

Cost and Wastage Estimates for an Oral Oncology Medication Split-Fill Option in a Patient Management Program.

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Payers have the opportunity to reduce pharmacy costs when patient's start on a new cancer medication. Statistical comparisons between the matched patients indicated the split fill process was effective on reducing pharmacy costs for all months, lowering initial copay, and increasing persistency in the second month. Modeled wastage was meaningfully lower for all months.

OBJECTIVES

- To compare patients with pharmacy benefit designs that include a split-fill option to similar patients without this option on patient discontinuation rates, patient reported side-effects, estimated pharmacy costs, and potential wastage. During the study period, 11 of the 15 medications were available, an extended list from prior studies.^{1,2,3,4}

METHODS

- Study design: Retrospective cohort.
- Patient exclusion criteria: < age 6; off-label pediatric medication; residing outside U.S. state territories; greater than a 40 day supply on initial dispense; starting with two concurrent oral oncolytic medications
- Propensity matching (1:1 greedy match algorithm) used to compare cohorts (split-fill vs. non split-fill). Propensity variables: Patient age, gender, state census areas, index medication, start date as historical half-year segmentation, and use of more than a single medication.
- Paired t-test were conducted on the outcomes of payer costs, copay cost, and persistency on matched patients.
- Wastage was modeled for only the non split-fill patients using the split-fill patient discontinuation rate, and no statistical analysis

Figure 1. Potential wastage in fill patterns for non split-fill based on rates of discontinuations in split-fill patterns applied to non split-fill discontinuations.

| Cohort | Pattern | First Month | | Second Month (discontinuations) | |
|----------------|---------|---------------------------------------|---------------------------------------|---------------------------------------|----------------------|
| | | 1 st half | 2 nd half | 1 st half | 2 nd half |
| Non Split Fill | A | 30 days fill -----> | | No fill | |
| | B | | 30 days fill -----> | | No fill |
| Split fill | A | 1 st 14 day fill -----> | 2 nd 14 day fill -----> | No fill | |
| | B | | 1 st 14 day fill -----> | 2 nd 14 day fill -----> | No fill |

RESULTS

- 2,363 of 2,473 patients within CMP met selection criteria
- 672 identified in the split-fill program and 1,691 in non split-fill
- One case did not match to a control patient, reducing the matched count to 671 per cohort
- Post-matched comparisons on propensity variables indicated differences within the accepted 10% value of standardized differences (see Table 1.)

Table 1. Standardized differences pre- and post-propensity matching, and post-propensity descriptive statistics.

| | Standardized Differences | | Post Propensity Descriptive Statistics | |
|----------------------------|--------------------------|------------|--|-------------|
| Variable | Pre-match | Post-match | Case | Control |
| Age | -0.047 | 0.002 | 57.6 (11.8) | 57.6 (13.9) |
| Female | -0.091 | -0.015 | 49.6% | 50.3% |
| Census area | 0.498* | 0.014 | | |
| Central EN,WN,ES | | | 50.3% | 49.7% |
| New England, Mid Atlantic | | | 50.2% | 49.8% |
| South Atlantic, WS Central | | | 49.2% | 50.8% |
| Mountain, Pacific | | | 49.8% | 50.2% |
| Index Medication | 0.144* | 0.082 | | |
| sorafenib | | | 20.2% | 20.2% |
| everolimus | | | 15.2% | 15.2% |
| sunitinib | | | 15.0% | 16.8% |
| pazopanib | | | 13.9% | 12.7% |
| erlotinib | | | 13.1% | 12.4% |
| dasatinib | | | 9.9% | 10.1% |
| nilotinib | | | 6.1% | 6.3% |
| crizotinib | | | 3.6% | 3.4% |
| ceritinib | | | 1.9% | 2.1% |
| vorinostat | | | 0.6% | 0.3% |
| bexarotene | | | 0.3% | 0.2% |
| Use 2+ medications | 0.057 | 0.015 | 48.8% | 51.2% |
| Historical 6 months | 0.144* | 0.023 | | |
| 1 st | | | 49.9% | 50.1% |
| 2 nd | | | 49.2% | 50.8% |
| 3 rd | | | 50.6% | 49.4% |
| 4 th | | | 50.5% | 49.5% |

*Difference exceeds 0.10 criterion

- Persistency (i.e., no discontinuation at 45 days): Significantly higher ($p < .0001$) for the split-fill group compared to non split-fill in the second month (71.6% vs 67.0%).
- Copay: In first month, split-fill had \$132.50 lower copay than non split-fill ($p < .007$).
- Pharmacy costs: Difference between cohorts in pharmacy costs was significantly lower for split-fill cohort per month, with the three month average being \$2,724.97 AWP (see Table 2).
- Split-fill rates applied to weight non split-fill rate in modeling wastage (pattern A rates multiplied + pattern B rates multiplied): 28.2% of non split-fill patients discontinued (wastage).

- The average wastage for non-split fill patients was \$2,782.29 AWP assuming that the latter 14 days of the month were not used as modeled (see Table 3).
- Patient reported side-effects did not differ significantly between cohorts, either in prevalence rates or time to first report.

Table 2. Monthly mean plan costs (AWP) difference for non split-fill compared to matched split-fill patients.

| Non Split Fill Cost Difference from Split Fill | | |
|--|----------|-----------|
| Month Filled | AWP \$* | Std. Dev. |
| 1 | 3,118.90 | 6,787.40 |
| 2 | 2,259.80 | 6,989.10 |
| 3 | 2,796.20 | 8,261.50 |
| 3 month average | 2,724.97 | |

*P<0.0001

Table 3. Non split-fill AWP for potential wastage due to discontinuation (weighted by split-fill discontinuation rates).

| Month Filled | AWP \$ PUPM* | AWP \$ 14 days |
|-----------------|--------------|----------------|
| 1 | 5,180.87 | 2,612.64 |
| 2 | 5,144.93 | 2,680.95 |
| 3 | 5,908.20 | 3,053.28 |
| 3 month average | 5,411.34 | 2,782.29 |

*PUPM=per utilizing patient per month

CONCLUSIONS

- Implementing a split-fill process within a pharmacy management program for the first three months of a new therapy may result in: lower discontinuation rates; lower copay; significantly reduced pharmacy costs; reduced potential wastage.
- By opting to adopt split-fill component, third party payers have a means of reducing the pharmacy cost for new therapy oncolytics
- Other researchers have also found similar benefits for modifying the quantity of initial fills for different therapies.^{5,6,7}

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