

PATIENT PACKAGE INSERT IN ACCORDANCE WITH THE PHARMACISTS' REGULATIONS (PREPARATIONS) – 1986

The medicine is dispensed
with a doctor's prescription only

Escitalopram Teva 10 mg Tablets

The active ingredient and its quantity:
Each tablet contains:

Escitalopram (as oxalate) 10 mg

Escitalopram Teva 20 mg Tablets

The active ingredient and its quantity:
Each tablet contains:

Escitalopram (as oxalate) 20 mg

For information on the inactive ingredients,
see section 2 – "Important information about
some of the ingredients of the medicine"
and section 6 – "Further Information".

**Read the leaflet carefully in its entirety
before using the medicine.** This leaflet
contains concise information about the
medicine. If you have further questions,
refer to the doctor or pharmacist.

This medicine has been prescribed for you.
Do not pass it on to others. It may harm
them even if it seems to you that their
medical condition is similar.

This medicine is not usually intended for
use in children and adolescents under the
age of 18.

Antidepressants and anti-anxiety
medicines increase the risk of suicidal
behavior and thoughts in children,
adolescents and young adults up to the
age of 25.
Upon commencement of treatment
with the medicine, patients of all
ages, and their relatives, must monitor
behavioral changes, such as: worsening
of depression, suicidal thoughts,
aggressiveness and the like.
If such changes occur, refer to a doctor
immediately.

1. WHAT IS THE MEDICINE INTENDED FOR?

Escitalopram Teva is an antidepressant
from the selective serotonin reuptake
inhibitors (SSRI) group. Medicines
belonging to this group act on the serotonin
system in the brain by increasing serotonin
levels. Disruptions in the serotonin system
are considered important factors in
development of depression and diseases
associated with depression. Escitalopram
Teva is used to treat the following
conditions:

depression
panic disorders
generalized anxiety
social anxiety
obsessive-compulsive disorder

A few weeks may pass until you start to feel
better. Continue taking the medicine even
if it takes time until your feeling improves.

Talk to the doctor if you do not experience
an improvement or if you feel worse.

2. BEFORE USING THE MEDICINE

Do not use the medicine if:

- You are sensitive (allergic) to the active
ingredient (escitalopram) or to any of
the additional ingredients contained
in the medicine (see list of inactive
ingredients in section 6 – "Further
Information").
- You are taking other medicines
from the non-selective, irreversible
monoamine oxidase inhibitors
(MAOI) group, such as: phenelzine,
iproniazid, isocarboxazid, nialamide,
tranylcypromine. Wait 14 days after
taking these medicines before
beginning to use Escitalopram Teva,
or wait 7 days after taking Escitalopram
Teva before starting treatment with
these medicines. Likewise, do not
concomitantly take Escitalopram
Teva with reversible monoamine
oxidase A inhibitors (MAO-AI) (e.g.,
moclobemide to treat depression)
or the non-selective and reversible
monoamine oxidase inhibitor (MAOI)
linezolid (an antibiotic), or irreversible
monoamine oxidase B inhibitors
(MAO-BI) (e.g., selegiline for treatment
of Parkinson's). The combination with
these medicines increases the risk of
serotonin syndrome (see "Drug
interactions" in section 2).
- You are suffering from heart rhythm
disorders (as tested by ECG, a heart
function test), or from a congenital
defect in the electrical activity of the
heart.
- You are taking medicines to treat heart
rhythm disorders or which may have an
effect on the heart rhythm (see "Drug
interactions" in section 2).
- You are taking the medicine pimozide.

