

Package leaflet: Information for the user

Comirnaty™ 10 micrograms/dose

concentrate for dispersion for injection

Children 5 to 11 years

**COVID-19 mRNA Vaccine (nucleoside modified)
tozinameran**

▼ This medicine is subject to additional monitoring. This will allow quick identification of new safety information. You can help by reporting any side effects your child may get. See the end of section 4 for how to report side effects.

Read all of this leaflet carefully before you receive this vaccine because it contains important information for you.

- Keep this leaflet. You may need to read it again.
- If you have any further questions, ask your doctor, pharmacist or nurse.
- If your child gets any side effects, talk to your doctor, pharmacist or nurse. This includes any possible side effects not listed in this leaflet. See section 4.

What is in this leaflet

1. What Comirnaty is and what it is used for
2. What you need to know before your child receives Comirnaty
3. How Comirnaty is given
4. Possible side effects
5. How to store Comirnaty
6. Contents of the pack and other information

1 What Comirnaty is and what it is used for

Comirnaty is a vaccine used for preventing COVID-19 caused by SARS-CoV-2 virus.

Comirnaty 10 micrograms/dose concentrate for dispersion for injection is given to children from 5 to 11 years of age.

The vaccine causes the immune system (the body's natural defences) to produce antibodies and blood cells that work against the virus, so giving protection against COVID-19.

As Comirnaty does not contain the virus to produce immunity, it cannot give your child COVID-19.

2 What you need to know before your child receives Comirnaty

Comirnaty should not be given

- if your child is allergic to the active substance or any of the other ingredients of this medicine (listed in section 6)

Warnings and precautions

Talk to your doctor, pharmacist or nurse before your child is given the vaccine if your child:

- has ever had a severe allergic reaction or breathing problems after any other vaccine injection or after having been given Comirnaty in the past.
- is feeling nervous about the vaccination process or has ever fainted following any needle injection.
- has a severe illness or infection with high fever. However, your child can have the vaccination if he/she have a mild fever or upper airway infection like a cold.
- has a bleeding problem, bruises easily or uses a medicine to prevent blood-clots.
- has a weakened immune system, because of a disease such as HIV infection or a medicine such as corticosteroid that affects the immune system.

There is an increased risk of myocarditis (inflammation of the heart muscle) and pericarditis (inflammation of the lining outside the heart) after vaccination with Comirnaty (see section 4). These conditions

can develop within just a few days after vaccination and have primarily occurred within 14 days. They have been observed more often after the second vaccination, and more often occurred in younger males. Following vaccination, you should be alert to signs of myocarditis and pericarditis, such as breathlessness, palpitations and chest pain, and seek immediate medical attention should these occur.

As with any vaccine, Comirnaty may not fully protect all those who receive it and it is not known how long you will be protected.

Your child may receive a third dose of Comirnaty. The third dose may still not provide full immunity to COVID-19 in people who are immunocompromised. In these cases, you should continue to maintain physical precautions to help prevent COVID-19. In addition, your close contacts should be vaccinated as appropriate. Discuss appropriate individual recommendations with your doctor.

Children

Comirnaty is not recommended for children aged under 5 years.

Other medicines and Comirnaty

Tell your doctor or pharmacist if your child is using, has recently used or might use any other medicines or has recently received any other vaccine.

Pregnancy and breast-feeding

If your child is pregnant or breast-feeding, ask your doctor or pharmacist for advice before your child receives this vaccine.

Driving and using machines

Some of the effects of vaccination mentioned in section 4 (Possible side effects) may temporarily affect your ability to use machines or undertake activities such as cycling. Wait until these effects have worn off before resuming activities that require your full attention.

3 How Comirnaty is given

Comirnaty is given after dilution as an injection of 0.2 mL into a muscle of the upper arm.

Your child will receive 2 injections.

It is recommended to receive the second dose of the same vaccine 3 weeks after the first dose to complete the vaccination course.

If your child is immunocompromised, he or she may receive a third dose of Comirnaty at least 28 days after the second dose.

If you have any further questions on the use of Comirnaty, ask your doctor, pharmacist or nurse.

4 Possible Side Effects

Like all vaccines, Comirnaty can cause side effects, although not everybody gets them.

Very common side effects: may affect more than 1 in 10 people

- injection site: pain, swelling, redness
- tiredness
- headache
- muscle pain
- chills
- joint pain
- diarrhoea
- fever

Some of these side effects were slightly more frequent in adolescents 12 to 15 years than in adults.

Common side effects: may affect up to 1 in 10 people

- nausea
- vomiting

Uncommon side effects: may affect up to 1 in 100 people

- enlarged lymph nodes (more frequently observed after the booster dose)
- feeling unwell
- arm pain

- insomnia
- injection site itching
- allergic reactions such as rash or itching
- feeling weak or lack of energy/sleepy
- decreased appetite
- excessive sweating
- night sweats

Rare side effects: may affect up to 1 in 1,000 people

- temporary one sided facial drooping
- allergic reactions such as hives or swelling of the face

Very rare side effects: may affect up to 1 in 10,000 people

inflammation of the heart muscle (myocarditis) or inflammation of the lining outside the heart (pericarditis) which can result in breathlessness, palpitations or chest pain

Not known (cannot be estimated from the available data)

- severe allergic reaction
- extensive swelling of the vaccinated limb
- swelling of the face (swelling of the face may occur in patients who have had facial dermatological fillers)
- a skin reaction that causes red spots or patches on the skin, that may look like a target or “bulls-eye” with a dark red centre surrounded by paler red rings (erythema multiforme)
- unusual feeling in the skin, such as tingling or a crawling feeling (paraesthesia)
- decreased feeling or sensitivity, especially in the skin (hypoesthesia)

Reporting of side effects

- If your child gets any side effects, talk to your doctor, pharmacist or nurse. This includes any possible side effects not listed in this leaflet. You can also report side effects directly via ADR Reporting, Website: www.medicinesauthority.gov.mt/adrportal and include batch/Lot number if available.

