

Government Gazette
Staatskoerant

REPUBLIC OF SOUTH AFRICA
REPUBLIEK VAN SUID AFRIKA

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government printing
Department:
Government Printing Works
REPUBLIC OF SOUTH AFRICA

HIGH ALERT: SCAM WARNING!!!

TO ALL SUPPLIERS AND SERVICE PROVIDERS OF THE GOVERNMENT PRINTING WORKS

It has come to the attention of the *GOVERNMENT PRINTING WORKS* that there are certain unscrupulous companies and individuals who are defrauding unsuspecting businesses disguised as representatives of the *Government Printing Works* (*GPW*).

The scam involves the fraudsters using the letterhead of *GPW* to send out fake tender bids to companies and requests to supply equipment and goods.

Although the contact person's name on the letter may be of an existing official, the contact details on the letter are not the same as the *Government Printing Works*'. When searching on the Internet for the address of the company that has sent the fake tender document, the address does not exist.

The banking details are in a private name and not company name. Government will never ask you to deposit any funds for any business transaction. *GPW* has alerted the relevant law enforcement authorities to investigate this scam to protect legitimate businesses as well as the name of the organisation.

Example of e-mails these fraudsters are using:

PROCUREMENT@GPW-GOV.ORG

Should you suspect that you are a victim of a scam, you must urgently contact the police and inform the *GPW*.

GPW has an official email with the domain as **@gpw.gov.za**

Government e-mails DO NOT have org in their e-mail addresses. All of these fraudsters also use the same or very similar telephone numbers. Although such number with an area code 012 looks like a landline, it is not fixed to any property.

GPW will never send you an e-mail asking you to supply equipment and goods without a purchase/order number. *GPW* does not procure goods for another level of Government. The organisation will not be liable for actions that result in companies or individuals being resultant victims of such a scam.

Government Printing Works gives businesses the opportunity to supply goods and services through RFQ / Tendering process. In order to be eligible to bid to provide goods and services, suppliers must be registered on the National Treasury's Central Supplier Database (CSD). To be registered, they must meet all current legislative requirements (e.g. have a valid tax clearance certificate and be in good standing with the South African Revenue Services - SARS).

The tender process is managed through the Supply Chain Management (SCM) system of the department. SCM is highly regulated to minimise the risk of fraud, and to meet objectives which include value for money, open and effective competition, equitability, accountability, fair dealing, transparency and an ethical approach. Relevant legislation, regulations, policies, guidelines and instructions can be found on the tender's website.

Fake Tenders

National Treasury's CSD has launched the Government Order Scam campaign to combat fraudulent requests for quotes (RFQs). Such fraudulent requests have resulted in innocent companies losing money. We work hard at preventing and fighting fraud, but criminal activity is always a risk.

How tender scams work

There are many types of tender scams. Here are some of the more frequent scenarios:

Fraudsters use what appears to be government department stationery with fictitious logos and contact details to send a fake RFQ to a company to invite it to urgently supply goods. Shortly after the company has submitted its quote, it receives notification that it has won the tender. The company delivers the goods to someone who poses as an official or at a fake site. The Department has no idea of this transaction made in its name. The company is then never paid and suffers a loss.

OR

Fraudsters use what appears to be government department stationery with fictitious logos and contact details to send a fake RFQ to Company A to invite it to urgently supply goods. Typically, the tender specification is so unique that only Company B (a fictitious company created by the fraudster) can supply the goods in question.

Shortly after Company A has submitted its quote it receives notification that it has won the tender. Company A orders the goods and pays a deposit to the fictitious Company B. Once Company B receives the money, it disappears. Company A's money is stolen in the process.

Protect yourself from being scammed

- If you are registered on the supplier databases and you receive a request to tender or quote that seems to be from a government department, contact the department to confirm that the request is legitimate. Do not use the contact details on the tender document as these might be fraudulent.
- Compare tender details with those that appear in the Tender Bulletin, available online at **www.gpwonline.co.za**
- Make sure you familiarise yourself with how government procures goods and services. Visit the tender website for more information on how to tender.
- If you are uncomfortable about the request received, consider visiting the government department and/or the place of delivery and/or the service provider from whom you will be sourcing the goods.
- In the unlikely event that you are asked for a deposit to make a bid, contact the SCM unit of the department in question to ask whether this is in fact correct.

Any incidents of corruption, fraud, theft and misuse of government property in the *Government Printing Works* can be reported to:

Supply Chain Management: Ms. Anna Marie Du Toit, Tel. (012) 748 6292.
Email: **Annamarie.DuToit@gpw.gov.za**

Marketing and Stakeholder Relations: Ms Bonakele Mbhele, at Tel. (012) 748 6193.
Email: **Bonakele.Mbhele@gpw.gov.za**

Security Services: Mr Daniel Legoabe, at tel. (012) 748 6176.
Email: **Daniel.Legoabe@gpw.gov.za**

Closing times for **ORDINARY WEEKLY** *GOVERNMENT GAZETTE* 2024

*The closing time is **15:00** sharp on the following days:*

- **28 December 2023,** Thursday for the issue of Friday **05 January 2024**
- **05 January,** Friday for the issue of Friday **12 January 2024**
- **12 January,** Friday for the issue of Friday **19 January 2024**
- **19 January,** Friday for the issue of Friday **26 January 2024**
- **26 January,** Friday for the issue of Friday **02 February 2024**
- **02 February,** Friday for the issue of Friday **09 February 2024**
- **09 February,** Friday for the issue of Friday **16 February 2024**
- **16 February,** Friday for the issue of Friday **23 February 2024**
- **23 February,** Friday for the issue of Friday **01 March 2024**
- **01 March,** Friday for the issue of Friday **08 March 2024**
- **08 March,** Friday for the issue of Friday **15 March 2024**
- **14 March,** Thursday for the issue of Friday **22 March 2024**
- **20 March,** Wednesday for the issue of Thursday **28 March 2024**
- **27 March,** Wednesday for the issue of Friday **05 April 2024**
- **05 April,** Friday for the issue of Friday **12 April 2024**
- **12 April,** Friday for the issue of Friday **19 April 2024**
- **19 April,** Friday for the issue of Friday **26 April 2024**
- **25 April,** Thursday for the issue of Friday **03 May 2024**
- **03 May,** Friday for the issue of Friday **10 May 2024**
- **10 May,** Friday for the issue of Friday **17 May 2024**
- **17 May,** Friday for the issue of Friday **24 May 2024**
- **24 May,** Friday for the issue of Friday **31 May 2024**
- **31 May ,** Friday for the issue of Friday **07 June 2024**
- **07 June,** Friday for the issue of Friday **14 June 2024**
- **13 June,** Thursday for the issue of Friday **21 June 2024**
- **21 June,** Friday for the issue of Friday **28 June 2024**
- **28 June,** Friday for the issue of Friday **05 July 2024**
- **05 July,** Friday for the issue of Friday **12 July 2024**
- **12 July,** Friday for the issue of Friday **19 July 2024**
- **19 July,** Friday for the issue of Friday **26 July 2024**
- **26 July,** Friday for the issue of Friday **02 August 2024**
- **01 August,** Thursday for the issue of Thursday **08 August 2024**
- **08 August,** Thursday for the issue of Friday **16 August 2024**
- **16 August,** Friday for the issue of Friday **23 August 2024**
- **23 August,** Friday for the issue of Friday **30 August 2024**
- **30 August,** Friday for the issue of Friday **06 September 2024**
- **06 September,** Friday for the issue of Friday **13 September 2024**
- **13 September,** Friday for the issue of Friday **20 September 2024**
- **19 September,** Thursday for the issue of Friday **27 September 2024**
- **27 September,** Friday for the issue of Friday **04 October 2024**
- **04 October,** Friday for the issue of Friday **11 October 2024**
- **11 October,** Friday for the issue of Friday **18 October 2024**
- **18 October,** Friday for the issue of Friday **25 October 2024**
- **25 October,** Friday for the issue of Friday **01 November 2024**
- **01 November,** Friday for the issue of Friday **08 November 2024**
- **08 November,** Friday for the issue of Friday **15 November 2024**
- **15 November,** Friday for the issue of Friday **22 November 2024**
- **22 November,** Friday for the issue of Friday **29 November 2024**
- **29 November,** Friday for the issue of Friday **06 December 2024**
- **06 December,** Friday for the issue of Friday **13 December 2024**
- **12 December,** Thursday for the issue of Friday **20 December 2024**
- **18 December,** Wednesday for the issue of Friday **27 December 2024**

LIST OF TARIFF RATES

FOR PUBLICATION OF NOTICES

COMMENCEMENT: 1 APRIL 2018

NATIONAL AND PROVINCIAL

Notice sizes for National, Provincial & Tender gazettes 1/4, 2/4, 3/4, 4/4 per page. Notices submitted will be charged at R1008.80 per full page, pro-rated based on the above categories.

Pricing for National, Provincial - Variable Priced Notices		
Notice Type	**Page Space**	**New Price (R)**
Ordinary National, Provincial	1/4 - Quarter Page	252.20
Ordinary National, Provincial	2/4 - Half Page	504.40
Ordinary National, Provincial	3/4 - Three Quarter Page	756.60
Ordinary National, Provincial	4/4 - Full Page	1008.80

EXTRA-ORDINARY

All Extra-ordinary National and Provincial gazette notices are non-standard notices and attract a variable price based on the number of pages submitted.

The pricing structure for National and Provincial notices which are submitted as **Extra ordinary submissions** will be charged at R**3026.32** per page.

GOVERNMENT PRINTING WORKS - BUSINESS RULES

The **Government Printing Works** (**GPW**) has established rules for submitting notices in line with its electronic notice processing system, which requires the use of electronic *Adobe* Forms. Please ensure that you adhere to these guidelines when completing and submitting your notice submission.

Closing Times for Acceptance of Notices

1. The *Government Gazette* and *Government Tender Bulletin* are weekly publications that are published on Fridays and the closing time for the acceptance of notices is strictly applied according to the scheduled time for each gazette.

2. Please refer to the Submission Notice Deadline schedule in the table below. This schedule is also published online on the Government Printing works website www.gpwonline.co.za

   All re-submissions will be subject to the standard cut-off times.
   All notices received after the closing time will be rejected.

Government Gazette Type	Publication Frequency	Publication Date	Submission Deadline	Cancellations Deadline
National Gazette	Weekly	Friday	Friday 15h00 for next Friday	Tuesday, 15h00 - 3 working days prior to publication
Regulation Gazette	Weekly	Friday	Friday 15h00 for next Friday	Tuesday, 15h00 - 3 working days prior to publication
Petrol Price Gazette	Monthly	Tuesday before 1st Wednesday of the month	One day before publication	1 working day prior to publication
Road Carrier Permits	Weekly	Friday	Thursday 15h00 for next Friday	3 working days prior to publication
Unclaimed Monies (Justice, Labour or Lawyers)	January / September 2 per year	Last Friday	One week before publication	3 working days prior to publication
Parliament (Acts, White Paper, Green Paper)	As required	Any day of the week	None	3 working days prior to publication
Manuals	Bi- Monthly	2nd and last Thursday of the month	One week before publication	3 working days prior to publication
State of Budget (National Treasury)	Monthly	30th or last Friday of the month	One week before publication	3 working days prior to publication
Extraordinary Gazettes	As required	Any day of the week	*Before 10h00 on publication date*	*Before 10h00 on publication date*
Legal Gazettes A, B and C	Weekly	Friday	One week before publication	Tuesday, 15h00 - 3 working days prior to publication
Tender Bulletin	Weekly	Friday	Friday 15h00 for next Friday	Tuesday, 15h00 - 3 working days prior to publication
Gauteng	Weekly	Wednesday	Two weeks before publication	3 days **after** submission deadline
Eastern Cape	Weekly	Monday	One week before publication	3 working days prior to publication
Northern Cape	Weekly	Monday	One week before publication	3 working days prior to publication
North West	Weekly	Tuesday	One week before publication	3 working days prior to publication
KwaZulu-Natal	Weekly	Thursday	One week before publication	3 working days prior to publication
Limpopo	Weekly	Friday	One week before publication	3 working days prior to publication
Mpumalanga	Weekly	Friday	One week before publication	3 working days prior to publication

GOVERNMENT PRINTING WORKS - BUSINESS RULES

Government Gazette Type	Publication Frequency	Publication Date	Submission Deadline	Cancellations Deadline
Gauteng Liquor License Gazette	Monthly	Wednesday before the First Friday of the month	Two weeks before publication	3 working days **after** submission deadline
Northern Cape Liquor License Gazette	Monthly	First Friday of the month	Two weeks before publication	3 working days **after** submission deadline
National Liquor License Gazette	Monthly	First Friday of the month	Two weeks before publication	3 working days **after** submission deadline
Mpumalanga Liquor License Gazette	Bi-Monthly	Second & Fourth Friday	One week before publication	3 working days prior to publication

Extraordinary Gazettes

3. *Extraordinary Gazettes* can have only one publication date. If multiple publications of an *Extraordinary Gazette* are required, a separate Z95/Z95Prov *Adobe* Forms for each publication date must be submitted.

Notice Submission Process

4. Download the latest *Adobe* form, for the relevant notice to be placed, from the **Government Printing Works** website www.gpwonline.co.za.

5. The *Adobe* form needs to be completed electronically using *Adobe Acrobat* / *Acrobat Reader*. Only electronically completed *Adobe* forms will be accepted. No printed, handwritten and/or scanned *Adobe* forms will be accepted.

6. The completed electronic *Adobe* form has to be submitted via email to submit.egazette@gpw.gov.za. The form needs to be submitted in its original electronic *Adobe* format to enable the system to extract the completed information from the form for placement in the publication.

7. Every notice submitted **must** be accompanied by an official **GPW** quotation. This must be obtained from the *eGazette* Contact Centre.

8. Each notice submission should be sent as a single email. The email **must** contain **all documentation relating to a particular notice submission**.

   8.1. Each of the following documents must be attached to the email as a separate attachment:

      8.1.1. An electronically completed *Adobe* form, specific to the type of notice that is to be placed.

         8.1.1.1. For National *Government Gazette* or *Provincial Gazette* notices, the notices must be accompanied by an electronic Z95 or Z95Prov *Adobe* form

         8.1.1.2. The notice content (body copy) **MUST** be a separate attachment.

      8.1.2. A copy of the official **Government Printing Works** quotation you received for your notice. *(Please see Quotation section below for further details)*

      8.1.3. A valid and legible Proof of Payment / Purchase Order: **Government Printing Works** account customer must include a copy of their Purchase Order**. Non-Government Printing Works** account customer needs to submit the proof of payment for the notice

      8.1.4. Where separate notice content is applicable (Z95, Z95 Prov and TForm 3, it should **also** be attached as a separate attachment. *(Please see the Copy Section below, for the specifications)*.

      8.1.5. Any additional notice information if applicable.

GOVERNMENT PRINTING WORKS - BUSINESS RULES

9. The electronic *Adobe* form will be taken as the primary source for the notice information to be published. Instructions that are on the email body or covering letter that contradicts the notice form content will not be considered. The information submitted on the electronic *Adobe* form will be published as-is.

10. To avoid duplicated publication of the same notice and double billing, Please submit your notice **ONLY ONCE.**

11. Notices brought to **GPW** by "walk-in" customers on electronic media can only be submitted in *Adobe* electronic form format. All "walk-in" customers with notices that are not on electronic *Adobe* forms will be routed to the Contact Centre where they will be assisted to complete the forms in the required format.

12. Should a customer submit a bulk submission of hard copy notices delivered by a messenger on behalf of any organisation e.g. newspaper publisher, the messenger will be referred back to the sender as the submission does not adhere to the submission rules.

QUOTATIONS

13. Quotations are valid until the next tariff change.

    13.1. ***Take note:*** **GPW**'s annual tariff increase takes place on ***1 April*** therefore any quotations issued, accepted and submitted for publication up to ***31 March*** will keep the old tariff. For notices to be published from 1 April, a quotation must be obtained from **GPW** with the new tariffs. Where a tariff increase is implemented during the year, **GPW** endeavours to provide customers with 30 days' notice of such changes.

