Estimated costs of production and potential prices for the WHO Essential Medicines List

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ABSTRACT

Introduction There are persistent gaps in access to affordable medicines. The WHO Model List of Essential Medicines (EML) includes medicines considered necessary for functional health systems.

Methods A generic price estimation formula was developed by reviewing published analyses of cost of production for medicines and assuming manufacture in India, which included costs of formulation, packaging, taxation and a 10% profit margin. Data on per-kilogram prices of active pharmaceutical ingredient exported from India were retrieved from an online database. Estimated prices were compared with the lowest globally available prices for HIV/AIDS, tuberculosis (TB) and malaria medicines, and current prices in the UK, South Africa and India.

Results The estimation formula had good predictive accuracy for HIV/AIDS, TB and malaria medicines. Estimated generic prices ranged from US$0.01 to US$1.45 per unit, with most in the lower end of this range. Lowest available prices were greater than estimated generic prices for 214/277 (77%) comparable items in the UK, 142/212 (67%) in South Africa and 118/298 (40%) in India. Lowest available prices were more than three times above estimated generic price for 47% of cases compared in the UK and 22% in South Africa.

Conclusion A wide range of medicines in the EML can be profitably manufactured at very low cost. Most EML medicines are sold in the UK and South Africa at prices significantly higher than those estimated from production costs. Generic price estimation and international price comparisons could empower government price negotiations and support cost-effectiveness calculations.

INTRODUCTION

Lack of access to affordable medicines continues to represent a major global health burden. A recent analysis found that, based on per-capita pharmaceutical expenditure, a basket of 201 essential medicines was unaffordable in nearly all low-income countries and 13 middle-income countries. An earlier estimate put the number of people lacking regular access to essential medicines at one-third of the global population. In low-income and middle-income countries (LMICs), only 58% of essential medicines are available in the public sector, and 67% in the private sector, according to surveys of pharmacies. Medicines account for a quarter of all health expenditures globally, and 100% of health expenditures for about half of households in LMICs. Recent high-level groups have recommended greater transparency in drug pricing. Data on the costs of production for medicines are not publicly available, and

Key questions

What is already known about this topic?

► High prices of essential medicines pose a barrier to their use in many different settings.

► We searched the PubMed database and grey literature using the term (cost* OR price*) AND (manufacture OR production) AND essential AND (medicine* OR drug*).

While the costs of production for a few products have previously been analysed, we did not find any studies on production costs for all medicines on the WHO Essential Medicines List, or any similarly broad group of products.

► This study was commissioned by WHO, who identified this as a gap in the literature.

What are the new findings?

► The methodology presented in this study can be used to reliably estimate the generic price that can be achieved if profit margins are competitive, for a wide range of medicines.

► We show that prices could fall for most essential medicines in the UK and South Africa, and for nearly half of essential medicines in India. This suggests that even for old and widely used medicines, continued efforts should be made to encourage competitive supply.

Recommendations for policy

► This methodology can be used by governments and international institutions to inform price negotiations.

► Cost of production analysis could be incorporated into government tenders, health technology assessments and medicine price negotiations, allowing a reduction of information asymmetry.
health systems have limited negotiating power when a medicine is sold in a monopoly situation. However, prices can fall substantially when generic competition is enabled, achieving, for example, price reductions of 99% in first-line HIV/AIDS medicines.6

The WHO Model List of Essential Medicines (EML) was created in 1977 to support national health systems in prioritising drugs for procurement, and includes medicines that are needed to ‘satisfy priority health needs of the population’,7 based on disease burden, efficacy, safety and cost-effectiveness compared with other medicines in the same therapeutic group.8

The price of the raw medicine substance, or active pharmaceutical ingredient (API), is generally the most significant component of pharmaceutical cost of production.9 This analysis used data of API exported from India to estimate generic prices for all medicines in solid oral dosage forms included in the EML. We have previously undertaken analyses of production costs for viral hepatitis (HBV), tuberculosis (TB)12 and cancer drugs,13 using similar methods.

**METHODS**

Additional references for this section are given in an online supplementary appendix. Medicines with solid oral formulations from the 2015 EML were included (see online supplementary appendix for excluded formulations). All monetary values are expressed in US dollars.