By reporting side effects you can help provide more information on the safety of this medicine.

5 How to store Comirnaty

Keep this medicine out of the sight and reach of children.

The following information about storage, expiry and use and handling is intended for healthcare professionals.

Do not use this medicine after the expiry date which is stated on the carton and label after EXP. The expiry date refers to the last day of that month.

Store in freezer at -90 °C to -60 °C for 9 months.

Store in the original package in order to protect from light.

The vaccine will be received frozen at -90 °C to -60 °C. Frozen vaccine can be stored either at -90 °C to -60 °C or 2 °C to 8 °C upon receipt.

When stored frozen at -90 °C to -60 °C, 10-vial packs of the vaccine can be thawed at 2 °C to 8 °C for 4 hours or individual vials can be thawed at room temperature (up to 30 °C) for 30 minutes.

Once removed from the freezer, the unopened vial may be stored refrigerated at 2 °C to 8 °C for a single period of up to 10 weeks within the 9-month shelf life. The outer carton should be marked with the new discard date at 2 °C to 8 °C. Once thawed, the vaccine cannot be re-frozen.

Thawed vials can be handled in room light conditions.

After dilution, store and transport the vaccine at 2 °C to 30 °C and use within 12 hours. Discard any unused vaccine.

Do not use this vaccine if you notice particulates in the dilution or discolouration.

Do not throw away any medicines via wastewater or household waste. Ask your pharmacist how to throw away medicines you no longer use. These measures will help protect the environment.

6 Contents of the pack and other information

What Comirnaty contains

- The active substance is COVID-19 mRNA Vaccine called tozinameran. After dilution, the vial contains 10 doses of 0.2 mL with 10 micrograms tozinameran each.
- The other ingredients are:
 - ((4-hydroxybutyl)azanediyl)bis(hexane-6,1-diyl)bis(2-hexyldecanoate) (ALC-0315)
 - 2-[(polyethylene glycol)-2000]-N,N-ditetradecylacetamide (ALC-0159)
 - 1,2-Distearoyl-sn-glycero-3-phosphocholine (DSPC)
 - cholesterol
 - trometamol
 - trometamol hydrochloride
 - sucrose
 - water for injections

What Comirnaty looks like and contents of the pack

The vaccine is a white to off-white dispersion (pH: 6.9 - 7.9) provided in a multidose vial of 10 doses in a 2 mL clear vial (type I glass), with a rubber stopper and an orange flip-off plastic cap with aluminium seal.

Pack sizes: 195 vials or 10 vials

Not all pack sizes may be marketed.

Marketing Authorisation Holder

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Manufacturers

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For any information about this medicine, please contact the local representative of the Marketing Authorisation Holder:

Malta

Vivian Corporation Ltd, Tel: +35621 344610

This leaflet was last revised in 02/2022

This medicine has been given ‘conditional approval’. This means that there is more evidence to come about this medicine. The European Medicines Agency will review new information on this medicine at least every year and this leaflet will be updated as necessary.



Scan the code with a mobile device to get the package leaflet in different languages.

URL: www.comirnatyglobal.com

Detailed information on this medicine is available on the European Medicines Agency web site: <http://www.ema.europa.eu>.

This package leaflet is available in all EU/EEA languages on the European Medicines Agency website.

The following information is intended for healthcare professionals only:

Administer Comirnaty intramuscularly after dilution as a course of 2 doses (0.2 mL each) 3 weeks apart.

A third dose may be given at least 28 days after the second dose to individuals who are severely immunocompromised.

Traceability

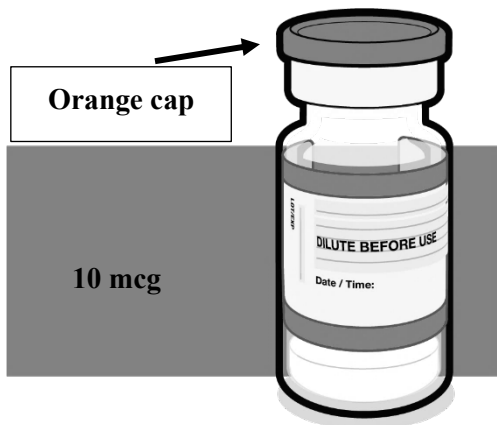
In order to improve the traceability of biological medicinal products, the name and the batch number of the administered product should be clearly recorded.

Handling instructions

Comirnaty 10 micrograms/dose should be prepared by a healthcare professional using aseptic technique to ensure the sterility of the prepared dispersion.

DOSE VERIFICATION OF COMIRNATY 10 MICROGRAMS/DOSE CONCENTRATE FOR DISPERSION FOR INJECTION (CHILDREN 5 TO 11 YEARS)

- Verify that the vial has an orange plastic cap.
- If the vial has a purple plastic cap, please make reference to the Summary of Product Characteristics for Comirnaty 30 micrograms/dose concentrate for dispersion for injection.
- If the vial has a grey plastic cap, please make reference to the Summary of Product Characteristics for Comirnaty 30 micrograms/dose dispersion for injection.



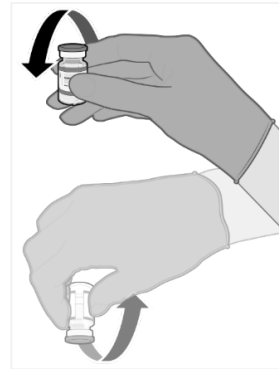
HANDLING PRIOR TO USE OF COMIRNATY 10 MICROGRAMS/DOSE CONCENTRATE FOR DISPERSION FOR INJECTION (CHILDREN 5 TO 11 YEARS)

- If the multidose vial is stored frozen it must be thawed prior to use. Frozen vials should be transferred to an environment of 2 °C to 8 °C to thaw; a 10 vial pack may take 4 hours to thaw. Ensure vials are completely thawed prior to use.
- Upon moving vials to 2 °C to 8 °C storage, update the expiry date on the carton.
- Unopened vials can be stored for up to 10 weeks at 2 °C to 8 °C within the 9-month shelf life.
- Alternatively, individual frozen vials may be thawed for 30 minutes at temperatures up to 30 °C.
- Prior to use, the unopened vial can be stored for up to 12 hours at temperatures up to 30 °C. Thawed vials can be handled in room light conditions.