14. Each quotation has a unique number.

15. Form Content notices must be emailed to the *eGazette* Contact Centre for a quotation.

    15.1. The *Adobe* form supplied is uploaded by the Contact Centre Agent and the system automatically calculates the cost of your notice based on the layout/format of the content supplied.

    15.2. It is critical that these *Adobe* Forms are completed correctly and adhere to the guidelines as stipulated by **GPW**.

16. **APPLICABLE ONLY TO GPW ACCOUNT HOLDERS**:

    16.1. **GPW** Account Customers must provide a valid **GPW** account number to obtain a quotation.

    16.2. Accounts for **GPW** account customers **must** be active with sufficient credit to transact with **GPW** to submit notices.

        16.2.1. If you are unsure about or need to resolve the status of your account, please contact the **GPW** Finance Department prior to submitting your notices. (If the account status is not resolved prior to submission of your notice, the notice will be failed during the process).

17. **APPLICABLE ONLY TO CASH CUSTOMERS:**

    17.1. Cash customers doing **bulk payments** must use a **single email address** in order to use the **same proof of payment** for submitting multiple notices.

18. The responsibility lies with you, the customer, to ensure that the payment made for your notice(s) to be published is sufficient to cover the cost of the notice(s).

19. Each quotation will be associated with one proof of payment / purchase order / cash receipt.

    19.1. This means that **the quotation number can only be used once to make a payment.**

GOVERNMENT PRINTING WORKS - BUSINESS RULES

COPY (SEPARATE NOTICE CONTENT DOCUMENT)

20. Where the copy is part of a separate attachment document for Z95, Z95Prov and TForm03

20.1. Copy of notices must be supplied in a separate document and may not constitute part of any covering letter, purchase order, proof of payment or other attached documents.

The content document should contain only one notice. (You may include the different translations of the same notice in the same document).

20.2. The notice should be set on an A4 page, with margins and fonts set as follows:

Page size = A4 Portrait with page margins: Top = 40mm, LH/RH = 16mm, Bottom = 40mm; Use font size: Arial or Helvetica 10pt with 11pt line spacing;

Page size = A4 Landscape with page margins: Top = 16mm, LH/RH = 40mm, Bottom = 16mm; Use font size: Arial or Helvetica 10pt with 11pt line spacing;

CANCELLATIONS

21. Cancellation of notice submissions are accepted by **GPW** according to the deadlines stated in the table above in point 2. Non-compliance to these deadlines will result in your request being failed. Please pay special attention to the different deadlines for each gazette. Please note that any notices cancelled after the cancellation deadline will be published and charged at full cost.

22. Requests for cancellation must be sent by the original sender of the notice and must accompanied by the relevant notice reference number (N-) in the email body.

AMENDMENTS TO NOTICES

23. With effect from 01 October 2015, **GPW** will not longer accept amendments to notices. The cancellation process will need to be followed according to the deadline and a new notice submitted thereafter for the next available publication date.

REJECTIONS

24. All notices not meeting the submission rules will be rejected to the customer to be corrected and resubmitted. Assistance will be available through the Contact Centre should help be required when completing the forms. (012-748 6200 or email info.egazette@gpw.gov.za). Reasons for rejections include the following:

24.1. Incorrectly completed forms and notices submitted in the wrong format, will be rejected.

24.2. Any notice submissions not on the correct *Adobe* electronic form, will be rejected.

24.3. Any notice submissions not accompanied by the proof of payment / purchase order will be rejected and the notice will not be processed.

24.4. Any submissions or re-submissions that miss the submission cut-off times will be rejected to the customer. The Notice needs to be re-submitted with a new publication date.

GOVERNMENT PRINTING WORKS - BUSINESS RULES

APPROVAL OF NOTICES

25. Any notices other than legal notices are subject to the approval of the Government Printer, who may refuse acceptance or further publication of any notice.

26. No amendments will be accepted in respect to separate notice content that was sent with a Z95 or Z95Prov notice submissions. The copy of notice in layout format (previously known as proof-out) is only provided where requested, for Advertiser to see the notice in final Gazette layout. Should they find that the information submitted was incorrect, they should request for a notice cancellation and resubmit the corrected notice, subject to standard submission deadlines. The cancellation is also subject to the stages in the publishing process, i.e. If cancellation is received when production (printing process) has commenced, then the notice cannot be cancelled.

GOVERNMENT PRINTER INDEMNIFIED AGAINST LIABILITY

27. The Government Printer will assume no liability in respect of—

27.1. any delay in the publication of a notice or publication of such notice on any date other than that stipulated by the advertiser;

27.2. erroneous classification of a notice, or the placement of such notice in any section or under any heading other than the section or heading stipulated by the advertiser;

27.3. any editing, revision, omission, typographical errors or errors resulting from faint or indistinct copy.

LIABILITY OF ADVERTISER

28. Advertisers will be held liable for any compensation and costs arising from any action which may be instituted against the Government Printer in consequence of the publication of any notice.

CUSTOMER INQUIRIES

Many of our customers request immediate feedback/confirmation of notice placement in the gazette from our Contact Centre once they have submitted their notice – While **GPW** deems it one of their highest priorities and responsibilities to provide customers with this requested feedback and the best service at all times, we are only able to do so once we have started processing your notice submission.
GPW has a 2-working day turnaround time for processing notices received according to the business rules and deadline submissions.
Please keep this in mind when making inquiries about your notice submission at the Contact Centre.

29. Requests for information, quotations and inquiries must be sent to the Contact Centre ONLY.

30. Requests for Quotations (RFQs) should be received by the Contact Centre at least **2 working days** before the submission deadline for that specific publication.

GOVERNMENT PRINTING WORKS - BUSINESS RULES

Payment of cost

31. The Request for Quotation for placement of the notice should be sent to the Gazette Contact Centre as indicated above, prior to submission of notice for advertising.

32. Payment should then be made, or Purchase Order prepared based on the received quotation, prior to the submission of the notice for advertising as these documents i.e. proof of payment or Purchase order will be required as part of the notice submission, as indicated earlier.

33. Every proof of payment must have a valid **GPW** quotation number as a reference on the proof of payment document.

34. Where there is any doubt about the cost of publication of a notice, and in the case of copy, an enquiry, accompanied by the relevant copy, should be addressed to the Gazette Contact Centre, **Government Printing Works**, Private Bag X85, Pretoria, 0001 email: info.egazette@gpw.gov.za before publication.

35. Overpayment resulting from miscalculation on the part of the advertiser of the cost of publication of a notice will not be refunded, unless the advertiser furnishes adequate reasons why such miscalculation occurred. In the event of underpayments, the difference will be recovered from the advertiser, and future notice(s) will not be published until such time as the full cost of such publication has been duly paid in cash or electronic funds transfer into the **Government Printing Works** banking account.

36. In the event of a notice being cancelled, a refund will be made only if no cost regarding the placing of the notice has been incurred by the **Government Printing Works**.

37. The **Government Printing Works** reserves the right to levy an additional charge in cases where notices, the cost of which has been calculated in accordance with the List of Fixed Tariff Rates, are subsequently found to be excessively lengthy or to contain overmuch or complicated tabulation.

Proof of publication

38. Copies of any of the *Government Gazette* or *Provincial Gazette* can be downloaded from the **Government Printing Works** website www.gpwonline.co.za free of charge, should a proof of publication be required.

39. Printed copies may be ordered from the Publications department at the ruling price. The **Government Printing Works** will assume no liability for any failure to post or for any delay in despatching of such *Government Gazette*(s)

GOVERNMENT PRINTING WORKS CONTACT INFORMATION

Physical Address:	**Postal Address:**	**GPW Banking Details:**
Government Printing Works	Private Bag X85	**Bank:** ABSA Bosman Street
149 Bosman Street	Pretoria	**Account No.:** 405 7114 016
Pretoria	0001	**Branch Code:** 632-005

For Gazette and Notice submissions: Gazette Submissions:	**E-mail:** submit.egazette@gpw.gov.za
For queries and quotations, contact: Gazette Contact Centre:	**E-mail:** info.egazette@gpw.gov.za
	Tel: 012-748 6200
Contact person for subscribers: Mrs M. Toka:	**E-mail:** subscriptions@gpw.gov.za
	Tel: 012-748-6066 / 6060 / 6058
	Fax: 012-323-9574

GOVERNMENT NOTICES • GOEWERMENTSKENNISGEWINGS

DEPARTMENT OF AGRICULTURE, LAND REFORM AND RURAL DEVELOPMENT

NO. 4478 **8 March 2024**

AMENDING GOVERNMENT NOTICE NO. 37334 OF 2014 PUBLISHED IN THE GOVERNMENT GAZETTE NO. 37334 DATED 21 FEBRUARY 2014 INTERMS OF SECTION 11(A) OF THE RESTITUTION OF LAND RIGHTS ACT, ACT NO. 22 OF 1994, AS AMENDED.

Notice is hereby given in terms of Section 11A (4) of the Restitution of Land Rights Act, No. 22 of 1994, as amended, that the Commissioner for Restitution of Land Rights is amending the said Gazette Notice to reflect the correct affected properties and total affected extent for the land claim lodged by the late Graham Nkosi ID No 470325 5226 081 registered as KRP 1067. The correct properties to be published is the Remaining Extent of the farm (Portion 0) and Portion 2 of the farm Belfast 492 IT situated at Mkhondo Local Municipality at Gert Sibande District Municipality, therefore this seeks to clarify that.

CURRENT PARTICULARS OF THE PROPERTY
BELFAST 492 IT

Description of property	Owner of Property	Title Deed Number	Extent of Property	Bonds	Bond Holder	Other Endorsements
The Remaining extent of the farm 492 IT	Mondi South Africa PTY [1967013038O7]	T89352/1999	327.7676 ha (116.6847 ha) extent claimed	None	None	I-740/2021C
Portion 2	Mondi South Africa PTY [1967013038O7]	T89352/1999	327.7691 ha (65.1332 ha) extent claimed	None	None	I/740/2021C K15/1961L

TOTAL EXTENT CLAIMED 181.7819 ha

Notice is hereby given in terms of Section 11A (4) of the Restitution of Land Rights Act, No. 22 of 1994, as amended, that the Commissioner for Restitution of Land Rights is amending the said Gazette Notice to reflect the correct affected properties and total affected extent for the land claim lodged by the late Graham Nkosi ID No 470325 5226 081 registered as KRP 1067. The correct properties to be published is the Remaining Extent of the farm (Portion 0) and Portion 2 of the farm Belfast 492 IT situated at Mkhondo Local Municipality at Gert Sibande District Municipality, therefore this seeks to clarify that.

The Regional Land Claims Commissioner, Mpumalanga Province will investigate all the claims in terms of the provisions of the Act, any party interested in the above mentioned property is hereby invited to submit within **30 [thirty days]** from the date of publication of this notice to submit any comments, or further information to:

Commission on Restitution of Land Rights
30 Samora Machel Drive
Nelspruit
1200

Tel No. 013 756 6000



MR. L.H. MAPHUTHA
REGIONAL LAND CLAIMS COMMISSIONER
MPUMALANGA PROVINCE
DATE: 2024/02/28

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DEPARTMENT OF AGRICULTURE, LAND REFORM AND RURAL DEVELOPMENT

NO. 4479 **8 March 2024**

GENERAL NOTICE IN TERMS OF THE RESTITUTION OF LAND RIGHTS ACT, NO.22 OF 1994, AS AMENDED

NOTICE IS HEREBY GIVEN IN TERMS OF SECTION 11(1) OF THE RESTITUTION OF LAND RIGHTS ACT, ACT NO.22 OF 1994 AS AMENDED, THAT CLAIM FOR RESTITUTION OF LAND RIGHTS ON.

Reference No : KRN6/2/2/C/3/0/3

Claimant : Mr. Daniel Johannes Botes

Property Description : Plot No.35, division of Rodenbeck "B" No.294 situated in the Registration Division of Bloemfontein, Mangaung Metropolitan Municipality, Motheo District Municipality of the Free State Province.

Extent of Land : 4,5896 ha

Current Title Deed : T23509/2001

Date Submitted : 03 May 1997

Current Owner : Mangaung Local Municipality

Current Land Use : Residential purpose

Has been submitted to the Regional Land Claims Commissioner for the Free State Province and that the Commission on Restitution of Land Rights will further investigate the claims in terms of the provision of the Act, as amended in due course.

Any party who has an interest in the abovementioned land claim is hereby invited to submit, within 30 days from the date of the publication of this Notice, any comments/information to:

The Regional Land Claims Commissioner
Free State Province
P O Box 4376
Bloemfontein,
9300
Tel: 051 430 0444

Lebjane Maphutha
Regional Land Claims Commissioner
Date: 14/11/2023

DEPARTMENT OF AGRICULTURE, LAND REFORM AND RURAL DEVELOPMENT

NO. 4480 **8 March 2024**

GENERAL NOTICE IN TERMS OF THE RESTITUTION OF LAND RIGHTS ACT, 22 OF 1994, AS AMENDED

Notice is hereby given in terms of Section 11(1) of the Restitution of Land Rights Act No. 22 of 1994 (as amended), that claims for the restitution of land rights on:

Reference No.	:	N6/2/2/0/C/413/0/0/05
Claimant	:	Mr Mmatli Bethuel Motaung
Description of Properties	:	The property under claim is described below as - The Farm Holhoek 1842, district Bethlehem Free State Province;
Date submitted	:	27 July 1997; 10 November 1997 and unknown date
Current Land Use	:	Hospitality Industry and Ecotourism
Total Hectorage	:	1001,0121 hectares

The Regional Land Claims Commissioner
Free State Province
P O Box 4376
Bloemfontein,
9300
Tel: 051 430 0444

Lebjane Maphutha
Regional Land Claims Commissioner
Date: 2023/09/20

DEPARTMENT OF AGRICULTURE, LAND REFORM AND RURAL DEVELOPMENT

NO. 4481 **8 March 2024**

GENERAL NOTICE IN TERMS OF THE RESTITUTION OF LAND RIGHTS ACT, 1994 (ACT 22 OF 1994) AS AMENDED

Notice is hereby given in terms of section 11(1) (c) of the Restitution of Land Rights Act, 1994 as amended) that a claim has been lodged for restitution of land rights on:

REF NO	CLAIMANT	PROPERTY DESCRIPTION & CLAIMED EXTENT	CURRENT LANDOWNERS	BONDS / NO BONDS	DEED OF TRANSFER	INTERESTED PARTIES
Z 0337 (KRP 7546)	Mr. Jim David Buta	Portion 9 of farm Rietfontein 470 JR	Republic of South Africa	None	T8224/1985 T96120/2006	Land Claimant, the current landowners, and the Ekangala District Municipality.
		Barnetein farm 619 JR	National Government of the Republic of South Africa	None	T64719/1988 T11790/2012	
		Claimed Extent = 9.5925 hectares				

Take further notice that the Commission on Restitution of Land Rights will investigate the claim in terms of the provisions of Rule 5 of the Rules Regarding Procedure of Commission Established in terms of section 16 of Restitution of Land Rights Act as amended. Any interested party on the claim is hereby invited to submit, representations in terms of section 11A of the Restitution of Land Rights Act 22 of 1994 as amended within 90 (ninety) working days from the publication date of this notice, any comments/information may be send to:

Chief Directorate: Land Restitution Support Gauteng Province
Private Bag X03
ARCADIA
0007
Tel: (012) 310-6500
Fax: (012) 324-5812

MR. L.H MAPHUTHA
REGIONAL LAND CLAIMS COMMISSIONER
DATE: 20-4/02/26

DEPARTMENT OF AGRICULTURE, LAND REFORM AND RURAL DEVELOPMENT

NO. 4482 **8 March 2024**

FERTILIZERS, FARM FEEDS, AGRICULTURAL REMEDIES AND STOCK REMEDIES ACT, 1947 (ACT NO. 36 OF 1947)

REGULATIONS RELATING TO THE TARIFFS FOR THE REGISTRATION OF FERTILIZERS, FARM FEEDS, AGRICULTURAL REMEDIES, STOCK REMEDIES, STERILIZING PLANTS AND PEST CONTROL OPERATORS, APPEALS AND IMPORTS: AMENDMENT

The Minister for Agriculture, Land Reform and Rural Development, has under Section 23 of the Fertilizers, Farm Feeds, Agricultural Remedies and Stock Remedies Act, 1947 (Act No. 36 of 1947), made the regulations in the Schedule.