Special warnings regarding use of the medicine

**Before treatment with Escitalopram
Teva, tell the doctor if:**

- You are pregnant or breastfeeding (see
"Pregnancy, breastfeeding and fertility"
section).
- You are suffering, or have suffered in
the past, from epilepsy. Discontinue
treatment with Escitalopram Teva if you
have an epileptic attack for the first time
or if there is an increase in the frequency
of attacks (also see section 4 – "Side
Effects").
- You are suffering, or have suffered in
the past, from impaired liver or kidney
function. The doctor may need to adjust
the dosage of the medicine for you.
- You have diabetes. Treatment with
Escitalopram Teva may affect your sugar
level balance. The dosage of the insulin
and/or other diabetes medicines may
need to be adjusted.
- You have reduced blood sodium levels.
- You have a tendency to bleed easily or
develop bruises or, if you are pregnant
(see Pregnancy, breastfeeding and
fertility" section).
- You are receiving electroconvulsive
therapy (ECT).
- You are suffering, or have suffered in
the past, from coronary heart disease.
- You are suffering, or have suffered in
the past, from heart rhythm disturbances or
if you have recently suffered from a heart
attack.
- You have a low heart rate at rest and/or
if you know you may develop a decrease
in blood salts, as a result of prolonged
diarrhea or vomiting or as a result of
taking diuretics.
- You are suffering, or have suffered in
the past, from symptoms which may be
indicative of heart rhythm disorders such
as: rapid or irregular heart rate, fainting,
collapse or dizziness when getting up
from sitting or lying down.
- You have or, previously had, eye
problems, such as certain types of
glaucoma (increased intraocular
pressure).

Attention:

Some patients suffering from manic
depression may enter a state of mania,
characterized by unusual ideas, which
rapidly change, unexplained happiness
and more physical activity than usual. If
you experience these symptoms, refer to
a doctor.

Symptoms such as restlessness or difficulty
sitting or standing still may also occur in
the first weeks of treatment. If they occur,
report them immediately to the doctor.

Medicines such as Escitalopram Teva
(called SSRI/SNRI) may cause symptoms
of sexual function disturbances (see
section 4 – "Side Effects"). In certain
cases, the symptoms persisted even after
discontinuation of the treatment.

Panic disorder

When treating panic disorders, 2-4 weeks
usually pass before any improvement is felt.
Some patients may experience increased
anxiety at the beginning of treatment,
which will disappear during the first or second
week of treatment. Therefore, in cases of
panic disorders, it is recommended to start
treatment with a low dosage.

It is very important to strictly follow the
doctor's instructions and not to stop
treatment or change the dose without
consulting the doctor.

Suicidal thoughts and worsening of depression or of anxiety disorder

Suicidal thoughts or thoughts of self-harm
are common in psychiatric disorders,
especially in patients suffering from
depression and/or anxiety disorder. These
thoughts may increase at the beginning
of treatment with antidepressants, as
the effect of these medicines takes time,
usually two weeks, but sometimes longer.
The chance of occurrence of such thoughts
is higher if:

- You have had suicidal thoughts or
thoughts of self-harm in the past.
- You are a young adult. Data from clinical
trials show that there is increased risk of
suicidal behavior among adults below the
age of 25 with a psychiatric condition who
were treated with antidepressants.

If you have suicidal thoughts or thoughts
of self-harm at any time, **refer to your doctor
or proceed to a hospital immediately.**

**It is recommended that you tell a relative
or close friend that you are suffering from
depression or from an anxiety disorder, and
ask them to read this leaflet.** You can ask
them to tell you when they think there has
been a worsening of your depression or your
anxiety disorder, or if they are concerned
about a change that has occurred in your
behavior.

Children and adolescents

Escitalopram Teva is not usually intended
for use in children and adolescents under
the age of 18.

In patients under the age of 18 who have
taken medicines from this group, there is
an increased risk of side effects, such as
suicide attempts, suicidal thoughts and
hostility (particularly aggressiveness,
rebellious behavior and anger). Despite
this, your doctor can prescribe this
medicine for patients under the age of 18
if he thinks that this is for their benefit.

If the doctor has prescribed the medicine
for a patient under the age of 18 and you
are interested in discussing it – refer to the
doctor again.

Inform your doctor if some of the above
listed side effects occurred or became
worse in patients under the age of 18.
Likewise, the long-term effects of
Escitalopram Teva on growth, maturation
and cognitive-behavioral development in
this age group have not yet been tested.

