APPROVED PATIENT INFORMATION LEAFLET

SCHEDULING STATUS: S4

SONKE LAMIVUDINE 150 Tablets

Lamivudine

Sugar-free

Read all of this leaflet carefully before you start taking SONKE LAMIVUDINE 150

- Keep this leaflet. You may need to read it again.
- If you have further questions, please ask your doctor, pharmacist, nurse or other health care provider.
- **SONKE LAMIVUDINE 150** 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 **SONKE LAMIVUDINE 150** is and what it is used for
2. What you need to know before you take **SONKE LAMIVUDINE 150**
3. How to take **SONKE LAMIVUDINE 150**
4. Possible side effects
5. How to store **SONKE LAMIVUDINE 150**
6. Contents of the pack and other information

1. What SONKE LAMIVUDINE 150 is and what it is used for

SONKE LAMIVUDINE 150 tablets are used as part of the combination treatment of HIV infection in adults and children above the age of 12 years old.

2. What you need to know before you take SONKE LAMIVUDINE 150

Do not take SONKE LAMIVUDINE 150 tablets if

- You have previously had an allergic reaction to lamivudine or to any of **SONKE LAMIVUDINE 150** ingredients listed above. (An allergic reaction may include rash, itching, swelling of the face, lips, tongue or hands/feet or breathing difficulties) (listed in section 6).
- You have blood disorders
- You are pregnant or breastfeeding

Warnings and precautions

Take special care with SONKE LAMIVUDINE 150:

- If you have a history of liver disease, including chronic active hepatitis. Patients with liver disease including chronic hepatitis B or C, who are treated with combination antiretrovirals, have a higher risk of severe and potentially life-threatening liver problems. Your doctor may conduct blood tests in order to check how well your liver is working or may switch you to another medicine. If you have severe liver disease, do not take SONKE LAMIVUDINE 150.
- If you have hepatitis B infection, your doctor will carefully consider the best treatment regimen for you. Symptoms of your hepatitis may become worse after discontinuation of SONKE LAMIVUDINE 150.
- If you develop stomach pain, nausea or vomiting. Check with your doctor, as these could be signs of an inflamed pancreas or a condition called lactic acidosis. If you are feeling unwell, consult your doctor.
- Liver disease has been known to occur.
- If you have kidney disease.
- If you have lipodystrophy (a problem with the way your body uses and stores fat). This condition is characterised by being overweight (obesity), build-up of fat on the back of the neck between the shoulders (buffalo hump), loss of fat under the skin (making the face look thin), breast enlargement and high levels of fat and glucose (sugar) in the blood. If you have any of these signs, you should be thoroughly examined by your doctor.
- In rare cases, as your immune system becomes stronger, it can also attack healthy body tissue (autoimmune disorders). The symptoms of autoimmune disorders may develop many months after you start taking medicine to treat your HIV infection.

- If you notice any of the symptoms of osteonecrosis (where parts of the bone tissue die, because of reduced blood supply to the bone): joint aches and pain, joint stiffness or difficulty in movement, tell your doctor.
- You may still develop infections and other complications of HIV infection while using **SONKE LAMIVUDINE 150.**
- You are still at a risk of transmission of HIV to other people through sexual contact or blood contamination, while taking **SONKE-LAMIVUDINE 150.**
- You should stop **SONKE-LAMIVUDINE 150** tablets immediately if you develop abdominal (intestinal) pain, nausea or vomiting, shortness of breath, tiredness and weight loss and you should contact your doctor.

SONKE-LAMIVUDINE 150 must be used with caution in patients with liver and kidney disorders as well as in the elderly.

Other medicines and SONKE-LAMIVUDINE 150

Always tell your health care provider if you are taking any other medicine. (This includes all complementary or traditional medicines).

If you are taking other medicines on a regular basis, including complimentary or traditional medicines, concomitant use of SONKE-LAMIVUDINE 150 with these medicines may cause undesirable interactions. Please consult your doctor, pharmacist or other health care professional for advice.

Care is needed if you are taking the following medicines along with **SONKE LAMIVUDINE 150:**

- Trimethoprim/sulphamethoxazole (a medicine to treat infections)
- Zalcitabine
- Emtricitabine (used to treat HIV infection).
- Cladribine (used to treat a certain type of cancer).
- Medicines (usually liquids) containing sorbitol and other sugar alcohols (such as xylitol,)

SONKE-LAMIVUDINE 150 with food or drink

SONKE LAMIVUDINE 150 can be taken with or without food.

