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### **Epic Order and Manual Enter/Edit of Point-of-Care Tests**

	Name, Position	Signature	Date
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Approving Authority:			
Approving Authority:			
Approval Date:			

#### **Bi-Annual Review/ Revisions**

Date	Signature	Date	Signature

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#### Epic Order and Manual Enter/Edit of Point-of-Care Tests

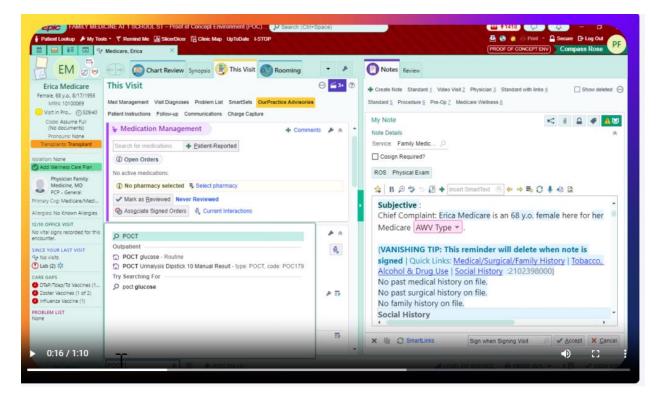
**Purpose:** This standard operating procedure (SOP) provides instructions for ordering and resulting Point-of-Care Tests in Epic for hospital and ambulatory sites.

This SOP governs the Enter/Edit features in Epic from ordering the Point-of-Care Tests (POCT) to entering patient results.

#### **Procedure:**

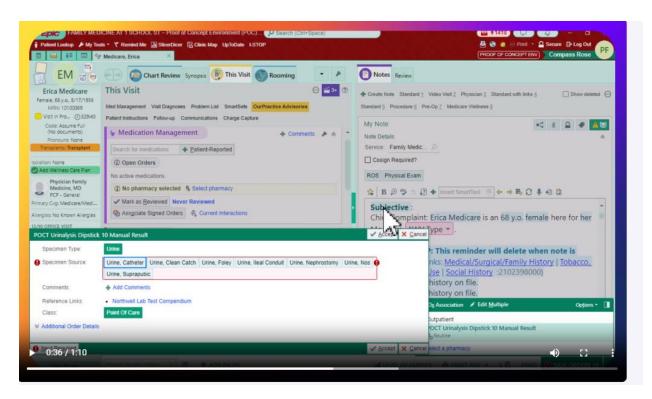
#### **Epic POC Orders**

- 1. When in the patient's chart go to Add Orders button and click in bottom left in the chart
- 2. The conduct search using key order words (Note: Point-of-Care orders begin with POC or POCT

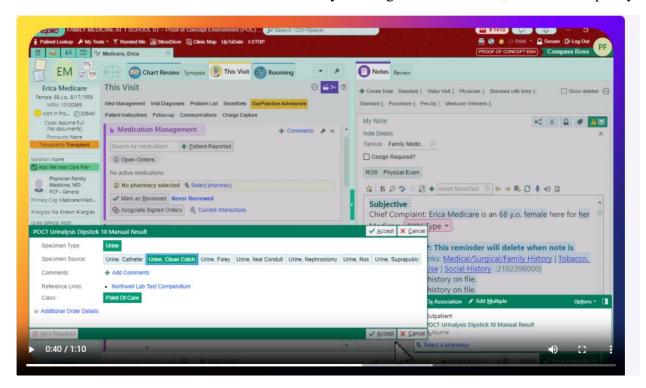


3. Select the test, then review any key items needed for the test to include, specimen type, specimen source, etc.. There is also a link to the Northwell Lab Test Compendium to review more in-depth information about the test

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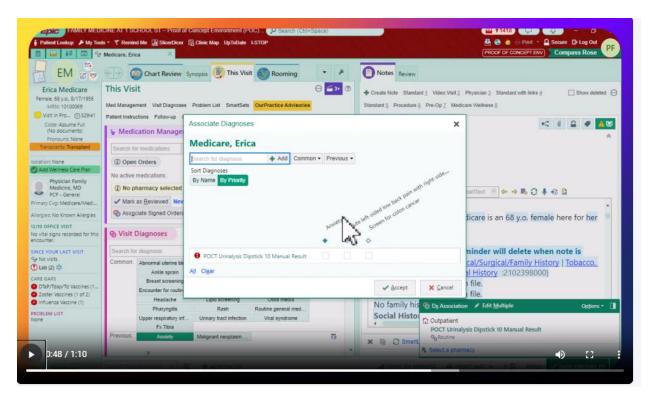