SCHEDULE

Definition

1. In this Schedule "the Regulations" means the regulations published by Government notice No. R. 1449 of 1 July 1983, as amended by government Notices Nos. R. 96 of January 1984, R. 2055 of 14 September 1984, R. 1053 of 3 June 1988, R. 1242 of 9 June 1990, r. 1409 of 6 August 1993, R.1592 of 30 September 1996, r. 1017 of 14 August 1998, R. 216 of 10 March 2000, R. 964 of 5 October 2001, R. 1096 of 30 August 2002, R. 1475 of 17 October 2003, R. 3448 of 15 April 2005, R.1139 of 2 December 2005, R. 225 of 17 March 2006, R935 of 22 September 2006, R. 956 of 29 September 2006, R. 1086 of 3 November 2006, R. 1087 of 3 November 2006, R. 250 of 23 March 2007, R. 483 of 8 June 2007, R.755 of 18 July 2008,R.112 of 13 February 2009, R.72 of 12 February 2010 and R.97 of 18 February 2011, R.75 of 8 February 2013, R259 of 5 April 2013, R 207 of 1 April 2014, R 285 of 31 March 2015,
R 372 of 29 March 2016, No R 310 of 31 March 2017, No R 394 of 28 March 2018,
No R 471 of 29 March 2019, No.R.395 of 27 March 2020, No. R. 265 of 26 March 2021
No R 1978 of 1 April 2022, No R 1752 of 21 April 2023

Substitution of Table 1 of the Regulations R 1752 of 21 April 2023

2. The Regulations are hereby amended by the substitution for Table 1 of the following table:

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Table 1
"FEES PAYABLE"

TARIFFS, RATES AND SCALES FOR SERVICES, GOODS AND SUPPLIES PROVIDED BY THE DEPARTMENT OF AGRICULTURE, LAND REFORM AND RURAL DEVELOPMENT

NATURE OF SERVICE, GOODS OR SUPPLIES PROVIDED	TARIFF APPLICABLE FROM 1 APRIL 2023	TARIFF APPLICABLE FROM 1 APRIL 2024	REMARKS / EXPLANATION
AGRICULTURE INPUTS CONTROL			
FERTILIZERS, FARM FEEDS, AGRICULTURAL REMEDIES AND STOCK REMEDIES ACT, 1947 (Act No. 36 of 1947)			
1 REGISTRATIONS: Application for registration of Farm Feed and Pet Food including application on lapsed registration, parallel and daughter registrations.			
Kinds of Farm Feeds and Pet Food (Groups and kinds)			
1.1 Importers for own use			
1.1 (a) Raw material of plant, animal origin and their by-products including blended raw materials of plant and animal origin and their by-products	R 1 809.00 per application/product	R 1 923.00 per application/product	Registration is valid for 3 years
1.1 (b) **Feed Additives**			
(i) Technological additives	R 3 292,00 per application/product	R 3 499.00 per application/product	Registration is valid for 3 years
(ii) Sensory additives	R 1 809,00 per application/product	R 1 923.00 per application/product	Registration is valid for 3 years.
(iii) Zootechnical additives	R 3 292,00 per application/product	R 3 499.00 per application/product	
(iv) Nutritional additives	R 1 809,00 per application/product	R 1 923.00 per application/product	

NATURE OF SERVICE, GOODS OR SUPPLIES PROVIDED	TARIFF APPLICABLE FROM 1 APRIL 2023	TARIFF APPLICABLE FROM 1 APRIL 2024	REMARKS / EXPLANATION
(v) Nutritional Additives Premixes	R 3 292,00 per application/product	R 3 499.00 per application/product	
(vi) Livestock feeds (complete concentrate and supplements)	R 3 292,00 per application/product	R 3 499.00 per application/product	
1.1 (c) **Pet foods**			
(i) Complete dog and cat foods	R 3 292,00 per application/product	R 3 499.00 per application/product	Registration is valid for 3 years.
(ii) Complete miscellaneous pet foods	R 874,00 per application/product	R 929.00 per application/product	Registration is valid for 3 years.
(iii) Complementary pet foods	R 1 316,00 per application/product	R 1 399.00 per application/product	Registration is valid for 3 years.
(iv) Pet-Neutraceuticals	R 3 292. 00 per application/product	R 3 499.00 per application/product	Registration is valid for 3 years.
(v) Herbal supplements (including horses)	R 874,00 per application/product	R 929.00 per application/product	
(vi) Nutritional supplements (including horses)	R 1 809.00 per application/product	R 1 923.00 per application/product	
(vii) Seed and Grain mixtures	R 874.00 per application/product	R 929.00 per application/product	
1.2 **Importers for retail / Local trader/ Distributor/ Seller/ Manufacture for retail/ Parallel registration/ Daughter registration**			
1.2 (a) Raw material of plant, animal origin and their by-products including blended raw materials of plant and animal origin and their by-products	R 2 965,00 per application/product	R 3 152.00 per application/product	

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NATURE OF SERVICE, GOODS OR SUPPLIES PROVIDED	TARIFF APPLICABLE FROM 1 APRIL 2023	TARIFF APPLICABLE FROM 1 APRIL 2024	REMARKS / EXPLANATION
1.2 (b) **Feed Additives**			
(i) Technological additives	R 5 813,00 per application/product	R 6 179.00 per application/product	
(ii) Sensory additives	R 3 456,00 per application/product	R 3 674.00 per application/product	
(iii) Zootechnical additives	R 5 813.00 per application/product	R 6 179.00 per application/product	
(iv) Nutritional additives	R 3 456,00 per application/product	R 3 674.00 per application/product	
(v) Nutritional additives premixes	R 5 813,00 per application/product	R 6 179.00 per application/product	
(vi) Livestock feeds (complete concentrate and supplements)	R 5 813,00 per application/product	R 6 179.00 per application/product	

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NATURE OF SERVICE, GOODS OR SUPPLIES PROVIDED	TARIFF APPLICABLE FROM 1 APRIL 2023	TARIFF APPLICABLE FROM 1 APRIL 2024	REMARKS / EXPLANATION
1.2 (c) **Pet food**			
(i) Complete dog and cat foods	R 5 813.00 per application/product	R 6 179.00 per application/product	
(ii) Complete miscellaneous pet foods	R 1 809.00 per application/product	R 1 923.00 per application/product	
(iii) Complementary pet foods	R 1 809.00 per application/product	R 1 923.00 per application/product	
(iv) Pet-Neutraceuticals	R 5 813.00 per application/product	R 6 179.00 per application/product	
(v) Herbal supplements (including horses)	R 1 809.00 per application/product	R 1 923.00 per application/product	
(vi) Nutritional supplements (including horses)	R 3 456.00 per application/product	R 3 674.00 per application/product	
(vii) Seed and Grain mixtures	R 1 809.00 per application/product	R 1 923.00 per application/product	
2. RENEWALS: **Application for the renewal of the registration of a Farm Feed and Pet Food** **(Groups and kinds of farm feeds and pet food)**			
2.1 **Importers for own use**			
2.1 (a) Raw material of plant, animal origin and their by-products including blended raw materials of plant and animal origin and their by-products	R 687.00 per application/product	R 730.00 per application/product	

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NATURE OF SERVICE, GOODS OR SUPPLIES PROVIDED	TARIFF APPLICABLE FROM 1 APRIL 2023	TARIFF APPLICABLE FROM 1 APRIL 2024	REMARKS / EXPLANATION
2.1 (b) **Feed Additives**			
(i) Technological additives	R 687,00 per application/product	R 730.00 per application/product	
(ii) Sensory additives	R 687,00 per application/product	R 730.00 per application/product	
(ii) Zootechnical additives	R 687,00 per application/product	R 730.00 per application/product	
(iv) Nutritional additives	R 687,00 per application/product	R 730.00 per application/product	
(v) Nutritional Additives Premixes	R 687,00 per application/product	R 730.00 per application/product	
(vi) Livestock feeds (complete, concentrate and supplements)	R 687,00 per application/product	R 730.00 per application/product	
2.1 (c) **Pet Food**			
(i) Complete dog and cat foods	R 687,00 per application/product	R 730.00 per application/product	-do-
(ii) Complete miscellaneous pet foods	R 687,00 per application/product	R 730.00 per application/product	-do-
(iii) Complementary pet foods	R 687,00 per application/product	R 730.00 per application/product	-do-
(iv) Pet-Neutraceuticals	R 687,00 per application/product	R 730.00 per application/product	-do-
(v) Herbal supplements (including horses)	R 687,00 per application/product	R 730.00 per application/product	
(vi) Nutritional supplements (including horses)	R 687.00 per application/product	R 730.00 per application/product	
(vii) Seed and Grain mixture	R 687,00 per application/product	R 730.00 per application/product	

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NATURE OF SERVICE, GOODS OR SUPPLIES PROVIDED	TARIFF APPLICABLE FROM 1 APRIL 2023	TARIFF APPLICABLE FROM 1 APRIL 2024	REMARKS / EXPLANATION
2.2. **Importers for retail/ Local trader/Distributor/ Seller/ Manufacture for retail/ Parallel registration/ Daughter registration**			
2.2 (a) Raw material of plant, animal origin and their by-products including blended raw material of plant and animal origin and their by-products	R 1 538.00 per application/product	R 1 635.00 per application/product	
2.2 (b) **Feed Additives**			
(i) Technological additives	R 2 787,00 per application/product	R 2 963.00 per application/product	
(ii) Sensory additives	R 1 657,00 per application/product	R 1 761.00 per application/product	
(iii) Zootechnical additives	R 2 787,00 per application/product	R 2 963.00 per application/product	
(iv) Nutritional Additives	R 1 657,00 per application/product	R 1 761.00 per application/product	
(v) Nutritional Additives Premixes	R 2 787,00 per application/product	R 2 963.00 per application/product	
(vi) Livestock feeds (complete, concentrate and supplements)	R 2 787,00 per application/product	R 2 963.00 per application/product	

NATURE OF SERVICE, GOODS OR SUPPLIES PROVIDED	TARIFF APPLICABLE FROM 1 APRIL 2023	TARIFF APPLICABLE FROM 1 APRIL 2024	REMARKS / EXPLANATION
2.2 (c) **Pet Food**			
(i) Complete dog and cat foods	R 2 787.00 per application/product	R 2 963.00 per application/product	
(ii) Complete miscellaneous pet foods	R 869.00 per application/product	R 924.00 per application/product	
(iii) Complementary pet foods	R 869.00 per application/product	R 924.00 per application/product	
(iv) Pet-Neutraceuticals	R 2 787.00 per application/product	R 2 963.00 per application/product	
(v) Herbal supplements including horses.	R 869.00 per application/product	R 924.00 per application/product	
(vi) Nutritional supplements (including horses)	R 1 657.00 per application/product	R 1 761.00 per application/product	
(vii) Seed and Grain mixture	R 869.00 per application/product	R 924.00 per application/product	

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3 LATE RENEWAL APPLICATIONS: **Payment additional to that mentioned in 2.1 (a)-(c) and 2.2 (a)-(c) above, in case of a late application for the renewal of animal feed and pet food.**			
3.1. **Importers for own use**			
(i) Raw material of plant, animal origin and their by-products including blended raw materials of plant and animal origin and their by product.	R 527.00 per application/product	R 560.00 per application/product	
(ii) Groups and kinds of farm feeds and pet food mentioned 2.1(b & c)	R 527.00 per application/product	R 560.00 per application/product	
(Groups and kinds of farm feeds and pet food for):			
3.2. **Importers for retail/ Local trader/ distributor/ Seller/ Manufacture for retail/Parallel registration/ Daughter registration**			
3.2. (a) Raw material of plant, animal origin and their by-products including blended raw materials of plant and animal origin and their by - products	R 835.00 per application/product	R 888.00 per application/product	

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NATURE OF SERVICE, GOODS OR SUPPLIES PROVIDED	TARIFF APPLICABLE FROM 1 APRIL 2023	TARIFF APPLICABLE FROM 1 APRIL 2024	REMARKS / EXPLANATION
3.2 (b) **Livestock Feed Additives**			
(i) Technological additives	R 1 640.00 per application/product	R 1 743.00 per application/product	
(ii) Sensory additives	R 976.00 per application/product	R 1 037.00 per application/product	
(iii) Zootechnical additives	R 1 640.00 per application/product	R 1 743.00 per application/product	
(iv) Nutritional Additives	R 976.00 per application/product	R 1 037.00 per application/product	
(v) Nutritional Additives Premixes	R 1 640.00 per application/product	R 1 743.00 per application/product	
(vi) Livestock feeds (complete concentrate and supplements	R 1 640.00 per application/product	R 1 743.00 per application/product	
3.2 (c) **Pet Food**			
(i) Complete dog and cat foods	R 1 640.00 per application/product	R 1 743.00 per application/product	
(ii) Complete miscellaneous pet food	R 510.00 per application/per product	R 542.00 per application/per product	
(iii) Complementary pet food	R 510.00 per application/product	R 542.00 per application/product	
(iv) Pet Neutraceuticals	R 1 640.00 per application/product	R 1 743.00 per application/product	
(v) Herbal supplements (including horses)	R 510.00 per application/product	R 542.00 per application/product	
(vi) Nutritional supplements (including horses)	R 976.00 per application/product	R 1 037.00 per application/product	
(vii) Seed and Grain mixture	R 510.00 per application/product	R 542.00 per application/product	
4. Any other minor amendment to the registered product requested by the registration holder, e.g. transfer of registration, company and	R 1 316.00 per application/product	R 1 399.00 per application/product	

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NATURE OF SERVICE, GOODS OR SUPPLIES PROVIDED	TARIFF APPLICABLE FROM 1 APRIL 2023	TARIFF APPLICABLE FROM 1 APRIL 2024	REMARKS / EXPLANATION
product name change, label amendment, additional manufacturer etc.			
4.1 **Payment for information and other services**.			
(i) Import Permit	R 756.00 per product	R 804.00 per product	
(ii) Advertisement approval	R 874.00 per request	R 929.00 per request	
(iii) Free Sale certificate	R 72.00 per certificate	R 77.00 per certificate	
(iv) Re-print of the certificate	R 72.00 per certificate	R 77.00 per certificate	
(v) Other documents	R 131 per request plus, R1.00 per page	R 139 per request plus, R1.00 per page	
(vi) Guideline documents	#	#	Guidelines are available for download on DALRRD website for free.
(vii) An appeal under section 6 of Act no 36 of 1947	R 7 458.00 per application/product	R 7 928.00 per application/product	
5. Application for sterilization plant			
5.1 Application for the registration of the Sterilization plant.	R 5 462.00 per application/product	R 5 806.00 per application/product	
5.2 Application for the renewal of the sterilization plant and rendering plant	R 2 808.00 per application/product	R 2 985.00 per application/product	Renewal is valid for 3 years.

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NATURE OF SERVICE, GOODS OR SUPPLIES PROVIDED	TARIFF APPLICABLE FROM 1 APRIL 2023	TARIFF APPLICABLE FROM 1 APRIL 2024	REMARKS / EXPLANATION
5.3 Payment additional to that mentioned in 5.2 in case of late application for the renewal of the sterilizing plant and rendering plant	R 2 091.00 per application/plant	R 2 223.00 per application/plant	
5.4 **Payments for information and other services.**			
(i) Import Permit	R 756.00 per product	R 804.00 per product	
(ii) Advertisement approval	R 874.00 per advertisement.	R 929.00 per advertisement.	
(iii) Free Sale certificate	R 72.00 per certificate	R 77.00 per certificate	
(iv) Re- print of the certificate	R 72.00 per certificate	R 77.00 per certificate	
(v) Other documents	R 131 per request plus, R1.00 per page	R 139 per request plus, R1.00 per page	
(vi) Any other amendment on registered product requested by the registration holder, e.g., transfer of registration, company, and product name change, etc.	R 1 316.00 per application/product.	R 1 399.00 per application/product.	
(vii) An appeal under section 6 of Act No 36 of 1947	R 7 458.00 per application/product.	R 7 928.00 per application/product.	