Pregnancy and breastfeeding

You must not use SONKE LAMIVUDINE 150 when pregnant or breastfeeding your infant.

If you are pregnant or breastfeeding your baby while taking this medicine, please consult your doctor, pharmacist or other health care professional for advice.

Driving and using machines

SONKE LAMIVUDINE 150 may cause dizziness or tiredness in some patients. Do not drive or operate any tools or machines until you know how **SONKE LAMIVUDINE 150** affects you because your concentration may be affected.

3. How to take SONKE-LAMIVUDINE 150

Do not share medicines prescribed for you with other persons.

Always take SONKE LAMIVUDINE TABLETS exactly as your doctor or pharmacist has told you. You should check with your doctor or pharmacist if you are unsure.

Adults and adolescents: Take two tablets once a day or one tablet twice daily in combination with zidovudine.

Children weighing between 14 kg to 20 kg:

Take one tablet once a day.

Children weighing at least 25 kg:

Take one tablet twice a day or take two tablets once a day.

If you have liver and kidney problems, your doctor may adjust the doses accordingly.

To help you remember to take **SONKE LAMIVUDINE 150**, try to get into the habit of taking it at the same time each day.

Take **SONKE LAMIVUDINE 150** as directed and for as long as directed; do not stop them, even if you feel better, as the symptoms may return.

*If you have the impression that the effect of **SONKE LAMIVUDINE 150** tablets is too strong or too weak, talk to your doctor or pharmacist.*

If you have taken more SONKE LAMIVUDINE 150 than you should

In the event of overdosage, consult your doctor or pharmacist. If neither is available, rush the patient to the nearest hospital or poison centre. Take this leaflet or some tablets with you so your doctor will know what you have taken.

If you forget to take SONKE LAMIVUDINE 150 tablets

Take them as soon as you remember. However, if it is almost time for your next dose, skip the missed dose and go back to your regular dosing schedule. Do not take a double dose to make up for forgotten individual doses.

4. Possible side effects

SONKE LAMIVUDINE 150 can have side effects.

Not all side-effects reported for **SONKE LAMIVUDINE 150** are included in this leaflet. Should your general health worsen while taking **SONKE LAMIVUDINE 150**, please consult your doctor, pharmacist or other health care professional for advice.

If any of the following happens, stop taking **SONKE LAMIVUDINE 150** and tell your doctor immediately or go to the casualty department at your nearest hospital:

- Rashes, hives, itching, chest constriction, shortness of breath or swelling of the face, lips, tongue, hands/ feet, fainting, high temperature (fever)

These are all very serious side effects. If you have them, you may have had a serious reaction to **SONKE LAMIVUDINE 150**. You may need urgent medical attention or hospitalisation.

Tell your doctor immediately or go to the casualty department at your nearest hospital if you notice any of the following:

- A breakdown of muscle tissue that releases a damaging protein into the blood (rhabdomyolysis); symptoms include: dark, reddish urine, a decreased amount of urine, weakness and muscle aches.

- Excess lactic acid in the blood (lactic acidosis); symptoms include: rapid breathing, excessive sweating, cool and clammy skin, sweet-smelling breath, stomach pain, nausea or vomiting, confusion and tiredness.
- Inflammation of the pancreas (pancreatitis); symptoms include severe stomach pain with nausea and vomiting and fever.
- Inflammation of the liver (hepatitis); symptoms include: yellow skin or eyes (jaundice), nausea, fatigue and fever.

These are all serious side effects. You may need urgent medical attention

Tell your doctor if you notice any of the following side effects:

Frequent side effects:

- Nausea, vomiting, stomach pain or cramps, diarrhoea
- Unusual weakness or tiredness
- Fever, general feeling of illness
- Difficulty in sleeping
- Headache, sensation of pins and needles, tingling, burning, numbness
- Muscle pain or cramping
- Joint pain
- Cough, irritated or runny nose

Side effects with frequency unknown:

- Loss of hair

Side effects that may show up in your blood tests:

- A reduction in the number of white blood cells (neutropenia)
- A reduction in the number of platelets (cells that make the blood clot) (thrombocytopenia)
- A reduction in the number of red blood cells (anaemia)
- A failure of the bone marrow to produce new red blood cells (pure red cell aplasia)
- An increase in liver enzymes

- An increase in the level of an enzyme called amylase in the blood

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, pharmacist or nurse. You can also report side effects to SAHPRA via the "**6.04 Adverse Drug Reaction Reporting Form**", found online under SAHPRA's publications: https://www.sahpra.org.za/Publications/Index/8. By reporting side effects, you can help provide more information on the safety of **SONKE LAMIVUDINE 150**

5. How to store SONKE LAMIVUDINE 150

Store all medicines out of reach of children.