MIXING PRIOR TO DILUTION OF COMIRNATY 10 MICROGRAMS/DOSE CONCENTRATE FOR DISPERSION FOR INJECTION (CHILDREN 5 TO 11 YEARS)

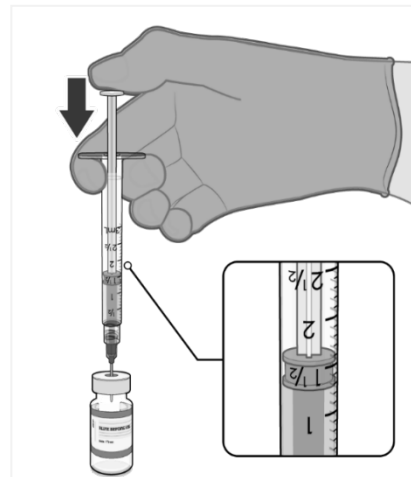
- Allow the thawed vial to come to room temperature and gently invert it 10 times prior to dilution. Do not shake.
- Prior to dilution, the thawed dispersion may contain white to off-white opaque amorphous particles.



Gently x 10

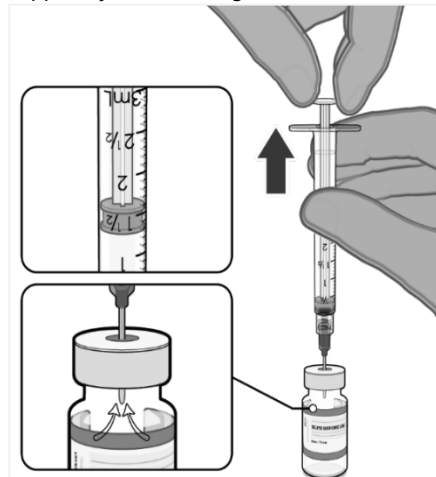
DILUTION OF COMIRNATY 10 MICROGRAMS/DOSE CONCENTRATE FOR DISPERSION FOR INJECTION (CHILDREN 5 TO 11 YEARS)

- The thawed vaccine must be diluted in its original vial with 1.3 mL sodium chloride 9 mg/mL (0.9%) solution for injection, using a 21 gauge or narrower needle and aseptic techniques.



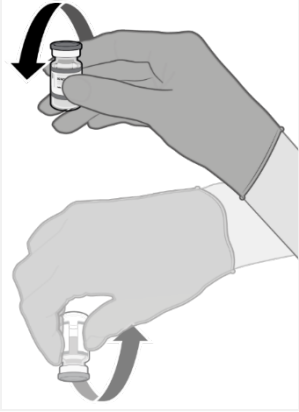
1.3 mL of 0.9% sodium chloride injection

- Equalise vial pressure before removing the needle from the vial stopper by withdrawing 1.3 mL air into the empty diluent syringe.



Pull back plunger to 1.3 mL to remove air from vial.

- Gently invert the diluted dispersion 10 times. Do not shake.
- The diluted vaccine should present as a white to off-white dispersion with no particulates visible. Do not use the diluted vaccine if particulates or discolouration are present.



Gently x 10

- The diluted vials should be marked with the appropriate date and time.
- After dilution, store at 2 °C to 30 °C and use within 12 hours.
- Do not freeze or shake the diluted dispersion. If refrigerated, allow the diluted dispersion to come to room temperature prior to use.



**Record appropriate date and time.
Use within 12 hours after dilution.**

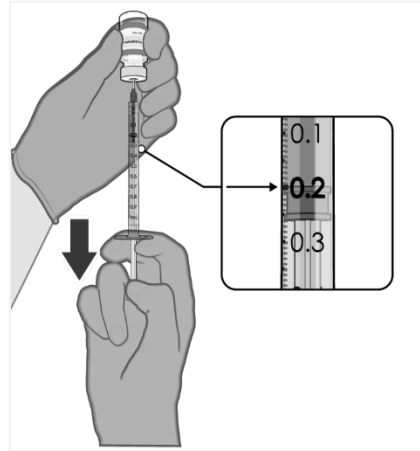
PREPARATION OF INDIVIDUAL 0.2 mL DOSES OF COMIRNATY 10 MICROGRAMS/DOSE CONCENTRATE FOR DISPERSION FOR INJECTION (CHILDREN 5 TO 11 YEARS)

- After dilution, the vial contains 2.6 mL from which 10 doses of 0.2 mL can be extracted.
- Using aseptic technique, cleanse the vial stopper with a single use antiseptic swab.
- Withdraw 0.2 mL of Comirnaty for children age 5 to 11 years.

Low dead-volume syringes and/or needles should be used in order to extract ten doses from a single vial. The low dead-volume syringe and needle combination should have a dead-volume of no more than 35 microlitres.

If standard syringes and needles are used, there may not be sufficient volume to extract ten doses from a single vial.

- Each dose must contain 0.2 mL of vaccine.
- If the amount of vaccine remaining in the vial cannot provide a full dose of 0.2 mL, discard the vial and any excess volume.
- Discard any unused vaccine within 12 hours after dilution.



0.2 mL diluted vaccine

Disposal

Any unused medicinal product or waste material should be disposed of in accordance with local requirements.

