Front Side

280 mm

PRIBEMAK 15
PRIBEMAK 30
PM04339448

PATIENT INFORMATION LEAFLET

SCHEDULING STATUS S5

PRIBEMAK 15
Cyclobenzaprine hydrochloride 15 mg (extended release capsule)
Contains 49,8 mg sugar (sucrose).

PRIBEMAK 30
Cyclobenzaprine hydrochloride 30 mg (extended release capsule)
Contains 99,60 mg sugar (sucrose).

Read all of this leaflet carefully before you start taking PRIBEMAK
- Keep this leaflet. You may need to read it again.
- If you have further questions, please ask your doctor, pharmacist, nurse or other healthcare provider.
- **PRIBEMAK** has been prescribed for you personally and you should not share your medicine with other people. It may harm them, even if their symptoms are the same as yours.

What is in this leaflet

1. What **PRIBEMAK** is and what it is used for
2. What you need to know before you take **PRIBEMAK**
3. How to take **PRIBEMAK**
4. Possible side effects
5. How to store **PRIBEMAK**
6. Contents of the pack and other information

1. What PRIBEMAK is and what it is used for

PRIBEMAK contains cyclobenzaprine.

PRIBEMAK is used along with rest and physiotherapy to help decrease muscle spasms and pain by relaxing the muscles associated with sprains, strains and/or other muscle injuries.
PRIBEMAK should only be used for up to 2 – 3 weeks.

2. What you need to know before you take PRIBEMAK

Do not take PRIBEMAK:
- If you are allergic to cyclobenzaprine or any of the other ingredients of this medicine, listed in **section 6**.
- are taking certain antidepressants, known as monoamine oxidase (MAO) inhibitors (such as isocarboxazid, phenelzine, selegiline or tranylcypromine) or it has been 14 days or less since you stopped taking a MAO inhibitor.
- have had a recent heart attack.
- have heart rhythm problems (dysrhythmias).
- have heart failure.
- have an overactive thyroid (hyperthyroidism).
- elderly.
- have liver problems.

Warnings and precautions
Talk to your doctor before taking **PRIBEMAK**
- Tell your doctor if you are taking other medicines used to treat depression, mood, anxiety, psychotic or thought disorders as they may cause a serious reaction when taken with **PRIBEMAK**. Agitation, restlessness, hallucinations, insomnia, feeling confused, a rapid heart rate and high blood pressure, dilated pupils or loss of muscle co-ordination or twitching muscles (these may be signs of a serious condition called serotonin syndrome).
- have a history of eye problems including glaucoma.
- have trouble emptying your bladder (urinary retention).
- if you suffer from epilepsy (seizures).

Use in children and adolescents
The use of **PRIBEMAK** in children and adolescents (below the age of 18 years) is not recommended.

Other medicines and PRIBEMAK
Always tell your healthcare professional if you are taking any other medicine. (This includes complementary or traditional medicines.)
Tell your doctor or pharmacist if you are taking any of the following medicines:
- A medicine used to treat depression, mood, anxiety, psychotic or thought disorders, as they may cause a serious and life threatening reaction when taken with **PRIBEMAK**.
- A pain medicine called tramadol or meperidine, as it may increase the risk of seizures.
- A medicine to help quit smoking called bupropion, as it may increase the risk of side effects.
- A blood pressure medicine called verapamil, as **PRIBEMAK** may reduce the effect of verapamil.

PRIBEMAK with food and drink:
PRIBEMAK may enhance the effects of alcohol and increase side effects. Alcohol should be avoided.

Pregnancy, breastfeeding and fertility

If you are pregnant or breastfeeding your baby, think you may be pregnant or are planning to have a baby please consult your doctor, pharmacist or other healthcare professional for advice before taking **PRIBEMAK**.

PRIBEMAK should not be used during pregnancy.
PRIBEMAK should not be used during breastfeeding.

Driving and using machines
PRIBEMAK frequently causes drowsiness and dizziness, patients using **PRIBEMAK** should not drive or operate machinery.
It is not always possible to predict to what extent **PRIBEMAK** may interfere with the daily activities of a patient. Patients should ensure that they do not engage in the above activities until they are aware of the measure to which **PRIBEMAK** affects them.

PRIBEMAK contains sucrose
If you have been told by your doctor that you have an intolerance to some sugars, contact your doctor before taking this medicinal product.