- Store at or below 25 °C.
- Store in the original package.
- Keep the blister in the outer carton.

- Protected from light/moisture.
- Do not store in a bathroom.
- Do not use after the expiry date stated on the label.
- Return all unused medicine to your pharmacist.
- Do not dispose of unused medicine in drains or sewerage systems e.g. toilets.

6. Contents of the pack and other information

What SONKE LAMIVUDINE 150 contains

Each SONKE LAMIVUDINE 150 tablet contains 150 mg lamivudine. (active substance).

The other ingredients are:

Tablet core: Magnesium stearate, microcrystalline cellulose; sodium starch glycollate

Flim coat: Hypromellose; titanium dioxide; propylene glycol.

What SONKE LAMIVUDINE 150 looks like and contents of the pack

White to off-white, oval shaped, biconvex, film-coated tablets with 'RX919' debossed on one side and a score line on the other side.

10 tablets are packed in blister strips of clear, transparent, PVC film with an aluminium foil backing. Cartons contain 10, 30 or 60 tablets.

56 or 60 Tablets packed in securitainers or in white opaque HDPE bottles.

Holder of Certificate of Registration

RANBAXY PHARMACEUTICALS (PTY) LTD

a Sun Pharma company

14 Lautre Road, Stormill Ext 1

Roodepoort, 1724

South Africa

This leaflet was last revised in

Date of registration: 10 August 2007

Date of revision: 25 January 2023

Registration number

A40/20.2.8/027

Marketed by Sonke Pharmaceuticals (Pty) Ltd

This product is for use only in the territories of South Africa, Namibia, Botswana, Lesotho and Swaziland, is not for resale and any other use is not authorised.

Botswana: S2 Reg. no.: BOT0801207

Namibia: NS2 Reg. no.: 05/20.2.8/0416

GOEDGEKEURDE PASIËNTINLIGTINGSTUK

SKEDULERINGSTATUS: S4

SONKE LAMIVUDINE 150 Tablette

Lamivudien

Suikervry

Lees hierdie hele inligtingstuk noukeurig deur voordat jy begin om SONKE LAMIVUDINE 150 te neem.

- Hou hierdie inligtingstuk. Dit mag vir jou nodig wees om dit weer te lees.
- Vra asseblief jou dokter, apteker, verpleegkundige of ander gesondheidsorgverskaffer as jy verdere vrae het.
- **SONKE LAMIVUDINE 150** is vir jou persoonlik voorgeskryf en jy moenie jou medisyne met ander persone deel nie. Dit kan skadelik vir hulle wees, selfs al is hulle simptome dieselfde as joune.

Wat is in hierdie inligtingstuk

1. Wat **SONKE LAMIVUDINE 150** is en waarvoor dit gebruik word
2. Wat jy behoort te weet voordat jy **SONKE LAMIVUDINE 150** neem
3. Hoe om **SONKE LAMIVUDINE 150** te neem
4. Moontlike newe-effekte
5. Hoe om **SONKE LAMIVUDINE 150** te bewaar
6. Inhoud van die verpakking en ander inligting

1. Wat SONKE LAMIVUDINE 150 is en waarvoor dit gebruik word

SONKE LAMIVUDINE 150 tablette word gebruik as deel van die kombinasie-behandeling van MIV-infeksie in volwassenes en kinders bo die ouderdom van 12 jaar.

2. Wat jy behoort te weet voordat jy SONKE LAMIVUDINE 150 neem

Moenie SONKE LAMIVUDINE 150 tablette neem nie as

- jy voorheen ‘n allergiese reaksie gehad het teenoor lamivudien of teenoor enige van die ander bestanddele van **SONKE LAMIVUDINE 150** (gelys in afdeling 6).