Fuljett ta' tagħrif: Informazzjoni għall-utent

Comirnaty™ 10 mikrogramma/doża

konċentrat għal dispersjoni għall-injezzjoni

Tfal minn 5 snin sa 11-il sena

Vaċċin tal-mRNA tal-COVID-19 (b'nucleoside modifikat)

tozinameran

▼ Dan il-prodott mediċinali huwa suġġett għal monitoraġġ addizzjonali. Dan ser jippermetti identifikazzjoni ta' malajr ta' informazzjoni ġdida dwar is-sigurtà. Inti tista' tgħin billi tirrapporta kwalunkwe effett sekondarju li jista' jkollok. Ara t-tmieni ta' sezzjoni 4 biex tara kif għandek tirrapporta effetti sekondarji.

Aqra sew dan il-fuljett kollu qabel tircievi dan il-vaċċin peress li fih informazzjoni importanti għalik.

- Żomm dan il-fuljett. Jista' jkollok bżonn terġa' taqrah.
- Jekk ikollok aktar mistoqsijiet, staqsi lit-tabib, lill-ispizjar jew lill-infermier tiegħek.
- Jekk it-tifel jew it-tifla tiegħek ikollu/ha xi effett sekondarju kellem lit-tabib, lill-ispizjar jew lill-infermier tiegħek. Dan jinkludi xi effetti sekondarju possibbli li mhuwiex elenkat f'dan il-fuljett. Ara sezzjoni 4.

F'dan il-fuljett

1. X'inhu Comirnaty u għalxiex jintuża
2. X'għandek tkun taf qabel ma tircievi Comirnaty
3. Kif jingħata Comirnaty
4. Effetti sekondarji possibbli
5. Kif taħžen Comirnaty
6. Kontenut tal-pakkett u informazzjoni oħra

1 X'inhu Comirnaty u għalxiex jintuża

Comirnaty huwa vaċċin użat għall-prevenzjoni tal-COVID-19 ikkawżata mill-virus SARS-CoV-2.

Comirnaty 10 mikrogrammi/doża konċentrat għal dispersjoni għall-injezzjoni jingħata lil tfal b'età minn 5 snin sa 11-il sena.

Il-vaċċin jikkawża s-sistema immuni (id-difiża naturali tal-ġisem) biex tipproduċi antikorpi u ċelluli tad-demem li jaħdmu kontra l-virus, u b'hekk tagħti protezzjoni kontra l-COVID-19.

Peress li Comirnaty ma fihx il-virus biex jipproduċi l-immunità, ma jistax jagħti COVID-19 lit-tifel jew lit-tifla tiegħek.

2 X'għandek tkun taf qabel ma t-tifel jew it-tifla tiegħek jircievi/tircievi Comirnaty

Comirnaty m'għandux jingħata

- jekk it-tifel jew it-tifla tiegħek huwa/hija allergiku/a għas-sustanza attiva jew għal xi sustanza oħra ta' din il-mediċina (imniżżla fis-sezzjoni 6)

Twissijiet u prekawzjonijiet

Kellem lit-tabib, lill-ispizjar jew lill-infermier tiegħek qabel ma t-tifel jew it-tifla tiegħek jingħata/tingħata dan il-vaċċin jekk it-tifel/tifla tiegħek:

- qatt kellu/kellha reazzjoni allergika severa jew problemi bin-nifs wara l-injezzjoni ta' kwalunkwe vaċċin ieħor jew wara li ngħata/ingħatat Comirnaty fil-passat.
- qed iħossu/iħossha nervuż/a dwar il-proċess tat-tilqim jew jekk xi darba intilef/intilfet minn sensih/ha wara kwalunkwe injezzjoni b'labra.
- għandu/għandha marda severa jew infezzjoni b'deni għoli. Madankollu, it-tifel/tifla tiegħek jista'/tista' jiehu/tiehu t-tilqima tiegħek jekk ikollu/ha deni ħafif jew infezzjoni ħafifa tal-parti ta' fuq tal-passaġġ tan-nifs bħal riħ.

- għandu/għandha problema ta' ħruġ ta' demm, jitbenġel/titbenġel malajr jew juża/tuża mediċina biex tipprevjeni emboli tad-demem.
- għandu/għandha sistema immuni mdgħajfa, minħabba marda bħal infezzjoni bl-HIV jew mediċina bħal kortikosteroidi li taffettwa s-sistema immuni.

Hemm żieda fir-riskju ta' mijokardite (infjammazzjoni tal-muskolu tal-qalb) u perikardite (infjammazzjoni tar-rita barra mill-qalb) wara tilqim b'Comirnaty (ara sezzjoni 4). Dawn il-kundizzjonijiet jistgħu jiżviluppaw fi żmien ftit jiem biss wara t-tilqim u seħħew primarjament fi żmien 14-il jum. Dawn ġew osservati aktar spiss wara t-tieni tilqima, u aktar spiss f'irġiel iżgħar fl-età. Wara t-tilqim, għandek toqgħod attent għal sinjali ta' mijokardite u perikardite, bħal qtugħ ta' nifs, palpitazzjonijiet u wġiġh fis-sider, u ftitex attenzjoni medika immedjata jekk dawn iseħħu.

Bħal kull vaċċin, Comirnaty jista' ma jiprotegix b'mod sħiħ lil dawk kollha li jirċevuh u mhux magħruf kemm ser iddum protett.

It-tifel jew it-tifla tiegħek jista'/tista' tircievi t-tielet doża ta' Comirnaty. It-tielet doża xorta tista' ma jipprovdi immunità sħiħa għal COVID-19 f'nies li huma immunokompromessi. F'dawn il-każijiet, għandek tkompli tiehu prekawzjonijiet fiżiċi biex tgħin tipprevjeni l-COVID-19. Barra minn hekk, persuni viċin tiegħek għandhom jiġu mlaqqma kif xieraq. Iddiskuti rakkomandazzjonijiet individwali xierqa mat-tabib tiegħek.

