demolished apartment with broken windows. How do we comfort the mother who is tucking her five-year-old son with leukemia in bed? She is terrified by the impending shortage of chemotherapy; even the cancer center he receives treatment at has been severely damaged by the blast. What do we tell the mother who has been laid off? She knows that, with losing her job, the entire family lost their healthcare coverage. What do we tell the hundreds of parents who lost their children and the hundreds of children who lost their parents as a result of incompetence, bureaucracy, and political corruption? When will we be angry enough with them and for them, with ourselves and for ourselves?

One of the rivers that run north of Beirut is believed to have carried the blood of the Phoenician god Adonis to the Mediterranean Sea. Legend has it that Adonis was killed by a wild boar in the forest. Like Adonis, Lebanon has been bleeding; COVID-19 and the Beirut explosion have certainly hastened the hemorrhage. But, isn’t the wild boar, in this situation, our inaction for years in the face of social injustice, political corruption, and shortsighted sectarianism? Have we failed to see the humanity in our neighbors until our very basic needs of food, shelter, medicine, physical safety, and education were at stake - just like them? When will we place our basic rights and wellbeing ahead of our sectarian and political affiliations? When will our health, mental and physical, become more important than perpetuating political and financial practices that have failed to deliver for decades?

Some of us, Lebanese people, are a sea, an ocean, and a thousand lakes away, waiting for our loved ones to pick up the phone and tell us they are safe. Some of us are living this reality firsthand every day in the heart of Beirut, opening our eyes every day to destruction and calamity that were avoidable, unjustified, and unnecessary. Will we emerge from this crisis with more debt accumulated, more lives lost, but no lessons learned? Will inequality be made only more extreme as the socioeconomically underprivileged lose their assets and as healthcare out-of-pocket payments increase? Or, will we grasp this opportunity to reform our political structure and our public healthcare system? We worry that we will be satisfied with a partially refurbished version of the current status quo. We nonetheless hope that we, the people of Lebanon living in the country and abroad, will find in the midst of this turmoil an opportunity for true and fundamental rebirth - just like Adonis.

DATA AVAILABILITY STATEMENT

Data sharing is not applicable to this article as no new data were created or analyzed in this study.

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Making hydroxyurea affordable for sickle cell disease in Tanzania is essential (HASTE): How to meet major health needs at a reasonable cost

To the Editor:

More than 300 000 babies with sickle cell disease are born every year worldwide, and more than three-quarters are in Africa.

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Tanzania is among the five countries in the world that rank highest with respect to this burden. In view of this, in 2004 the Muhimbili University of Health and Allied Sciences (MUHAS) and the Muhimbili National Hospital (MNH) established the Muhimbili Sickle Cell (MSC) program, a systematic and comprehensive program aimed to integrate research, training, education and advocacy into the care of SCD in the country.1

Definitive cure of SCD can be achieved only by bone marrow transplantation, a procedure not free of risk that requires ad hoc facilities and is currently out of reach for most patients living in Africa; or by gene therapy that holds great promise, but must be regarded as being still at an experimental stage. Recently new drugs have been approved in high income countries (glutamine, voxelotor, crizanlizumab), but their role in the management of SCD patients at large is still to be defined.2

At the moment, the one medicine that has been universally established as part of the standard of care for SCD is hydroxyurea (HU). In view of its proven efficacy and long-term safety HU was listed by the World Health Organization (WHO) as an essential medicine; and in a formal trial it has been shown to be valuable, as expected, in Africa as it is elsewhere3; however, the sad reality is that the majority of patients in this continent are in fact not receiving this medicine.

In the aim to correct this serious shortcoming in the management of SCD, we first wished to assess the availability and affordability of HU in Dar-es-Salaam, Tanzania. The collaborative partnership between WHO and the international non-governmental organization Health Action International (WHO/HAI) developed a methodology for measuring the availability and affordability of medicines worldwide. The WHO/HAI defines a drug as available when found in a particular pharmacy on the day of data collection; as affordable when the out-of-pocket cost of a one-month course of treatment is less than 1 day salary of the lowest-paid unskilled government worker (LPGW); and as accessible when it is available and affordable.4 In two surveys (carried out in 2016 and in 2018), we found that HU was not available in three out of five pharmacies (Figure 1A). From interviews with pharmacists we learned that HU was not in stock because level of demand was low. The selling price ranged from USD 0.27 per 500 mg capsule in the MNH Hospital Pharmacy, to USD 0.91 in one of the nearby retail pharmacies. Therefore the average cost for a monthly course of treatment (1 g/day for 30 days) was USD 35.16 (range: 16.50-54.60 USD). Since in Tanzania the estimated daily salary of the LPGW is USD 1.4,5 to procure 1 month of HU treatment would require on average 25.11 days, rather than one (Figure 1A).