**Country of production**

We assumed API procurement and formulation of finished pharmaceutical product (FPP) in India due to its large generic industry and historical importance in treatment of the HIV epidemic.6 The main alternative country to consider as a focus for the analysis was China; for pharmaceutical manufacture, India and China are similar in cost of labour, infrastructure and tax. However, Indian manufacturers have more experience with the WHO prequalification programme (an initiative that certifies the quality of medicines), which currently lists 335 products manufactured in India, but only 23 manufactured in China.15

**Estimation of API cost**

Data on price per kilogram of API exported from India was collected from an online database (infodriveindia.com) that collects data published pursuant to Indian customs regulations. Prices for exported goods in the database are given in both Indian rupees and US dollars, with the US dollar value calculated based on the exchange rate on the day of the transaction. Search terms used and full exclusion criteria are available in the online supplementary appendix. The default timeframe for API shipments used in the database search was 1 July 2014 to 1 July 2016, and was extended backwards by 2 years until at least 100 records were available, up to a maximum 6-year timeframe.

Linear regression was used to estimate the per-kilogram price on 1 July 2016 of exported API. API price per dose was calculated, and multiplied by a ‘salt-factor’ for drugs whose dose is expressed in the EML in terms of the active molecule, but for which the API is manufactured in a salt form (eg, an ‘erythromycin 250mg’ tablet contains 360 mg of erythromycin estolate). Stata/IC V.14.0 for Mac was used for statistical analysis of the API dataset.

**Costs of excipients**

Typical costs of excipients used in tablets and typical proportion of the FPP made up by each excipient were combined, and excipients were assumed to represent 50% of total tablet weight. This yielded an estimated average excipient cost of US$2.63 per kilogram of FPP (see online supplementary appendix).

**Conversion costs**

A number of sources were used to inform an assumed per-unit cost for converting raw API and excipients into an FPP (‘conversion cost’). We reviewed reports of capital and operating costs for pharmaceutical tablet formulating plants and contacted large generic manufacturers for confidential estimates. In addition, we identified the product with the lowest price in the UK, South Africa and India: with API cost and profit margins approaching zero for the cheapest products, the total per-unit cost can be seen as a high estimate of the conversion cost. Estimated conversion costs per tablet extracted from these sources are summarised in table 1, with details and full references available in the online supplementary appendix.

Per-unit conversion costs reported in analyses in the last 10 years (operating cost including depreciated capital cost) ranged from US$0.0002 to US$0.013 per tablet. The estimated conversion cost by Chaudhuri and West (US$0.01 per tablet) included costs of environmental protection and compliance with current Good Manufacturing Practice (cGMP) standards.16 The costs attributed to cGMP compliance are similar to costs estimated in an Indian government white paper (see online supplementary appendix). We therefore used a conversion cost of US$0.01 per tablet in this study.

**Tax and profit margin**

The average profit margin for the pharmaceutical industry in India was 8.8% in 1995 and 15.4% in 2005. We assumed a profit margin of 10%. Tax rates in India range from 18.5% to 34.6%; we assumed a midpoint value of 26.6% tax charged on net profits. The final price estimation algorithm is shown in figure 1. References are mentioned in the online supplementary appendix.

**Current prices in the UK, South Africa and India**

Lowest currently available prices (LCP) were collected from publicly accessible databases. For South Africa, we used public procurement prices published by the National Department of Health. For Tamil Nadu, India, prices were extracted from a list of a medicine tenders, or, if not available from that source, from an online database.
database of Indian Maximum Retail Prices. For the UK, prices were extracted from the British National Formulary (BNF) and the drugs and pharmaceutical electronic market information tool (eMit), with the lower of the two used to represent the UK price. The BNF reports ‘indicative’ prices, while eMit prices represent actual government purchases. In addition, the availability of more than one manufacturer in the UK was noted, as an indicator for generic availability and competition. Details on price sources and exchange rates use are available in the online supplementary appendix.

**Validation**

We compared estimated generic prices to global lowest current prices of medicines for which there are global market-managing initiatives, such as the Global Fund. Details on price sources are available in the online supplementary appendix.

**RESULTS**

Estimated API price per kilogram, generic price estimates and current prices are given in online supplementary appendix table 2.

