Great South African Molecules: The Case For Mycothiol

Comfort M. Nkambule

Department of Chemistry, Tshwane University of Technology, Pretoria, 0001, South Africa.

Received 13 September 2016, revised 15 March 2017, accepted 17 March 2017.

Abstract

South Africa has one of the oldest chemical societies in the world and has a long history of natural products, synthetic, and medicinal chemistry yet the visibility of molecules discovered or synthesized in South Africa is very low. Is this because South African scientists are incapable of discovering influential and celebrated molecules, or is there inadequate publicity of such discoveries? Perspective profiles on the discovery and scope of research on ‘Great South African Molecules’ should be a good start to redress this state of affairs. One such molecule deserving publicity is the antioxidant mycothiol which is produced by mycobacteria. This is a molecule of interest not only because of its medicinal potential in the fight against tuberculosis, but also from synthetic chemistry on the African continent. This is indeed the story of many molecules from South Africa which have come from other organisms, terrestrial and marine alike, including plants, fungi, parasites, sponges and bacteria. In many cases these natural compounds have been modified into derivatives which have vastly different properties to the original, either by humans or by exposure to the elements. The impact of molecules covers all spheres of life including the key sectors of medicine (active pharmaceutical agents), nutrition (vitamins and supplements), clothing (polymers and fabrics), energy (light and energy-harnessing molecules and fuels) and security (fabrics for personal protection against trauma, chemical and fire damage). The most influential molecules are linked directly to Nobel Prize awards in recognition of their significance (Fig. 1). These prizes have been awarded in the different categories of Chemistry [Chlorophyll (1915), haemin and chlorophyll (1930), vitamin A, vitamin B2 and vitamin C (1937), fullerences (1996)], Physiology or Medicine [Prontosil (1939), vitamin K (1943), penicillin (1945), dichlorodiphenyltrichloroethane or DDT (1948), streptomycin (1952), prostaglandins (1982), artemisinin (2015) and the avermectins (2015)] and Physics [graphene (2010)].

1. What is a Great Molecule?

The diversity of organic chemical entities that have influenced and shaped human civilization is marvellous and includes synthetic molecules designed purely from the imagination of chemists or through serendipitous discoveries, but the majority have come from other organisms, terrestrial and marine alike, including plants, fungi, parasites, sponges and bacteria. In many cases these natural compounds have been modified into derivatives which have vastly different properties to the original, either by humans or by exposure to the elements. The impact of molecules covers all spheres of life including the key sectors of medicine (active pharmaceutical agents), nutrition (vitamins and supplements), clothing (polymers and fabrics), energy (light and energy-harnessing molecules and fuels) and security (fabrics for personal protection against trauma, chemical and fire damage). The most influential molecules are linked directly to Nobel Prize awards in recognition of their significance (Fig. 1). These prizes have been awarded in the different categories of Chemistry [Chlorophyll (1915), haemin and chlorophyll (1930), vitamin A, vitamin B2 and vitamin C (1937), fullerences (1996)], Physiology or Medicine [Prontosil (1939), vitamin K (1943), penicillin (1945), dichlorodiphenyltrichloroethane or DDT (1948), streptomycin (1952), prostaglandins (1982), artemisinin (2015) and the avermectins (2015)] and Physics [graphene (2010)].

A common theme behind the success of these great molecules is the determination and perseverance of the scientists behind their discovery or application to be active advocates of their utility, significance and value to science or commerce. Recently Berhanu Abegaz wrote an insightful and stirring commentary on the challenges of doing chemistry in Africa, but on a positive note he also emphasized the opportunities for endeavours in chemistry on the African continent. This is indeed the story of the future of chemistry in Africa and the South African government must be acknowledged for the positive role they are playing in advancing support for these opportunities. The government has engaged various strategies and concepts to diversify the South African economy away from being intensive on resource supply to focus on the so-called beneficiation and knowledge.
economic pursuits. In 2012 the Department of Science and Technology (DST) released the much anticipated committee report on the review of the National System of Innovation (NSI) wherein innovation was broadly defined as “the capacity to generate, acquire and apply knowledge to advance economic and social purposes”. Such a definition is intended to encompass research and development, plus education and social marketing of knowledge that might inspire the emergence and growth of small and medium enterprises. The objectives of the NSI are that this type of translation of research should be more socially transformative and sustainable in the long term, so what are the challenges or barriers that have hindered these goals?