4. You can make a selection for the source by clicking on that selection, then hit Accept key



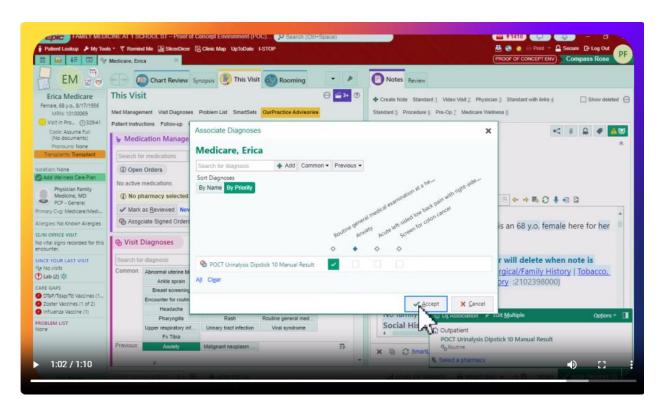
5. Then click Assign orders in the bottom right on the screen

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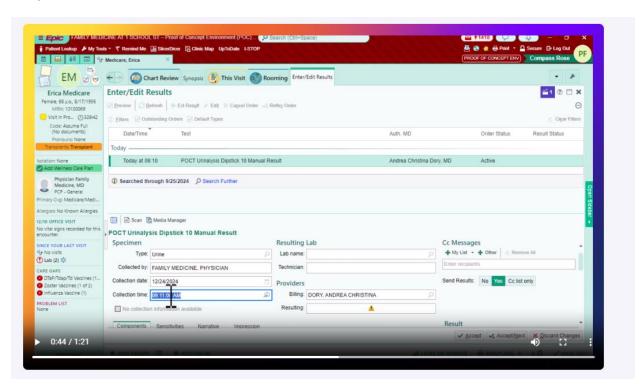


- 6. This will lead to assigning your diagnosis field (see above)
- 7. Add diagnosis then click Accept button

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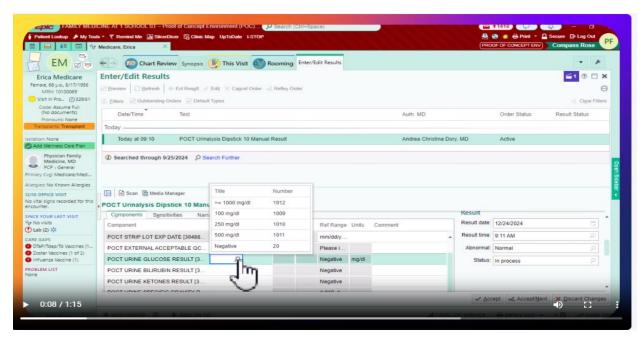
8. Afterwards the test can now be resulted



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#### **Epic POC Result Entry**

1. When ready to result in the patient's chart go to Enter/Edit Results tab

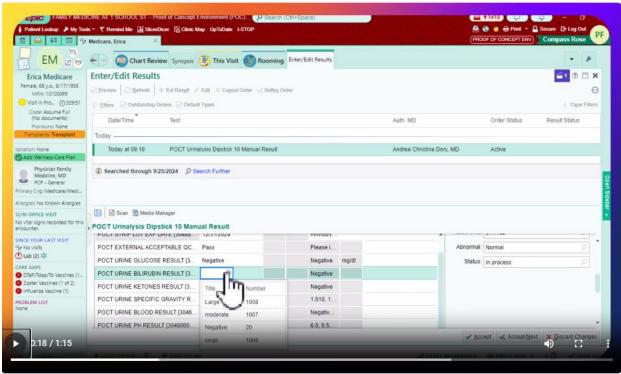


- 2. There are various fields to be completed in the results area (entries highlighted with a red marker are mandatory fields and must have an entry to be able to finalize test results). Here are some of the fields which require information:
  - a. Result date/time
  - b. Test kit lot#s (lot #s can be scanned into this field with a barcode scanner to ensure accurary)
  - c. Test kit expiration date
  - d. Internal or External QC checks (for results to be valid and internal and/or external QC has to pass successfully)
  - e. Patient test results (results can be in multiple fields of entry, e.g., fecal occult blood can have 1-3 fields on entry if more than one card is tested per order)
    - i. Qualitative result entry from test kits of "Negative", "Positive", "Reactive", "Non Reactive", "Invalid" or other results
    - ii. Quantitative result entry is usually from a device that measures test levels (e.g., AccuChek Inform II or CoaguChek XS Plus/Pro)
  - f. Instrument number field (required for sites that have multiple test devices)
    - i. Enter serial number of analyzer or
    - ii. 1,2,3, etc to denote which analyzer is in use