Thus, by WHO/HAI definition, HU was poorly available, not affordable, and therefore not accessible. We also note that the prices in the pharmacies visited were between 1.2 and 4.1 times higher than the international reference price (IRP: USD 0.22 per capsule in 2018).

Next, we explored pharmacy compounding as a modality that might make HU more accessible to SCD patients in Tanzania. Compounding is the act of preparing, mixing, assembling, packaging, and labelling a drug in response to a practitioner’s prescription drug order or initiative based on the practitioner-patient-pharmacist relationship in the course of professional practice.6 We recalled suggestions made one century ago by Robert P Fischelis: he pointed out that in certain cases health care professionals can use their own know-how efficiently, and he exhorted the “pharmacist to utilize all means at his command in an effort to meet the

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**FIGURE 1** Curbing the price of HU to SCD patients by offering a galenic preparation. A, Availability, prices and affordability of HU in Dar es Salaam. A, Available; NA, Not Available. The price at which HU would have sold, whether available or not on the day of the interview, is expressed in USD (average exchange rate: 1 USD = 2200 Tanzanian shillings - TZS). The Median price ratio (MPR) - the price of HU from the survey divided by the international reference price (IRP) - was used to contextualize local prices at international level. B, Stepwise approach to the compounding process for hydroxyurea at MUHAS
demands of physicians, and at the same time supply the remedies prescribed at as reasonable a price as possible. We took heed from this seminal paper, which was published at about the same time that James B. Henrick reported the first case of SCD. Since the HU patent expired a long time ago, there is no direct legal impediment to the galenic trajectory.

As a first step, we obtained permission from the Tanzania Food and Drug Authority (TFDA) to administer compounded HU to named SCD patients at the Hematology Clinic of MNH. Next we purchased raw materials, that is, HU Active Pharmaceutical Ingredient, microcrystalline cellulose excipient, and hard gelatine capsules, from which, at the School of Pharmacy, MUHAS. Compounding of HU was carried out in compliance with international Pharmacopeia standards, including quality controls by high performance liquid chromatography through the comparison of the purchased HU API with the standard obtained from the European Directorate for the Quality of Medicines & Healthcare (EDQM), (Figure 1B). Finally, HU capsules were dispensed directly to 41 adult patients during clinic visits, providing each patient with a supply of HU sufficient for 1.5 months. We had no reason to carry out a formal trial; however, we have reviewed data from clinic visits on 26 out of 41 patients. Before galenic HU the mean hemoglobin level was 7.10 g/dL (SD 1.55; range 3.2-9.6), and the mean MCV was 88.2 fL (SD 6.36; range 73-102). After galenic HU the mean hemoglobin level was 7.93 g/dL (SD 1.50; range 3.7-10.1), and the mean MCV was 97.5 fL (SD 8.44; range 80-115). In 11 out of 26 patients the Hb increment was greater than 1 g/dL.

In this pilot experiment HU was issued free of charge to the patients, thanks to a donation from an outside source and to the good will of all MUHAS staff concerned. Given that we have established practical feasibility of this approach, we proceeded to work out a budget required to produce a 1-year HU supply for the treatment of 1000 patients - such as may be needed for a cohort of SCD patients at a referral center in Africa. We estimate the total cost at USD 66 613, of which paid personnel would be 40%, raw materials 27%, amortization of equipment 13%, add-on costs (import permits, energy, supplies, etc.) 20%. By this estimate, the final cost per daily dose of 1000 mg would be USD 0.18. Thus, compared with the products from our survey, pharmacy compounded HU from HASTE would be 3.6 times cheaper than commercially available HU from the hospital pharmacy, and 5 to 10 times cheaper than HU from the retail market in Tanzania (Figure 1A).

When a person in Tanzania has an acute illness, for example, from bacterial infection, that person and family will make sacrifices to buy a course of an antibiotic - even if not affordable by the WHO/HAIR definition. The situation is obviously quite different with a chronic, lifelong disorder like SCD. Given a poverty rate in Tanzania of 26.8%, many patients would be unable to purchase HU for a lifetime.

