

Using the dotLab System in a 21 CFR Part 11 Compliant Environment

- For users who are working in a FDA Regulated Environment



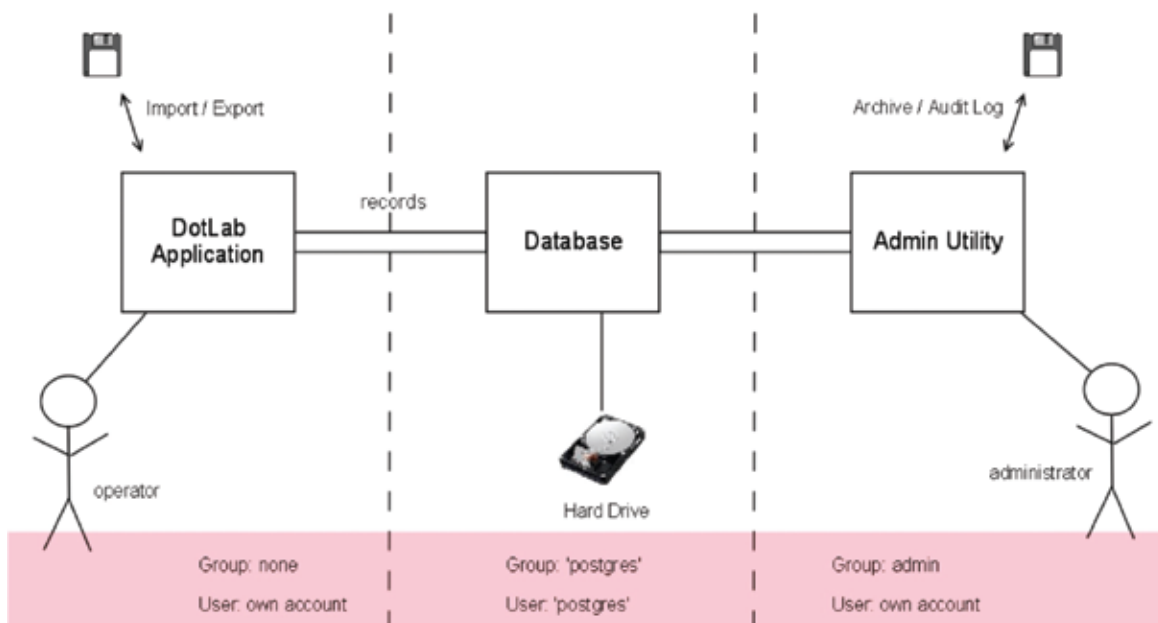
Title 21 CFR Part 11 of the Code of Federal Regulations presents the requirements of the U.S. FDA regarding electronic records and electronic signatures and defines the criteria under which electronic records and electronic signatures are considered to be trustworthy, reliable and equivalent to paper records.

dotLab Software is now available with support for 21 CFR Part 11 Compliance

Features Include:

- Secure Database
- Audit Trail
- Signature controls
- User/Password Controls

Full details in: "Guide to Using the dotLab System in a 21 CFR Part 11 Compliant Environment" found at: <http://www.axela.com/products/literature.php>



Architecture: The architecture drawing above shows the data flow and database structure within the dotLab software, along with inputs and outputs of the dotLab System.

To inquire about purchasing the 21 CFR Part 11 option contact Axela at:

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