From a chemist’s perspective it is puzzling why a country that boasts an incredible diversity of flora and fauna does not have...
molecules that feature amongst the world’s greatest molecules? After all, South African chemists have access to the warm Indian and frigid Atlantic Oceans and all the assorted marine compounds they harbour, in addition to the incredible geographical landscape that stretches from the tropical north to the temperate southwest and desert-like interior regions of the Karoo and the Northern Cape. While South African botanists have taken advantage of their proximity to the Cape Floral Kingdom, one of the six floral kingdoms of the world, to carve out a position of prestige in world botany, chemists seem to be lagging behind. Michael Davies-Coleman already made a ‘call to arms’ to all chemists in his article ‘Natural Products Research in South Africa: End of an Era on Land or the Beginning of an Endless Opportunity in the Sea?’ when he cautioned that ‘a perception from current allocation of research funding in SA would suggest that the future of natural products research will increasingly become the responsibility of botanists and ethnobotanists and this new development must be challenged by SA’s organic chemists who have been the custodians of natural products chemistry in this country for over a century’.5

Even the National Development Plan (NDP), the very ambitious strategic vision for the country in 2030, denounces South African teachers, supervisors and research investigators for not being sufficiently innovative. Considering that we have one of the oldest world chemical societies (The South African Chemical Society was founded in 1912), is it that South African scientists have not, or are incapable of, discovering great molecules, or is it that we have not been good champions of the molecules we have discovered? Given the long history of natural products, synthetic and medicinal chemistry in the country, it is surprising that there is such low awareness of any ‘great molecules’ from South Africa.

To start the redress of this state of affairs it seems appropriate to start profiling ‘Great South African Molecules’ in the South African Journal of Chemistry. The objective is to educate society about the possibilities of how fundamental research is an avenue for entrepreneurs within the bio-economy to counter the persistent perception that a shortage of intellectual capital/capacity is to be blamed for failures in the NSI. What may be lacking is targeted marketing, coordination and incubation of the nascent projects whose long-term nurturing may lie beyond the current management of performance in academia. Thus a concerted effort to spin these ideas in small or medium enterprises is a worthwhile pursuit. Therefore, as part of the response to this challenge, in December 2015 I presented a talk at the National Convention of the South African Chemical Institute (SACI 2015) to profile the first of what I consider is a great South African molecule, a pseudo-disaccharide called mycothiol (1) that is produced by mycobacteria, including one of humanity’s greatest nemeses, Mycobacterium tuberculosis (M. tb) (Fig. 3).

Some of the better known South African molecules whose history is well documented include combretastatin (2) which was isolated from the African bush willow of the Eastern Cape.
This compound causes the rapid collapse and necrosis of a tumour’s vascular structure and is thus a biologically active antiproliferative since the endothelial cells of tumours are more susceptible to its activity.6 A phosphate analogue of combretastatin called fosbretabulin (3) is currently under drug development as an anti-cancer drug (Zybrestat) presumably due to its action as a vascular disruptor agent (VDA).7