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**Table 1** Per-unit cost component values available from published sources in the last 10 years

<table>
<thead>
<tr>
<th>Source (year, location)</th>
<th>Cost components included in estimate</th>
<th>Conversion cost per tablet (US$)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Most recent conversion cost ‘norms’ published under India’s Drug (Price Control) Order 1995 (2012, India)</td>
<td>CC not including distribution or packaging</td>
<td>0.0002–0.001</td>
</tr>
<tr>
<td>Lowest-priced product in UK, South Africa, India (2016)</td>
<td>Cost per unit of lowest-priced solid oral formulation FPP</td>
<td>0.0011–0.0043</td>
</tr>
<tr>
<td>‘Can local producers compete with low-cost imports? A simulation study of pharmaceutical industry in low-income Africa’, Chaudhuri and West (2015 India)</td>
<td>CC including depreciation of capital and packaging, but not including sales and distribution</td>
<td>0.0056</td>
</tr>
<tr>
<td>Confidential discussion by authors with large generic companies (2016, India)</td>
<td>CC including depreciation of capital and packaging, and including sales and distribution</td>
<td>0.0105</td>
</tr>
<tr>
<td>‘Outlook on pharma operations’ McKinsey &amp; Company (2014, no specific territory)</td>
<td>CC including packaging, but not depreciation of capital, sales or distribution</td>
<td>0.006</td>
</tr>
<tr>
<td></td>
<td>Total production costs, lowest-cost plants (not further specified)</td>
<td>0.013</td>
</tr>
</tbody>
</table>

CCC, conversion cost.

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**Figure 1** Algorithm used to estimate generic prices. API, active pharmaceutical ingredient.
There were sufficient API data for 148 medicines on the EML (including 20 fixed-dose combinations or medicines listed as combinations), and for 375 finished pharmaceutical products overall when including various dose forms. For 49 medicines on the EML, production costs could not be calculated due to a lack of data on exported API. Lowest current prices were identified for 73% of items in the UK, 47% in South Africa and 75% in India.


Estimated generic prices ranged from US$0.011 per tablet (glyceryl trinitrate 500 μg) to US$1.447 per tablet (darunavir 800mg), and were heavily skewed towards lower prices in this range. There were 186 estimated generic prices <2.5 cents per tablet, versus only 51 prices above 10 cents per tablet. There was a strong correlation between the estimated generic prices with current global lowest prices for medicines used for HIV, TB and malaria (figure 2). Current lowest UK prices were median 171% above estimated generic price (IQR 4%–1230%), South African prices were median 39% above estimated generic price (IQR −24% to 183%) and Indian prices were median 40% below estimated generic price (IQR −70% to 59%).

In the UK, prices were above estimated generic price in 214 of 277 (77%) comparable cases, and more than three times above estimated price in 47% of cases. LCP were 0.2–387 times the estimated price where multiple suppliers existed, and 3.5–8803 where only one supplier was available. The items with highest LCP to estimated price ratio in the UK were daclatasvir 30mg (8803x), daclatasvir 60mg (5063x), sofosbuvir 400mg (958x), ledipasvir/sofosbuvir 90/400mg FDC (593x) and dexamethasone 1.5mg (387x). Prices were available only in the BNF in 109 cases, only in eMIT in 5 cases. The eMIT price was lower than BNF price in 185 of 209 cases where a price was available from both sources.

In South Africa, 142 of 212 (67%) comparable prices were above the estimated generic price. LCP were 0.2–139 times the estimated price. The items with highest price ratios were ondansetron 24mg (159x), mercapto- purine 50mg (106x), ondansetron 8mg (90x), omeprazole 40mg (32.8x) and haloperidol 2mg (25x).

In India, 118 of 298 (40%) comparable prices were above estimated Indian generic price. The items with highest Indian price to estimated price ratios in were zidovudine 250mg (45x), praziquantel 150mg (15.5x), capetabine 150mg (13.8x), efavirenz/emtricitabine/tenofovir 600/200/300mg FDC (10.7x) and efavirenz 200mg (10.5x). Among prices collected from the database of tenders, LCP to estimated price ratios ranged from 0.04 to 4.3. Among private market prices, lowest Indian price to estimated price ratios ranged from 0.03 to 45.1.

Large price variations were seen for many medicines that are presently under patent protection, or were under protection until recently. In many cases, generic medicines had relatively consistent prices, while for some there was notable variation in prices (figure 3). The therapeutic group with most price variation was antiretroviral medicines.

