



# What is Electronic Investigator Site File (eISF)?

eISF is a web-based platform that investigational sites can use as a replacement for the typical paper-based Investigator Site File (ISF) for a clinical trial.

Site participation in eISF is voluntary. However, because of the anticipated site benefits, we want to ensure that sites fully understand eISF in order to make an informed decision.

When a site utilizes eISF—they have full ownership of the contents—just as they do when managing essential study documentation via a paper binder.

Florence is the eISF vendor that we are working with.

We are offering the use of eISF for new Sponsor studies, at no cost to sites.

## Potential benefits for research sites utilizing eISF

Cost effective, minimizes site document storage costs

Green solution: reduces use of paper

Removes paper burden for managing essential documents (no need to print and scan)

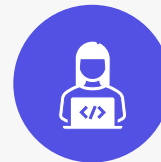
Easy to search and locate documents

eSignatures for Study Investigators

Remote monitoring of the site file

25-year electronic archiving

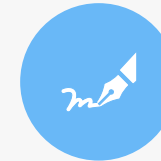
## Key features of Florence eISF to aid research sites



Simple Site Account Set-up Process



Intuitive Navigation & Document Management



Streamlined Signatory Process



Audit Trail

## Next Steps

If you are interested in utilizing Florence eISF to help manage your investigator site file, please notify your Sponsor CRA as soon as possible.

If you have already confirmed your interest in using Florence eISF for this study, it is good to consider who will be your eISF Account Administrator at your site. Once identified, please inform your CRA.

If you have further questions about Florence eISF, please visit [Why Florence | MSD - Florence \(florencehc.com\)](https://www.florencehc.com).