

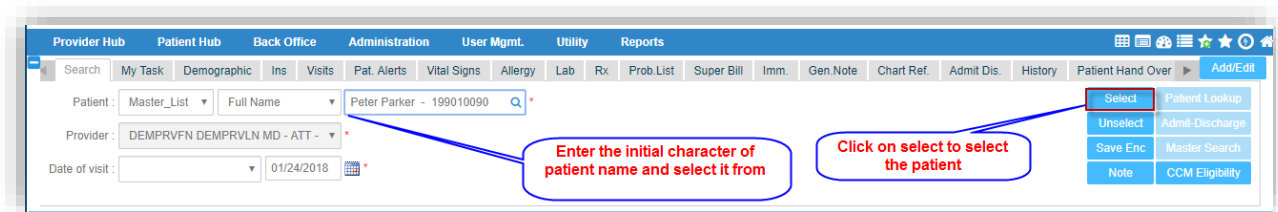
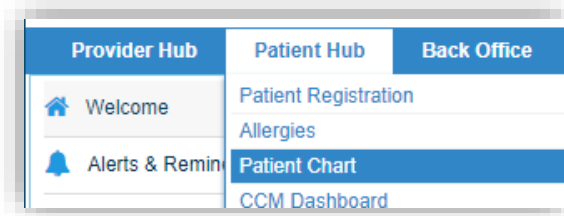


Geesemed Implant Device Manual-2020

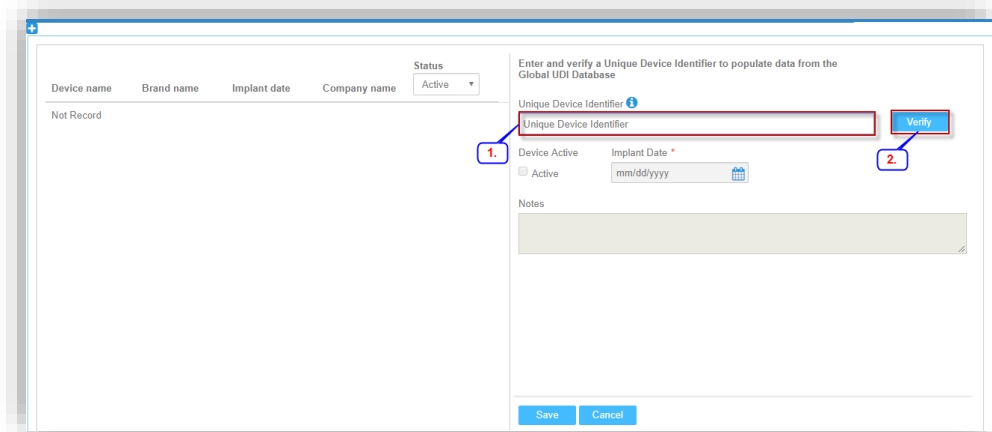


Implant Device

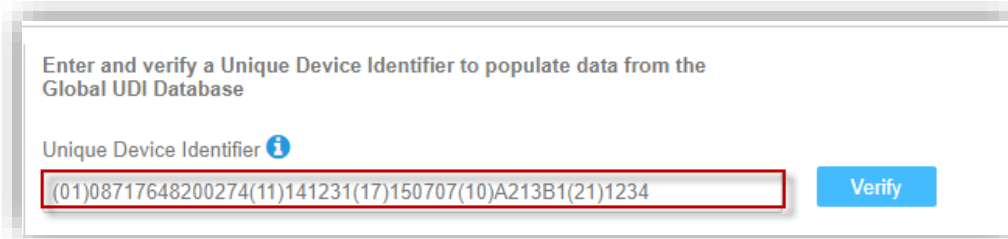
- Go to Patient chart menu for select a patient through below path.
- Path: “**Patient Hub>>Patient Chart**”



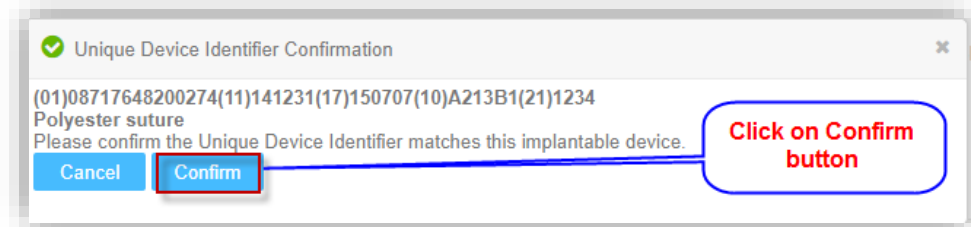
- Go to Implantable device menu through below path.
- Path: “**Patient Hub>>Implantable Device**”