3. How to take PRIBEMAK

Do not share medicines prescribed for you with any other person.

Always take **PRIBEMAK** exactly as your doctor has instructed you. You should check with your doctor or pharmacist if you are not sure.

The recommended dose is 15 mg (1 capsule) daily.
Some patients may require up to 30 mg/day given as one 30 mg capsule once daily or 2 x 15 mg capsules taken once daily.
PRIBEMAK must be taken orally, at the same time each day.
PRIBEMAK should only be taken for short periods of time.
Your doctor will tell you how much to take and for how long you should take it.
If you have the impression that the effect of **PRIBEMAK** is too strong or too weak, talk to your doctor or pharmacist.

If you take more PRIBEMAK than you should

In the event of over dosage, consult your doctor or pharmacist. If neither is available, contact the nearest hospital or poison control centre.

If you forget to take PRIBEMAK
It is important to take your **PRIBEMAK** regularly at the same time each day. If you forget to take a dose, take it as soon as you remember unless it is time for your next dose. Do not take two doses at the same time, to make up for a missed dose.

If you stop taking PRIBEMAK
If **PRIBEMAK** is stopped earlier than your doctor has prescribed, your symptoms may return or worsen. Always discuss with your doctor before stopping **PRIBEMAK**.

4. Possible side effects

PRIBEMAK can have side effects.
Not all side effects reported for **PRIBEMAK** are included in this leaflet. Should your general health worsen or if you experience any untoward effects while taking **PRIBEMAK**, please consult your doctor, pharmacist or other healthcare professional for advice.

If any of the following happens, stop taking **PRIBEMAK** and tell your doctor immediately or go to the casualty department at your nearest hospital:
- Difficulty breathing, hives, swelling of your face or tongue, itching.
- Agitation, restlessness, hallucinations, insomnia, feeling confused, a rapid heart rate and high blood pressure, dilated pupils or loss of muscle co-ordination or twitching muscles (these may be signs of a serious condition called serotonin syndrome).
- A faster heartbeat, chest pain, dizziness or lightheadedness.

These are all very serious side effects. If you have them, you may have had a serious reaction to **PRIBEMAK**. You may need urgent medical attention or hospitalisation.

Tell your doctor if you notice any of the following:
Frequently occurring:
- Dry mouth
- Dizziness
- Fatigue
- Constipation
- Nausea
- Upset stomach
- Drowsiness
- Headache

Less frequent
- Feeling anxious or depressed, abnormal thoughts or dreams, hallucinating, becoming excited
- Fits, muscle twitches, inability to sleep
- Sensation of ringing in the ears

Frequency unknown:
- Unexplained bleeding or bruising
- Increased or decreased sugar levels (you may experience increased thirst, tiredness, dry mouth, blurred vision or feeling shaky, irritable, sweating, chills, lightheaded, dizziness, confusion and hungry).
- Weight gain or loss
- Increased or decreased blood pressure
- Swelling of ankles or feet
- Hepatitis, jaundice, sweating, light sensitivity, itching, facial swelling, rash, frequent urination, enlarged breasts.

If you notice any side effects not mentioned in this leaflet, please inform your doctor or pharmacist.

Reporting of side effects
If you get side effects, talk to your doctor or pharmacist or nurse. This includes any possible side effects not listed in this leaflet. You can also report side effects to SAHPRA via the Med Safety APP (Medsafety X SAHPRA) and eReporting platform (who-umc.org) found on SAHPRA website. By reporting side effects, you can help provide more information on the safety of **PRIBEMAK.**

5. How to store PRIBEMAK

Store at or below 25 °C.
KEEP OUT OF THE REACH OF CHILDREN.
Store in the original package / container.
Keep the blisters in the carton until required for use.
Do not store in a bathroom.
Do not use after the expiry date stated on the blister / label / carton bottle.
Return all unused medicine to your pharmacist.
Do not dispose of unused medicine in drains or sewerage systems (e.g. toilets).

