Cost-minimization analysis of generic equivalents (bortezomib, decitabine and capecitabine) in comparison to the originator brand medicines in Colombia

Shyam Akku, MD; Amit Garg, MD; Suhas Khandarkar, MPharm

Introduction: Majority of the Colombian population has become health insured leading to improved healthcare access in the country. Healthcare expenses have risen drastically, and cancer is responsible for the major disease burden. An option to reduce cost burden is to replace the use of originator drug products with generic formulations which are considered to be bioequivalent to the originator drugs and cost much less. We conducted a cost-minimization study for Colombia to assess the savings that could be achieved with the use of generic formulations of three anticancer drugs, namely bortezomib, decitabine and capecitabine as compared to their originator products.

Methods: We compared the prices of originator and Dr Reddy’s Laboratories Ltd generic equivalents of bortezomib, decitabine and capecitabine in Colombia for the year 2015. Prices of originator drugs were obtained from SISMED (Sistema de información de Precios de Medicamentos – Drug Price Information System), the prices of generic drugs were provided by Dr Reddy’s Laboratories Ltd. Cost savings were estimated in value and percentages.

Results: The study showed the substantial savings of 63% (US$4.68 million) with bortezomib, 26% (US$0.29 million) with decitabine, and 46% (US$1.50 million) with capecitabine from Dr Reddy’s Laboratories Ltd generics as compared to their originator branded medicines.

Conclusion: Considering the high economic burden of cancer and high prices of originator anticancer drugs, our study highlights the need to replace the originator branded medicines with the generics. This will not only improve the patient access to medications but also be beneficial for the insurance payers in Colombia, thereby improving the overall healthcare system in the country.

Keywords: Anticancer drugs, cost-minimization, generics, pharmacoeconomics

Introduction

Colombia is an upper-middle income country with over 48 million people, as reported by the World Health Organization (WHO) in the year 2015 [1, 2]. Over the past few years, Colombia has undergone various demographic and epidemiological changes viz. fall in total fertility rate from 3.24 children per woman in 1985 to 2.3 in 2013 while life expectancy has increased by four years over the period of 2000 to 2012; mortality rates have also declined from 1990 to 2012, e.g. under-five mortality rate: 35 vs 17 per 1,000 live births, maternal mortality ratio: 100 vs 83 per 100,000 live births [2, 3]. Overall, healthcare expenditure has increased drastically in Colombia. According to WHO, around US$962 per capita were spent on health expenditure in 2014 which accounted for 7.2% of gross domestic product (GDP) in the country [1, 3].

Non-communicable diseases account for 71% of total deaths in Colombia, and cancers constitute 17% of the total deaths [4]. Out of all the cancers, breast, stomach and colorectal cancer are three of the five most prevalent forms in Colombia accounting for 7%, 13.1% and 8.5% of total cancer mortality, respectively. Besides these, blood/bone marrow cancers (9.9%) are also responsible for high burden of cancer-related deaths in the country (leukaemia 4.9%, multiple myeloma 1.2%, non-Hodgkin lymphoma 3.4%, and Hodgkin lymphoma 0.4%) [5].

However, launch prices for the anticancer drugs have substantially increased over time. Anticancer medicines rank first in terms of global spending in the pharmaceutical market (US$91 billion in 2013). According to a recent study in the US, the average price of an anticancer drug is US$65,900 in 2013, while the average survival benefit is 0.46 years [6]. Overall, the economic burden due to cancer cost US$290 billion in 2010 with the greatest share of the medical costs (53%). In fact, economic burden of cancer is estimated to be US$88.3 trillion between the period 2011–2030 [7]. The economic burden of cancer per patient ranged from US$0.54 in India to US$4.32 in China, US$7.92 in South America, US$244 in Japan, and US$460 in the US [8].