Among the lesser known compounds which are nonetheless significant in the history of South African molecules is rosmarinine (4), a pyrrolizidine alkaloid whose structure was elucidated by the doyen of South African organic chemistry, Frank Warren, after whom the national meetings of the Organic Chemistry Division of SACI are named.8 Other South African compounds of interest include the extremely toxic monofluoroacetic acid (MFA) (5) which is the first known natural compound to contain fluorine,9 and the guaiane sesquiterpene geigerin (6) which was suspected to be responsible for over 1 million sheep deaths from so-called ‘vomiting disease’ in the Karoo in the 1930s because it is one of the bitter components found in the Geigeria shrub species.10 The real toxic culprit is now thought not to be geigerin, but rather the elusive vermeeric acid precursor of the bis-lactone vermeerin (7).11 Lastly, a discussion of famous South African molecules cannot exclude the mycotoxins even though these are not ‘a molecule’, but rather a diverse group (>300) of fungitoxic compounds including the aflatoxins (8–12) and fumonisins (13, 14). These compounds are produced by fungi that spoil post-harvest maize and wheat grain in storage; many South African chemists working primarily out of the Council for Scientific and Industrial Research (CSIR), including Pieter Steyn, Robert Vleggaar and Fanie van Heerden, were very active in the 1980s on investigations of the isolation and biosynthesis of mycotoxins.12

So which contemporary compounds deserve to join the list of celebrated South African molecules? There will obviously be dispute, contestation and disagreement about what qualifies a molecule to be classified as South African and why it should be considered great if it is not yet an economic powerhouse or indeed a Nobel Prize winner. As to what qualifies a molecule to be South African, this will be easier to resolve if we set the follow-

![Figure 4](http://journals.sabinet.co.za/sajchem/).}
ing two reasonable restrictions, that a compound is or may be considered to be South African if:

- it was **discovered** by scientist(s) working in a South African university, research institute or company; this will be the default definition of a South African researcher, not necessarily immigration or citizenship status.
- **South African researchers** are at the forefront of highlighting the significance of the molecule, even if they were not the initial discoverers.

As to what qualifies a non-Nobel Prize winning or blockbuster molecule to ‘greatness’ will always be a topic of great discussion, but a molecule could be classified as great if it has a scientific impact and is well-known and researched. This may be indicated by the number of publications, citations, h-index, or even that the molecule has a common or trivial name. For example, the h-index is a factor that seeks to gauge the long-term value of a topic based not only on how well cited that topic might be, but also how well cited are those doing the citation. Thus a high h-index value suggests that the topic is highly valued by other scientists whose work is also well regarded by other scientists.

The most important criterion, however, should be the potential and scope for continued and further investigation of the utility of the compound. This point is important to generate local research or commercial interest in the molecule, which might in turn stimulate economic activity or has the capacity of generating translation that addresses the needs of the society. While it might be true that molecules may have some inherent potential for greatness, that potential can only be unlocked by scientists exploring, reporting and exploiting the knowledge embedded in the molecule and its properties. Therefore, it should not just be a matter of whether a molecule exceeds some minimum threshold metrics, but advocates of the molecule must make an argument to justify the research capacity of the molecule. Ideally this case should be subjected to peer and public scrutiny via publication in the South African Journal of Chemistry.

### 2. Is Mycothiol a Great Molecule?

Does mycothiol qualify for consideration as a great molecule? To be transparent in this evaluation, mycothiol is compared to the other well-known South African compounds as shown in Table 1. To further illustrate the reasonableness of the criteria, Table 1 also includes a few international ‘great molecules’ for comparison. While the international molecules all have superior metrics (Table 1, entries 8–12), some of the well-known South African molecules like MFA (Table 1, entry 1), geigerin (Table 1, entry 2) and rosmarinine (Table 1, entry 3) have very low publication and citation metrics. However, fosfretubulin (Table 1, entry 4), combretastatin (Table 1, entry 5) and mycothiol (Table 1, entry 7) have reasonable publication presence to be considered as great molecules, thus there is little dispute that mycothiol passes the first hurdle. Therefore we may proceed to the determination of whether the molecule is South African, before delving into its potential for further research or societal impact.