6. Contents of pack and other information

What PRIBEMAK contains
The active ingredient is cyclobenzaprine.
The other ingredients are:
Sugar Spheres
Hypromellose
Isopropyl Alcohol
Methylene Chloride
Colloidal Silicon dioxide (Aerosil 200)
Ethyl Cellulose

PRIBEMAK 15 contains gelatin, water, titanium dioxide, Sodium lauryl sulfate, D & C Yellow 10 (E104), FD & C Red 40 (E129) and FD & C Green 3 (E143)
PRIBEMAK 30 contains gelatin, Sodium lauryl sulfate, water, titanium dioxide, FD & C Red 40 (E129), FD & C Yellow 6 (E110) and FD & C Blue 1 (E133)

What PRIBEMAK looks like and contents of the pack
PRIBEMAK 15: Each extended-release capsule contains 15 mg cyclobenzaprine.
PRIBEMAK 30: Each extended-release capsule contains 30 mg cyclobenzaprine.

Contents of the pack:
HDPE pack:
Capsules are packed in 60 ml round, white, HDPE container, 33 mm neck finish with 33 mm polypropylene child-resistant closure with Pulp and white-printed Heat Seal Liner.
Blister pack:
Capsules are packed in aluminium foil laminated on one side and laminated to clear PVC on the other side. Pack sizes include 10s, 15s, 30s and 60s capsules.
Not all pack sizes may be marketed.

Holder of Certificate of Registration
MACLEODS PHARMACEUTICALS SA (PTY) LTD
Office Block 1,
Bassonia Estate Office Park (East),
1 Cussonia Drive,
Bassonia Rock Ext 12
Alberton
Gauteng

This leaflet was last revised in 20 May 2025

Registration numbers
PRIBEMAK 15: 57/17.3/0304
PRIBEMAK 30: 57/17.3/0305

Access to the corresponding Professional Information
The corresponding professional information can be accessed at https://www.macleodssaproducts.co.za or email safety@macleodspharma.com or contact 011 682 1169.

335 mm

Back Side

PASIËNTINLIGTINGSTUK

SKEDULERINGSTATUS: S5

PRIBEMAK 15
Siklobensaprienhidrochloried 15 mg (verlengde vrystellings kapsule)
Bevat 49,8 mg suiker (sukrose).

PRIBEMAK 30
Siklobensaprienhidrochloried 30 mg (verlengde vrystellings kapsule)
Bevat 99,60 mg suiker (sukrose).

Lees hierdie hele inligtingstuk noukeurig deur voordat jy PRIBEMAK begin neem
- Hou hierdie inligtingstuk. Dit mag nodig wees dat jy dit weer moet lees.
- Indien jy verdere vrae het, vra asseblief jou dokter, apteker, verpleegkundige of ander gesondheidsorgverskaffer.
- **PRIBEMAK** is vir jou persoonlik voorgeskryf, en jy moet nie jou medisyne met ander persone deel nie. Dit mag skadelik vir hulle wees, selfs al is hul simptome dieselfde as joune.

Wat is in hierdie inligtingstuk

1. Wat **PRIBEMAK** is en waarvoor dit gebruik word
2. Wat jy behoort te weet voordat jy **PRIBEMAK** neem
3. Hoe om **PRIBEMAK** te neem
4. Moontlike newe-effekte
5. Hoe om **PRIBEMAK** te bewaar
6. Inhoud van die pak en ander inligting

1. Wat PRIBEMAK is en waarvoor dit gebruik word

PRIBEMAK bevat siklobensaprien.

PRIBEMAK word saam met rus en fisioterapie gebruik om spierspasmas en pyn te verminder deur die spiere te ontspan wat geassosieer word met verstuitings, verrekkings en/of ander spierbeserings.
PRIBEMAK moet slegs vir tot en met 2 – 3 weke gebruik word.

2. Wat jy behoort te weet voordat jy PRIBEMAK neem

Moet nie PRIBEMAK neem nie:
- Indien jy allergies is vir siklobensaprien of enige van die ander bestanddele van hierdie medisyne, gelys in afdeling 6.
- sekere antidepressante neem, bekend as monoamien-oksidase (MAO) inhibeerders (soos isokarboksasied, fenelsien, selegilien of tranielsipromien) of dit is 14 dae of minder sedert jy opgehou het om 'n MAO-inhibeerder te neem.
- 'n onlangse hartaanval gehad het.
- hartritme probleme het (disritmieë).
- hartversaking het.
- 'n ooraktiewe skildklier (hipertireose) het.
- 'n Bejaarde is.
- Lewerprobleme het.