1. Site has to keep a written list of the analyzer serial numbers and which ones are labeled as 1,2,3, etc on file

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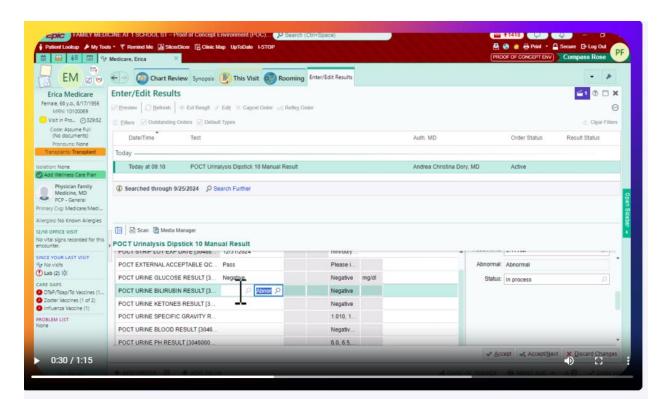
3. Type in result(s) by selecting magnifying glass and selecting from list or manually start typing results in open field which will initiate auto populate resulting



**NOTE:** When an abnormal or critical value is added, it will flag in entry area and total test areas

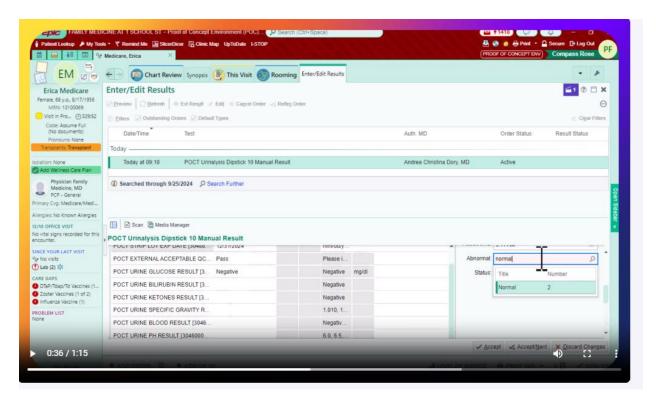
4. If an error is made, correct by editing first in the component (main entry area), then clear out component in box to the right of the main entry area

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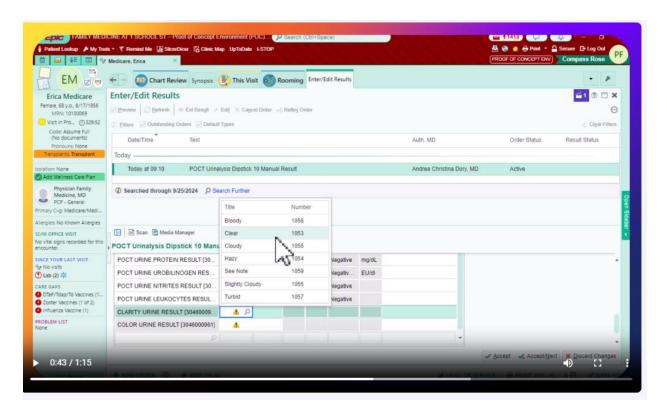
5. Then you can go to the far right area on the page for the main test and change that result to corrected result

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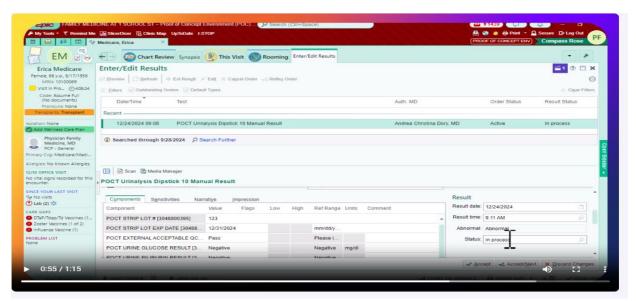