#### 2.1. Is Mycothiol a South African Molecule?

Mycothiol is a low-molecular-weight thiol (LMWT) weighing only 486.49 g mol⁻¹ with a molecular formula of C₆H₁₂N₂O₂S. It was first discovered, but not characterized, by Robert Fahey and Gerald Newton and their group at the University of California San Diego (UCSD, USA) as the ‘surprisingly previously unknown thiol’ present as the majority thiol in their study of ‘low-molecular-weight thiols in streptomyces and their potential role as antioxidants’. While acknowledging the discovery of U17 (unknown thiol eluting at 17 minutes) as a potential antioxidant, they concluded their report with the prophetic words that ‘the first step in testing the protective role of U17 is to fully elucidate its structure and studies to achieve this goal are in progress’. This paper was accepted on 23 February 1993 (submitted on 26 October 1992) while across both the Atlantic and Pacific oceans two other research groups were independently also hot on the trail of this thiol. Yasuhiro Yamada and his co-workers at Osaka University (Japan) were investigating the biosynthetic pathway towards allosamidin (15) which was the first known chitinase inhibitor produced by streptomyces sp. (Fig. 5). In 1994 they published their report (submitted on 4 February 1994) on the structure of a dimeric disulfide metabolite found in the mycelial extracts of streptomyces whose monomer was determined to be 2-(N-acetyl-L-cysteynil)amido-2-deoxy-α-D-glucopyranosyl-(1→1)-D-myo-inositol (1). At the same time Daniel Steenkamp was working in chemical pathology at the University of Cape Town (South Africa) investigating the ‘thiols of intracellular pathogens’ focussing on the protozoa, especially the *Leishmania* species. After having confirmed that ovthiol A (16) is indeed produced by *Leishmania donovani* which causes visceral leishmaniasis (Fig. 5), Steenkamp decided to ‘isolate and perform a structural analysis’ of any of the then uncharacterised thiols produced by Gram-positive bacteria. The choice of *Mycobacterium bovis* BCG which was moti...
vated in part by its close genetic relationship to pathogenic mycobacteria, namely *Mycobacterium tuberculosis* (*M. tb*), proved fortuitous because the isolated major thiol was structurally determined to be none other than 1-S-steenkamp and Hendrik Spies (University of Stellenbosch, South Africa) who performed the NMR structural elucidation, coined the name mycothiol for this cysteiny1-y pseudo-disaccharide and published their report in 1994 (submitted on 15 April 1994).17 In their article Spies and Steenkamp speculated that the unknown thiol U17 discovered by Newton *et al.* the previous year could be mycothiol and this was later confirmed in a follow-up paper by the UCSD group published in 1995 (submitted on 27 December 1994).18 In 1996 the abbreviation MSH for mycothiol was proposed by Robert Fahey’s group in their seminal paper on the distribution of thiols in microorganisms which confirmed that MSH is the major thiol in the actinomycetes.19

Thus, while Spies and Steenkamp may not have been the first to discover mycothiol, along with Yamada’s group they simultaneously proposed a structure of this thiol and indeed it was they who first proposed the common name for the molecule. Significantly, as the name suggests, Spies and Steenkamp were the first to indelibly link this compound to mycobacteria and by extension to *M. tb*, which subsequently opened up a very vibrant and diverse research enterprise because of the significance of *M. tb*.1 It is therefore justified to claim MSH as a South African molecule.

To further emphasize the critical link to South Africa, Steenkamp and co-workers were the pioneers in investigation and elucidation of the biosynthesis of MSH,20 but given the smaller number of researchers in South Africa working on the topic, faster progress was made elsewhere. The elucidated biosynthetic pathway to date is shown in Scheme 1. This is the composite picture from the efforts of a diverse group of researchers, but the central role of Fahey and his collaborators in the USA and Canada is unmistakable.21 Several informative reviews have been written that highlight the identification, isolation, mechanisms and biochemistry of MSH biosynthesis.22

3. Why is Mycothiol a Molecule of Great Potential?

Since we have established that mycothiol is a South African molecule that surpasses the minimum requirements for greatness, as the advocate for the molecule I will now discuss its virtues as to the potential that exists for further investigation to sustain local research interest in its utility and capacity for the development of commercial outputs as part of research translation to address the needs of the South African society.