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NATURE OF SERVICE, GOODS OR SUPPLIES PROVIDED	TARIFF APPLICABLE FROM 1 APRIL 2023	TARIFF APPLICABLE FROM 1 APRIL 2024	REMARKS / EXPLANATION
6 Application for the registration of fertilizers			
6.1 Application for registration of fertilizer products including lapsed registration, parallel and daughter registration.	R 5 462.00 per application/product	R 5 806.00 per application/product	Registration is valid for 3 years.
6.2 Application for the renewal of the Fertilizer product	R 2 809.00 per application/product	R 2 986.00 per application/product	Renewal is valid for 3 years.
6.3 Payment additional to that mentioned in 6.2 in case of late application for the renewal of the fertilizer product	R 2 091.00 per application/product	R 2 223.00 per application/product	
6.4 Application for a group 3 fertilizer product including lapsed registration, parallel and daughter registration.	R 8 008.00 per application/product	R 8 513.00 per application/product	
6.5 **Payment for information and other services.**			
(i) Import Permit	R 756.00 per product	R 804.00 per product	
(ii) Advertisement approval	R 874.00 per advertisement	R 929.00 per advertisement	
(iii) Free Sale certificate	R 72.00 per certificate	R 77.00 per certificate	
(iv) Re- print of the certificate	R 72.00 per certificate	R 77.00 per certificate	
(v) Other documents	R 131 per request plus, R1.00 per page	R 139 per request plus, R1.00 per page	
(vi) Guideline documents	#	#	
(vii) Any other amendment requested by the registration holder, e.g., change	R 1 316.00 per application/product	R 1 399.00 per application/product	

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NATURE OF SERVICE, GOODS OR SUPPLIES PROVIDED	TARIFF APPLICABLE FROM 1 APRIL 2023	TARIFF APPLICABLE FROM 1 APRIL 2024	REMARKS / EXPLANATION
in product name, change in company name, address, etc.			
(viii) An appeal in terms of section 6 of Act 36 of 1947	R 7 458.00 per application	R 7 928.00 per application	
7 Application for Registration of Pest Control Operator			
7.1 Application for registration of Pest Control Operator including lapsed registration.	R 2 809.00 per application	R 2 986.00 per application/product	Registration is valid for 3 years.
7.2 Application for the renewal of the Pest Control Operator.	R 1 484.00 per application	R 1 577.00 per application/product	Renewal is valid for 3 years.
7.3 Payment additional to that mentioned in 7.2 in case of late application for the renewal of Pest Control Operator certificate.	R 788.00 per application	R 838.00 per application/product	
7.4 An appeal in terms of section 6 of Act No.36 of 1947	R 7 458.00 per application/product	R 7 928.00 per application/product	
7.5 Payment for information and documentation.	R 131 per request plus, R1.00 per page	R 139 per request plus, R1.00 per page	
7.6 Re-print of the certificate	R 72.00 per certificate	R 77.00 per certificate	
8 Application for Registration of an Agricultural Remedy /Stock Remedy			
8.1 Application for the registration of an Agricultural Remedy or Stock Remedy including lapsed registration, parallel and daughter registration.	R 12 140.00 per application/product	R 12 905.00 per application/product	Registration is valid for 3 years.
8.2 Application for the renewal of the registration of an Agricultural Remedy or Stock Remedy	R 6 085.00 per application/product	R 6 468.00 per application/product	Renewal is valid for 3 years.

NATURE OF SERVICE, GOODS OR SUPPLIES PROVIDED	TARIFF APPLICABLE FROM 1 APRIL 2023	TARIFF APPLICABLE FROM 1 APRIL 2024	REMARKS / EXPLANATION
8.3 Payment additional to that mentioned in 8.2 in case of late application for renewal of an Agricultural Remedy or Stock Remedy.	R 4 247,00 per application/product	R 4 515.00 per application/product	
8.4 **Other services payments**:			
(a) Approval of additional or new source of active ingredient(s), for Agricultural Remedies	R 6 421.00 per application/product	R 6 826.00 per application/product	
(b) Change of active ingredient purity specification/ notification of new impurity of technical material/ manufacturing process change	R 6 421.00 per application/product	R 6 826.00 per application/product	
(c) Change of manufacturer or additional manufacturer for Stock Remedy or Agricultural Remedy.	R 6 421.00 per application/product	R 6 826.00 per application/product	
(d) Major change in the formulation	R 6 421.00 per application/product	R 6 826.00 per application/product	
(e) Minor change in the formulation	R 1 316.00 per application/product	R 1 399.00 per application/product	
(f) Amendment of shelf life/packaging material for Stock Remedy or Agricultural Remedy	R 6 421.00 per application/product	R 6 826.00 per application/product	
(g) Additional claim(s) and withdrawal period requested by the registration holder of an Agricultural Remedy or Stock Remedy	R 12 140.00 per application/product	R 12 905.00 per application/product	
(h) Amendment requested by the office of the Registrar in relation to restricted or controlled substance.	Free	Free	

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NATURE OF SERVICE, GOODS OR SUPPLIES PROVIDED	TARIFF APPLICABLE FROM 1 APRIL 2023	TARIFF APPLICABLE FROM 1 APRIL 2024	REMARKS / EXPLANATION
(i) Change in product name, change in company name, address, spelling mistakes, species scientific name changes excluding technical changes.	R 1 316.00 per application/product	R 1 399.00 per application/product	
(j) Any other minor amendments excluding technical on registered product requested by the registration holder e.g., manufacture name change, supplier name change, transfer of registered valid registration, administrative amendments, etc.	R 1 316.00 per application/product	R 1 399.00 per application/product	
(k) Dossier updates or notifications for Stock Remedies and Agricultural Remedies	R 1 316.00 per application/product	R 1 399.00 per application/product	
(l) Approval of change of artwork on approved label, e.g., artwork approval and minor label amendment	R 1 316.00 application/product	R1 399.00 application/product	
(m) Protocol approval	R 3 866.00 per application/product	R 4 110.00 per application/product	
(n) Data waiver application: scientific data analysis	R 3 866.00 per application/product	R 4 110.00 per application/product	
(o) Fulfillment of conditional registration in case of emergency registration, e.g., minor crops/ minor species in Agricultural remedies and Stock remedies.	R 1 973.00 per application/product	R 2 097.00 per application/product	
(p) Technical/Data evaluation in case of application for an exemption.	R 1 973.00 per application/product	R 2 097.00 per application/product	

NATURE OF SERVICE, GOODS OR SUPPLIES PROVIDED	TARIFF APPLICABLE FROM 1 APRIL 2023	TARIFF APPLICABLE FROM 1 APRIL 2024	REMARKS / EXPLANATION
8.5 **Payment for information and other Services.**			
(i) Import Permit	R 756.00 per product	R 804.00 per product	
(ii) Advertisement approval	R 874.00 per advertisement	R 929.00 per advertisement	
(iii) Free Sale certificate	R 72 00 per certificate	R 77.00 per certificate	
(iv) Re- print of the certificate	R 72.00 per certificate	R 77.00 per certificate	
(v) Other documents	R 131 per request plus, R1.00 per page	R 139 per request plus, R1.00 per page	
(vi) Guideline documents	#	#	Guidelines are available free on DALRRD website.
(vii) An appeal in terms of section 6 of Act no 36 1947	R 7 458.00 per application	R 7 928.00 per application	

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DEPARTMENT OF CORRECTIONAL SERVICES

NO. 4483 **8 March 2024**

correctional services
Department:
Correctional Services
REPUBLIC OF SOUTH AFRICA

AMENDMENTS TO DCS DELEGATIONS OF AUTHORITY: ANNEXURE A TO THE TREASURY REGULATIONS, PUBLIC SERVICE REGULATIONS AND CORRECTIONAL SERVICES ACT

I, **Makgothi Samuel Thobakgale, National Commissioner (and accounting officer) of the Department of Correctional Services** in accordance with the powers vested in me by:-

- Section 97(2) of the Correctional Services Act, 1998 (act 111 of 1998) as amended;
- The Public Finance Management Act, Act 1 of 1999, as amended by Act 29 of 1999 and (b) the Treasury Regulations, 2005, promulgated in terms of Section 76 of the said Act;
- The Public Service Act, 1994, as amended by Act 30 of 2007, and (b) the Public Service Regulations, 2016, promulgated in terms of Section 41 of the said Act, as set;

hereby amend the following delegations that formed part of the delegations published in the Government gazette no 49108 on 11 August 2023. The amendments becomes effective on the date of this publication:

1. **Delegation number 19**: Delegations by the Accounting officer, Treasury Regulations: Annexure A: Delegations related to expenditure control
2. **Delegation numbers 10; 15; 17, 19 and 103**: Delegations by the National Commissioner: Correctional Services Act 111 of 1998 as amended.
3. **Delegation number 41**: PART B, APPENDIX B: Delegations Register: National Commissioner delegations to performer levels in terms of the Public Service Regulations 2016.

The amendments is set out in the attached amended pages.

MS Thobakgale
National Commissioner
Date: 12/02/2024

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ANNEXURE A

DEPARTMENT OF CORRECTIONAL SERVICES

AMENDMENT TO DELEGATIONS RELATED TO EXPENDITURE CONTROL

Note: All general abbreviations and conditions applicable to the principal delegations is applicable to this Annexure A

NO	DESCRIPTION	DELEGATION			
		HEAD OFFICE	REGIONS	MANAGEMENT AREAS / CORRECTIONAL CENTRES	REMARKS/FINANCIAL LIMITS (PER TRANSACTION)
19	1. Processing and payments of arbitration awards on PERSAL and BAS in Head Office.	1. DC: ER DC: HRM DC: FMA	1. Not delegated	1. Not delegated	• The delegation and limitation under section 96(2)(a) of the CSA delegation number 103 **must** be adhered to. • Regional Co-coordinators Employee Relations in the regions and Head Office must facilitate and consolidate all arbitration awards and submit under the signature of RC and relevant CDC to Head Office: DC Employee Relations for validation and confirmation. • DC Employee Relations must register and validate all the awards and certify in writing to DC HRM and DC FMA for calculations and processing of the payments.
	2. Furnishing of security in terms section 145 (7) of the LRA if approved that an arbitration award can be taken on review by the delegated authority (refer to delegation no 103: Decision to review arbitration award-Correctional Services Act delegations)	2. DC LS	2. Not delegated	2. Not delegated	

MS Thobakgale
National Commissioner
Date: 12/02/2024

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AMENDMENTS TO DELEGATIONS: CORRECTIONAL SERVICES ACT AND REGULATIONS: DEPARTMENT OF CORRECTIONAL SERVICES

A. CORRECTIONAL SERVICES ACT 111 OF 1998 AS AMENDED

	POWER OR DUTY BEING DELEGATED		Principal	DESIGNATION/POST LEVEL DELEGATED TO PER TIER			DELEGATION CONDITIONS AND LIMITATIONS
No	Section	Topic Description	Functionary	Head Office	Region	Management Area	
10	Section 3(5)(c) and (g)	1. Approval of transport for school children, transport between residence and place of work	NC	1. CDC Human Resources	1. Regional Commissioner	Not delegated	Compliance to DPSA financial manual
		2. Approval of transport to funeral of an official	NC (HOD)	2. Relevant CDC where official was	2. Regional Commissioner	Not delegated	Compliance to prescripts issued by the DPSA and the Department of transport, as well as all relevant prescripts on cost containment
15	Section 3(5)(g) read with Section 96(3)(d)	Approve all placements and transfers/temporary transfers on state cost.	NC	Levels 2-7 Director HR Administration & Utilization Levels 8-10 (**Excluding** officials who are Heads of Correctional or Remand Detention Centres or Community Correction offices) DC HR Management Levels 11-12 (**Excluding** officials who are Heads of Correctional or Remand Detention Centres or Community Correction offices) DC HR Management) CDC Human Resources	No delegation	No delegation	Subject to the Policy on transfers and prescripts on finances and logistics. Note 1: Transfers and placement of all SMS members: Levels 13-15 not delegated. Decision maker: National Commissioner Note 2: **Officials who are Heads of Correctional or Remand Detention Centres or Community Correction offices not delegated. Decision maker remains the National Commissioner**
17	Section 3(5)(g) read with Section 96(3)(d)	Approve transfers/temporary transfers and placements of officials on own time and cost from level 2 (SCO or equivalent) to level 12 (DD or equivalent), including interdepartmental transfers from DCS to other departments/ institutions.	NC	Level 2-7 Relevant Director after consultation with the Director HR A&U Level 8-12 Relevant DC after consultation with Director HR A&U	Within RC Office Levels 2-7: Relevant RH Corporate Services after consultation with the DRC Across Management Areas: Deputy Regional Commissioner after consultation with the RH Corporate Services	Levels 2-7: Within Management Area only, Area Commissioner (including interdepartmental transfers from DCS)	Compliance to transfer policy and procedures. To be processed/managed by HR In regional offices and Head Office the application for transfer must be submitted to Personnel Office that will process the application to the relevant Directorate. Note: **Officials who are Heads of Correctional or Remand Detention Centres or Community**

MS Thobakgale
National Commissioner
Date: 12/02/2024

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AMENDMENTS TO DELEGATIONS: CORRECTIONAL SERVICES ACT AND REGULATIONS: DEPARTMENT OF CORRECTIONAL SERVICES

	POWER OR DUTY BEING DELEGATED		Principal	DESIGNATION/POST LEVEL DELEGATED TO PER TIER			DELEGATION CONDITIONS AND LIMITATIONS
No	Section	Topic Description	Functionary	Head Office	Region	Management Area	
					Levels 8-10: (**Excluding** officials who are Heads of Correctional or Remand Detention Centres or Community Correction offices) Deputy Regional Commissioner (Excluding Heads of Centres and Community Correction offices) Levels 11-12: (**Excluding** officials who are Heads of Correctional or Remand Detention Centres or Community Correction offices) Regional Commissioner		**Correction offices not delegated. Decision maker remains the National Commissioner**
19	Section 3(5)(g) read with Section 96(3)(d) and Public Service Regulations 2016, Regulation 63	Appoint employees in acting capacity (read with delegations in terms of the Public Service Regulations)	NC	Level 2-12 Relevant Director Level 13 Relevant DC in consultation with the CDC of the Branch. Level 14 Relevant CDC in consultation with the NC.	Level 2-12 Relevant Director Level 13 DRC in consultation with the RC Level 14 RC in consultation with the NC.	Level 2-12 Area Commissioner Level 13 DRC in consultation with the RC Level 14 RC in consultation with the NC.	1. Level 15 in consultation with the Minister. 2. Compliance to relevant policy and Public Service Regulation 63 at all times. 3. The appointment in acting capacity of SMS members for longer than 6 months must be submitted to the Minister for approval in terms of DPSA policy via the National Commissioner. 4. The delegation to approve any level in acting capacity must comply with the prescripts on acting and the National Commissioner may at any time revoke such acting appointment. 5. HR Head Office will provide template letters to be used for appointment of officials who must act. 6. HR must certify that a post is vacant on PERSAL before any acting appointment where acting allowance is involved. 7. CDC HR to issue a directive on the administrative conditions for acting appointments.
103	Section 96(2)(a) read with Chapter VII of the Labour Relations Act 66 of 1995 as	Decision to review all arbitration awards (Section 145 of the LRA) or to withdraw such review.	NC	Director Employee Relations for Head Office to decide if matter must be	Regional Head Human Resources for region to decide if matter must be	No delegation	**NB: If the award is not referred for a decision to review there is no requirement for a mandate to implement and must be implemented, since**

MS Thobakgale
National Commissioner
Date: 12/02/2024

AMENDMENTS TO DELEGATIONS: CORRECTIONAL SERVICES ACT AND REGULATIONS: DEPARTMENT OF CORRECTIONAL SERVICES

	POWER OR DUTY BEING DELEGATED		Principal	DESIGNATION/POST LEVEL DELEGATED TO PER TIER			DELEGATION CONDITIONS AND LIMITATIONS
No	Section	Topic Description	Functionary	Head Office	Region	Management Area	
	amended read together with PSCBC Resolution 5/2005			implemented or not. **If not to be implemented it must be referred to the DC Legal Services for a decision to review.** Director CEU for Section 95B cases to decide if award must be implemented or not. If not to be implemented, it must be referred to the **DC Legal Services** for a decision to review. **Decision to review in all cases is with the DC Legal Services.**	implemented or not. **If not to be implemented it must be referred to the DC Legal Services for a decision to review.** **Regions have no delegation to review arbitration awards.** **No delegation for Section 95B matters**		**there is already an award that is enforceable in terms of Section 143 of the LRA. Failure to implement may lead to contempt proceedings in the Labour Court.** The Director Employee Relations, Director CEU (latter in Section 95B cases only) and the Regional Head Human Resources are responsible to ensure implementation and to issue instruction/s to ensure such implementation.