Waarskuwings en voorsorgmaatreëls
Praat met jou dokter voordat jy **PRIBEMAK** neem
- Vertel jou dokter as jy ander medisyne neem wat gebruik word om depressie, buie, angstigheid, psigotiese- of denkversteurings te behandel, aangesien dit 'n ernstige reaksie kan veroorsaak wanneer dit saam met **PRIBEMAK** geneem word. Opgewondenheid, rusteloosheid, hallusinasies, slapeloosheid, verward voel, 'n vinnige harttempo en hoë bloeddruk, verwyde pupille of verlies aan spierkoördinasie of spiertrekkings (dit kan tekens wees van 'n ernstige toestand genaamd serotoniensindroom).
- 'n geskiedenis van oogprobleme het, insluitend gloukoom.
- sukkel om jou blaas leeg te maak (urinêre retensie).
- as jy aan epilepsie ly (aanvalle).

Gebruik in kinders en adolessente
Die gebruik van **PRIBEMAK** in kinders en adolessente (onder die ouderdom van 18 jaar) word nie aanbeveel nie.

Ander medisyne en PRIBEMAK
Vertel altyd jou professionele gesondheidsorgwerker as jy enige ander medisyne gebruik. (Dit sluit komplementêre of tradisionele medisyne in.)
Vertel jou dokter of apteker as jy enige van die volgende medisyne neem:
- 'n Medisyne wat gebruik word om depressie, buie, angstigheid, psigotiese- of denkversteurings te behandel, aangesien dit 'n ernstige en lewensbedreigende reaksie kan veroorsaak wanneer dit saam met **PRIBEMAK** geneem word.
- 'n Pynmedisyne genaamd tramadol of meperidien.
- 'n Medisyne om jou te help om op te hou rook naamlik bupropioon, aangesien dit die risiko van newe-effekte kan verhoog.
- 'n Bloeddrukmedisyne genaamd verapamil, aangesien **PRIBEMAK** die effek van verapamil kan verminder.

PRIBEMAK met voedsel en drank:
PRIBEMAK kan die effekte van alkohol versterk en newe-effekte verhoog. Alkohol moet vermy word.

Swangerskap, borsvoeding en fertiliteit
Indien jy swanger is of jou baba borsvoed, vermoed dat jy moontlik swanger is of beplan om 'n baba te hê, raadpleeg asseblief jou dokter, apteker of ander professionele gesondheidsorgwerker vir advies voordat jy **PRIBEMAK** neem.

PRIBEMAK moet nie tydens swangerskap gebruik word nie.
PRIBEMAK moet nie tydens borsvoeding gebruik word nie.

Bestuur en gebruik van masjinerie

PRIBEMAK veroorsaak gereeld lomerigheid en duiseligheid, pasiënte wat **PRIBEMAK** gebruik, moet nie bestuur of masjinerie hanteer nie.
Dit is nie altyd moontlik om te voorspel tot watter mate **PRIBEMAK** met die daaglikse aktiwiteite van 'n pasiënt kan inmeng nie. Pasiënte moet seker maak dat hulle nie aan die bogenoemde aktiwiteite deelneem nie, totdat hulle bewus is van die mate waartoe **PRIBEMAK** hulle affekteer.

PRIBEMAK bevat sukrose
Indien jy deur jou dokter ingelig is dat jy 'n onverdraagsaamheid teenoor sommige suikers het, kontak jou dokter voordat jy hierdie medisinale produk neem.

3. Hoe om PRIBEMAK te neem

Moet nie medisyne wat vir jou voorgeskryf is met enige ander persoon deel nie.

Neem altyd **PRIBEMAK** presies soos wat jou dokter vir jou gesê het. Bevestig met jou dokter of apteker indien jy nie seker is nie.

Die aanbevole dosis is 15 mg (1 kapsule) daagliks.
Sommige pasiënte benodig moontlik tot en met 30 mg/dag gegee as een 30 mg kapsule een keer per dag of 2 x 15 mg kapsules wat een keer per dag geneem word.
PRIBEMAK moet mondelings, op dieselfde tyd elke dag ingeneem word.
PRIBEMAK moet net vir kort tydperke geneem word.
Jou dokter sal jou vertel hoeveel om te neem en vir hoe lank jy dit moet neem.
Indien jy die indruk kry dat die effek van **PRIBEMAK** te sterk of te swak is, praat met jou dokter of apteker.

Indien jy meer PRIBEMAK neem as wat jy moet
In die geval van oordosering, raadpleeg jou dokter of apteker. Indien nie een beskikbaar is nie, kontak die naaste hospitaal of gifbeheersentrum.