  (‘n Allergiese reaksie kan uitslag, gejeuk, swelling van die gesig, lippe, tong of hande/voete of asemhalingsprobleme insluit).
- jy bloedversteurings het
- jy swanger is of borsvoed

Waarskuwings en voorsorgmaatreëls

Tref spesiale voorsorg met **SONKE LAMIVUDINE 150:**

- As jy ‘n geskiedenis van lewersiekte het, insluitend chroniese aktiewe hepatitis. Pasiënte met lewersiekte, insluitend chroniese hepatitis-B of -C, wat met kombinasie-antiretrovirale middels behandel word, het ‘n hoër risiko van ernstige en potensieel lewensbedreigende lewerprobleme. Jou dokter mag bloedtoetse uitvoer om vas te stel hoe goed jou lewer funksioneer of mag jou na ‘n ander medisyne omskakel. As jy ernstige lewersiekte het moet jy nie **SONKE LAMIVUDINE 150** neem nie.
- As jy hepatitis-B-infeksie het, sal jou dokter die beste behandelingsregimen vir jou oorweeg. Simptome van jou hepatitis mag vererger nadat jy opgehou het om **SONKE LAMIVUDINE 150** te neem.
- As jy maagpyn, naarheid of braking ontwikkel. Kontroleer met jou dokter, aangesien hierdie tekens van ‘n ontsteekte pankreas of ‘n toestand wat melksuurasidose genoem word, mag wees. Raadpleeg jou dokter as jy nie goed voel nie.
- Dit is bekend dat lewersiekte kan voorkom.
- As jy niersiekte het.
- As jy lipodistrofie het (‘n probleem met die wyse waarop jou liggaam vet gebruik en stoor). Hierdie toestand word gekenmerk deur oorgewig (vetsugtig) te wees, opbou van vet op die agterkant van die nek tussen die skouers (buffelskof), verlies van vet onder die vel (dit veroorsaak dat die gesig maer vertoon), borsvergroting en hoë vlakke van vet en glukose (suiker) in die bloed. Indien jy enige van hierdie tekens het, moet jy deeglik deur jou dokter ondersoek word.

- In seldsame gevalle, aangesien jou immuunstelsel versterk het, kan dit ook gesonde liggaamsweefsel aanval (outo-immuunversteurings). Die simptome van outo-immuunversteurings kan baie maande nadat jy begin het om medisyne te neem om jou MIV-infeksie te behandel, ontwikkel.
- As jy enige simptome van osteonekrose opmerk (waar gedeeltes van die beenweefsel afsterf as gevolg van verminderde bloedvoorsiening aan die been): gewrigsongemak en pyn, gewrigstyfheid of moeilike beweging, moet jy jou dokter inlig.
- Jy kan steeds infeksies en ander komplikasies van MIV-infeksie ontwikkel terwyl jy **SONKE LAMIVUDINE 150** gebruik**.**
- Daar is steeds ‘n risiko dat jy MIV aan ander persone deur seksuele kontak of bloedkontaminasie kan oordra terwyl jy **SONKE-LAMIVUDINE 150** neem**.**
- Staak **SONKE-LAMIVUDINE 150** tablette onmiddellik as jy abdominale (maag-) pyn, naarheid of braking, kortasemigheid, moegheid en gewigsverlies ontwikkel en jy moet jou dokter kontak.

SONKE-LAMIVUDINE 150 moet met omsigtigheid gebruik word by pasiënte met lewer- en niervers teurings asook by bejaardes.

Ander medisyne en SONKE-LAMIVUDINE 150

Lig jou gesondheidsorgverskaffer altyd in as jy enige ander medisyne gebruik. (Dit sluit komplementêre of tradisionele medisyne in.)

As jy ander medisyne op ‘n gereelde basis gebruik, insluitend komplementêre of tradisionele medisyne, kan die gelyktydige gebruik van SONKE LAMIVUDINE 150 saam met hierdie medisyne ongunstige interaksies veroorsaak. Raadpleeg asseblief jou dokter, apteker of ander gesondheidsorgverskaffer vir advies.

Versigtigheid is nodig as jy die volgende medisyne saam met **SONKE LAMIVUDINE 150** neem**:**

- Trimetoprim/sulfametoksasool (‘n medisyne om infeksies te behandel)
- Salsitabien
- Emtrisitabien (gebruik om MIV-infeksie te behandel).
- Kladribien (gebruik om sekere tipes kanker te behandel).
- Medisyne (gewoonlik vloeistowwe) wat sorbitol en ander suikeralkohole bevat (soos xilitol)

SONKE-LAMIVUDINE 150 saam met voedsel of drinkgoed

SONKE LAMIVUDINE 150 kan met of sonder voedsel geneem word.

Swangerskap en borsvoeding

Jy moenie **SONKE LAMIVUDINE 150** gebruik wanneer jy swanger is of jou suigeling borsvoed nie.

Konsulteer asseblief jou dokter, apteker of ander gesondheidsorgverskaffer vir advies as jy swanger is of jou baba borsvoed terwyl jy hierdie medisyne gebruik.