MS Thobakgale
National Commissioner
Date: 12/02/2024

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AMENDMENT: PART B, APPENDIX B: DELEGATIONS REGISTER - NATIONAL COMMISSIONER
HEAD OF DEPARTMENT DELEGATIONS TO PERFORMER LEVELS IN TERMS OF THE PUBLIC SERVICE REGULATIONS, 2016

ROW NO	POWER OR DUTY BEING DELEGATED		Principal Functionary	DESIGNATION/POST LEVEL DELEGATED TO PER TIER			DELEGATION CONDITIONS AND LIMITATIONS
	Regulation	Topic Description		Head Office	Region	Management Area	Indicators for Execution/Reporting/ Consultation/ Informing
	CHAPTER 1	GENERAL PROVISIONS					
41	48	Leave A head of department shall-- (a) encourage an employee to fully utilize his or her annual leave entitlement in the leave cycle; (b) ensure that all leave taken by an employee is recorded accurately and in full; and (c) ensure that an employee does not abuse sick leave.	NC	Approval of application for all types of leave **1-30** days (except leave without pay and Incapacity leave): Supervisor (minimum level 7) and next level manager. **31- 60 Days** Minimum level DD **61 – 89 Days** Relevant Director **Leave without pay** **1-7 Days** Relevant Director **8-29 Days** Director HR Administration & Utilization **30-89 Days** DC HR Management **90 days+** (all types of leave except incapacity leave) Relevant Branch Head in Head Office More than **184** calendar days of unpaid leave in a period of 18 months DC HR Management	Approval of application for all types of leave **1-30** days (except leave without pay and Incapacity leave): Supervisor (minimum level 7) and next level manager **31- 60 Days** Minimum level DD **61 – 89 Days** RH Corporate Services **Leave without pay** **1-7 Days** Relevant Director **8-29 Days** RH Corporate Services **30-89 Days** Deputy Regional Commissioner all types of leave except incapacity leave) Regional Commissioner More than **184** calendar days of unpaid leave in a period of 18 months DC HR Management	Approval of application for all types of leave **1-30** days (except leave without pay and Incapacity leave): Supervisor (minimum level 7) and next level manager **31- 60 Days** Minimum level DD **61 – 89 Days** Area Commissioner **Leave without pay** **1-7 Days** Head Correctional/ Remand Centre/ Community Corrections (minimum level DD) **8-29 Days** Area Commissioner **30-89 Days** DRC or Area Commissioner on DC level **90 days+** (all types of leave except incapacity leave) Regional Commissioner More than **184** calendar days of	DPSA determination on leave must be adhered to. In Management Areas where the AC is on the level of a Director the leave forms of DD's must be recommended by the Area Coordinator: Corporate Services to verify correctness and availability of leave. Leave forms for Area Commissioner (including on DC level) in regions must be recommended by the DRC. Leave forms of DRC's must be recommended by the Regional Head: Human Resources to verify correctness and availability of leave. Leave forms of CDC's and RC's must be recommended by the CDC HR to verify correctness and availability of leave

MS Thobakgale
National Commissioner
Date: 12/02/2024

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AMENDMENT TO PART B, APPENDIX B: Delegations Register - NC delegations to performer levels in terms of the PSR
HEAD OF DEPARTMENT DELEGATIONS TO PERFORMER LEVELS IN TERMS OF THE PUBLIC SERVICE REGULATIONS, 2016

				Incapacity leave (including 'conditional' approval) Director HRM & A Permanent incapacity leave DD Retention & Termination Deviation from the recommendation of the Health Risk Manager on temporary incapacity leave and ILL health retirement Director HR A&U in consultation with the relevant Director Approval of special leave not specified in the special leave policy DC HRM	Incapacity leave (including 'conditional' approval) Regional Coordinator: HR Permanent incapacity leave DD Retention & Termination Deviation from the recommendation of the Health Risk Manager on temporary incapacity leave and ILL health retirement Director HR A&U in consultation with the RH HR Approval of special leave not specified in the special leave policy No delegation	unpaid leave in a period of 18 months DC HR Management Incapacity leave (including 'conditional' approval) Area Coordinator: Corporate Services Permanent incapacity leave DD Retention & Termination Deviation from the recommendation of the Health Risk Manager on temporary incapacity leave and ILL health retirement Director HR A&U in consultation with the RG HR Approval of special leave not specified in the special leave policy No delegation	Policy and procedure on incapacity leave and ILL- Health Retirement (PILIR) and DPSA determination on leave must be adhered to
		Approval of payment of leave gratuity on service termination	**HOD**	**DD Auxiliary Services**	**Regional Coordinator Human Resources**	**Area Coordinator Human Resources**	**Compliance to DPSA directives and all applicable prescripts.**

MS Thobakgale
National Commissioner
Date: 12/02/2024

DEPARTMENT OF FORESTRY, FISHERIES AND THE ENVIRONMENT

NO. 4484 **8 March 2024**

NATIONAL ENVIRONMENTAL MANAGEMENT: BIODIVERSITY ACT (10/2004)

NOTICE OF INTENTION TO DECLARE ERF 5159, ERF 3023 AND ERF 3009 BETTYS BAY AS AN EXTENSION TO THE HAROLD PORTER NATIONAL BOTANICAL GARDEN

I, Barbara Dallas Creecy, Minister of Forestry, Fisheries and the Environment hereby, in terms of Section 33(2)(b) of the National Environmental Management: Biodiversity Act, 2004 (Act No. 10 of 2004), give notice of the intention to declare Erf 5159, Erf 3023 and Erf 3009 Bettys Bay as an extension to the Harold Porter National Botanical Garden.

The Harold Porter National Botanical Garden is one of the national botanical gardens managed by the South African National Biodiversity Institute (SANBI) and forms part of the Kogelberg Biosphere Reserve, a UNESCO World Heritage Site. The garden covers an area of 200.5 hectares and showcases the rich diversity of the Cape Floral Kingdom.

Erf 5159, Erf 3023 and Erf 3009 Bettys Bay are three adjacent pieces of land located near the garden, measuring approximately 11.3 hectares in total. Erfs 3023 and 3009 are state-owned and are being managed by SANBI with the intention of incorporating them into the garden. The lands contain natural vegetation that is representative of the coastal fynbos region, as well as some alien invasive plants that need to be cleared. The lands also serve as a natural buffer zone between the garden and the surrounding urban development.

The declaration of these land portions as an extension to the garden will enhance its conservation value, protect its biodiversity, provide opportunities for restoration, research, education and recreation, and contribute to the objectives of the Act and other relevant policies and legislation.

Members of the public are invited to submit written representations on or objections to this notice within 60 days from the date of publication of this notice in the Gazette.

By post to: The Director-General

Department of Forestry, Fisheries and the Environment

Attention: Mr Tshilidzi Netshisaulu

Private Bag X447

PRETORIA

0001

By hand at: Environment House
478 Steve Biko Road
ARCADIA
0083

By email: PKershaw@dffe.gov.za

Comments received after the closing date may not be considered.

The Department of Forestry, Fisheries and the Environment complies with the protection of Personal Information Act, 2013 (Act No. 4 of 2013). Comments received on the draft regulations, and the responses thereto, are collated in a comments and response report which will be made available to the public as part of the consultation process. If a commenting party has any objection to his or her name, or the name of the represented organisation being made publicly available in the comments and response report, such objection should be highlighted in bold as part of the submission.

Any enquiries in connection with this notice can be directed to Ms Pamela Kershaw at 012 399 9585.

BARBARA CREECY

MINISTER OF FORESTRY, FISHERIES AND ENVIRONMENT

SCHEDULE

Region: situated in Western Cape Province in the Bettys Bay Township in the Hangklip-Kleinmond Local Municipality

Description	Diagram	Extent (hectares)
Erf 3023 Bettys Bay	T3376/1980	1,6910
Erf 3009	T11618/1938	7,4286
Erf 5159	T52202/1989	2,1868

TOTAL: 11.3064 Hectares

DEPARTMENT OF TRADE, INDUSTRY AND COMPETITION

NO. 4485 **8 March 2024**

STANDARDS ACT, 2008
STANDARDS MATTERS

In terms of the Standards Act, 2008 (Act No. 8 of 2008), the Board of the South African Bureau of Standards has acted in regard to standards in the manner set out in the Schedules to this notice.

SECTION A: DRAFTS FOR COMMENTS

The following draft standards are hereby issued for public comments in compliance with the norm for the development of the South Africa National standards in terms of section 23(2)(a) (ii) of the Standards Act.

Draft Standard No. and Edition	Title, scope and purport	Closing Date
SANS 10400-O Ed 4	*The application of the National Building Regulations – Part O: Lighting and ventilation*. Provides deemed-to-satisfy requirements for compliance with part O (Lighting and ventilation) of the National Building Regulations.	2024-04-04
SANS 2070 Ed 1	*Disinfectants and Detergent-disinfectants for use in domestic and industrial environments*. Specifies the requirements for disinfectants and detergent-disinfectants for use in domestic, institutional and industrial environments (i.e. non-food preparation environments).	2024-04-08

SCHEDULE A.1: AMENDMENT OF EXISTING STANDARDS

The following draft amendments are hereby issued for public comments in compliance with the norm for the development of the South African National Standards in terms of section 23(2)(a) (ii) of the Standards Act.

Draft Standard No. and Edition	Title	Scope of amendment	Closing Date
SANS 657-1 Ed 3.4	*Steel tubes for non-pressure purposes – Part 1: Sections for scaffolding, general engineering and structural applications.*	Amended to update referenced standards, and to delete the annex on notes to purchasers.	2024-04-04
SANS 1321-1 Ed 1.2	*Non-metallic waste traps – Part 1: Plastics waste traps.*	Amended to remove the annex on notes to purchases.	2024-04-09
SANS 1077 Ed 1.3	*Sealing compounds for the building and construction industry, two-component, polyurethane-base.*	Amended to delete the annex on notes to purchasers.	2024-04-09
SANS 62-1 Ed 3.4	*Steel pipes – Part 1: Pipes suitable for threading and of nominal size not exceeding 150 mm.*	Covers the requirements for two classes (medium and heavy-based on wall thickness) of welded and seamless steel pipes with screwed or plain ends and of nominal size in the range 8 mm to 150 mm.	2024-04-09
SANS 3001-CO3-2 Ed 1.2	*Civil engineering test methods – Part CO3-2: Concrete durability index testing – Oxygen permeability test.*	Describes the method to test for oxygen permeability on concrete specimens that have been prepared by coring and cutting concrete cubes in the laboratory, or by taking cores from concrete elements on site.	2024-04-23

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SCHEDULE A.2: WITHDRAWAL OF THE SOUTH AFRICAN NATIONAL STANDARDS

In terms of section 24(1)(C) of the Standards Act, the following published standards are issued for comments with regard to the intention by the South African Bureau of Standards to withdrawn them.

Draft Standard No. and Edition	Title	Reason for withdrawal	Closing Date

SCHEDULE A.3: WITHDRAWAL OF INFORMATIVE AND NORMATIVE DOCUMENTS

In terms of section 24(5) of the Standards Act, the following documents are being considered for withdrawal.

Draft Standard No. and Edition	Title	Reason for withdrawal	Closing Date

SECTION B: ISSUING OF THE SOUTH AFRICAN NATIONAL STANDARDS

SHEDULE B.1: NEW STANDARDS

Standard No. and year	Title, scope and purport
SANS 8201:2024 Ed 2	*Alarm systems – Audible emergency evacuation signal – Requirements*. Specifies the requirements for an audible emergency evacuation signal intended to indicate without ambiguity, to all persons within the reception area of the signal, that an emergency situation (fire, gas leaks, explosion, nuclear radiation, etc.) requires immediate evacuation.
SANS 24511:2024 Ed 1	*Activities relating to drinking water and wastewater services – Guidelines for the management of wastewater utilities and for the assessment of wastewater services*. Provides guidelines for the management of wastewater utilities and for the assessment of wastewater services.
SANS 24512:2024 Ed 1	*Activities relating to drinking water and wastewater services – Guidelines for the management of drinking water utilities and for the assessment of drinking water services*. Provides guidelines for the management of drinking water utilities and for the assessment of drinking water services.
SANS 20166-4:2024 Ed 1	*Molecular in vitro diagnostic examinations – Specifications for pre-examination processes for formalin-fixed and paraffin-embedded (FFPEJ tissue – Part 4: In situ detection techniques*. Specifies requirements and gives recommendations for the collection, handling, documentation, transport, storage and processing during the pre-examination phase of formalin-fixed and parraffin-embedded (FFPE) tissue specimens intended for qualitative and/or (semi-) quantitative in situ examination of the morphology and of biomolecules, such as metabolites, protein, DNA and or RNA, on FFPE tissue section by using different in situ detection technique.
SANS 17664-1:2024 Ed 1	*Processing of health care products – Information to be provided by the medical device manufacturer for the processing of medical devices – Part 1: Critical and semi-critical medical devices*. Specifies requirements for the information to be provided by the medical device manufacturer for the processing of critical or semi-critical medical devices or medical devices that are intended to be sterilized.
SANS 60086-3:2024 Ed 3	*Primary batteries – Part 3: Watch batteries*. Specifies dimensions, designation, methods of tests and requirements for primary batteries for watches.
SANS 60086-5:2024 Ed 5	*Primary batteries – Part 5: Safety of batteries with aqueous electrolyte.* Specifies tests and requirements for primary batteries with aqueous electrolyte to ensure their safe operation under intended use and reasonably foreseeable misuse.

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Standard No. and year	Title, scope and purport
SANS 60335-2-16:2024 Ed 4	*Household and similar electrical appliances – Safety – Part 2-16: Particular requirements for food waste disposers.* Deals with the safety of electric food waste disposers for household and similar purposes, their rated voltage being not more than 250 V including direct current (DC) supplied appliances and battery-operated appliances.
SANS 62115:2024 Ed 2	*Electric toys – Safety.* Specifies safety requirements for electric toys that have at least one function dependant on electricity, electric toys being any product designed or intended, whether or not exclusively, for use in play by children under 14 years of age.
SANS 62264-3:2024 Ed 2	*Enterprise-control system integration – Part 3: Activity models of manufacturing operations management.* Defines activity models of manufacturing operations management that enable enterprise system to control system integration.
SANS 62264-5:2024 Ed 2	*Enterprise-control system integration – Part 5: Business to manufacturing transactions.* Defines transactions in terms of information exchanges between applications performing business and manufacturing activities associated with Levels 3 and 4.
SANS 10400-T:2024 Ed 5	*The application of the National Building Regulations – Part T: Fire protection.* Provides deemed-to-satisfy requirements for compliance with part T (Fire Protection) of the National Building Regulations.
SANS 14764:2024 Ed 3	*Software Engineering – Software Life Cycle Processes – Maintenance.* Provides guidance for the maintenance of software, based on the maintenance process and its activities and tasks defined in ISO/IEC/IEEE 12207:2017, 6.4.13.
SANS 23081-2:2024 Ed 2	*Information and documentation – Metadata for managing records – Part 2: Conceptual and implementation issues.* Outlines a framework for defining metadata elements consistent with the principles and Implementations outlined in SANS 23081-1.

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SCHEDULE B.2: AMENDED STANDARDS

The following standards have been amended in terms of section 24(1)(a) of the Standards Act.