3.1. Mycothiol’s Significance to the TB Problem

Quite obviously due its association with *M. tb*, MSH has a significant potential role as a druggable target in the development of new treatments to alleviate the dread and burden of tuberculosis disease (TB). In the contemporary South Africa envisioned in the NDP, the most important aspect to investing in MSH research would be its potential to address a key South African or African societal need which is particularly urgent for a Third World country at the confluence of an increasing burden of First World ‘lifestyle’ diseases like hypertension, cardiovascular and diabetes, with the Third World threats of TB and HIV.23

MSH plays a role in the self-defensive antioxidant response of mycobacteria to an elevated oxidative intracellular environment,24 and also a part in the detoxification mechanism of xenobiotics. Thus MSH is a druggable target in the search and development of drugs against TB, particularly new drugs to counter bacterial resistance against the current therapies.21–e,24–25 The need for new antitubecular drugs, even from the semi-synthetic derivatizations of naturally occurring products has recently been highlighted as part of the urgency in the search for new treatments.26 There is an urgent need to develop more efficacious treatments against a disease that affects over 9 million people per year worldwide and is the second deadliest by a single infectious agent, trailing only HIV with which it most often occurs as a co-infection.27 For a country like South Africa where the public primary healthcare infrastructure is weak the problem is exacerbated by haphazard compliance with the current lengthy TB treatment regimens leading to the development of drug resistance,28 including the emergence of multiple and extensively drug-resistant *M. tb* strains (MDR and XDR, respectively) and even some totally drug-resistant (TDR) forms of the disease.29–31

3.1.1. Mycothiol as an Antioxidant and a Co-factor

MSH chemistry provides a research opportunity for the search of new treatments based on the complex biochemistry of this
molecule. While the enzymes involved in MSH biosynthesis have been identified, many aspects of the role of MSH are incomplete and are still open for further exploration.\textsuperscript{20,21a,21d} These include understanding the specific mechanism of MSH’s role as an antioxidant because it has been established that MSH is more resistant to auto oxidation compared to glutathione and cysteine,\textsuperscript{22} and that MSH-deficient mutants of \textit{M. smegmatis} have been shown to be hypersensitive to oxidative stress.\textsuperscript{23,24} However, MSH deficiency seemed to imbue mutants with resistance to the commonly used first-line bactericidal drugisoniazid, yet it has also been reported that vitamin C is a very effective mycobacterial antioxidant, even against MDR strains.\textsuperscript{25} It therefore seems likely that the development of MSH-inhibiting drugs used in combination with vitamin C could be the basis of a new treatment for TB.

To further highlight its potential as a target in the development of antibiotics against TB, MSH has also been demonstrated to enhance the activity of other antibiotics like rifampin,\textsuperscript{26} or that it of antibiotics against TB, MSH has also been demonstrated to be a co-factor for other detoxification and antioxidant systems.\textsuperscript{33} Given the recent reigniting of interest in MSH biosynthesis it is quite evident that plenty of opportunities are available for local chemists and biochemists to play their role.

3.1.2. Mycothiol Analogues as Potential Biosynthetic Targets

There have been a few reports on the chemical synthesis of MSH and its intermediates going back to the first total synthesis of Steenkamp and co-workers in 2002, up to the most recent efforts to improve glycosylation stereochemistry via endocyclic cleavage anomerization of \( \beta \)-glycosides to \( \alpha \)-glycosides reported by Manabe and Ito.\textsuperscript{34} These reports highlight how MSH is not only a medicinal target, but also that since it is a formidable system there are opportunities for further research by carbohydrate and synthetic chemists. It is reasonable to hypothesize that competitive substrate inhibition of the biosynthesis of MSH could also be a gateway to possible antitubecular drug targets.\textsuperscript{35}

