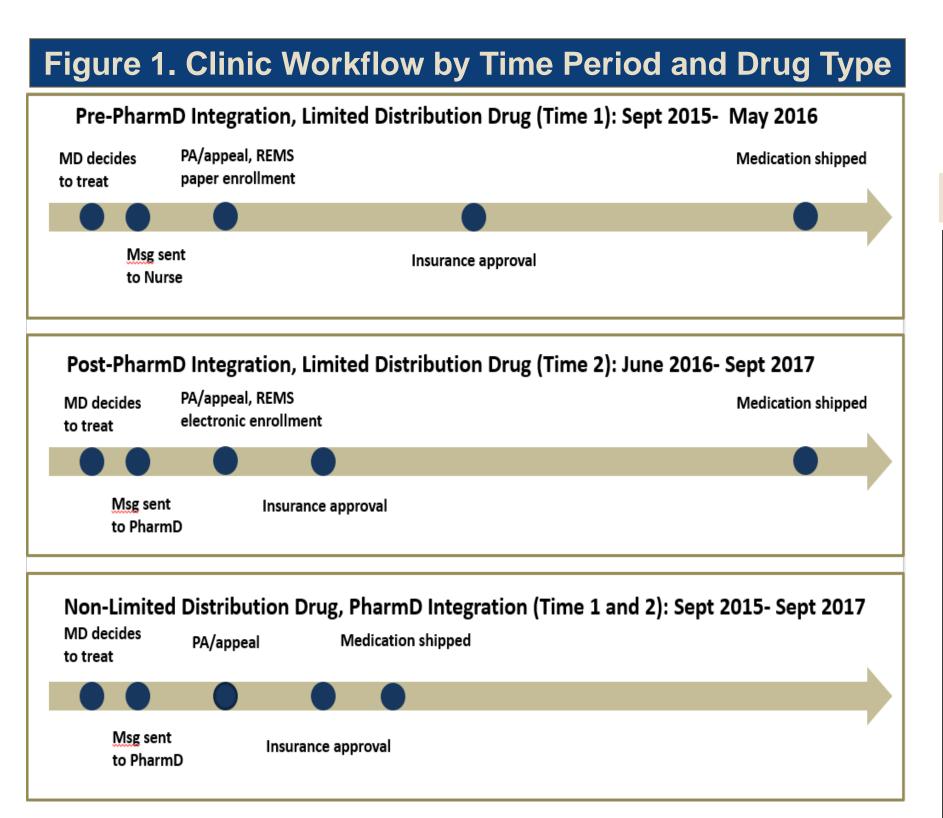


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BACKGROUND

- Oral anti-neoplastic therapy can be difficult to access due to insurance authorization, out of pocket costs, and limited distribution drugs (LDDs).¹
- In September 2015, a clinical pharmacist joined the Hematology Clinic at Vanderbilt-Ingram Cancer Center to facilitate timeliness of medications dispensed by Vanderbilt Specialty Pharmacy (non-LDDs).
- The pharmacist's scope expanded to manage LDDs in June 2016 (Workflow shown in Figure 1).



OBJECTIVES

- Compare access time for LDD vs. non-LDD prescriptions
- Assess whether integrating a clinical pharmacist into clinic decreased access time to LDD medications

Inclusion criteria:

recipients.

Primary outcome:

shipment

Statistical analysis:

	Time 1 (n=119) n (%)	Time 2 (n=291) n (%)			
Insurance					
Commercial	70 (59%)	143 (49%)			
Government	49 (41%)	148 (51%)			
Combination Therapy					
Yes	9 (8%)	31 (11%)			
No	110 (92%)	260 (89%)			
Off Label					
Yes	10 (8%)	36 (12%)			
No	109 (92%)	255 (88%)			
Drug Type					
Non-LDD	89 (75%)	196 (67%)			
LDD	30 (25%)	95 (33%)			
Common Medications					
LDD:					
Revlimid®	23 (19%)	60 (21%)			
Pomalyst®	7 (6%)	35 (12%)			
Non-LDD:					
Imbruvica®	30 (25%)	41 (14%)			
Ninlaro®	16 (13%)	39 (13%)			
Jakafi®	17 (14%)	36 (12%)			

PREDICTING TIME TO MEDICATION ACCESS FOR HEMATOLOGIC MALIGNANCIES: THE IMPACT OF AN INTEGRATED SPECIALTY PHARMACY AND LIMITED DISTRIBUTION DRUG NETWORKS

METHODS

 Oral anti-neoplastic therapy prescribed by a hematology provider to an adult patient between Sept 2015-Sept 2017, excluding uninsured patients or free drug sample

• Time (in days) from treatment decision to medication

 Proportional odds logistic regression to test whether access time was associated with drug type (LDD vs. non-LDD), Time Period (Time 1: 9/2015-5/2016; Time 2: 6/2016-9/2017), and Drug Type* Time Period, controlling for off-label use and insurance type.

