



What is it?

It is a standardized electronic format for reporting suspected **adverse drug reactions** (discomfort caused by medications) developed to facilitate exclusive reporting by **patients, consumers and health professionals**.

Contains the variables that comply with national and international regulations. Notifications are transmitted directly and immediately to the corresponding State Pharmacovigilance Centre (CEFV) and to the National Pharmacovigilance Centre (CNFV).

VigiRam Key Features :

- Electronic notifications save the notifier time and facilitate the capture data tower
- Available via QR code on any browser, mobile devices and tablets
- Installs as an application on the device
- Does not require user registration
- Allows data capture without an internet connection
- Requires very little data for notification transfer
- All information sent over the Internet is encrypted keeping
Your data is protected
- Features drop-down menus for easy and standardized data entry.
- The user has full control over the shipping process
- The notifier's data is saved, so it is not necessary to enter it again.
- It is possible to save copies of the notifications sent on the device

VigiRam should not be used by the pharmaceutical industry, health research establishments or distributors to comply with the notification or otherwise establish an official provision. , unless the CNFV