**DISCUSSION**

This study calculated production costs and estimated generic prices for 148 medicines on the WHO EML, showing that most essential medicines can be manufactured at low cost. Calculation was possible for 148/197 (75%) of the medicines meeting inclusion criteria. Despite most medicines on the EML being off-patent, 214 of 277 comparable prices in the UK, 142 of 212 comparable prices in South Africa and 118 of 298 comparable prices in India exceeded the price that would be expected based on cost of production and a 10% profit margin.

This study used prices of actual, recent sales of API exported from India as the main data input for price estimation. For most medicines, at least 100 export shipments of API were available. Prices of API were generally stable over time, or had a slight decreasing trend (see online supplementary appendix). The estimation formula accounted for capital and operating expenses including labour costs, land and utilities costs, costs of running equipment, costs associated with environmental protection and compliance with cGMP standards, taxation and a profit margin. Validation exercises demonstrated a good ability of the estimation algorithm to predict current lowest global prices for treatments of
HIV, TB and malaria—diseases with large international treatment programmes (Figure 2). Lowest current prices in India were higher than the estimated generic price in 118 cases and lower in 180 cases, and median 40% lower than estimated generic price. This distribution suggests that the algorithm used to calculate minimum costs of production errs on the conservative side (Table 1 and online supplementary appendix).

Most of the high-priced medicines in India were found only in the private market price source, and not the (Tamil Nadu) government tender list, suggesting a lack of availability in public facilities. Over 75% of health expenditure is out-of-pocket in India, of which the majority is spent on medicines.17 While we found Indian prices to be below our estimated generic price in many cases, Indian prices were mostly government tender prices, which are likely to be significantly lower than the private market prices more often experienced by those needing medicines in India. Further analysis of the Indian market would be necessary to determine prices available to various facilities, provinces and patient groups.

Generic competition achieved massive price reductions for antiretroviral drugs in the early 2000s.6 However, non-communicable diseases (NCDs) now represent a larger disease burden than communicable disease, and similar price reductions in medicines for cancer, type 2 diabetes and anticoagulation may be valuable in enabling wider treatment of NCDs in LMICs.18

To protect the right to health, all countries need to ensure affordable access to medicines. This requires avoiding and tackling monopolies, including through legislation, as well as encouraging competition, ensuring a robust supply chain and monitoring shortages and stock-outs. Mechanisms to overcome intellectual property
restrictions in the public interest include governments issuing compulsory licences and originator companies offering voluntary licences. Both are legal mechanisms that allow the use of patented products (including production, import and use) before patent expiry. One successful example of voluntary licensing is embodied in the Medicines Patent Pool, which negotiates licences with originator companies, and then sub-licenses production rights to generic manufacturers in resource-limited countries.

The World Trade Organization’s Agreement on Trade-Related Aspects of Intellectual Property Rights (TRIPS), which sets minimum requirements and limitations on intellectual property laws globally, additionally provides (in article 31bis) for compulsory licences predominantly for export—meaning that compulsory licences can be granted in countries with large manufacturing capacity (such as India) in order to supply other countries where there is a pressing need.

The TRIPS agreement does not prevent individual patients importing medicines for personal use, regardless of patent status, although national legislation must also permit importation for personal use to make this a viable route for access, and this is not uniformly true. This route is being used by patients to access affordable HIV pre-exposure prophylaxis and hepatitis C drugs, with evidence of good clinical outcomes. However, personal-use importation as a method to overcome access barriers requires the patient population to be organised and assertive, have a high level of access to information and financial resources and cannot be considered a feasible large-scale solution.

Comparative clinical and cost-effectiveness is assessed both by national regulators such as the UK’s National Institute for Health and Care Excellence and the WHO Expert Committee reviewing proposed additions to the EML. Cost-effectiveness calculation depends on the efficacy and available price for the drug and the comparator. Where a novel drug is being compared with a drug that is near patent expiry, a cost estimation exercise can help anticipate the generic price of the latter. Recent examples where this analysis could have been applied include novel oral anticoagulants compared with warfarin, dasatinib as a second-line after imatinib for chronic myeloid leukaemia and dolutegravir compared with efavirenz for the treatment of HIV. Generic market entry of comparators should trigger recalculation of cost-effectiveness for the treatment of HIV. Successful countries (such as India) in order to supply other countries where there is a pressing need.

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