Standard No. and year	Title, scope and purport
SANS 868-1-1:2024 Ed 1.2	*Compression-ignition engine systems and machines powered by such engine systems, for use in mines and plants with explosive gas atmospheres or explosive dust atmospheres or both – Part 1-1: Hazardous locations in underground mines – Basic explosion protected engines. Consolidated edition incorporating amendment No. 2.* Amended to delete reference to certification mark scheme.
SANS 868-3-3:2024 Ed 1.2	*Compression-ignition engine systems and machines powered by such engine systems, for use in mines and plants with explosive gas atmospheres or explosive dust atmospheres or both – Part 3-3: Hazardous locations on surface – Machines. Consolidated edition incorporating amendment No. 2.* Amended to update the clause on testing and certification of surface-based machines with compression-ignition engines, and the clause on marking.
SANS 1515-3-2:2024 Ed 1.1	*Gas measuring equipment primarily for use in mines – Part 3-2: Fixed, transportable, and vehicle-mounted toxic gas measuring and warning sensor heads, instruments and devices. Consolidated edition incorporating amendment No. 1.* Amended to update referenced standards, the definitions, and the clause on general requirements, to remove references to certification and certification mark, and to update the annex on sampling and assessment of compliance with this part of SANS 1515.
SANS 238:2024 Ed 3.1	*Liquid toilet soap. Consolidated edition incorporating amendment No. 1.* Amended to delete the annex on notes to purchasers.
SANS 511:2024 Ed 2.1	*Women's workwear. Consolidated edition incorporating amendment No. 1.* Amended to update referenced standards, and to delete the annex on notes to purchasers.
SANS 1437:2024 Ed 2.2	*Footwear soling material and footwear sole and heel components of rubber and plastics. Consolidated edition incorporating amendment No. 2.* Amended to update referenced standards, and to delete the annex on notes to purchasers.
SANS 1828:2024 Ed 2.3	*Cleaning chemicals for use in the food industry. Consolidated edition incorporating amendment No. 3.* Amended to update the requirements on suitability for purpose and pH value, referenced standards, the requirements on method of test for effect on corrosion-resistant steel, and on marking and labelling.
SANS 1411-7:2024 Ed 2.3	*Materials of insulated electric cables and flexible cords – Part 7: Polyethylene (PE). Consolidated edition incorporating amendment No. 3.* Amended to update referenced standards, definitions, and to include sheath shrinkage requirements.
SANS 1464-22:2024 Ed 2.1	*Safety of luminaires – Part 22: Luminaires for emergency lighting. Consolidated edition incorporating amendment No. 1.* Amended to update referenced standards, the clauses on changeover operation, high-temperature operation, and on battery chargers for self-contained emergency luminaires, and the annex on rated relative lamp light output.
SANS 633:2024 Ed 1.1	*Soil profiling and rotary percussion borehole logging on dolomite land in Southern Africa for engineering purposes. Consolidated edition incorporating amendment No. 1.* Amended to delete a trade name.
SANS 665-1:2024 Ed 1.3	*Wedge gate and resilient seal valves for general purposes – Part 1: General. Consolidated edition incorporating amendment No. 3.* Amended to delete the annex on notes to purchasers.
SANS 1223:2024 Ed 2.4	*Fibre-cement pressure pipes and couplings. Consolidated edition incorporating amendment No. 4.* Amended to delete the annex on notes to purchasers.
SANS 1420:2024 Ed 1.4	*Bed bases (spring upholstered). Consolidated edition incorporating amendment No. 4.* Amended to delete the appendix on notes to purchasers.
SANS 1808-31:2024 Ed 1.4	*Water supply and distribution system components – Part 31: Automatic control valves. Consolidated edition incorporating amendment No. 4.* Amended to delete the annex on notes to purchasers.
SANS 949:2024 Ed 3.1	*Strongroom doors or Vault doors. Consolidated edition incorporating amendment No. 1.* Amended to delete reference to fire resistance, to update the clause on marking, the table on tools for testing resistance to attack, and to delete the annex on notes to purchasers.
SANS 15:2024 Ed 2.4	*Wax emulsion polish for floors and furniture. Consolidated edition incorporating amendment No. 4.* Amended to delete the appendix on notes to purchasers.

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SCHEDULE B.3: WITHDRAWN STANDARDS

In terms of section 24(1)(C) of the Standards Act, the following standards have been withdrawn.

Standard No. and year	Title
SANS 1781:2018 Ed 1	*Biological based treatment products.*

SCHEDULE B4: ESTABLISHMENT OF TECHNICAL COMMITTEES

In terms of the South African Norm for the development of South African National Standards, the following technical committee has been established:

Committee No.	Title	Scope

If your organization is interested in participating in these committees, please send an e-mail to Dsscomments@sabs.co.za for more information.

SCHEDULE B5: RETRACTION OF PREVIOUSLY GAZETTED ITEMS

Notice is hereby given that the following standards gazetted for public enquiry have been retracted.

Standard No.	Title	Scope	Date gazetted

SCHEDULE B6: GENERAL

Notice is hereby given that the following standards/draft standard have been renumbered.

Standard/draft No.	Title	Scope	New number/designation

SCHEDULE B7: ADDRESS OF THE SOUTH AFRICAN BUREAU OF STANDARDS HEAD OFFICE

Copies of the standards mentioned in this notice can be obtained from the Head Office of the South African Bureau of Standards at 1 Dr Lategan Road, Groenkloof, Private Bag X191, Pretoria 0001.

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DEPARTMENT OF HEALTH

NO. 4486 **8 March 2024**

HEALTH PROFESSIONS ACT, 1974 (ACT NO.56 OF 1974)

REGULATIONS RELATING TO THE QUALIFICATIONS FOR REGISTRATION OF OPTOMETRISTS: AMENDMENT, 2024

The Minister of Health intends, in terms of section 24 of the Health Professions Act, 1974 (Act No. 56 of 1974), and on the recommendation of the Health Professions Council of South Africa, to make the regulations in the Schedule.

Interested persons are invited to submit any substantiated comments or representations on the proposed regulations to the Director-General of Health, Private Bag X828, Pretoria, 0001 (for attention of the Director: Public Entities Governance; mihloti.mushwana@health.gov.za and paul.tsebe@health.gov.za), within three month of the date of publication of this notice.

DR M.J. PHAAHLA, MP
MINISTER OF HEALTH
DATE: 15/02/2024

SCHEDULE

Definitions

1. In these regulations and any word or expression to which a meaning has been assigned in the regulations shall bear such meaning, unless the context indicates otherwise –
"the Act" means the Health Professions Act, 56 of 1974; and
"the Regulations" means the regulations relating to the qualifications for the registration of optometrists as published under Government Notice No. 2554 in *Government Gazette* No. 46991 of 30 September 2022 and Government Notice No. 2599 in *Government Gazette* No. 47257 of 07 October 2022.

Substitution of regulation 2 of the regulations

2. The following regulation is hereby substituted for regulation 2 of the regulations –

"2. The Registrar may register as an Optometrist any person who –

(a) is in possession of the following qualifications; and
(b) has passed the national examination determined by the professional board for optometry and dispensing opticians.

Examining Authority	***Qualification***	***Abbreviation for registration***
Technikon Witwatersrand	Bachelor of Technology (Optometry) Diploma in Optometry SA	B Tech (Optom) Wits Tech Dip Optom Wits Tech
University of Durban-Westville	Bachelor of Optometry	B Optom Durban-Westville
University of the North	Bachelor of Optometry	B Optom North
Rand Afrikaans University	Bachelor of Optometry	B Optom RAU
University of the Free State	Bachelor of Optometry	B Optom Free State

Examining Authority	*Qualification*	*Abbreviation for registration*
University of Johannesburg	Bachelor of Optometry	B Optom Johannesburg
University of KwaZulu Natal	Bachelor of Optometry	B Optom KwaZulu Natal
University of Limpopo	Bachelor of Optometry	B Optom Limpopo

Short title

3. These Regulations are called Regulations Relating to the Qualifications for Registration of Optometrists: Amendment, 2024.

DEPARTMENT OF HEALTH

NO. 4487 **8 March 2024**

MEDICINES AND RELATED SUBSTANCES ACT, (ACT NO. 101 OF 1965)

ANNUAL SINGLE EXIT PRICE ADJUSTMENT [SEPA] OF MEDICINES AND SCHEDULED SUBSTANCES FOR THE YEAR 2024

I, DR MJ PHAAHLA, the Minister of Health, has determined on recommendation of the Pricing Committee, in terms of Regulation 8(1) of the Regulations relating to a Transparent Pricing System for Medicines and Scheduled Substances published in terms of the Medicines and Related Substances Act, (Act 101 of 1965), that the Single Exit Price (SEP) of Medicines and Scheduled Substances may be adjusted by not greater than **6.79%** of the SEP of medicines and their related pack sizes that was available as at 20th December 2023 regardless of how that SEP was arrived at in the 2021 cycle. Applications for adjustments of the SEP may only be submitted for the first time in 2024 from 12th January 2024 and by no later than 08th March 2024.

All medicines and their related pack sizes with SEP approved with an effective date later than 20th December 2023 are not eligible for SEPA 2024. An applicant may only submit once in the 2024 cycle unless a re-submission is made for eligible medicines that have not been previously approved for an adjustment in the 2024 period in which an application was made. The final date for re-submissions will be 25th March 2024. Applicants with any other type of SEP-related submission in the system must either wait for the finalization of such a submission before lodging the SEPA 2024 submission or withdraw the submission within the system before the SEPA 2024 submission is lodged. The lodgment of a SEPA 2024 submission shall automatically terminate the processing of any SEP-related submission already in the system.

An adjustment in the Single Exit Price in terms of this Notice may only be implemented by the manufacturer or importer of the relevant medicine or scheduled substance, no later than 32 working days after the date that the manufacturer or importer has communicated the information requested by the Director-General in terms of the Notice published in terms of Regulation 21 of the Regulations Relating to a Transparent Pricing System for medicines and scheduled substances.

DR MJ PHAAHLA, MP
MINISTER OF HEALTH
DATE: 11/01/2024

MEDICINES AND RELATED SUBSTANCES ACT, (ACT NO. 101 OF 1965)

INFORMATION TO BE PROVIDED BY MANUFACTURERS AND OR IMPORTERS OF MEDICINES AND SCHEDULED SUBSTANCES WHEN LODGING SUBMISSIONS FOR THE SINGLE EXIT PRICE ADJUSTMENT (SEPA) FOR 2024

I, DR SSS BUTHELEZI, Director General, have determined in accordance with Regulation 21 of the Regulations Relating to a Transparent Pricing System for Medicines and Scheduled Substances published in Government Gazette number 28214 of 11 November 2005, that the information required in the submissions for the 2021 SEP adjustment as determined by the Minister be submitted to the Directorate: Pharmaceutical Economic Evaluation (PEE) within the National Department of Health by a manufacturer or importer of the medicine or scheduled substance, who is the applicant of the medicine, in accordance to the information and instruction document appended to this Notice.

Such information should be submitted electronically to sepupdates@health.gov.za The submission should include information containing the applicant's entire portfolio; including the medicines for which the applicant is not requesting an adjustment of the SEP.



DR SSS BUTHELEZI
DIRECTOR-GENERAL: HEALTH
DATE: 11/01/2024

health
Department:
Health
REPUBLIC OF SOUTH AFRICA

2030 NDP

INFORMATION AND INSTRUCTIONS FOR THE 2024 SINGLE EXIT PRICE ADJUSTMENT (SEPA) SUBMISSIONS

ALL 2024 SEPA SUBMISSIONS MUST BE SUBMITTED ELECTRONICALLY VIA EMAIL ADDRESS sepupdates@health.gov.za WITH ALL SUPPORTING DOCUMENTS SAVED ON A ZIPPED FOLDER

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PREAMBLE

This document provides information and instructions on how to present the required information when communicating the 2024 SEP adjustment (SEPA) for medicines prices adjusted in terms of Section 22G of Medicines and Related Substances Act (101 of 1965) as amended, and Regulation 8(1) of the Regulations Relating to a Transparent Pricing System for Medicines and Scheduled Substances. The applicants are required to comply with all the requirements and instructions in this document, failure to do so will result in the submission being considered incomplete. Incomplete submissions shall be regarded as ineligible for processing on the basis of non-compliance with the requirements of these guidelines. These guidelines must be read together with the relevant sections of the Medicines Pricing regulations.

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1. ACRONYMS

CFO – Chief Financial Officer
DoH – Department of Health
NDoH – National Department of Health
DoP – Database of Single Exit Prices
MCC – Medicines Control Council
MPR – Medicine Pricing Registry
NAPPI – National Pharmaceutical Product Interface
PEE – Pharmaceutical Economic Evaluations
PI – Package Insert
SAHPRA – South African Health Products Regulatory Authority
SEP – Single Exit Price
SEPA – Single Exit Price Adjustment
VAT – Value Added Tax
WHO ATC – World Health Organisation Anatomical Therapeutic Chemical

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2. APPLICANT INFORMATION

2.1 APPLICANT REQUIREMENTS

All registered applicants for medicines sold in SA, who are eligible in terms of the notice as signed by the Minister of Health, may forward submissions for the Single Exit Price Adjustment (SEPA) for 2024 for all scheduled medicines appearing on the Database of Medicines Prices (DoP) published on 20th December 2023.

The medicines that form part of the applicant's portfolio that do not appear on the DoP of 20th December 2023 must be included at the bottom of tab 1 of the excel spreadsheet as part of the 2024 SEPA submission and the supporting evidence in the form of a communicated email that had a signed letter of approval and its associated excel spreadsheet both reflecting the effective date of the approved Single Exit Price (SEP) must be provided for such medicines as confirmation that the medicines were previously processed and allocated an official SEP by the department of Health.

a) The following medicines are legible for 2024 SEPA implementation:
    i. Medicines whose SEP's were approved for implementation on or before the 20th December 2023.
b) The following medicines are **NOT** legible for 2024 SEPA implementation:
    i. Medicines for which 2024 SEPA adjustment is not applicable
    ii. Medicines for which 2024 SEPA implementation is not required
    iii. Discontinued medicines
    iv. Medicines whose SEP's were approved for implementation after the 20th December 2023.

NB: Applicants must note that medicines that are **NOT** eligible for 2024 SEPA implementation under 2.1 (b) must still be included in the submission for 2024 SEPA implementation under tab 1.

c) The information contained in the published gazette with respect to the 2024 SEPA should be read carefully and the contents thereof must be complied with as required.
d) The dates and timelines contained in the published gazette with respect to the implementation period for 2024 SEPA must be read carefully and complied with as required and the dates are as follows:
    i. 20 December 2023 – reference Dop base SEP's for applying 2024 SEPA
    ii. 12 January 2024 – First day of receiving 2024 SEPA submissions
    iii. 8 March 2024 – Last day of receiving 2024 first time SEPA submissions
    iv. 25 March 2024 - Last day of receiving 2024 SEPA resubmissions.

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e) Read carefully the information and instructions contained in this document before completing all the fields of both tabs (Tab 1 and Tab 2) of the latest 2024 excel SEPA template which is available on the website https://www.health.gov.za/nhi/.

NB: A submission, which has been signed off by the Chief Financial Officer is presumed to be free from calculation errors. Similarly, a submission, which has been signed off by the Responsible Pharmacist is presumed to be free from any errors which are regulatory in nature.

f) Provide the required information on the cover page (**Annexure A**).

g) Sign the declaration annexed to this document (**Annexure B**).

NB: Do NOT amend the declaration form AND no information appearing on the submission shall be changed post facto if the declaration form is found to be completed and signed by all the relevant officials responsible for lodging the 2024 SEPA submission.

h) Complete the checklist that is also annexed to this document (**Annexure C**).

i) Complete **all** sections of all tabs (Tab 1 and Tab 2) of the latest 2024 SEPA template in the fields provided (**Annexure D**).

j) Include a signed covering letter on a company letterhead, stating the purpose of your submission, with every submission or re-submission where applicable.

k) A complete submission should include a fully completed latest SEPA template for 2024, annexure A, B, C and D (A fully completed SEPA template) and a signed covering letter on the applicant's letterhead.

l) Ensure all the SEPA template fields are completed in full and the base Single Exit Prices, to be used as a reference for adjustment purposes are those which were applicable on 20th December 2023 and that they have an effective date of 20th December 2023 or earlier. The base SEP's for each submitted medicine must be verified as correct and that it is appropriate for the relevant medicine prior to lodging the submission.

m) Ensure that all fields have been completed as per DoP of 20th December 2023.