Tfal

Comirnaty mhux rakkomandat għal tfal b'età ta' inqas minn 5 snin.

Mediċini oħra u Comirnaty

Għid lit-tabib jew lill-ispizjar tiegħek jekk it-tifel/tifla tiegħek qed juża/tuża, uża/użat dan l-aħħar jew jista'/tista' juża/tuża xi mediċini oħra jew jekk riċentement irċieva/irċieviet xi vaċċin ieħor.

Tqala u tredidigh

Jekk it-tifla tiegħek hija tqila jew qed tredida' itlob il-parir tat-tabib jew tal-ispizjar tiegħek qabel it-tifla tiegħek tircievi dan il-vaċċin.

Sewqan u tħaddim ta' magni

Uħud mill-effetti tat-tilqima msemmija fis-sezzjoni 4 (Effetti sekondarji possibbli) jistgħu jaffettwaw b'mod temporanju l-hila tiegħek li tħaddem magni jew li twettaq attivitajiet bħal issuq rota. Stenna sakemm dawn l-effetti jgħaddu qabel ma tkompli b'attivitajiet li jeħtieġu l-attenzjoni sħiħa tiegħek.

3 Kif jingħata Comirnaty

Comirnaty jingħata wara d-dilwizzjoni bħala injezzjoni ta' 0.2 mL ġo muskolu fin-naħa ta' fuq tad-driegħ.

It-tifel/tifla tiegħek ser jircievi/tircievi 2 injezzjonijiet.

Huwa rakkomandat li tircievi t-tieni doża tal-istess vaċċin 3 ġimgħat wara l-ewwel doża biex tlesti l-kors ta' tilqim.

Jekk it-tifel/tifla tiegħek huwa/hija immunokompromess/a, jista'/tista' jircievi/tircievi t-tielet doża ta' Comirnaty mill-inqas 28 jum wara t-tieni doża.

Jekk għandek aktar mistoqsijiet dwar l-użu ta' Comirnaty, staqsi lit-tabib, lill-ispizjar jew lill-infermier tiegħek.

4 Effetti sekondarji possibbli

Bħal kull vaċċin ieħor, Comirnaty jista' jikkawża effetti sekondarji, għalkemm ma jidhrux f'kulhadd.

Effetti sekondarji komuni ħafna: jistgħu jaffettwaw aktar minn persuna waħda minn kull 10

- sit tal-injezzjoni: uġiġh, neffa, ħmura
- għeja
- uġiġh ta' ras
- uġiġh fil-muskoli
- tkexkix ta' bard
- uġiġh fil-gogi
- dijarea

- deni

Uħud minn dawn l-effetti sekondarji kienu kemmxejn aktar frekwenti f'adolessenti b'età minn 12 sa 15-il sena milli fl-adulti.

Effetti sekondarji komuni: jistgħu jaffettwaw sa persuna waħda minn kull 10

- dardir
- rimettar

Effetti sekondarji mhux komuni: jistgħu jaffettwaw sa persuna waħda minn kull 100

- glandoli limfatiċi minfuħa (osservati b'mod aktar frekwenti wara d-doża *booster*)
- tħossok ma' tiflaħx
- uġiġħ fid-dirgħajn
- insomnja
- ħakk fis-sit tal-injezzjoni
- reazzjonijiet allergiċi bħal raxx jew ħakk
- tħossok dgħajjef jew b'nuqqas ta' enerġija/bi ngħas
- tnaqqis fl-aptit
- għaraq eċċessiv
- għaraq matul il-lejl

Effetti sekondarji rari: jistgħu jaffettwaw sa persuna waħda minn kull 1,000

- naħa waħda tal-wiċċ tiddendel b'mod temporanju
- reazzjonijiet allergiċi bħal horriqija jew nefħa fil-wiċċ

Effetti sekondarji rari ħafna: jistgħu jaffettwaw sa persuna waħda minn kull 10,000

- infjammazzjoni tal-muskolu tal-qalb (mijokardite) jew infjammazzjoni tar-rita barra mill-qalb (perikardite) li tista' tirriżulta fi qtugħ ta' nifs, palpitazzjonijiet jew uġiġħ fis-sider

Mhux magħruf (ma tistax tittieħed stima mid-*data* disponibbli)

- reazzjoni allergjika severa
- nefħa estensiva fid-driegħ li fih ingħata l-vaċċin
- nefħa fil-wiċċ (nefħa fil-wiċċ tista' sseħħ f'pazjenti li kellhom fillers dermali tal-wiċċ)
- reazzjoni tal-ġilda li tikkawża tikek jew rqajja' ħomor fuq il-ġilda, li jistgħu jidhru bħal targit jew "bull's-eye" b'ċentru aħmar skur imdawwar bi ċrieki ħomor ċari (eritema multiforme)
- sensazzjoni mhux tas-soltu fil-ġilda, bħal tneħħim jew sensazzjoni ta' xi ħaġa miexja fuq il-ġilda (parestesija)
- tnaqqis fis-sensazzjoni jew fis-sensittività, speċjalment fil-ġilda (ipoestesija)

Rappurtar tal-effetti sekondarji

Jekk it-tifel/tifla tiegħek ikollu/ha xi effett sekondarju, kellew lit-tabib, lill-ispizjar jew lill-infermier tiegħek. Dan jinkludi xi effetti sekondarju possibbli li mhux elenkat f'dan il-fuljett. Tista' wkoll tirrapporta effetti sekondarji direttament permezz tas-sit elettroniku tar-rappurtar tar-reazzjonijiet avversi suspettati: www.medicinesauthority.gov.mt/adportal u inkludi n-numru tal-lott/Lott jekk disponibbli.

Billi tirrapporta l-effetti sekondarji tista' tgħin biex tiġi pprovduta aktar informazzjoni dwar is-sigurtà ta' din il-medicina.