Bestuur van voertuie en gebruik van masjinerie

SONKE LAMIVUDINE 150 kan duiseligheid of moegheid by sommige pasiënte veroorsaak. Moenie voertuie bestuur of enige gereedskap of masjinerie gebruik totdat jy weet hoe **SONKE LAMIVUDINE 150** jou affekteer nie, omdat jou konsentrasie geaffekteer mag wees.

3. Hoe om SONKE-LAMIVUDINE 150 te neem

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

Neem **SONKE LAMIVUDINE** tablette altyd presies soos wat jou dokter of apteker dit aan jou voorgeskryf het. Jy moet jou dokter of apteker raadpleeg as jy onseker is.

Volwassenes en adolessente: Neem twee tablette een keer per dag of een tablet twee keer per dag in kombinasie met sidovudien.

Kinders wat tussen 14 kg tot 20 kg weeg:

Neem een tablet een keer per dag.

Kinders wat ten minste 25 kg weeg:

Neem een tablet twee keer per dag of neem twee tablette een keer per dag.

Indien jy lewer- en nierprobleme het, sal jou dokter die dosis dienooreenkomstig aanpas.

Om jou te help onthou om **SONKE LAMIVUDINE 150** te neem, probeer om in die gewoonte te kom om dit op dieselfde tyd elke dag te neem.

Neem **SONKE LAMIVUDINE 150** soos voorgeskryf en vir so lank as wat dit voorgeskryf is; moenie ophou om dit te neem nie, selfs as jy beter voel, aangesien die simptome kan terugkeer.

*Praat met jou dokter pf apteker as jy onder die indruk is dat die effek van **SONKE LAMIVUDINE 150** tablette te sterk of te swak is.*

As jy meer SONKE LAMIVUDINE 150 geneem het as wat jy behoort te neem

Raadpleeg jou dokter of apteker in die geval van oordosering. Indien nie een van hulle beskikbaar is nie, neem die pasiënt dadelik na die naaste hospitaal of gifbeheersentrum. Neem hierdie inligtingstuk of sommige van die tablette saam met jou sodat jou dokter kan weet wat jy geneem het.

As jy vergeet het om SONKE LAMIVUDINE 150 tablette te neem

Neem dit gou as wat jy onthou. As dit egter amper tyd is vir jou volgende dosis, slaan die dosis oor wat jy nie geneem het nie en gaan terug na jou normale dosisskedule. Moenie ‘n dubbele dosis neem om te vergoed vir die individuele dosisse wat jy vergeet het nie.

4. Moontlike newe-effekte

SONKE LAMIVUDINE 150 kan newe-effekte hê.

Nie alle newe-effekte wat vir **SONKE LAMIVUDINE 150** aangemeld word, is in hierdie inligtingstuk ingesluit nie. Konsulteer asseblief jou dokter, apteker of ander gesondheidsorgverskaffer vir advies indien jou algemene gesondheid verswak terwyl jy **SONKE LAMIVUDINE 150** neem.

Indien enige van die volgende gebeur, staak die gebruik van **SONKE LAMIVUDINE 150** en lig jou dokter onmiddellik in of gaan na die noodgevalle afdeling by jou naaste hospitaal:

- Uitslag, galbulte, gejeuk, gehyg, kortasemigheid of swelling van die gesig, lippe, tong, hande/ voete, floute, hoë koors

Hierdie is almal baie ernstige newe-effekte. As jy hulle het, kon jy ‘n ernstige reaksie teenoor **SONKE LAMIVUDINE 150** gehad het. Jy mag dringende mediese aandag of hospitalisasie benodig.

Lig jou dokter onmiddellik in of gaan na die noodgevalle-afdeling by jou naaste hospitaal as jy

enige van die volgende opmerk:

- ‘n Afbraak van spierweefsel wat ‘n skadelike proteïen in die bloed vrystel (rabdomiolise); simptome sluit in: donker, rooierige urine, ‘n afname in die hoeveelheid van die urine, swakheid en spierpyne.
- Oormatige melksuur in the blood (melksuurasidose); simptome sluit in: vinnige asemhaling, oormatige sweet, koue en klam vel, asem wat soet ruik, maagpyn, naarheid of braking, verwarring en moegheid.
- Ontsteking van die pankreas (pankreatitis); simptome sluit erge maagpyn met naarheid en braking en koors in.
- Ontsteking van die lewer (hepatitis); simptome sluit in: geel vel en oë (geelsug), naarheid, moegheid en koors.