6. Components with yellow yield sign are recommended entry fields but not required (result entry is the same as all other manual results)

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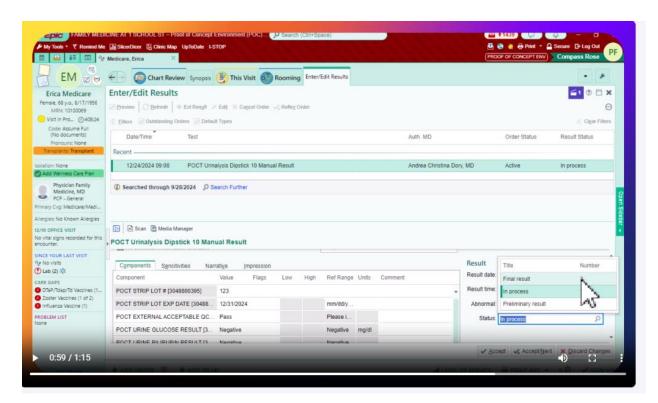


7. Once all results are added, go to review tab, select, then change Status in Result from In process to completed

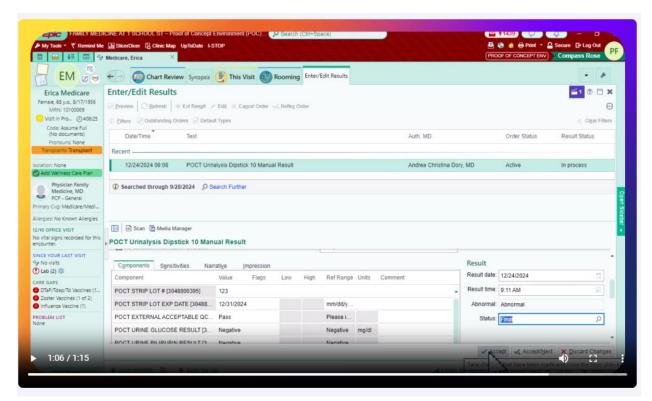


8. Click magnifying glass and choose Final result or enter Final result manually

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9. Click Accept, the test will finalize and drop from pending POCT queue test list



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- 10. Any issues with resulting any tests must be reported immediately to site supvs/mgrs, who will vet out any issues to ensure accurate test results are ordered and completed for each patient.
- 11. Site supvs/mgrs, must keep a log of any unusual ordering/resulting issues to include the issues, who was contacted to assist with resolving issues (e.g., Epic IT Help Desk, POCT Supv/Mgr, etc).
- 12. Discrepancies must be reported to the POCT Dept as part of the Quality Assurance program to ensure the highest accuracy of testing compliance as deemed by multiple accrediting agencies (NYS DoH, College of American Pathologists, etc).
- 13. Site staff will be trained on Ordering and Enter/Edit procedures by site management teams or be assisted by the POCT departments overseeing any sites by completing a competency. Competencies will be initial, then annual
  - a. Annual competencies will be completed by site management teams
  - b. Competencies relating to staff member(s) needing retraining due to needing more training or non-compliance issues should be completed by site management and documented as re-training on competency.
- 14. Critical values for the following tests are as follows:

	Test	Critical Low	Critical High	Age (if specified)
	POCT Glucose Accucheck Manual Result	<= 54 mg/dL	>= 450 mg/dL	1 to adults
	POCT Glucose Accucheck Manual Result	<= 45 mg/dL	>= 300 mg/dL	Neonates to < 4 weeks of age
es	POCT Glucose Accucheck Manual Result	<= 45 mg/dL	>= 450 mg/dL	4 weeks to 1 year of age
ases	POCT Whole Blood Hemoglobin Hemocue Manual Result (POC WB Hemoglobin)	<7.0 g/dL	>19 g/dL	
	POCT Ph Right Spot		>= 6.0 Placement not confirmed	
inks	Cholestech Manual Result (POC Glucose)	<45	>450	
	POCT Glycosylated Hemoglobin DCA Vantage (HGB A1C) Manual Result		High Risk (prediabetes): 5.7% - 6.4% Diabetic, diagnostic: > 6.5%	
	POCT I-STAT Creatinine Abbott Manual Result		>10.0	

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Distribution List:		
Hospital staff		
Ambulatory site staff		
Revision/Version His	·	
Revision/Version His Version No.	Date	Additions/Amendments
	·	Additions/Amendments NEW
	Date	