Drug interactions

**If you are taking, or have recently
taken, other medicines, including non-
prescription medicines and nutritional
supplements, tell the doctor or
pharmacist. In particular, inform the
doctor or pharmacist if you are taking:**

- Do not use with non-selective monoamine
oxidase inhibitors (MAOI) (such as:
phenelzine, iproniazid, isocarboxazid,
nialamide, tranylcypromine). If you took
one of these medicines, you should wait
14 days before starting treatment with
Escitalopram Teva. After discontinuing
treatment with Escitalopram Teva,
you should wait 7 days before starting
treatment with these medicines.
- Do not use together with selective,
reversible monoamine oxidase A
inhibitors (MAO-AI), such as
moclobemide (for treatment of
depression).
- Do not use together with irreversible
monoamine oxidase B inhibitors
(MAO-BI), such as selegiline (to treat
Parkinson's disease). They may increase
the risk of side effects.
- Do not use together with the antibiotic
linezolid.
- Lithium (for treatment of manic
depression) and tryptophan.
- Imipramine and desipramine (for
treatment of depression).
- Sumatriptan and similar medicines (for
treatment of migraine) and also tramadol
and other similar medicines (opioids
used to relieve severe pain). These
medicines may increase the risk of side
effects.
- Cimetidine, lansoprazole and
omeprazole (for treatment of gastric
ulcer), fluconazole (for treatment of fungal
infections), flvoxamine (for treatment
of depression) and ticlopidine (to
reduce risk of stroke). These medicines
may increase the concentration of
Escitalopram Teva in the blood.
- St. John's wort (*Hypericum perforatum*,
a herbal medicine for treatment of
depression).
- Aspirin (acetylsalicylic acid) and
non-steroidal anti-inflammatory drugs
(NSAIDs) (medicines to relieve pain or
thin the blood, anticoagulants). These
medicines may increase the tendency
for bleeding.
- Warfarin, dipyridamole and
phenprocoumon (blood thinners,
anticoagulants): the doctor will monitor
your blood coagulation time at the start
of treatment and at the end of treatment
with Escitalopram Teva, to makes sure
that the dosage of the blood-thinner
medicine you are taking is appropriate.
- Medicines that may lower your sensitivity
threshold for convulsions: meprobolone
(for treatment of malaria), mefloquine
(for treatment of depression) and tramadol
(for treatment of severe pain), neuroleptic
preparations (medicines for treatment
of schizophrenia and psychoses), and
antidepressants (from the tricyclic
antidepressant group, and SSRIs).
- Flecainide, propafenone and
metoprolol (for treatment of heart and
vascular diseases), clomipramine,
nortriptyline (antidepressants) and
risperidone, thioridazine and haloperidol
(antipsychotics). It may be necessary to
adjust the dosage of Escitalopram Teva.
- Medicines that lower blood potassium or
magnesium levels, as such conditions
increase the risk of life-threatening
arrhythmias.

**Do not take Escitalopram Teva
concomitantly with medicines to treat
heart rhythm disorder or with medicines
that may have an effect on heart rhythm,
such as class IA and III anti-arrhythmias,
antipsychotics (e.g., phenothiazine
derivatives, pimozide, haloperidol), tricyclic
antidepressants, certain antimicrobial
agents (e.g., sparflloxacin, moxifloxacin,
erythromycin administered intravenously,
pentamidine, anti-malarials particularly
halofantrine), certain antihistamines
(for the treatment of allergy, astemizole,
hydroxyzine, mizolastine). If you have
further questions on this matter, refer to
the doctor.**

• SNRI and SSRI antidepressants may
cause serotonin syndrome (rare),
which includes symptoms such as
nervousness, restlessness, confusion,
sweating, high fever, increased reflexes,
muscle spasms, tremors, increase in
heart rate (see section 4 – "Side Effects").
The syndrome may occur at a higher
frequency if you are concomitantly
taking additional medicines that affect
the nervous system (see above).
Therefore, inform the doctor about any
other medicine you are taking.

Use of the medicine and food

The medicine can be taken with or without
food.

