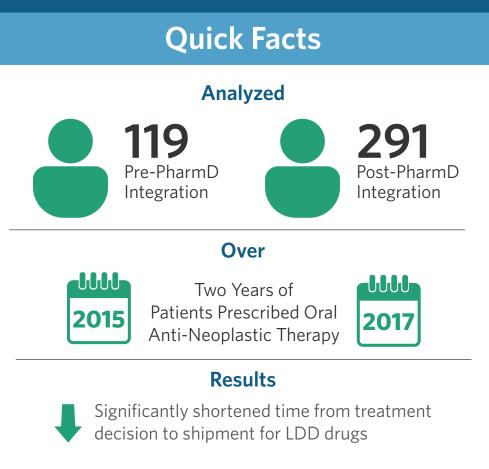
Predicting Time to Medication Access for Hematologic Malignancies: The impact of an integrated specialty pharmacy and limited distribution drug networks

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Increased value to patient access and outperformed LDDs, challenging the value of LDD networks beyond medical economics

100% of insurance appeals were approved

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The impact of an integrated specialty pharmacy and limited distribution drug networks

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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 of certain agents (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).

Figure 1: Clinic Workflow by Time Period and Drug Type

Pre-PharmD Integration, Limited Distribution Drug (Time 1): 09/15 - 05/16



Post-PharmD Integration, Limited Distribution Drug (Time 2): 06/16 - 09/17



Non-Limited Distribution Drug, PharmD Integration (Time 1 & 2): 09/15 - 09/17



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: 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 recipients.
- Primary outcome: Time (in days) from treatment decision to medication shipment

Methods

Results

Table 2: Proportional Odds Logistic Regression

Testing Predictors of Medication Access Time

 Statistical analysis: 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.

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

Primary Outcomes

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

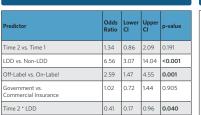


Figure 2. Mean Days between Treatment Decision, PA Completion, Insurance Approval, and Drug Shipment



Note: LDD=Limited Distribution Drug, PA=Prior Authorization,Tx Dec=Treatment Decision

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

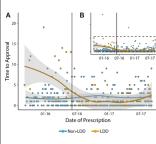


Figure 3. Time from Treatment Decision

to Insurance Approval: Time 1 {A} vs. Time 2 {B}

Conclusions-

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