At present, majority of the Colombian population is covered under health insurance (85% in 2008 and 97% in 2010) [3, 9]. This has secured the general population and increased the access to health care. However, health care in Colombia is in jeopardy as it faces financial constraints. One of the reasons attributed for this condition is high drug expenditure in the country. Colombia is believed to be one of the highest payers for drugs in the world due to extensive use of expensive originator drugs rather than generic molecules. Thus, there is a need to replace the originators with generic molecules in Colombia to bring down the overall healthcare economic expenditure [9].

Author for correspondence: Shyam Akku, MD, Manager – Global Medical Affairs, Dr Reddy’s Laboratories Ltd, 8-2-337, Road No. 3, Banjara Hills, Hyderabad – 500034 AP, India

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A recent cost-minimization study conducted by Alexandra et al. (Essential Medicines and Pharmaceutical Policies, World Health Organization) across 17 countries showed that substantial savings can occur by switching the purchases from originator medicines to generics. These savings ranged from 11% for beclomethasone inhaler to 73% for ceftriaxone injection [10]. Keeping in mind the high cost of anticancer drugs, we conducted a pharmacoeconomic study to estimate the savings that could be achieved with the use of generic formulations of anticancer drugs – bortezomib, decitabine and capecitabine – in comparison to their originator brands in Colombia.

**Methods**

**Rationale**

Pharmacoeconomics identifies, measures and compares the costs of various drug therapies to the society and the healthcare system. It helps the clinicians, health-payers, and other decision-makers to assess the costs and outcomes of various treatment options via cost-minimization, cost-benefit, cost-effectiveness and cost-utility analyses [11-13]. The choice of the analysis depends upon the clinical situation to be evaluated and the question being asked. Cost-minimization analysis compares the costs of treatments having similar outcomes and aims at identifying the most economic option available. On the other hand, cost-benefit analysis compares the costs and benefits of treatments that have different outcomes and the outcome is measured in monetary value. Cost-effectiveness analysis compares the costs per standardized units of effectiveness for treatments that have different outcomes. Cost-utility analysis compares the costs per quality-adjusted life years for treatments that have different outcomes. Thus, of the different pharmacoeconomic methods, cost-minimization analysis is the one that compares the two interventions when the clinical outcomes of both are equivalent [12]. This means that if the two drugs are equivalent to each other in terms of clinical and humanistic outcomes, e.g. generics and originator molecules, this method can be viable in making a decision with respect to the cost of the drug in order to choose the economical one [11].

Literature evidence clearly shows that generic molecules are equivalent to the originator drugs in terms of outcomes. As per the US Food and Drug Administration (FDA), generic drugs are approved only when they are bioequivalent to the branded drugs, and have demonstrated similar quality and performance [14]. Several systematic reviews have shown that cardiovascular and neurological generics are as clinically effective as the branded drug products [15-17]. Therefore, we chose the cost-minimization study to compare the costs of generic and originator products of three anticancer drugs namely bortezomib, decitabine and capecitabine in Colombia. Our study is further supported by a recent cost-minimization study that was conducted across 17 countries to estimate the savings that could be achieved by switching the use from the originator to generic drugs [10].

**Study methods**

In our cost-minimization study, we compared the prices of originator and Dr Reddy’s Laboratories Ltd (DRL) generic drugs of bortezomib, decitabine and capecitabine in Colombia. For originator products, price data was obtained from the SISMED (Sistema de información de Precios de Medicamentos – Drug Price Information System), Colombia. SISMED is an information system that provides the necessary data to analyse and control the drug prices in Colombia, and thus, helps in the regulation of the pharmaceutical market in the country.

The mean unit price of originator drugs was calculated for the year 2015. On the other hand, prices of generic drugs were provided by DRL. To estimate the maximum cost savings that could be generated if the originator branded medicines consumed were purchased at the price of generics, mean prices of originator brands and the price of generics were applied to the volume of originator branded medicines consumed. All the prices were calculated in US dollars to avoid any fluctuations in the currency value throughout the year 2015.