Thus the synthesis of MSH analogues has been a vibrant area of research in the last 22 years. David Gammon and his co-workers at the University of Cape Town (South Africa) have sustained the local interest in this area (Fig. 6).\textsuperscript{36} Several MSH analogues, especially the conjugates of plumbagin\textsuperscript{21a–21d} have shown promising inhibitory activity and good binding specificity to some of the enzymes involved in mycobacterial self-defence. In the spirit of continuing to mine this local resource, our group at the Tshwane University of Technology (Pretoria, South Africa) is also actively involved in the pursuit of mycothiol chemistry as part of our programme on tuberculosis and cancer research.\textsuperscript{37}

Other groups that have been very active in the synthesis and biological evaluation of MSH analogues include Spencer Knapp at Rutgers University (New Jersey, USA),\textsuperscript{38} and Carol Bewley at the National Institutes of Health (NIH) (Maryland, USA).\textsuperscript{39} Two compounds that have shown promising biological activity from the large libraries which have been screened by these groups are shown in Fig. 7. The Knapp-inspired compound 27 was found to be a very good substrate inhibitor of the deacetylase enzymes MshB and Mca,\textsuperscript{40} while compound 28 showed moderate inhibition of MshC, but surprisingly results in a 60 % decline in the production of MSH at only 30 \( \mu \)M and a 99 % loss of viability of non-replicating cultures of \textit{M. tb} at 90 \( \mu \)M.\textsuperscript{21c,40}

Since South Africa has limited resources and a smaller critical mass of researchers, it makes reasonable sense for local chemists to scavenge information about promising lead compounds from the literature for further derivatization and ‘molecular editing’ to enhance the efficacy of these molecules. The experience of the Chinese in their search for an anti-malarial candidate in the 1960s, leading to the discovery of artemisinin, should be a model for South Africa in scouring the literature about what is already known and how we could exploit that resource pool. Project 523 was a Chinese government initiative begun in 1967 to expeditiously find a treatment for chloroquine resistant \textit{Plasmodium falciparum} to aid their North Vietnamese allies who were now battling both the Americans and malaria.\textsuperscript{41} Over 500 scientists and 60 laboratories were recruited for the initiative and by 1969 three treatments were available. The discovery of artemisinin by Tu. The lesson for us in South Africa and Africa in general it should be emphasized, must be that a painstaking evaluation and modification of what is already known may open the door to great treasures just beneath the surface. I think that MSH chemistry possesses such a potential to be exploited.

3.1.3. Mycothiol Analogues as Trojan Carriers

A recent exciting development in mycothiol chemistry is the exploitation of the promiscuous activity of MSH enzymes, espe-
cially the deacetylases MshB and Mca. For example, Teesdale-Spittle and co-workers (New Zealand) reported how benign mycothiol conjugate analogues (29) could be used to smuggle pro-reactive electrophiles \[33\] into the intracellular space where a toxic mercapturic species is generated via a cascade of reactions initiated by the action of MshB or Mca in the deacetylation/detoxification of cysteinyl S-conjugates \[42\] (Scheme 3). In this work the mycothiol analogues are proposed to be progenitors of a new generation of anticancer cytotoxic agents. This is a wonderful expansion of the possible scope of work arising from the deepening exploration of MSH chemistry which bodes well for the growing stature of this molecule.

Similarly, Janata and co-workers (Czech Republic) have shown that the action of MshB and Mca triggers a downstream production of lincosamide antibiotics, with the potential to expedite the synthesis of new libraries of analogues of the natural antibiotics. Compound 34 is a mycothiol S-conjugation electrophilic adduct formed from 4-propyl-L-proline, an octose amino sugar and ergothioneine which reacts with MSH to form an S-conjugate (35) that is ‘detoxified’ by Mca. However, the excreted mercapturic acid 36 can be consumed by the lincomycin biosynthesis pathway to form the antibiotic lincomycin (37) (Scheme 4). In their concluding remarks these authors highlighted how finding that a key detoxification step in one process can be switched seamlessly towards a biosynthetic mode in another process bodes well for more exciting research in this area. I concur and that is why a key objective of this article is to churn the interest of South African chemists to explore more aspects of MSH chemistry!

