SEARLE

PHARMACOVIGILANCE

Searle Pharmacovigilance Philosophy

We at Searle believe that Patient safety is one of our core business principles. The aim of Pharma-covigilance is to promote public health by identifying, evaluating and minimizing safety issues in all possible ways. All of our medicines undergo thorough safety monitoring and evaluation processes at every stage of a medicine's lifecycle.

Searle encourages consumers & healthcare professionals to report any discomfort or adverse event experienced following consumption or administration of any of the Searle's product.

SEARLE Pharmacovigilance

Let's Make Medication Use Safer Together

PHARMACOVIGILANCE

ADVERSE EVENT (AE)
An untoward medical occurrence in a patient who has been or is being administered a medicinal product/medical device and the occurrence does not necessarily have a causal relationship with this treatment.

ADVERSE DRUG REACTION (ADR)
A response to a drug which is noxious and unintended, and which occurs at doses normally used in man for the prophylaxis, diagnosis, or therapy of disease, or for the modifications of physiological function.

Observe/Experience Adverse Event

- Doctor
- Pharmacist
- Nurse

Report to

SEARLE Pharmacovigilance

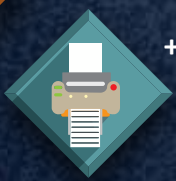
Reporting Helps

- ***Make*** medicine safer
- ***Prevent*** future harm to patients
- ***Safer*** prescribing and use of medicines
- ***Improve patient*** safety and protect public health
- ***Improve medicines*** information and education

“Safety First” is “Safety Always”

All AEs should be reported IMMEDIATELY or within One Calender Day to Searle Pharmacovigilance via following details

+ 92 21 371 70 200 - 1



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Pharmacovigilance unit:
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For more details, please refer to the Searle Website:
www.searlecompany.com