Over the past several decades attempts have been made to negotiate lower prices of drugs through tendering or purchasing agreements, but these have not been very successful. In principle, access to health care can be provided in one of two ways. On the one hand, the so-called vertical programmes provide medicines for specific diseases from international agencies. Unfortunately, SCD, although recognized as a major public health issue in Sub-Saharan Africa, has not yet received from international donors the funding that has accrued to HIV, TB or malaria: we think this anomaly should be corrected. On the other hand, horizontal programs are aimed at improving the capacity of the pharmaceutical system as a whole, and ultimately to provide universal health care (as in Rwanda).

In conclusion, our test setting of pharmacy compounded HU has proven to be economically advantageous with respect to this essential medicine. The TFDA regulatory approval for a named patient approach has been essential; and the tasks of ensuring compounding capacity, quality control management, and trust building among prescribers, pharmacists and patients have passed the test. However, it is clear that, in the long run, pharmacy compounding cannot be a long-term solution. Ultimately, for the SCD patient population to have full benefit from HU will depend on policy and political decisions by individual governments. We hope that this pilot study may encourage drug developers, manufacturers and indigenous entrepreneurs to invest in African countries, especially where necessary local expertise is already present. This is also in line with the recommendations of the UN High-Level Panel on Access to Medicines, recommending strengthening of manufacturing capacity as a measure to improve accessibility to essential medicines.

CONFLICT OF INTEREST

The authors declare no conflict of interest.

AUTHOR CONTRIBUTIONS

E.C. conceived of the study, participated in its design, carried out the survey and the budget impact analysis and wrote the paper. P.T. performed the compounding. E.S. participated in the design of the study. H.L. participated in the planning of analyses and interpretation of results. J.M. participated in the coordination of the study, and in the recruitment and follow up of patients. E.K. conceived of the study, participated in its design and performed the quality control analysis. L.L. conceived of the study, participated in its design, carried out the survey and the budget impact analysis and wrote the paper.

All authors helped with the manuscript: they read and approved the final version.

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DATA AVAILABILITY STATEMENT

The data that support the findings of this study are available from the corresponding author upon reasonable request.

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Overall survival of patients with triple-class refractory multiple myeloma treated with selinexor plus dexamethasone vs standard of care in MAMMOTH

To the Editor:
Multiple myeloma (MM) is the second most common hematologic malignancy worldwide, with an estimated annual incidence of 1.5-6.8 per 100,000 population. Proteasome inhibitors (PI) and immunomodulatory agents (IMiDs) have advanced the management of MM by improving the depth and duration of response. However, when patients become refractory to both drug classes, outcomes are poor, with median overall survival (OS) of 9 months in patients refractory to bortezomib and an IMiD. Anti-CD38 and anti-SLAMF7 antibodies have further advanced the field; despite this, most patients develop MM refractory to available therapies and have a particularly poor prognosis, with survival measured in months. Patients who receive multiple lines of treatment, including the five major agents lenalidomide, pomalidomide, bortezomib, carfilzomib, and daratumumab, are classed as penta-treated. The use of three-drug and four-drug combinations in the early treatment of MM means that there are also an increasing population of patients who become triple-class refractory (TCR; refractory to IMiDs, PIs, monoclonal antibodies [mAbs]) early in treatment. These patients require novel therapies in the face of clonal diversity and multi-refractoriness. There is currently no clear consensus on the optimal treatment sequence for patients with relapsed and/or refractory MM (RRMM), nor a standard of care for patients with penta-treated TCR MM.

As a first-in-class oral selective inhibitor of nuclear export (SINE), selinexor targets exportin 1, which is overexpressed in MM cells. Inhibition of exportin 1 restores nuclear retention and functional activation of MM tumor suppressor proteins, inhibition of NF-kB signaling, and induction of translational suppression of several oncoprotein mRNAs. The efficacy of selinexor in combination with low-dose dexamethasone (sel-dex) was demonstrated in the pivotal Phase 2b Selinexor Treatment of Refractory Myeloma (STORM) Part two study (NCT02336815). In 122 patients with penta-treated/TCR-MM treated with sel-dex, the overall response rate (ORR) was 26% and overall survival (OS) was 8.6 months. Based on these results, selinexor was approved in the US for the treatment of patients with MM refractory to at least two PIs, at least two IMiDs, and an anti-CD38 mAb.