available.

GIVE US A TRY

To learn more about our adverse event translation program, contact us at lionbridge.com/moreinfo and check out our Life Sciences blog. blog.lionbridge.com/life-sciences/

HOW WELL ARE YOU RESPONDING TO ADVERSE EVENTS ?

When pharmaceutical companies launch clinical trials in markets around the world, translation is a key requirement. Although translation currently represents less than .5% of overall spend in a trial, it is absolutely on the critical path to clean data and the trial's success.

Given that adverse events can derail a trial if not properly managed, and in-market regulatory bodies require timely reporting of all adverse events, pharmaceutical companies need an established translation strategy to communicate negative episodes in a timely manner, in the correct number of languages.

SPEED IS CRITCAL

Drug companies typically choose one of two methods for adverse event translation: do-it-yourself or outsourcing. Companies that translate the reports themselves often experience long delays because they don't have translators dedicated to the effort; and the employees who do the translating have other job responsibilities. A long translation process postpones the reporting and the release of the adverse event to the regulatory body (typically required within 48 hours).

Companies choosing to outsource to a third-party are usually hoping to avoid burdening their team with translation work, but still need to translate and report the event as quickly as possible. Unfortunately, not all outsourcers are equipped to support the time-sensitive requirements of adverse event translation.

YOU NEED A PROVEN GLOBAL PARTNER

Lionbridge Life Sciences is a leading provider of expert language translation for clinical trial success. Our reputable and tested operations are particularly well-suited to time-sensitive, multilingual adverse event reporting.

- Our **global footprint** allows us to take advantage of different time zones so translation work is ongoing and efficient.
- Our operation is **highly scalable** so we can handle a variable number of AEs; whatever your needs, we can ramp up or down to accommodate them.
- We specialize in all aspects of **clinical trial translation**, covering everything from pre-clinical and clinical to post-approval pharmacovigilance. Wherever you are in the process, we have a solution to help your trial succeed globally.