**Results**

Our study showed high costs of the originator products as compared to the generic molecules for all the three anticancer drugs. The originator drug of bortezomib cost US$7.48 million, decitabine US$1.12 million and capecitabine US$3.27 million. On the other hand, the generic bortezomib costed only US$2.80 million, decitabine costed US$0.83 million and capecitabine costed US$1.78 million, see Figure 1.

In total, a saving of 63% was achieved with the use of the generic form of bortezomib, 26% with decitabine, and 46% with capecitabine as compared to the originator drugs. Complete results for the three drugs are depicted in Table 1.

![Figure 1: Total costs with the use of originator versus DRL generic of bortezomib, decitabine and capecitabine (estimated for the year 2015)](image-url)
The study highlights the substantial cost savings that could be achieved by replacing the originator drugs of bortezomib, decitabine, and capecitabine with their DRL generic forms. All these three drugs have gained a rational place in the treatment of various cancers and thus are commonly used. Bortezomib (as a single agent or in combination) is the preferred regimen for the treatment of multiple myeloma, both in the primary therapy as well as maintenance therapy, and is used in relapsed patients. Also, it is approved as a second-line treatment for mantle cell lymphoma [18, 19]. Decitabine has been approved and used for the treatment of myelodysplastic syndrome and acute myeloid leukaemia [20, 21]. Capecitabine is used in the management of gastric, breast and colorectal cancer [22-25]. Moreover, all these three drugs have shown to confer cost savings in comparison to other anticancer drugs in their respective indications [26-35]. Therefore, using these drugs will not only help in the treatment of various cancers but can lead also to substantial cost savings.

Regarding the efficacy of generic medicines, it is well-proven that generics are bioequivalent to the originator drugs [14-17]. In fact, the Clinical Guidelines Committee of the American College of Physicians advises that clinicians should prescribe generic medicines in place of the expensive branded drugs. Generic medicines are not only associated with substantial cost savings, but also provide long-term adherence to the treatment [36].

Taking into account the field of oncology where medications are highly expensive, generics can play a very important role [37]. Generic imatinib is estimated to cost less than US$1,000 per year [38]. Decitabine 1,053 1,059 1,115,049 784 825,263 289,785 26

## Table 1: Total cost savings with the use of generic molecules of DRL bortezomib, decitabine and capecitabine as compared to the originator drugs in the year 2015

<table>
<thead>
<tr>
<th>Year 2015</th>
<th>Originator</th>
<th>DRL</th>
<th>Savings</th>
</tr>
</thead>
<tbody>
<tr>
<td>Drug</td>
<td>Units</td>
<td>Price per unit</td>
<td>Total value</td>
</tr>
<tr>
<td>Bortezomib</td>
<td>9,586</td>
<td>781</td>
<td>7,482,740</td>
</tr>
<tr>
<td>Decitabine</td>
<td>1,053</td>
<td>1,059</td>
<td>1,115,049</td>
</tr>
<tr>
<td>Capecitabine</td>
<td>12,930</td>
<td>253</td>
<td>3,274,733</td>
</tr>
</tbody>
</table>

*Prices in USD. DRL: Dr Reddy’s Laboratories Ltd.

## Discussion

Our study is based on estimating the total savings that could be obtained by replacing the originator drugs of bortezomib, decitabine, and capecitabine with their DRL generic forms.

## Conclusion

This study highlights the substantial cost savings that could be achieved by replacing the originator drugs of bortezomib, decitabine, and capecitabine with the DRL generic formulations in Colombia. This can be very helpful for the insurance payers and bring a positive reform in the Colombian health care.

## Competing interests

All authors are employees of Dr Reddy's Laboratories Ltd. Financial support was provided by Dr Reddy's Laboratories, Hydrabad, India.

## Provenance and peer review

Not commissioned; externally peer reviewed.

## Authors

Shyam Akku, MD
Amit Garg, MD
Suhas Khandarkar, MPharm

Dr Reddy’s Laboratories Ltd, 8-2-337, Road No. 3, Banjara Hills, Hyderabad – 500034 AP, India

## References