RESULTS

Table 1. Characteristics of Prescriptions (n=410)

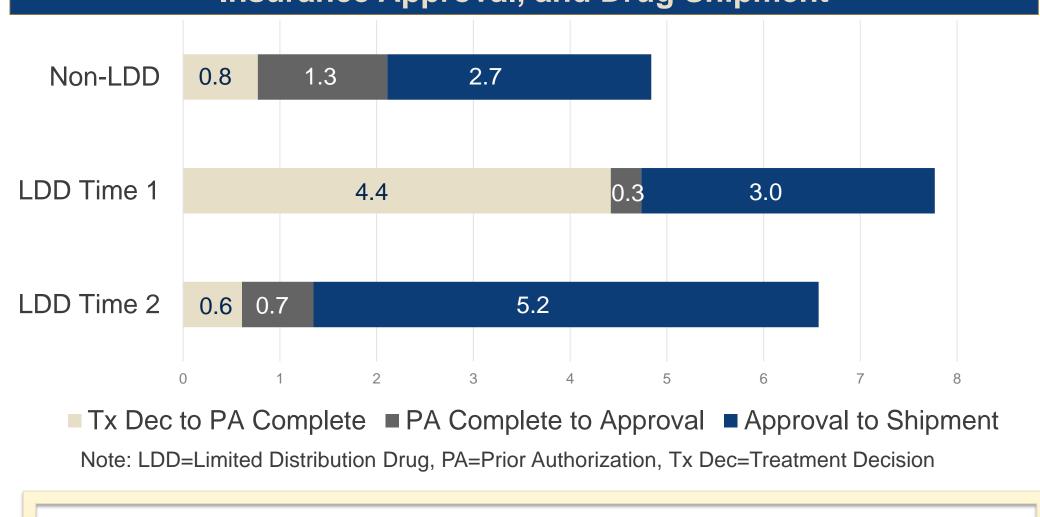
Primary Outcome

Median time from treatment decision to shipment:

- 6 days (IQR: 3-9) for LDD
- 3 days (IQR: 1-6) for non-LDD
- Predictors of Medication Access Time (Table 2)
- Longer access time for off-label than on-label indications
- In Time 1, time from treatment decision to shipment was significantly longer for LDD than non-LDD drugs
- For LDD drugs, access time reduced from Time 1 to Time 2

Table 2. Proportional Odds Logistic Regression testing predictorsof Medication Access Time					
Predictor	Odds Ratio	Lower Cl	Upper Cl	p-value	
Time 2 vs. Time 1	1.34	0.86	2.09	0.191	
LDD vs. Non-LDD	6.56	3.07	14.04	<0.001	
Off-label vs. on-label Government vs. Commercial	2.59	1.47	4.55	0.001	
Insurance	1.02	0.72	1.44	0.905	
Time 2 * LDD	0.41	0.17	0.96	0.040	

Figure 2. Mean Days between Treatment Decision, PA Completion, Insurance Approval, and Drug Shipment



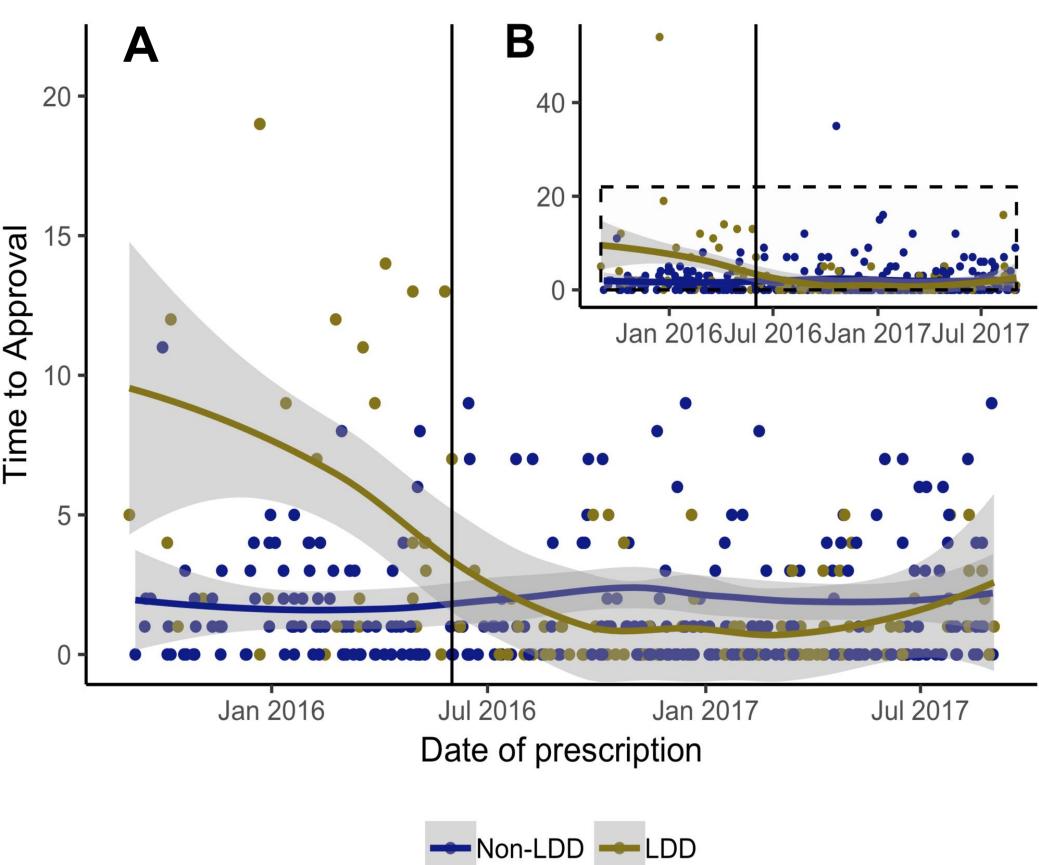
100% of insurance appeals were approved (5 in Time 1, 23 in Time 2)

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RESULTS







CONCLUSION

- Integrating a pharmacist into clinic significantly shortened time from treatment decision to shipment for LDD drugs, partially overcoming access barriers.
- Access to these drugs is still slower than non-LDD medications as they cannot be fully integrated into clinic workflow. The integrated specialty pharmacy program at VSP adds value to patient access and outperforms LDD medications, challenging the value of LDD networks beyond medical economics.

References:

1.Schwartz RN, Eng KJ, Frieze DA, et. al. NCCN Task Force Report: Specialty Pharmacy: J Natl Compr Canc Netw.2010 v. 8, p. S-1-S-12.