Lastly, a very interesting approach to the ‘hijacking’ of MSH biosynthetic enzymes comes from Padhye and co-workers (India) who have linked the results on how plumbagin analogues inhibit MshB activity by Gammon et al. to conjugate plumbagin to isoniazid to form a synergistic attack on the bacterial defences. The plumbagin-isoniazid analogue 39 (Fig. 8) showed promising inhibition even at low iron concentrations; this was a significant result because it implied that the mode of action of the new compound bypassed the mechanism of resistance to isoniazid which is inactive at low iron concentration. So while MSH-deficient mutants are more resilient to isoniazid, when MshB is selectively inhibited there is a decline in resistance and an enhancement of the isoniazid activity.

This reaffirms that there are many opportunities to explore for MSH chemistry, especially in the area of combination of natural products and other compounds of known bacteriostatic or bactericidal activity.45

### 3.2. Mycothiol Synthesis as a Synthetic Methodology Challenge

A significant challenge in MSH chemistry which is yet to be fully resolved is a simplified method for Gram-scale synthesis of MSH. Thus the synthesis of MSH has been a ‘proving ground’ for carbohydrate chemists and methodology gurus and junkies alike. Twenty two (22) years after its discovery, there are still new reports on attempts to improve the total synthesis of MSH.\[^{34e,34f}\]

### Figure 7 Substrate inhibitors of mycothiol biosynthesis.

19
20
21a - 21d \((n = 2-5)\)
22 \(X = O\)
23 \(X = S\)
24
25a \(X = NHAc; Z = O\)
25b \(X = NHAc; Z = NH\)
26a \(X = H; Z = O\)
26b \(X = H; Z = NH\)

**Table 1**

<table>
<thead>
<tr>
<th>Compound</th>
<th>IC(_{50}) (µM)</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>27</td>
<td>7µM for MshB</td>
<td>IC(_{50}) = 33 µM for Mca</td>
</tr>
<tr>
<td>28</td>
<td>100 µM for MshC</td>
<td></td>
</tr>
</tbody>
</table>

---

**Figure 6** Substrate analogues of mycothiol synthesized in South African laboratories.

---

**Figure 8** Substrate inhibitors of mycothiol biosynthesis.
The molecule is deceptively simple comprising as it does of only three subunits, a D-\textit{myo}-inositol that is glycosidically linked to a D-glucosamine whose amine is acetylated by N-acetyl cysteine. We recently wrote a comprehensive review on the advances made over the last five years in overcoming some of the synthetic challenges.\textsuperscript{46} The two paramount challenges are firstly, the desymmetrization and regioselective protection of \textit{myo}-inositol and secondly, the stereoselective glycosylation to give a 1,2-\textit{cis}-glycoside. Both of these are non-trivial synthetic challenges that have led to some of the most creative and ingenious solutions.

So far the most efficient method for the regioselective protection of \textit{myo}-inositol for application to the synthesis of C1 glycosylated derivatives is due to the work of Shashidhar (India) and co-workers to make \textit{myo}-inositol-1,3,5-orthoformate \textsuperscript{40} (the so-called Kishi’s triol) which is now also commercially available.\textsuperscript{47} This orthoformate can easily be regioselectively derivatized to give the meso diol \textsuperscript{44}. Shang-Cheng Hung (Taiwan) has shown that this diol can readily and efficiently be desymmetrized by reaction with a chiral ketopinyl ester as the chiral derivatizing agent to give the appropriately protected \textit{myo}-inositol \textsuperscript{45} as shown in Scheme 5.\textsuperscript{48} Other research groups in Australia, the UK and Sweden have also contributed creative solutions to \textit{myo}-inositol resolution and regioselective protection, including...
The glycosyl sulfide donor 53, which is an N-arylsulfonylamide, was alkylated by a suitably protected \textit{myo}-inositol acceptor 56 which was formed \textit{in situ} from the methylthiomethyl ether 55. The phenylsulfenyl chloride activation of the glycosyl sulfide, in the presence of catalytic AgOTf gave the AAA-stereoselective glycosylation in 93% yield. The subsequent deprotection steps of 58 to give $\alpha$-GlcN-Ins (29) are straightforward and therefore recommends this intramolecular aglycon transfer method as one of the preferred routes for the synthesis of 1,2-cis glucosamine glycosides.