Use of the medicine and alcohol consumption

As with many medicines, it is recommended
not to consume alcohol with Escitalopram
Teva, although no interaction between
Escitalopram Teva and alcohol is expected.

Pregnancy, breastfeeding and fertility

If you are pregnant or breastfeeding,
think you are pregnant or are planning
a pregnancy, consult with the doctor or
pharmacist before using this medicine.

Do not use Escitalopram Teva if you are
pregnant or breastfeeding, unless you have
consulted your doctor and discussed with
him the risk versus benefit of taking this
medicine.

If you took Escitalopram Teva during the
last 3 months of pregnancy, the following
signs may occur in a newborn baby:
breathing problems, blue skin, seizures,
change in body temperature, eating
problems, vomiting, low blood sugar level,
stiff or flaccid muscles, increased reflexes,
tremor, jitteriness, irritability, exhaustion,
persistent crying, sleepiness, sleeping
difficulties. If your baby shows these signs,
refer to the doctor immediately.

Make sure your midwife and/or doctor
know that you are taking Escitalopram
Teva. When taking medicines such as
Escitalopram Teva during pregnancy,
especially in the last 3 months of the
pregnancy, there is an increased risk of
a serious condition in babies called
persistent pulmonary hypertension of the
newborn (PPHN), which causes the baby
to breathe faster and to appear blue. These
symptoms usually appear during the first 24
hours after birth. If these symptoms occur
in your baby, refer to the midwife and/or
doctor immediately.

If you take Escitalopram Teva near the
end of your pregnancy, there may be an
increased risk of heavy vaginal bleeding
shortly after birth, especially if you have
a history of bleeding disorders. Your doctor
or midwife should be aware that you are
taking Escitalopram so they can advise you.

If you used Escitalopram Teva during
pregnancy, do not discontinue use abruptly.
Escitalopram may pass into breast milk
and therefore, it is not recommended to
breastfeed while using the medicine.

Animal studies have shown that citalopram,
a medicine similar to escitalopram, reduces
sperm quality. Theoretically, this may affect
fertility; however, no effect on human fertility
has yet been shown.

Driving and use of machines

It is recommended not to drive a vehicle
or operate dangerous machinery, or to
engage in any activity that require alertness
before knowing how Escitalopram Teva
affects you.

**Important information about some of
the ingredients of the medicine**

This medicine contains less than 1 mmol
(23 mg) sodium per tablet and is therefore
considered sodium-free.

3. HOW SHOULD YOU USE THE MEDICINE?

Always use the preparation according to
the doctor's instructions. Check with the
doctor or pharmacist if you are not
sure about the preparation dosage and
treatment regimen.

**The dosage and treatment regimen will
be determined by the doctor only. The
usual dosage is generally:**

Adults

Depression: The recommended dosage
is generally 10 mg once a day (once-daily
dosage). Your doctor may increase the
dosage up to 20 mg per day.

Panic disorder: The starting dosage is
5 mg per day (once-daily dosage) for the
first week, and afterwards, an increase in
the dosage to 10 mg per day. Your doctor
can increase the dosage up to 20 mg per
day.

Social anxiety: The recommended dosage
is usually 10 mg once a day (once-daily
dosage). Your doctor can lower the dosage
to 5 mg per day or increase the dosage
up to 20 mg per day, according to your
response to treatment with the medicine.

Anxiety disorder: The recommended
dosage is usually 10 mg once a day
(once-daily dosage). Your doctor can
increase the dosage up to 20 mg per day.

Obsessive-compulsive disorder: The
recommended dosage is usually 10 mg
once a day (once-daily dosage). Your
doctor can increase the dosage up to 20 mg
per day.

Elderly (above the age of 65)

The recommended dosage is usually
5 mg once a day (once-daily dosage). Your
doctor can increase the dosage up to 10 mg
per day.

Children and adolescents

Escitalopram Teva is not usually intended
for use in children and adolescents (see
section 2 – "Before Using The Medicine").

Impaired kidney function

It is recommended that caution be
exercised in patients with severe kidney
function impairment. Take the medicine as
prescribed by the doctor.