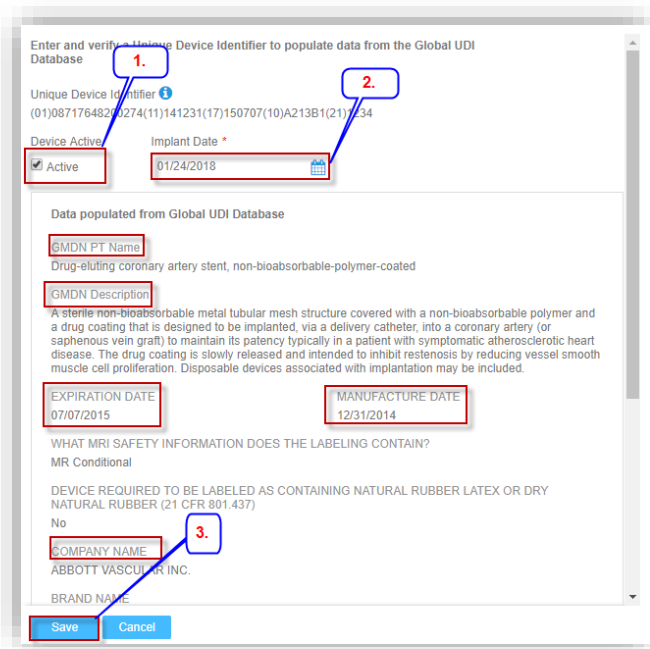
- Enter the Unique device identifier number which is available on device.



- Click on Verify then user will get unique device identifier confirmation window as shown in below screen shot.



- Now user will get all information related to Device like, name, description, Expiration date, Manufacturer date, Company name, Brand name, serial number, LOT batch, version model number etc...



Enter and verify a Unique Device Identifier to populate data from the Global UDI Database

Unique Device Identifier
(01)08717648200274(11)141231(17)150707(10)A213B1(21)1234

Device Active Active Implant Date * 01/24/2018

Data populated from Global UDI Database

GMDN PT Name
Drug-eluting coronary artery stent, non-bioabsorbable-polymer-coated

GMDN Description
A sterile non-bioabsorbable metal tubular mesh structure covered with a non-bioabsorbable polymer and a drug coating that is designed to be implanted, via a delivery catheter, into a coronary artery (or saphenous vein graft) to maintain its patency typically in a patient with symptomatic atherosclerotic heart disease. The drug coating is slowly released and intended to inhibit restenosis by reducing vessel smooth muscle cell proliferation. Disposable devices associated with implantation may be included.

EXPIRATION DATE: 07/07/2015 MANUFACTURE DATE: 12/31/2014

WHAT MRI SAFETY INFORMATION DOES THE LABELING CONTAIN?
MR Conditional

DEVICE REQUIRED TO BE LABELED AS CONTAINING NATURAL RUBBER LATEX OR DRY NATURAL RUBBER (21 CFR 801.437)
No

COMPANY NAME: ABBOTT VASCULAR INC.

BRAND NAME:

Save Cancel

- Select the active check box if device is implanted in patient body.
- Select the implementation date of device in patient.
- Click on '**Save**'.

- After saving the record that record will be displayed in left side grid.

Device name	Brand name	Implant date	Company name	Status
Drug-eluting coronary artery stent, non-bioabsorbable-polymer-coated	XIENCE ALPINE	01/24/2018	ABBOTT VASCULAR INC.	Active

Enter and verify a Unique Device Identifier to populate data from the Global UDI Database

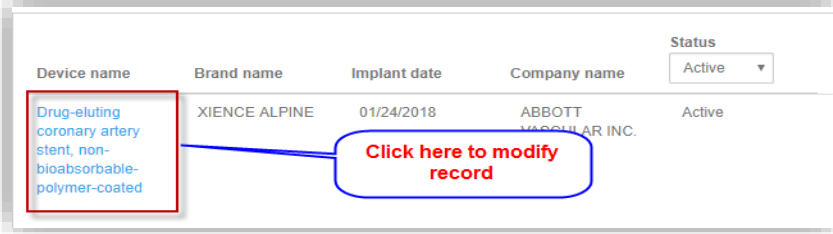
Unique Device Identifier Verify

Device Active Active Implant Date *

Notes



- If you want to modify existing record then click on device name link as shown on below screen.



- If user wants to in active implemented device then they need to uncheck the active check box.

