OnePacs Online Documentation

User Guides

Accessing the System

- Guide to Basic Functionality
- Setting User Preferences
- Using OnePacs on the iPad (or other mobile devices)

Transmitting or Uploading Cases to OnePacs

- How to Upload Studies to OnePacs
- Confirming Studies (Online Study Requisition System)
- Attaching Files to Cases; Receiving Case-related Faxes or Emails
- Study Notes and Flags
- Modifying Study Data after Transmission to OnePacs

Viewing Cases

- Viewer Options with the OnePacs system
  - OnePacs Workstation for Windows
  - OnePacs Workstation for macOS
  - OnePacs Web Viewer
  - RemotEye Viewer
  - Simple web viewer
  - Viewing cases on mac OS
  - Osirix HD for iPad
- Study Retriever (for use with the OnePacs workstation, or other viewers)

Sharing Cases

- Case Access Web Links

Reporting Cases

- Classic Report Editor
- Cloud Report Editor and VR
- Dictation with OnePacs
- Proofreading-based workflow ("Review workflow")
- Peer Review
- Result Coding Schemes
- Using Standard Report Texts
- Report Editor Tokens
- Using the Online QA Issue Management Solution
- Case Alerts
- Results Reporting
- Report keyboard shortcuts

Miscellaneous Features

- Secure, HIPAA-compliant email system
- Facility Directory Information system
- Burning Studies to CD or DVD
- Data Analysis Features
- Case Notifications
- Study History Audit
- Export
- Interface Management

Administrator's Guides

Installation Guides

- OnePacs Desktop for Windows installation guide (diagnostic viewer)
- OnePacs Desktop for macOS installation guide (diagnostic viewer)
- Study Retriever
  - Windows Study Retriever
  - Mac OS Study Retriever
  - Using the Windows Study Retriever with OnePacs Workstation for macOS, or OsiriX
- OnePacs Gateway (DICOM transmission system) Installation Guide

Administrative Guides

- Administrative features guide
- System architecture overview
- Capabilities of the free and paid versions

User Management

- Managing users
- Managing facilities
- Facility Subgroups
- System permissions
- Facility privileges
- Study filters

Workflow Management

- Managing access to cases
- Reporting and report distribution management
- Incoming Documents
- Scheduling

Other Administrative Features

- Billing functions
- Data analysis features
- Facility Radiologist IDs
- System Configuration

Transmission Computer (OnePacs Gateway) Management

- OnePacs Gateway User Interface
- Configuring the Gateway for Direct Downloads
- Configuring Compression
- Configuring the Gateway to route studies to more destinations
- Changing the AE Title of the Gateway
- Configuring the Gateway for Multiple Facilities
- Administrative Interface

Interfacing with Other Systems

- DICOM
  - Modality Worklist
  - Automated Prefetching of Prior Exams

FAQ

Frequently Asked Questions

Support

OnePacs support options

OnePacs User Forums

Access OnePacs User Forums

Miscellaneous

- Conformance statements
- Security considerations
- Indications for Use and Safety Information

Certifications and Registrations

The OnePacs System is 510(k) cleared by the US Food and Drug Administration as a class II medical device. OnePacs satisfies the requirements of the FDA for manufacturers of Class II devices, including Establishment Registration, Device Listing, U.S. Agent and Quality Systems (QS) regulation (21 CFR Part 820), and other regulations.

CE According to the guidelines stated in Directive 93/42/EEC of the European Community, the OnePacs system distributed in the EU is a Class I Medical Device. OnePacs satisfies the requirements for bearing the CE mark on its labeling.

ANVISA

The OnePacs system is cleared for medical use in Brazil under the ANVISA program.

O sistema OnePacs é aprovado pela ANVISA para uso clínico no Brasil.

We self-certify compliance with:

U.S.•EU SAFEHARBOR U.S. DEPARTMENT OF COMMERCE

- Related Study Matching
- HL7
  - Reports Outbound
  - Reports Inbound
  - Orders Inbound