Impaired liver function

These patients should receive no more
than 10 mg of the medicine per day. Take
the medicine as prescribed by the doctor.

**Patients known to be poor metabolizers
of the CYP2C19 enzyme**

Patients known to have this genotype
should receive no more than 10 mg of the
medicine per day. Take the medicine as
recommended by the doctor.

Do not exceed the recommended dose.

How to use

Swallow the medicine with some water.
The medicine can be taken with or without
food. Do not chew or crush the tablet, since
it has a bitter taste.

The tablet can be halved on the scoreline.

Duration of treatment

A few weeks may pass until you start to
feel better. Continue taking the medicine
even if it takes time until you feel better.
Do not change the dosage without consulting
the doctor.

Complete the treatment as recommended
by the doctor. Premature discontinuation
of treatment may cause the symptoms to
recur. It is recommended to adhere to
the treatment for at least 6 months after
you feel better.

**If you accidentally took an overdose
or if a child accidentally swallowed the
medicine, refer immediately to the doctor
or proceed to a hospital emergency room,
and bring the package of the medicine with
you. Do so even if you do not feel bad.**

Possible symptoms of an overdose are:
dizziness, tremor, nervousness, convulsions,
coma, nausea, vomiting, changes in heart
rate, decreased blood pressure, changes in
body salt/fluid balance.

**If you forgot to take this medicine at
the required time, do not take a double
dose.** If you forgot to take the medicine
and you remembered before going to sleep,
take the forgotten dose immediately and
continue as usual the following day. If you
only remembered during the night or the
following day, do not take the forgotten dose
and continue as usual.

Adhere to the treatment as recommended
by the doctor. Use this medicine at set
intervals as determined by the attending
doctor. Even if there is an improvement in
your health, do not stop treatment with the
medicine without consulting with the doctor
or pharmacist.

If you stop taking the medicine:

Do not stop treatment with Escitalopram
Teva before being instructed to do so by
the doctor. When completing treatment,
it is usually recommended to gradually
lower the dosage of the medicine over
several weeks. Abrupt discontinuation
of the treatment may cause the common
symptoms with discontinuation of
Escitalopram Teva treatment. The risk of
these symptoms occurring is higher when
Escitalopram Teva is used for a long time
or at a high dosage, or if the treatment is
reduced too quickly. In most patients, these
symptoms are mild or go away on their
own within two weeks. However, in some
patients, the symptoms can be severe or
can persist for a longer period of time (2-3
months and more). If you experience the
symptoms of treatment discontinuation on
completion of treatment with Escitalopram
Teva, consult the doctor. The doctor will
probably ask you to take the medicine
again and to lower the dosage more slowly.

Symptoms of treatment discontinuation
include: dizziness (unsteadiness or loss
of balance), feeling pins and needles
("prickling"), sensation of burning and
electric current (less common), including
in the head, sleep disturbances (vivid
dreams, nightmares, inability to sleep),
feeling anxious, headaches, nausea,
sweating (including night sweats), a
feeling of restlessness or nervousness,
tremor, confusion or disorientation, feeling
emotional or irritable, diarrhea, visual
disturbances, palpitations.

**Do not take medicines in the dark!
Check the label and the dose
each time you take the medicine. Wear
glasses if you need them.**

**If you have further questions regarding
use of the medicine, consult the doctor
or pharmacist.**

4. SIDE EFFECTS

**As with any medicine, use of
Escitalopram Teva may cause side
effects in some users. Do not be alarmed
when reading the list of side effects. You
may not suffer from any of them.**

These effects usually pass after a few
weeks of treatment. Note that some of
the side effects may be symptoms of your
illness and will therefore improve when you
begin to feel better.

**Refer to the doctor or proceed to a
hospital immediately if the following
symptoms occur:**

**Uncommon, frequency of up to 1 in 100
patients:**

Unusual bleeding, including gastrointestinal
bleeding.

Rare, frequency of up to 1 in 1,000 patients:
Swelling of the skin, tongue, lips, throat or
face, urticaria or breathing or swallowing
difficulties (signs of a severe allergic
reaction); High fever, nervousness or
restlessness, confusion, tremor, strong
muscle contraction may be symptoms of
a rare condition called serotonin syndrome.

