I'm unable to add my Investigator to one of our clinical sites. What do I do?

- Investigator (IVR), or Registration Coordinator (RC) on their behalf, accesses Registration and Credential Repository (RCR)
- Jump to Form FDA 1572 tab
- Add missing Practice Site to "Draft" RCR profile
- Jump to < Validate and View Documents >
- Generate, review, and sign just the Form FDA 1572
- Submit to CTEP
- When approval email is received, return to RUMS or NCORP-SYS and add the investigator to the required clinical site