NB: All the applicants must verify the correctness of the information which appears under the SEPA template excel spreadsheet Tab 1 column titled (Originator or Generic). Should this information be declared incorrect after the 2024 SEPA implementation process, the applicant will be required to provide evidence to support their claims i.e. should the column details be changed at a later stage after 20th December 2023.

n) Wherever the date is required, it should be stated in full (e.g. 14 March 2024).

o) Applicants are required to submit **ONLY** the electronic version of the entire submission via e-mail and the submission must include:

i. Signed cover letter on the official letter head of the applicant;
ii. Completed latest 2024 SEPA template;
iii. Completed Annexure A;
iv. Completed Annexure B;

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v. Completed Annexure C;
vi. Completed Annexure D (A fully completed latest 2024 SEPA template) and
vii. Supporting documents where applicable.

p) Applicants must ensure that all relevant documents such as the covering letter and the declaration form in Annexure B are signed prior to lodging the submission.

q) The responsible officials sign the declaration form (Annexure B) to verify and certify that the submission is complete and that the information contained in the submission is true, correct error-free, and every aspect of the 2024 SEPA gazette its guidelines is complied with in totality as prescribed.

r) The signed declaration form (Annexure B) also confirms that the submission in its entirety has been checked by all the persons whose signatures are appended under Annexure B, in addition to the person responsible for compiling the submission. **NB: A submission, which has been signed off by the Chief Financial Officer is presumed to be free from calculation errors. Similarly, a submission, which has been signed off by the Responsible Pharmacist is presumed to be free from any errors which are regulatory in nature.**

2.2 SEPA SUBMISSION REQUIREMENTS

a) The submissions lodged in terms of these guidelines are solely for the purpose of 2024 SEPA. For other medicine details amendments, applicants must use Template G of the SEP updates as published on the website https://www.health.gov.za/nhi/ or emailed to sepupdates@health.gov.za

b) For a submission to be considered complete, **ALL** sections of the 2024 SEPA template, inclusive of all excel spreadsheet fields, must be fully completed. A fully completed template must have all tabs (Tab 1 and Tab 2) and all the fields of the relevant worksheets completed. Within each tab, all the required fields must be completed for every medicine in the applicant's schedule as published on DoP of 20th December 2023.

c) **ALL** scheduled medicines that make up the applicant's portfolio on the date of the submission, **MUST** be presented in the latest SEPA template i.e. completed Annexure D.

d) **ALL** the medicines (and their respective prices) that have an SEP update which was approved, communicated and effected by the department in 2024, before the date of the applicant's 2024 SEPA submission, including those communicated to the applicant after the 20th December 2023, must be included in the 2024 SEPA submission. Both the letter and excel schedule received from the Directorate: Pharmaceutical Economic Evaluations (PEE) must be submitted for this category of items. The information will serve as SEPA 2024 supporting documents and it will be used to verify the information that is included as part of the 2024 SEPA submission (on the 2024 SEPA excel templates i.e. Tab 1 and Tab 2) but which may not be appearing on the 20th December 2023 database. Failure to provide these supporting documents will render the 2024 SEPA submission incomplete.

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e) Only the rightful applicant as recorded on the DoP of 20th December 2023 for the medicine as per the SAHPRA (formerly MCC) manufacturing license and MCC / SAHPRA medicines registration certificate must lodge the submission for the medicine(s) concerned.

f) Only those applicants whose manufacturing licenses have not expired may submit 2024 SEPA submissions.

g) In cases where an applicant name change occurred after the 20th December 2023 but before lodging the 2024 SEPA submission, only the applicant whose applicant name is reflected on the DoP of 20th December 2023 shall be considered for purposes of the 2024 SEPA submissions.

2.3 NOTES FOR APPLICANTS

a) The submission of 2024 SEPA is not obligatory. The eligible applicants are not compelled to compile and submit 2024 SEPA submissions.

b) The 2024 SEPA is only applicable on the medicines with a SEP that was already effective on the 20th December 2023, regardless of how these SEP's were arrived at. This includes the SEP approvals granted after a Non-Permanent Price Reduction submission. These non-permanent SEPs shall be regarded as permanent reductions at the point of lodging the 2024 SEPA submission.

c) Therefore, if the SEP of a medicine that appears on the 20th December 2023 database was arrived at after the applicant submitted a Template B submission, then such an SEP shall automatically become the SEP for applying 2024 SEPA on the first day of implementation of 2024 SEPA i.e. 12th January 2024.

d) Applicants must note that in terms of the Medicines Pricing Regulations, there shall only be one Single Exit Price at any given point in time.

e) Applicants are advised to compile their own list of reference medicine Single Exit Prices to enable the verification of prices during SEPA implementation. The schedule of 20th December 2023 may be found on the website https://www.health.gov.za/nhi/ under the folder for databases. Click on the excel spreadsheet titled *database of medicine prices 20th December 2023*.

f) There can only be one SEP submission launched at any given point in time. The applicant cannot request for an update on the SEP or lodge a Regulation 9 application, whilst the submission for SEPA is still in process. Similarly, the applicant cannot submit a SEPA or Regulation 9 application whilst the submission for an SEP update is still in process. In an event where the applicant has made a SEPA submission and any other SEP submissions and/or a Regulation 9 application the SEPA will not be considered. Should the applicant wish to re-submit, a new submission may be made once the other outstanding SEP submissions and/or Regulation 9 applications have been concluded.

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g) Each submission should include all the applicant's scheduled medicines, including discontinued medicines. Discontinued medicines should be indicated as such, as per the DoP under the status column. SEPA will not be allowed on officially declared discontinued medicines.

h) **The row order of all the applicant's medicines, as they appear on the DoP of 20th December 2023 must be maintained**.

i) Any medicines not appearing on the 20th December 2023 list should appear at the bottom of the 2024 SEPA template in an alphabetical order.

j) All medicines with related pack sizes that are presented on the template for 2024 SEPA must be unit priced unless the medicine, unless the medicines are exempt by the Minister of Health from unit pricing subsequent to submitting an exemption request to the Pricing Committee. In the case of unit pricing exemption, the applicant must provide a signed and valid exemption letter by the Minister of Health. When computing the unit prices, the resulting SEPs should not exceed the maximum allowable SEP after the adjustment on the SEP that existed on 20th December 2023 (i.e. SEP applicable as of 20th December 2023 + maximum allowable SEPA % as per the Minister's Notice).

k) All medicines including those with multiple pack sizes are required by law to be unit priced i.e. all same ingredient and dosage form medicines with related pack sizes must have the same unit price, unless the medicines are exempt by the Minister of Health from unit pricing subsequent to submitting an exemption request to the Pricing Committee and such exemption signed by the Minister of Health must be provided as supporting documentation. Non-compliance with unit pricing will result in the entire submission not being considered.

l) Where a new pack size is introduced after 20th December 2023, it is expected that this will result in a unit price that is no greater than the unit price that existed on a related pack sizes on 20th December 2023 unless a related pack size has been exempt by the Minister of Health from unit pricing subsequent to submitting an exemption request to the Pricing Committee and such exemption signed by the Minister of Health must be provided as supporting documentation. (Note that the newly launched medicines and/or pack sizes should be included in the portfolio of medicines in the submission for SEPA and should also be unit priced with their related pack sizes).

m) All submissions for SEPA will be processed within 32 working days (excluding weekends and holidays) upon receipt of the submission by the Directorate: PEE of the National Department of Health (NDoH).

n) The outcome of each processed submission will be communicated to the applicant within 32 working days of the date of your submission. Applicants are required to take note of this 2024 SEPA implementation time frame prior to following up on a submission status.

o) All processed and approved SEPs will be communicated to submitting applicants, price file managers and all the stakeholders that are registered on the Directorate: PEE of NDoH's emailing list. The approved prices will be published on the website https://www.health.gov.za/nhi/ at a later stage.

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p) All correspondence(s) concerning a submission will only be communicated to the applicant of the medicines applied for.

q) The electronic version of the submitted 2024 SEPA template (Annexure D) must be in excel (not pdf format) and should be saved with a file name extension "xls". Submissions containing password-protected documents and files in a version that is not accessible when using NDoH systems, such as those with the file extensions xlsx, docx and PDF, will not be considered.

r) 2024 SEPA can only be submitted on the published latest SEPA template for 2024 including both Tab 1 and Tab 2. **ANY** modification to the template will result in the entire submission not being considered. This also applies to re-submissions.

s) The final date for all 2024 SEPA submissions will be those as determined in the Minister's 2024 SEPA notice. No submission shall be received and reviewed outside of the dates that are stipulated in the 2024 SEPA notice.

t) An applicant may only submit once in the 2024 SEPA cycle. This does not apply to resubmissions under point (u) below.

i. Where no adjustment is requested, the existing SEP will be applicable for the 2024 SEPA cycle. The SEPA cycle is the period between two consecutive SEPA announcements by the Minister of Health. The applicant may not at a later stage re-submit a different SEPA request for the same medicine. The SEPA submission and the approval thereof for the 2024 cycle implies that previously approved non-permanent reductions automatically become the official SEP for applying 2024 SEPA at the end of the 2024 SEPA implementation cycle.

ii. An applicant's portfolio may not be divided into multiple submissions.

iii. The maximum allowable adjustment may not be divided into multiple submissions. Should an applicant request less than the maximum published adjustment, the balance will be forfeited for the 2024 cycle.

u) Resubmissions;

i. Resubmissions shall only be considered if submitted within the timelines stipulated in the 2024 SEPA notice.

ii. Will **only** be reviewed for medicines that had SEPs that were previously not adjusted in terms of the 2024 SEPA quantum, as a result of discrepancies identified in the first 2024 SEPA submission.

iii. All the requirements for the SEP submissions as stated in this document shall be applicable to re-submissions.

iv. A resubmission of the not-approved medicines may not be split into multiple re-submissions.

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v. MUST contain ALL the medicines listed under the Not-Approved sheet of Annexure E which is communicated to the applicant in response to the initial submission.
vi. Resubmissions must contain only medicines listed in the Not-Approved sheet of Annexure E communicated to the applicant in response to the initial submission.
vii. Re-submissions must only be submitted on the official and latest 2024 SEPA template.
viii. Must only be on the 2024 SEPA template, by the close off date as specified by the Minister of Health and reflected in the 2024 SEPA notice.

2.4 LODGING OF SUBMISSIONS

a) Submissions must be lodged electronically via the department of health email sepupdates@health.gov.za. No hard copies submissions will be considered.
b) The cover letter must reflect the following information:
   i. Applicant Name
   ii. 2024 SEPA Submission
   iii. Number of Medicines in TAB 1 of the submission template (e.g. Tab 1 = 50 medicines /75 line items
   iv. Number of Medicines in TAB 2 of the submission template (e.g. Tab 2 = 10 medicines /15 line items- Tab 2 is for originator medicines.
c) Each submission **MUST** be lodged on the latest 2024 SEPA template and must be accompanied by annexure A, B, C and D included in this document as well as the applicant's covering letter on the official letterhead of the applicant. All these documents must be saved on a zipped folder and submitted as such.
d) Where an applicant is uncertain about the contents of the submission being lodged, clarity must be solicited from the PEE directorate prior to lodging the submission and this must be done by no later than the closing dates for the lodging of the 2024 SEPA submissions. Queries relating to approved submissions that may contain information that was not corrected timeously will not be tolerated.
e) Only e-mail submissions will be accepted for SEPA 2024.
f) Every submission must be on a new e-mail with the heading that is specific to the content i.e. 2024 SEPA submission.
g) The 2024 SEPA submission email via sepupdates@health.gov.za must be addressed to:

2024 SEP Adjustment
Director: Pharmaceutical Economic Evaluations (PEE)
ATT: Ms Ntobeko Mpanza
National Department of Health

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Room C6-18A Dr Xuma Building
1112 Voortrekker Road, Pretoria Townlands 351 - JR
0001

Email submissions must be lodged made between 09:00 and 12:00 Monday to Friday excluding public holidays. All submissions will be allocated a reference number and no late submissions will be considered. Where the reference number is not received within five working days from the date of lodging the submission, the applicant must resubmit since their submission will not be reflecting on the list of submissions received by the department of Health.

For any enquiries regarding 2024 SEPA implementation, you may contact Ms Mahlogonolo Ledwaba between 13:00 and 15:00 at (012) 395 8186 or by e-mail at mahlogonolo.ledwaba@health.gov.za and Frieda.Seete@heath.gv.za at 012 395 8210 from Monday to Friday excluding public holidays.

All queries must include the reference number provided to the applicant as an acknowledgement of receipt of the SEPA submission

Note that the Department of Health will not be held responsible for submissions that were incorrectly submitted. A reference number reflected on the acknowledgment notice should be quoted in every communication made to NDoH by the applicant.

2.5 DOCUMENTS TO BE SUBMITTED

Applicants are required to submit **all** the following documents to ensure completeness of the submissions:

(a) Signed cover letter on the official letter head of the applicant; (The cover letter should include details of the number of medicines being submitted: see point 2.4 (b) above.
(b) Completed annexure A
(c) Completed annexure B
(d) Completed annexure C and
(e) Completed annexure D or latest 2024 SEPA template with both Tab 1 and Tab 2
(f) Supporting documents where necessary

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2.6 ACKNOWLEDGEMENT OF RECEIPT

Upon receipt of a submission, an acknowledgement notice will be provided to the representative of the applicant by the Directorate: PEE official. All applicants should retain their acknowledgment notice, for reference purposes.

3. HOW TO COMPLETE TEMPLATE COLUMNS

The details must be copied from the 20th December 2023 DoP for all the medicines for the applicant. All details and formatting must remain as it appears on DoP of 20th December 2023.

Failure to comply with the prescribed requirements under this section 3 below will result in the entire submission not being considered.

3.1 SEPA 2024 TEMPLATE TAB 1

3.1.1 For the information required under the following listed columns labels (headings) in the Template, applicants are required to copy such information from the DoP published on 20th December 2023 for all medicines that sought 2024 SEPA. All the information and formats and the order of medicines must remain as they appear on the DoP of 20th December 2023.

- APPLICANT SAHPRA/MCC LICENCE NUMBER
- APPLICANT NAME AS REGISTERED WITH SAHPRA/MCC
- SAHPRA/MCC MEDICINE REGISTRATION NUMBER
- NAPPI CODE (9-digit)
- ATC 4 CODE (WHO)
- SCHEDULE
- MEDICINE PROPRIETARY NAME
- ACTIVE INGREDIENT
- STRENGTH
- UNIT
- DOSAGE FORM
- PACK SIZE
- QUANTITY
- MANUFACTURER PRICE AS AT 20th December 2023
- LOGISTICS FEES AS AT 20th December 2023
- VAT

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- SEP AS AT 20th December 2023
- UNIT PRICE AS AT 20th December 2023
- EFFECTIVE DATE
- STATUS
- ORIGINATOR OR GENERIC

3.1.2 VOLUME OF SALES

This must be the total quantity of sales of each medicine for the period 01 January 2023 to 31 December 2023. Where the medicine is not being sold this should be indicated in the column. A blank will result in submission not being considered.

3.1.3 REQUESTED MANUFACTURER PRICE

This is the requested VAT exclusive manufacturer price of the medicine in South African Rands. This is a numerical field displayed at 2 decimal places, with no currency symbols. This column should be indented to the right.

3.1.4 REQUESTED LOGISTICS FEE

This is the requested VAT exclusive logistics fee for the medicine in South African Rands. This is a numerical field displayed at 2 decimal places, with no currency symbols. This column should be indented to the right.

3.1.5 VAT ON REQUESTED COMPONENTS

This column is the VAT component of the SEP, calculated at 15% to the sum of the requested manufacturer price and the requested logistics fee. This is a numerical field displayed at 2 decimal places with no currency symbols. This column should be indented to the right.

3.1.6 REQUESTED SEP

This is the requested Single Exit Price for the medicine in South African Rands. It is the sum of the requested ex-manufacturer price, the requested logistics fee and VAT. This is a numerical field displayed at 2 decimal places with no currency symbols. This column should be indented to the right.