5 Kif taħzen Comirnaty

Żomm din il-medicina fejn ma tidhix u ma tintlaħaqx mit-tfal.

It-tagħrif li ġej dwar il-ħażna, d-data ta' meta tiskadi u l-użu u l-immaniġġar huwa maħsub għall-professionisti tal-kura tas-saħħa.

Tużax din il-medicina wara d-data ta' meta tiskadi li tidher fuq il-kartuna u t-tikketta wara JIS. Id-data ta' meta tiskadi tirreferi għall-aħħar ġurnata ta' dak ix-xahar.

Aħżen fil-frیža f'temperatura ta' -90 °C sa -60 °C għal 9 xhur.

Aħżen fil-pakkett oriġinali sabiex tilqa' mid-dawl.

Il-vaċċin se jiġi riċevut ffrیżat f'temperatura ta' - 90 °C sa - 60 °C. Wara li jiġi riċevut vaċċin iffrیżat jista' jinħażen f'temperatura ta' - 90 °C sa - 60 °C jew f'temperatura ta' 2 °C sa 8 °C.

Meta jinħażnu ffrیżati f'temperatura ta' - 90 °C sa - 60 °C, pakketti b'10 kunjetti tal-vaċċin jistgħu jinħallu mis-silġ f'temperatura ta' 2 °C sa 8 °C għal 4 sigħat jew kunjetti individwali jistgħu jinħallu f'temperatura tal-kamra (sa 30 °C) għal 30 minuta.

Ladarba jitneħħa mill-frیža, il-kunjett mhux miftuħ jista' jinħażen fi frیġġ f'temperatura ta' 2 °C sa 8 °C għal perjodu wieħed sa 10 ġimgħat fi żmien is-9 xhur li fihom jibqa' tajjeb il-prodott. Il-kartuna ta' barra għandha tiġi mmarkata bid-data l-ġdida li fiha għandu jintrema jekk jinħażen f'temperatura ta' 2 °C sa 8 °C. Ladarba jinħall mis-silġ, il-vaċċin ma jistax jiġi ffrیżat mill-ġdid.

Kunjetti li nħallu mis-silġ jistgħu jiġu mmaniġġjati f'kondizzjonijiet ta' dawl tal-kamra.

Wara d-dilwizzjoni, aħżen u ttrasporta l-vaċċin f'temperatura ta' 2 °C sa 30 °C u użax fi żmien 12-il siegħa. Armi kwalunkwe vaċċin mhux użat.

Tużax dan il-vaċċin jekk tinnota partiċelli fid-dilwizzjoni jew bidla fil-kulur.

Tarmix medicini mal-ilma tad-dranagġ jew mal-iskart domestiku. Staqsi lill-ispizjar tiegħek dwar kif għandek tarmi medicini li m'għadex tużax. Dawn il-mizuri jgħinu għall-protezzjoni tal-ambjent.

6 Kontenut tal-pakkett u informazzjoni oħra

X'fih Comirnaty

- Is-sustanza attiva hi Vaċċin tal-mRNA tal-COVID-19 imsejħa tozinameran. Wara d-dilwizzjoni, il-kunjett ikun fih 10 dozi ta' 0.2 mL b'10 mikrogrammi ta' tozinameran kull wieħed.
- Is-sustanzi mhux attivi l-oħra huma:
 - ((4-hydroxybutyl)azanediyl)bis(hexane-6,1-diy)bis(2-hexyldecanoate) (ALC-0315)
 - 2-[(polyethylene glycol)-2000]-N,N-ditetradecylacetamide (ALC-0159)
 - 1,2-Distearoyl-sn-glycero-3-phosphocholine (DSPC)
 - kolesterol
 - trometamol
 - trometamol hydrochloride
 - sucrose
 - ilma għall-injezzjonijiet

Kif jidher Comirnaty u l-kontenut tal-pakkett

Il-vaċċin huwa dispersjoni ta' lewn abjad sa abjad maħmuġ (pH: 6.9-7.9) ipprovdut f'kunjett b'aktar minn doża waħda ta' 10 dozi f'kunjett trasparenti ta' 2 mL (ħġieg tat-tip I), b'tapp tal-lastku u għatu tal-plastik oranġjo li jitneħħa b'daqqa ta' saba' b'siġill tal-aluminju.

Daqsijiet tal-pakkett: 195 kunjett jew 10 kunjetti

Jista' jkun li mhux il-pakketti tad-daqsijiet kollha jkun fis-suq.

Detentur tal-Awtorizzazzjoni għat-Tqegħid fis-Suq

BioNTech Manufacturing GmbH

An der Goldgrube 12, 55131 Mainz, Il-Ġermanja

Phone: +49 6131 9084-0, Fax: +49 6131 9084 2121

service@biontech.de

Manifatturi

- BioNTech Manufacturing GmbH, Kupferbergterrasse 17 – 19, 55116 Mainz, Il-Ġermanja

Għal kull tagħrif dwar din il-medicina, jekk jogħġbok ikkuntattja lir-rappreżentant lokali tad-Detentur tal-Awtorizzazzjoni għat-Tqegħid fis-Suq:

Malta

Vivian Corporation Ltd, Tel: +35621 344610

Dan il-fuljett kien rivedut l-aħħar fl-02/2022



Skennja l-kowd b'*mobile* biex tikseb il-fuljett ta' tagħrif f'lingwi differenti.

URL: www.comirnatyglobal.com

Din il-medicina ngħatat approvazzjoni kondizzjonali'. Dan ifisser li għad trid tingħata aktar evidenza dwar din il-medicina. L-Aġenzija Ewropea għall-Medicini ser tirrevedi l-informazzjoni l-ġdida dwar din il-medicina mill-anqas kull sena u ser taġġorna dan il-fuljett kif meħtieġ.

