

0910-0312 eSubmitter Screenshots

The screenshot shows the eSubmitter application window. The title bar reads 'eSubmitter'. The menu bar includes 'File', 'Edit', 'View', 'Table', 'Output', 'Tools', and 'Help'. The status bar shows 'Submission Name: test' and 'Report Type: CTP: Transmittal Form'. The main content area has a tab labeled 'Introduction: Overview'. A green notification box at the top contains a lightbulb icon and text: 'Notice: eSubmitter submission templates are updated monthly. Using updated templates for your submission helps ensure the latest fields and structure of each submission can be received and processed by the FDA. When you create a new submission, eSubmitter will provide the newer template. If you want to create a submission from a prior submission, you can use the "Save As" option and eSubmitter will load the newer template. However, if eSubmitter updates are blocked by your IT infrastructure then eSubmitter cannot load the newer template. If eSubmitter updates are being blocked, check the FDA eSubmitter application history on the eSubmitter webpage to locate important CTP template updates so that you can perform manual reinstalls as needed. http://www.fda.gov/forindustry/FDAeSubmitter'. Below the notification, the section header is 'CTP Submission'. The text below reads: 'Use the arrows to navigate the submission screens. Please note, there are several icons within the application to help guide you. Most importantly, the light bulbs indicate additional instructions, definitions from the guidance document, and other helpful hints. Blue dots indicate required fields.'

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The screenshot shows the eSubmitter application window. The title bar reads 'eSubmitter'. The menu bar includes 'File', 'Edit', 'View', 'Table', 'Output', 'Tools', and 'Help'. The status bar shows 'Submission Name: BeYourself_01102022_PRA_TEST' and 'Report Type: CTP: Transmittal Form'. The main content area has a tab labeled 'Contact: Instructions'. The section header is 'Introduction > Identification > Submission Information'. Below the header, the text reads: 'You are in the Identification section. This section requests contact and address information for those responsible for the submission.' A red-bordered box highlights a section titled 'Paperwork Reduction Act' with the following text: 'According to the Paperwork Reduction Act of 1995, an agency may not conduct or sponsor, and a person is not required to respond to a collection of information unless it displays a valid OMB Control Number. The valid OMB Control Number for this information collection is 0910-0312 and expiration date XX/XX/20XX. The time required to complete this information collection is estimated to average 1 hour per response, including the time for reviewing instructions, searching existing data sources, gathering and maintaining the data needed, and completing and reviewing the collection of information. Send comments regarding this burden estimate or any other aspects of this information collection, including suggestions for reducing burden, to PRASStaff@fda.hhs.gov'.