Hierdie is almal ernstige newe-effekte. Jy mag dringende mediese aandag benodig.

Lig jou dokter in as jy enige van die volgende newe-effekte opmerk:

Dikwelse newe-effekte:

- Naarheid, braking, maagpyn of krampe, diaree
- Ongewone swakheid of moegheid
- Koors, algemene siekgevoel
- Sukkel om te slaap
- Hoofpyn, sensasie van naalde en spelde, tinteling, branderigheid, dooie gevoel
- Spierpyn of krampe
- Gewrigspyn
- Hoes, geïrriteerde of loopneus

Newe-effekte met onbekende voorkoms:

- Haarverlies

Newe-effekte wat deur jou bloedtoetse aangetoon kan word:

- ‘n Afname in die aantal witbloedselle (neutropenie)

- ‘n Afname in die aantal plaatjies (selle wat veroorsak dat bloed stol) (trombositopenie)
- ‘n Afname in die aantal rooibloedselle (anemie)
- ‘n Gebrek van die beenmurg om nuwe rooibloedselle te produseer (suiwer rooiselaplasie)
- ‘n Toename in lewerensieme
- ‘n Toename in die vlak van ‘n ensiem in die bloed wat amilase genoem word

Lig asseblief jou dokter of apteker in as jy enige newe-effekte opmerk wat nie in hierdie inligtingstuk genoem word nie.

Aanmelding van newe-effekte

Praat asseblief met jou dokter, apteker of verpleegkundige as jy newe-effekte ervaar. Jy kan ook newe-effekte by SAHPRA aanmeld *via* die “**6.04 Adverse Drug Reaction Reporting Form**”, wat aanlyn beskikbaar is onder SAHPRA se publikasies: https://www.sahpra.org.za/Publications/Index/8. Deur newe-effekte aan te meld, kan jy help om meer inligting te verskaf oor die veiligheid van **SONKE LAMIVUDINE 150.**

5. Hoe om SONKE LAMIVUDINE 150 te bewaar

Bewaar alle medisyne buite die bereik van kinders.

- Bewaar by of benede 25 °C.
- Bewaar in die oorspronklike verpakking.
- Hou die stulpstrook in die buitenste karton.
- Beskerm teen lig/vog.
- Moenie in ‘n badkamer bewaar nie.
- Moenie ná die vervaldatum, soos op die etiket aangedui, gebruik nie.
- Bring alle ongebruikte medisyne terug na jou apteker toe.
- Moenie van ongebruikte medisyne in dreinerings- of rioleringstelsels, bv. toilette, ontslae raak nie.

6. Inhoud van die verpakking en ander inligting

Wat SONKE LAMIVUDINE 150 bevat

Elke **SONKE LAMIVUDINE 150** tablet bevat 150 mg lamivudien (aktiewe bestanddeel).

Die ander bestanddele is:

Tabletkern: Magnesiumstearaat, mikrokristallyne sellulose; natrium-styselglikolaat

Filmbedekking: Hipromellose; titaniumdioksied; propileenglikool.

Hoe SONKE LAMIVUDINE 150 lyk en inhoud van die verpakking

Wit tot naaswit, ovaalvormige, bikonvekse, filmbedekte tablette met 'RX919' ingepers op een kant en 'n breeklyn op die ander kant.

10 tablette word verpak in stulpstroke van helder, deurskynende, PVC-film met 'n aluminiumfoelie rugkant. Kartonne bevat 10, 30 of 60 tablette.

56 of 60 tablette verpak in securitainers of in wit ondeursigtige HDPE-bottels.

Houer van die Registrasiesertifikaat

RANBAXY PHARMACEUTICALS (PTY) LTD

'n Sun Pharma maatskappy

Lautreweg 14, Stormill Uitbreiding 1

Roodepoort, 1724

Suid-Afrika

Hierdie inligtingstuk was laas hersien in

Datum van registrasie: 10 Augustus 2007

Datum van hersiening: 25 Januarie 2023

Registrasienommer

A40/20.2.8/027

Bemark deur Sonke Pharmaceuticals (Pty) Ltd

Hierdie produk is slegs vir gebruik in die gebiede van Suid-Afrika, Namibië, Botswana, Lesotho en Swaziland, is nie goedgekeur vir herverkope en enige gebruik nie.

Botswana: S2 Reg. nr.: BOT0801207

Namibië: NS2 Reg. nr.: 05/20.2.8/0416