Informazzjoni dettaljata dwar din il-medicina tinsab fuq is-sit elettroniku tal-Aġenzija Ewropea għall-Medicini:

<http://www.ema.europa.eu>.

Dan il-fuljett huwa disponibbli fil-lingwi kollha tal-UE/ŻEE fis-sit elettroniku tal-Aġenzija Ewropea għall-Medicini.

It-tagħrif li jmiss qed jingħata biss għall-professjonisti tal-kura tas-saħħa biss:

Agħti Comirnaty ġol-muskoli wara d-dilwizzjoni bħala kors ta' 2 doži (0.2 mL kull waħda) 3 ġimgħat bogħod minn xulxin.

It-tielet doża tista' tingħata mill-inqas 28 jum wara t-tieni doża lil individwi li huma immunokompromessi b'mod sever.

Traċċabilità

Sabiex tittejjeb it-traċċabilità tal-prodotti mediċinali bijoloġiċi, l-isem u n-numru tal-lott tal-prodott amministrat għandhom jiġu rrekordjati b'mod ċar.

Istruzzjonijiet dwar l-immaniġġar

Comirnaty 10 mikrogrammi/doża għandu jiġi ppreparat minn professjonist tal-kura tas-saħħa permezz ta' teknika asettika biex tiġi żgurata l-isterilità tad-dispersjoni ppreparata.

VERIFIKA TAD-DOŽA TA' COMIRNATY 10 MIKROGRAMMI/DOŽA KONĊENTRAT GĦAL DISPERSJONI GĦALL-INJEZZJONI (TFAL MINN 5 SNIN SA 11-IL SENA)

- Ivverifika li l-kunjett għandu għatu tal-plastik oranġjo.
- Jekk il-kunjett għandu għatu tal-plastik vjola, jekk jogħġbok irreferi għas-Sommarju tal-Karatteristiċi tal-Prodott għal Comirnaty 30 mikrogramma/doża konċentrat għal dispersjoni għall-injezzjoni.
- Jekk il-kunjett għandu għatu tal-plastik griż, jekk jogħġbok irreferi għas-Sommarju tal-Karatteristiċi tal-Prodott għal Comirnaty 30 mikrogramma/doża dispersjoni għall-injezzjoni.

Ghatu oranġjo

10 mcg

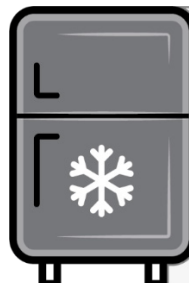


IMMANIĠĠAR QABEL L-UŽU TA' COMIRNATY 10 MIKROGRAMMI/DOŽA KONĊENTRAT GĦAL DISPERSJONI GĦALL-INJEZZJONI (TFAL MINN 5 SNIN SA 11-IL SENA)

- Jekk il-kunjett b'aktar minn doża waħda jinħażen iffriztat għandu jinħall mis-silġ qabel l-użu. Kunjetti ffrizati għandhom jiġu ttrasferiti għal ambjent b'temperatura ta' 2 °C sa 8 °C biex

jinħallu mis-silġ; pakkett ta' 10 kunjetti jista' jiehu 4 sigħat biex jinħall mis-silġ. Kun ċert li l-kunjetti jkunu nħallu kompletament mis-silġ qabel l-użu.

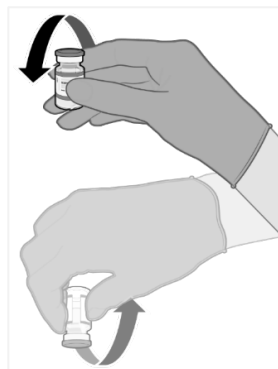
- Meta l-kunjetti jitmexxew għal hażna f'temperatura ta' 2 °C sa 8 °C, aġġorna d-data ta' meta jiskadi fuq il-kartuna.
- Kunjetti mhux miftuħa jistgħu jinħażnu għal massimu ta' 10 ġimgħat f'temperatura ta' 2 °C sa 8 °C fi żmien is-9 xhur li fihom idum tajjeb il-prodott.
- Inkella, kunjetti individwali ffrizati jistgħu jinħallu mis-silġ għal 30 minuta f'temperaturi sa 30 °C.
- Qabel l-użu, il-kunjett mhux miftuħ jista' jinħażen għal perjodu sa 12-il siegħa f'temperaturi sa 30 °C. Kunjetti li nħallu mis-silġ jistgħu jiġu mmaniġġjati f'kondizzjonijiet ta' dawl tal-kamra.



Ahžen għal perjodu sa 10 ġimgħat f'temperatura ta' 2 °C sa 8 °C

TAHLIT QABEL ID-DILWIZZJONI TA' COMIRNATY 10 MIKROGRAMMI/DOŽA KONĊENTRAT GĦAL DISPERSJONI GĦALL-INJEZZJONI (TFAL MINN 5 SNIN SA 11-IL SENA)

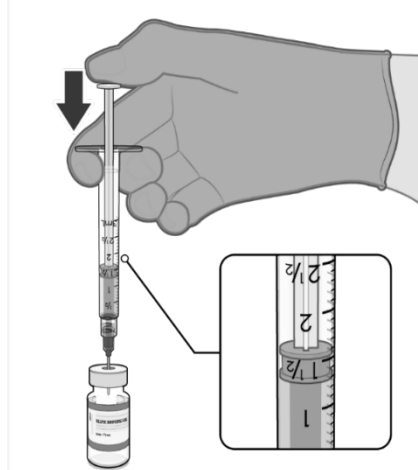
- F'alli l-kunjett li jkun inħall mis-silġ jilhaq it-temperatura tal-kamra u aqilbu bil-mod ta' taħt fuq għal 10 darbiet qabel id-dilwizzjoni. Thawwadx.
- Qabel id-dilwizzjoni, id-dispersjoni li tkun inħallet mis-silġ jista' jkun fiha partiċelli amorfi opaki ta' lewn abjad sa abjad maħmuġ.



