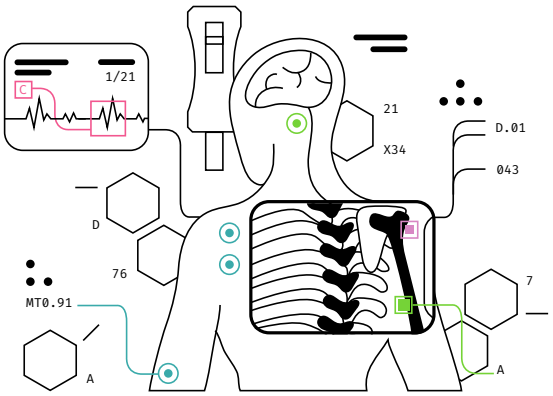


Your Medical Device Data

EirMed (@ Eudamed.com)

Under the European medical device regulations, all medical devices placed on sale inside the European Union must be certified and medical device data must be added to EUDAMED, the European Medical Device Database.

We have two solutions to help you to publish your UDI data in EUDAMED. We manage the complexity for you. To find out more visit **Eudamed.com**

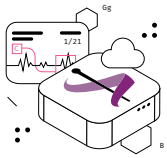
EudaMed SaaS — In the cloud

OR

EudaMed+ — Installed on your servers

Adding your data to EUDAMED is a multi-step process and we have automated the process as much as possible for you.

1

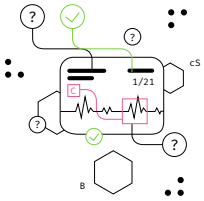


DATA ENTRY

EudaMed SaaS is populated directly from the EudaMed.com spreadsheet templates or from your data filled JSON files.

EudaMed+ can be populated directly from spreadsheet templates and/or web-based forms which include many functionalities and data management tools to help users.

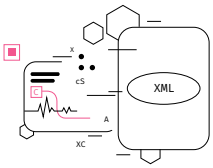
2



DATA VALIDATION

Both **EudaMed+** and **EudaMed SaaS** validate your data against 100s of data a nd complex business rules. You receive PDF files of your device data highlighting errors and any failed business rules which allow you to easily validate your data.

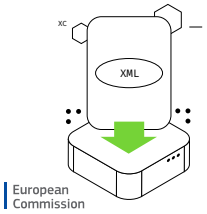
3



DATA CONVERSION

After error-free data validation, your data is converted to the EUDAMED required XML formats. Includes full batch management.

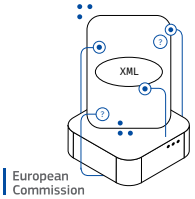
4



DATA SUBMISSON

When the XML files are ready EudaMed+ and EudaMed SaaS submit your data to EUDAMED.

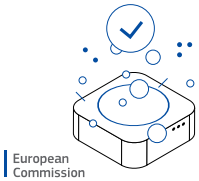
5



EUDAMED RESPONSE

EUDAMED will return messages of success or failure.

6



SUCCESS

Your data is added to EUDAMED, you can now sell products in the EU.

*If unsuccessful you fix the data and we send it again for you

EudaMed+ and **EudaMed SaaS** are always up to date.

If the European Commission changes the data requirements **EirMed** will rapidly implement the changes.

Using our software services you have...

- No in-house development costs
- No access point costs or worries
- No upgrade worries