Overall then, the challenge of synthesizing MSH has led to the development of new strategies to address a generic problem in glycosylation chemistry.

### 3.3. Mycothiol Monitoring and Bioassays

Finally, the last class of opportunities that arises out of MSH chemistry which I wish to highlight is the need for the development of faster, more accurate and specific bioassays to accelerate the search for new antitubercular agents. As early as 1998 Fahey and co-workers emphasized the need for a sensitive and specific immunoassay for MSH as part of screening for infections by the actinomycetes and \textit{M. tb} in particular. The current method to confirm \textit{M. tb} infection by culturing sputum is uneconomical with respect to duration and also in that it requires sophisticated infrastructure for the safety and expertise to grow cultures of \textit{M. tb}. An ELISA (enzyme-linked immunosorbent assay) microtitre plate test or a readable strip would result in time and expense savings and more importantly widespread testing at the primary health care facilities, before communicable infections spread. This remains an open challenge for South African chemists to develop affordable assays for use in this country.

Meanwhile Julie Leary and co-workers at the University of California Davis (USA) reported on a mass spectrometric method for the analysis (identification and quantification) of MSH that uses minimal sample preparation and is well suited for laboratories doing MSH biosynthetic inhibition studies. This method was checked and validated to be a suitable alternative to the traditional HPLC methods of analysis that require the prior derivatization of MSH with a UV active group; normally bromobimane is the choice derivatizing agent. However, some of the most exciting developments in this area of real-time monitoring of the biosynthetic activity of potential inhibitors have come from Chris Hamilton at the University of East Anglia (UK), Marcy Hernick at Virginia Tech (USA), and Anwar Jardine at the University of Cape Town (South Africa). Interest-
Scheme 6
Mechanistic rationale for nickel catalysed stereoselective glycosylation on N-benzylidene glucosamine donors.

Scheme 7
Complete diastereoselective glycosylation to form D-GlcN-Ins.

Scheme 8
Intramolecular aglycon transfer in 1,2-cis-glycosylation.
ingly, while the Hamilton and Hernick groups specifically set out to demonstrate new assays to monitor the mycothiol disulphide reductase (Mtr) and MshB enzymes respectively, Jardine and co-workers discovered a new assay for MshB by sheer serendipity (Scheme 9). When compound 59 which is a good substrate for MshB was exposed to this enzyme, N-deacetylation ensued followed by a spontaneous Smiles rearrangement to give N-aryl glycosylsulfide 61, which on reaction with Ellman’s reagent gave the anemic disulfide 62 together with thechromophore 63 whose concentration is easy to monitor. The discovery of this alternative assay for MshB activity may accelerate the search for new inhibitors of MshB due to its ease of application and monitoring.

4. Conclusion

This discovery of a new MshB assay by Jardine and his group, who himself is a graduate of the early days of MSH chemistry, is the perfect conclusion to this perspective article because it provides the best message for the South African chemist reader. That message is that MSH chemistry is rich in opportunities provided that we adopt an attitude of painstaking evaluation and modification of what is already known and that we remain open to new challenges, which may open the doors to unexpected treasures just beneath the surface.

I hope that this passionate discussion of all the chemistry around this simple yet intriguing molecule has been sufficient to convince the reader that indeed mycothiol is a great molecule, a Great South African Molecule.

References