Indien jy vergeet om PRIBEMAK te neem
Dit is belangrik om jou **PRIBEMAK** gereeld op dieselfde tyd elke dag te neem. As jy vergeet om 'n dosis te neem, vat dit sodra jy daarvan onthou, tensy dit tyd is vir jou volgende dosis. Moet nie twee dosisse op dieselfde tyd neem, om op te maak vir 'n vergete dosis nie.

Indien jy ophou om PRIBEMAK te neem
Wanneer **PRIBEMAK** vroeër gestaak word as wat jou dokter voorgeskryf het, kan jou simptome terugkeer of vererger. Bespreek altyd met jou dokter voordat jy **PRIBEMAK** staak.

4. Moontlike newe-effekte

PRIBEMAK kan newe-effekte hê.
Nie alle newe-effekte wat vir **PRIBEMAK** gerapporteer is, is in hierdie inligtingstuk ingesluit nie. Sou jou algemene gesondheid vererger of indien jy enige ongunstige effekte ervaar terwyl jy **PRIBEMAK** neem, raadpleeg asseblief jou dokter, apteker of ander professionele gesondheidsorgwerker vir advies.

Indien enige van die volgende gebeur, hou op om **PRIBEMAK** te neem en kontak onmiddellik jou dokter of gaan na die ongevalle-afdeling van jou naaste hospitaal:
- Probleme met asemhaling, galbulte, swelling van jou gesig of tong, gejeuk.
- Opgewondenheid, rusteloosheid, hallusinasies, slapeloosheid, verward voel, 'n vinnige harttempo en hoë bloeddruk, verwyde pupille of verlies aan spierkoördinasie of spiertrekkings (dit kan tekens wees van 'n ernstige toestand genaamd serotoniensindroom).
- 'n Vinniger hartklop, borskaspyn, duiseligheid of lighoofdigheid.

Hierdie is alles baie ernstige newe-effekte. Indien jy dit kry, het jy moontlik 'n ernstige reaksie op **PRIBEMAK** gehad. Jy benodig dalk dringende mediese aandag of hospitalisering.

Vertel jou dokter indien jy enige van die volgende opmerk:
Kom dikwels voor:
- Droë mond
- Duiseligheid
- Moegheid
- Hardlywigheid
- Naarheid
- Omgekrapte maag
- Lomerigheid
- Hoofpyn

Minder gereeld
- Voel angstig of depressief, abnormale gedagtes of drome, hallusineer, raak opgewonde
- Stuipe, spiertrekkings, onvermoë om te slaap
- Sensasie van suising in die ore

Frekwensie onbekend:
- Onverklaarbare bloeding of kneusplekke
- Verhoogde of verlaagde suikervlakke (jy kan verhoogde dors, moegheid, droë mond, dowwe visie ervaar of wankelrig voel, prikkelbaar, sweet, koue rillings, lighoofdig, duiseligheid, verwarring en honger voel).
- Gewigstoename of -verlies
- Verhoogde of verlaagde bloeddruk
- Swelling van enkels of voete
- Hepatitis, geelsug, sweet, ligsensitiwiteit, gejeuk, gesigswelling, uitslag, gereelde urinering, vergrote borste.

Indien jy enige newe-effekte opmerk wat nie in hierdie inligtingstuk genoem word nie, stel asseblief jou dokter of apteker in kennis.

Aanmelding van newe-effekte
Indien jy newe-effekte kry, praat met jou dokter of apteker of verpleegkundige. Dit sluit enige moontlike newe-effekte in wat nie in hierdie inligtingstuk gelys word nie. Jy kan ook newe-effekte aan SAHPRA rapporteer via die Med Safety APP (Medsafety X SAHPRA) en eReporting platform (who-umc.org) wat op SAHPRA se webwerf gevind word.
Deur newe-effekte aan te meld kan jy help om meer inligting beskikbaar te stel aangaande die veiligheid van **PRIBEMAK.**

5. Hoe om PRIBEMAK te bewaar

Berg by of benede 25 °C.
HOU BUITE BEREIK VAN KINDERS.
Bewaar in die oorspronklik verpakking / houer.
Hou die blister in die karton totdat dit vir gebruik benodig word.
Moet nie in 'n badkamer stoor nie.
Moet nie gebruik na die vervaldatum wat op die stulptrook / etiket / karton / bottel aangedui word nie.
Neem alle ongebruikte medisyne terug na jou apteker.
Moet nie ongebruikte medisyne in dreine of rioolstelsels (bv. toilette) weggooi nie.