3.1.7 REQUESTED UNIT PRICE

This is the resulting unit SEP of the medicine, considering its pack size and quantity of presentation as per the SAHPRA (formerly MCC) approved package insert (PI). The unit price

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should be obtained by; dividing the requested SEP by the pack size and then further divided by the quantity.

(a) This is the price of a unit of the medicine, e.g. one tablet, capsule, millilitre, gram, etc. The unit price as described in the Regulations Relating to a Transparent Pricing System for Medicines and Scheduled substances (section 22G of the Medicines and Related Substances Act) is the SEP divided by the number of units of the product. Note that unit pricing applies to all medicines with the same proprietary name, strength and dosage form.

(b) For injections the unit price shall be calculated per ml of reconstituted volume, even where the total volume of the medicine administered to a single patient is less than 1 ml.

(c) For inhalers, where the pack size is described in the SAHPRA (formerly MCC) approved PI as doses or puffs the unit price will be for 1 dose or puff.

(d) The unit price is the SEP divided by the pack size and then further divided by the quantity [the "quantity" represents the multiples in which the medicine is packed/the number of pack sizes e.g. for injections, the "quantity" for 50 vials containing 500mg powder for injection packed in 20ml vial to be reconstituted with 10ml of diluents is 50].

This is a numerical field displayed at decimal places with no currency symbols. This column should be indented to the right.

NOTE: The template with TAB 1 must always be maintained in the font and formats as it appears on Dop of 20th December 2023. Applicants should only make use of space, dashes or any other character if these are represented as such in official documentation.

3.2 SEPA 2024 TAB 2

Any blanks on Tab 2 will result in the submission not being considered. Where the medicine is a generic the applicant must comment. Where there is no price available the applicant must indicate this as well as measures taken to obtain the price. Proof of this communication must be supplied.

3.2.1 For the following columns:

- APPLICANT SAHPRA/MCC LICENCE NUMBER
- APPLICANT NAME AS REGISTERED WITH SAHPRA/MCC
- SAHPRA/MCC MEDICINE REGISTRATION NUMBER
- NAPPI CODE (9-digit)

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- ATC 4 CODE (WHO)
- SCHEDULE
- MEDICINE PROPRIETARY NAME
- ACTIVE INGREDIENT
- STRENGTH
- UNIT
- DOSAGE FORM
- PACK SIZE
- QUANTITY
- MANUFACTURER PRICE AS AT 20th December 2023
- LOGISTICS FEES AS AT 20th December 2023
- VAT
- SEP AS AT 20th December 2023
- UNIT PRICE AS AT 20th December 2023
- EFFECTIVE DATE
- STATUS
- ORIGINATOR OR GENERIC

The details must be copied from the 20th December 2023 DoP for all the medicines for the applicant. All details and formatting must remain as it appears on DoP of 20th December 2023.

3.2.2 For all medicines that are labelled originator, the following columns must be completed; Closest Australian Pack Size, Related Australia Quantity, Australian Manufacturer Price in AUS Dollars, AUS Dollar Exchange Rates, Australian Price in Rands, Australian matching pack size in Rands, Comment on Australian Price Provided, Closest Canada Pack Size, Related Canada Quantity, Canada Manufacturer Price in CAN Dollars, CAN Dollar Exchange Rates, CAN Price in Rands, Canadian matching pack size in Rands, Comment on Canadian Price Provided, Closest New-Zealand Pack Size, Related NZ Quantity, New-Zealand Manufacturer Price in NZ Dollars, NZ Dollar Exchange Rates, New-Zealand Price in Rands, New Zealand matching pack size in Rands, Comment on New Zealand Price Provided, Closest Spain Pack Size, Related Spain Quantity, Spain Manufacturer Price in EURO, EURO Exchange Rates, Spain Price in Rands, Spanish matching pack size in Rands, Comment on Spanish Price Provided, Closest Alternate Country Pack Size, Related Alternate Country Quantity, Manufacturer Price alternate currency , Alternate Currency Exchange Rates, Alternate Country Price in Rand, Alternate Country matching pack

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size in Rands, Comment on Alternate Country Price Provided. Where a medicine does not have a comparator product from Australia, Canada, New Zealand & Spain **all** other countries where the medicine is being sold must be listed and provided as alternate countries.

3.2.3 Where the exact pack size does not exist in the international market, the closest pack size will be used e.g. if there is 30 pack size in South Africa and only 28's and 100's in Spain the 28 pack size will be used as the closest pack to 30's. The related quantity refers to the quantity in which the pack size of the medicine is being sold in that country and allows for a like comparison of the South African medicine.

3.2.4 The exchange rate will be the average over the 12-month period (i.e. 01 November 2022 to 31 October 2023. These values will be published in the template for consistency. The following are the for the conversion to Rands:

AUS$ 12.1562
CAN$ 13.5342
NZD$ 11.2302
EUR€ 19.5976

NOTE: The template with Tab 1 and 2 must always be maintained in the font and format as it appears on DoP. Applicants should only make use of space, dashes or any other characters if these are represented as such in official documentation.

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4. ANNEXURES

4.1 ANNEXURE A: COVER PAGE

TO BE COMPLETED BY THE APPLICANT	
APPLICANT NAME *As it appears on the MCC / SAHPRA license*	
CONTACT PERSON Name: E-mail: Fax No: *(Person responsible for this submission)*	
NUMBER OF MEDICINES IN THE SUBMISSION *(Also include medicines for which SEP adjustment is not requested, rows which contain multiple active ingredients should not be counted.)*	
NUMBER OF ROWS BEING SUBMITTED *(Rows which contain only active ingredients should also be counted.)*	

FOR OFFICE USE ONLY (as per acknowledgement notice)	
Date received: (dd/month/yyyy)	
Received by **(Name and Surname):**	
Signature:	

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4.2 ANNEXURE B: DECLARATION

SEPA DECLARATION

I ....., (full name and surname) in my capacity as.....and having the authority to sign and enter into legally binding agreements on behalf of..... (Name of applicant) hereby certify that:

1. I have read and understood the information and instructions contained in the 2024 SEPA information and instruction document.
2. I have followed the instructions contained in the 2024 information and instruction document in completing the SEPA template.
3. I have correctly calculated unit pricing for all medicines in the applicant's portfolio.
4. I have requested only the SEPA and not any other medicine details amendments for the scheduled medicines in the applicant's portfolio.
5. I have enclosed a signed covering letter on the company letterhead, stating the purpose of this submission.
6. The information supplied in this submission is true and correct. (NB: please provide proof of authorization to sign on behalf of the company)

SIGNATURE (DEPONENT)

1. **.....(CFO name and signature)**
2. **.....(Responsible Pharmacist name and signature)**

The Deponent has acknowledged that he/she knows and understands the contents of this affidavit, which was signed and sworn to before me at .....on this the.....day of..... 2024 and that the regulations contained in Government Gazette Notice No. R 1258 of 21 July 1972 (as amended) has been complied with.

COMMISSIONER OF OATHS

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4.3 ANNEXURE C: CHECKLIST

SEPA CHECKLIST

Tick the appropriate box (√)

HAVE YOU:	YES	NO
a) Read and understood the entire instruction document for 2024 SEPA?		
b) Read, understood, and followed all the instructions in Section 2 and Section 3?		
c) Provided a signed covering letter on a company letterhead stating the purpose of the submission?		
d) Correctly completed the SEPA 2024 template?		
e) Completed the required fields of the covering page (Annexure A)?		
f) Signed the declaration as required, indicating that the information supplied with this application is true and correct (Annexure B)?		
g) Answered yes to all questions in this checklist (Annexure C)?		
h) There are no blanks on Tab 1 and Tab 2		

NOTE: *If any of the answer(s) to the question(s) above is **NO**, the submission will not be considered.*

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GENERAL NOTICES • ALGEMENE KENNISGEWINGS

STATISTICS SOUTH AFRICA

NOTICE 2359 OF 2024

STATISTICS SOUTH AFRICA

THE HEAD: STATISTICS SOUTH AFRICA notifies for general information that the Consumer Price Index is as follows:

Consumer Price Index, Rate **(Base Dec 2021=100)**

Rate: **December 2023 – 5.1**

PROCLAMATION 159 OF 2024

by the

President of the Republic of South Africa

AMENDMENT OF SCHEDULE 3 TO THE PUBLIC SERVICE ACT, 1994: JUDICIAL INSPECTORATE FOR CORRECTIONAL SERVICES

In terms of section 7(5)*(c)* of the Public Service Act, 1994 (promulgated under Proclamation No. 103 of 1994), I hereby amend, at the request of the Minister of Justice and Correctional Services and on the advice of the Minister for Public Service and Administration and the Minister of Finance, with effect from 1 April 2024, Part A of Schedule 3 to the said Act, by the insertion in columns 1, 2 and 3 of Part A of Schedule 3, after the words "Government Technical Advisory Centre", "Head: Government Technical Advisory Centre" and "National Treasury" of the words "Judicial Inspectorate for Correctional Services", "Chief Executive Officer: Judicial Inspectorate for Correctional Services" and "Department of Correctional Services", respectively.

Given under my Hand and the Seal of the Republic of South Africa at CAPE TOWN this 21 day of FEBRUARY Two Thousand and Twenty Four.

President

By Order of the President-in-Cabinet:

Minister of the Cabinet

PROKLAMASIE 159 VAN 2024

van die

President van die Republiek van Suid-Afrika

WYSIGING VAN DEEL A VAN BYLAE 3 BY DIE STAATSDIENSWET, 1994: REGTERLIKE INSPEKTORAAT VIR KORREKTIEWE DIENSTE

Ingevolge artikel 7(5)*(c)* van die Staatsdienswet, 1994 (gepromulgeer deur Proklamasie No. 103 van 1994), wysig ek hierby, op versoek en advies van die Minister van Justisie en Korrektiewe Dienste en op advies van die Minister vir die Staatsdiens en Administrasie en die Minister van Finansies, met ingang van 1 April 2024, Deel A van Bylae 3 by vermelde Wet deur in kolomme 1, 2 en 3, van Deel A van Bylae 3 voor die woorde "Munisipale Infrastruktuur Ondersteunings Agentskap", "Hoof: Munisipaliteit Infrastuktuur Ondersteunings Agentskap" en "Departement van Kooperatiewe Bestuur" onderskeidelik die woorde "Regterlike Inspektoraat vir Korrektiewe Dienste", "Hoof Uitvoerende Beampte: Regterlike Inspektoraat vir Korrektiewe Dienste" en "Departement van Korrektiewe Dienste", in te voeg.

Gegee onder my Hand en die Seel van die Republiek van Suid-Afrika te KAAPSTAD, op hede die 21 dag van FEBRUARIE Tweeduisend Vier en Twintig.

President

Op las van die President-in-Kabinet:

Minister van die Kabinet

BOARD NOTICE 573 OF 2024

DEFINING OF PRODUCTION AREA: BOT RIVER/BOTRIVIER (AMENDMENT)

The Wine Certification Authority, acting under section 6 of the Wine of Origin Scheme published by Government Notice No. R. 1434 of 29 June 1990 hereby –

(a) defines the area specified in the Schedule as a production area (ward) under the name Bot River; and

(b) repeals herewith Board Notice No 111 of 30 November 2007.

OLIVIA POONAH
EXECUTIVE MANAGER: WINE CERTIFICATION AUTHORITY

SCHEDULE

DEFINING OF PRODUCTION AREA: BOT RIVER

That portion of land situate within the following boundaries:

Start at the point where the western boundary of The Draay 563, just east of the township area of Kleinmond, meets the Atlantic Ocean; thence first north and then generally east with the western and northern boundaries of said The Draay 563 to the north-eastern beacon thereof; thence further east with an imaginary straight line to the south-western beacon of Lekker Water 545; thence further east with the southern boundary of said Lekker Water 545 and the northern boundary of The Draay 563 to the north-eastern beacon thereof; thence north-east with the north-western boundary of Hermanus River 542 to the northern beacon thereof; thence generally north-east with the general western boundaries of farms 474, 781, 473, 820 and Compagnies Drift 435 to a point where the N2-freeway intersects the western boundary of said Compagnies Drift 435; thence north-east with the north-western boundary of

said Compagnies Drift 435 to the northern beacon thereof; thence further north-east with an imaginary straight line, which is an extension of the north-western boundary of Compagnies Drift 435, to the northern beacon of farm 791; thence north-west with an imaginary straight line to the south-western beacon of farm 867; thence north with the western boundaries of said farm 767 and farm 866, Krakeel Poort 274 and Goedvertrouw 264 to the north-western beacon thereof; thence north-east for about 1925 meters with the north-western boundary of said Goedvertrouw 264 to a point where an imaginary straight line, drawn from the junction of the secondary roads (south-east of the farm Kanaän) joins it; thence south-east with the secondary road to where it joins the R43-road; thence south-east with said R43-road to where it joins the N2-freeway; thence south-west with said N2-freeway to a point where it intersects the north-eastern boundary of farm 362; thence generally south with said boundary and the eastern boundary of Avontuur 429 to where it joins the north-western boundary of Heuwelkraal 843; thence south-west with said north-western boundary of Heuwelkraal 843 and an imaginary straight line, which is an extension of said boundary, to where it joins the south-eastern beacon of farm 880; thence south-west with the south-eastern boundary of said farm 880 to a point where the Swart River intersects said boundary; thence south-east with said Swart River to a point where it intersects the western boundary of farm 475; thence south with an imaginary straight line to the northern beacon of Paarde Poort 539; thence west to where it joins the eastern boundary of Josephs Kraal 540; thence first north and then generally south with the eastern and general western boundaries of said Josephs Kraal 540 to where it joins the north-eastern beacon of Afdaks Rivier 575; thence generally south-west with the general eastern boundaries of said Afdaks Rivier 575 to a point where the R43-road intersects the south-western boundary thereof; thence north with said R43-road to the township Fisherhaven; thence north-west with the eastern boundary of said township to where it joins the Bot River Lagoon; thence generally south-west with the south eastern boundary of said lagoon to the Atlantic Ocean; thence north-west with the coastline to the point where the western boundary of The Draay 563 meets the Atlantic Ocean, the point of beginning mentioned above.

BOARD NOTICE 574 OF 2024

DEFINING OF PRODUCTION AREA: NIEUWOUDTVILLE (AMENDMENT)

The Wine Certification Authority, acting under section 6 of the Wine of Origin Scheme published by Government Notice No. R. 1434 of 29 June 1990 hereby –

(a) defines the area specified in the Schedule as a production area (ward) under the name Nieuwoudtville; and

(b) repeals herewith Board Notice No 12 of 8 February 2019.

OLIVIA POONAH
EXECUTIVE MANAGER: WINE CERTIFICATION AUTHORITY

SCHEDULE

DEFINING OF PRODUCTION AREA: NIEUWOUDTVILLE

The area situated within the following boundaries:

Start at the northeastern beacon of the farm Cloudskraal 644/1; thence generally west with the general northern boundaries of said Cloudskraal 644/1 to the southwestern beacon of farm 637; thence generally south with an imaginary straight line, drawn from said beacon of farm 637 to the escarpment of the Bokkeveld Mountain; thence generally south with the escarpment to the northwestern beacon of farm 648; thence generally north with the northern boundary of said farm 648, the eastern boundary of farm 645/1, the eastern, northern and western boundaries of farm 645/4, the eastern boundary of farms 644/2, 644 and 644/1 to the north-eastern beacon thereof, the point of beginning mentioned above.

BOARD NOTICE 575 OF 2024

DEFINING OF PRODUCTION AREA: KAROO-HOOGLAND

The Wine Certification Authority, acting under section 6 of the Wine of Origin Scheme published by Government Notice No. R. 1434 of 29 June 1990 hereby –

defines the area specified in the Schedule as a production area (region) under the name Karoo-Hoogland.

OLIVIA POONAH
EXECUTIVE MANAGER: WINE CERTIFICATION AUTHORITY

SCHEDULE

DEFINING OF PRODUCTION AREA: KAROO-HOOGLAND

The area situated within the following boundaries:

The municipal boundary of Karoo-Hoogland was used as the basis for the proposed Karoo-Hoogland (Region) and all the farms that are on average below 1000 m above sea level were taken out.

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