Bil-mod × 10

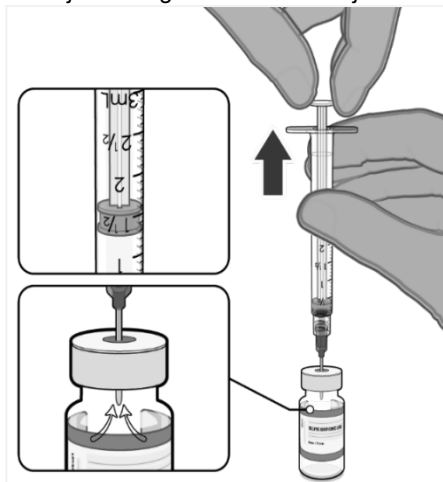
DILWIZZJONI TA' COMIRNATY 10 MIKROGRAMMI/DOŽA KONĊENTRAT GĦAL DISPERSJONI GĦALL-INJEZZJONI (TFAL MINN 5 SNIN SA 11-IL SENA)

- It-tilqima li tkun inħallet mis-silġ għandha tiġi dilwita fil-kunjett oriġinali ta' 1.3 mL ta' soluzzjoni għall-injezzjoni ta' 9 mg/mL (0.9%) sodium chloride, bl-użu ta' labra b'daqs ta' 21 gauge jew idjaq u tekniki asettiki.



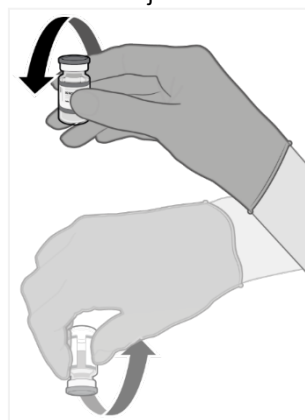
1.3 mL ta' injezzjoni ta' 0.9% sodium chloride

- Ugwalizza l-pressjoni fil-kunjett qabel ma tneħhi l-labra mit-tapp tal-kunjett billi tiġbed 1.3 mL ta' arja fis-siringa tad-dilwent vojta.



Iġbed il-plaġer lura sa 1.3 mL biex tneħhi l-arja mill-kunjett.

- Aqleb id-dispersjoni dilwita ta' taħt fuq bil-mod għal 10 darbiet. Thawwad.
- Il-vaċċin dilwit għandu jidher bħala dispersjoni ta' lewn abjad għal abjad maħmuġ mingħajr l-ebda partiċelli viżibbli. Tużax il-vaċċin dilwit jekk ikun hemm fraż jidur fil-kulur.



Bil-mod x 10

- Il-kunjetti dilwiti għandhom jiġu mmarkati bid-data u l-ħin xierqa.
- Wara d-dilwizzjoni, aħżen f'temperatura ta' 2 °C sa 30 °C u uża fi żmien 12-il siegħa.

- Tiffriżax u thawwad id-dispersjoni dilwita. Jekk titpoġġa fi friġġ, halli id-dispersjoni dilwita tilhaq it-temperatura tal-kamra qabel l-użu.



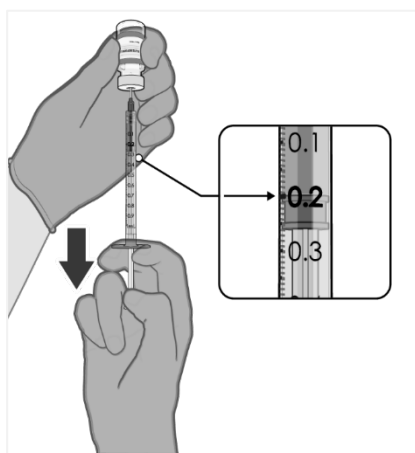
Irreġistra d-data u l-ħin xierqa. Uża fi żmien 12-il siegħa wara d-dilwizzjoni.

PREPARAZZJONI TA' DOŽI INDIVIDWALI TA' 0.2 mL TA' COMIRNATY 10 MIKROGRAMMI/DOŽA KONĊENTRAT GĦAL DISPERSJONI GĦALL-INJEZZJONI (TFAL MINN 5 SNIN SA 11-IL SENA)

- Wara d-dilwizzjoni, il-kunjett ikun fih 2.6 mL li minnhom jistgħu jinġibdu 10 doži ta' 0.2 mL.
- Bl-użu ta' teknika aseptika, naddaf it-tapp tal-kunjett b'imselha antiseptika li tintuża darba u tintrema.
- Iġbed 0.2 mL ta' Comirnaty għal tfal b'età minn 5 snin sa 11-il sena.

Sabiex jinġibdu 10 doži minn kunjett wieħed għandhom jintużaw siringi u/jew labar b'volum li ma jistax jintuża żgħir. Il-kombinazzjoni tas-siringa u tal-labra b'volum li ma jistax jintuża żgħir għandu jkollha volum li ma jistax jintuża ta' mhux aktar minn 35 mikrolitru.

- Jekk jintużaw siringi u labar standard, jista' ma jkunx hemm volum suffiċjenti biex jinġibdu għaxar doži minn kunjett wieħed.
- Kull doża għandu jkun fiha 0.2 mL ta' vaċċin.
- Jekk l-ammont ta' vaċċin li jifdal fil-kunjett ma jistax jipprovdi doża sħiħa ta' 0.2 mL, armi l-kunjett u kwalunkwe volum żejjed.
- Armi kwalunkwe vaċċin mhux użat fi żmien 12-il siegħa wara d-dilwizzjoni.



Vaċċin dilwit ta' 0.2 mL

Rimi

Kull fdal tal-prodott mediċinali li ma jkunx intuża jew skart li jibqa' wara l-użu tal-prodott għandu jintrema kif jitolbu l-liġijiet lokali.