6. Inhoud van die pak en ander inligting

Wat PRIBEMAK bevat
Die aktiewe bestanddeel is siklobensaprien.
Die ander bestanddele is:
Suiker Sfere
Hipromellose
Isopropielalkohol
Metileenchloried
Kolloïdale silikondioksied (Aerosil 200)
Etielsellulose

PRIBEMAK 15 bevat gelatien, water, titaandioksied, natriumlaurielsulfaat, D & C Geel 10 (E104), FD & C Rooi 40 (E129) en FD & C Groen 3 (E143)
PRIBEMAK 30 bevat gelatien, natriumlaurielsulfaat, water, titaandioksied, FD & C Rooi 40 (E129), FD & C Geel 6 (E110) en FD & C Blou 1 (E133)

Hoe PRIBEMAK lyk en inhoud van die pak
PRIBEMAK 15: Elke verlengde-vrystellings kapsule bevat 15 mg siklobensaprien.
PRIBEMAK 30: Elke verlengde-vrystellings kapsule bevat 30 mg siklobensaprien.

Inhoud van die pak:
HDPE-pak:
Kapsules word verpak in 60 ml ronde, wit, HDPE-houer, 33 mm nekafwerking met 33 mm polipropileen kinderbestande sluiting met Pulp en wit gedrukte hitteseëlvoering.
Stulpstrook pak:
Kapsules word verpak in aluminiumfoelie wat aan die een kant gelamineer is en gelamineer is tot deursigtige PVC aan die ander kant.
Verpakkingsgroottes sluit 10's, 15's, 30's en 60's kapsules in.
Nie alle verpakkingsgroottes word noodwendig bemark nie.

Houer van Sertifikaat van Registrasie
MACLEODS PHARMACEUTICALS SA (EDMS) BPK
Kantoorblok 1,
Bassonia Estate Kantoorpark (Oos),
1 Cussonia-rylaan ,
Bassonia Rock, Uitbr. 12,
Alberton
Gauteng

Hierdie inligtingstuk was laas hersien in 20 Mei 2025

Registrasienommers
PRIBEMAK 15 mg: 57/17.3/0304
PRIBEMAK 30 mg: 57/17.3/0305

Toegang tot die ooreenstemmende Professionele Inligting
Die ooreenstemmende professionele inligting kan verkry word by https://www.macleodssaproducts.co.za of e-pos safety@macleodspharma.com of kontak 011 682 1169.

280 mm

335 mm

PACKAGING RESEARCH & DEVELOPMENT

Restricted Circulation

(Front Page)

ARTWORK / PRINTPROOF APPROVAL FORM

1. Product: PRIBEMAK 15/30 (Cyclobenzaprine ER 15/30 mg capsules)	
2. Country: South Africa(Private South Africa)	**3. Packaging component: Patient Information**
4. No. of Colours: 1 Colour	**5. Pantone Shade No.: Black**
6. Specification: 40 gsm bible paper	
7. Machine Layout No.: NA	**8. Dimensions: 335 (L) x 280 (W) mm**
9. Design / Style: Booklet	**10. Item code No.: PM04339448**
11. Artwork has been made: Actual Size	**12. Pharma code No.: 40307 mini**
13. Barcode/NDC No: NA	**14. Supersedes Item code No: NA**
15. Revision no: 00	**16. A/W File name:Z:\South Africa\Cyclobenzaprine ER Capsules\Cyclobenzaprine ER Capsules PIL - PM04339448.cdr**
Approved By:	**17. Location: Baddi**

PRD	M.A. HOLDER	RAD	QC	PRODN	QA
	30/07/2025 				

Note: 1. To follow overleaf checklist for approval of Artwork / Print proof.

Instructions to Vendor :

1. Before processing, please ensure First pull out / Commercial supply should match with approved Artwork with respect to Design, Size, Fonts & Text.
2. Print proof to represent full printing area with Station Numbers.

FRM/CQA/021/04-01

Reason for revision :	Item code	Artwork Send for Printing Date	Artwork Implementation Date
1) New Artwork for South Africa (Private South Africa)	PM04339448		