Unknown frequency:
Difficulty passing urine; Convulsions (also
see "Special warnings regarding use of
the medicine" in section 2); Yellowing
of the skin and white area of the eyes,
which are signs of liver function problems/
hepatitis; Rapid and irregular heartbeat,
and fainting may be symptoms of a
life-threatening condition called torsade de
pointes (heart rate disorder); Thoughts of
self-harm or suicidal thoughts (also see
"Special warnings regarding use of the
medicine" in section 2); Sudden swelling of
the skin or mucous tissues (angioedema).

Additional side effects:

**Occur very frequently (frequency of more
than 1:10):**

Nausea, headache.

Occur frequently (frequency of up to 1:10):
Nasal congestion (sinusitis), reduced or
increased appetite, anxiety, restlessness,
abnormal dreams, difficulty falling
asleep, sleepiness, dizziness, yawning,
tremor, tingling sensation in the skin,
diarrhea, constipation, vomiting, dry
mouth, increased sweating, muscle and
joint pain, sexual function disturbances
(delayed ejaculation, problem with
erection, decreased sexual desire, difficulty
achieving orgasm in women), fatigue, fever,
weight gain.

**Occur infrequently (frequency of up to
1:100):**

Rash, itchy skin, teeth grinding, excitement,
nervousness, panic attack, confusion,
sleep disturbances, disturbances in sense
of taste, fainting, enlarged pupils, visual
disturbances, ringing in the ears (tinnitus),
hair loss, heavy menstrual bleeding,
abnormal menstrual cycle, reduction in
weight, fast heart rate, swelling of the limbs,
nosebleed.

Occur rarely (frequency of up to 1:1,000):
Aggressiveness, depersonalization,
hallucinations, slow heart rate.

Side effects of unknown frequency:
Decrease in blood sodium level (the
symptoms are malaise and nausea with
muscle weakness or confusion), dizziness
when getting up from sitting or from lying
down due to low blood pressure, abnormal
liver functions (increased liver enzymes in
the blood), movement disorders (involuntary
muscle movements), painful erection,
bleeding (including bleeding of the skin and
mucous tissues), abnormal ADH secretion,
which causes the body to retain water and
for the blood to thin, and to a reduction in
the amount of sodium, milk discharge in
men and in women that are not breastfeeding,
mania, an increased risk of fractures has
been observed in patients being treated with
this type of medicine, changes in the heart
rhythm (called prolongation of QT interval,
which can be seen in an ECG test of the
electrical activity of the heart). Heavy vaginal
bleeding shortly after birth, see further
information in "Pregnancy, breastfeeding
and fertility" section that can be found in
section 2 of the leaflet.

In addition, several side effects are known
to occur when taking medicines which act
similarly to escitalopram (the active ingredient
of Escitalopram Teva), which are: akathisia
(motor restlessness), loss of appetite.

**If a side effect occurs, or if one of the side
effects worsens or if you suffer from a
side effect not mentioned in the leaflet,
consult with the doctor.**

Reporting side effects

Side effects can be reported to the
Ministry of Health by clicking on the link
"Report Side Effects of Drug Treatment"
found on the Ministry of Health homepage
(www.health.gov.il) that directs you to the
online form for reporting side effects, or by
entering the link:
<https://sideeffects.health.gov.il>

5. HOW SHOULD THE MEDICINE BE STORED?

• **Avoid poisoning!** This medicine and
any other medicine must be kept in a
safe place out of the reach and sight of
children and/or infants in order to avoid
poisoning. Do not induce vomiting without
an explicit instruction from the doctor.

• Do not use the medicine after the expiry
date (exp. date) that appears on the
package. The expiry date refers to the
last day of that month.

• **Storage conditions:** Store the medicine
in the original package in order to protect
from light and moisture, below 25°C.

• Do not discard medicines into the
wastewater or waste bin. Ask the
pharmacist how to dispose of medicines
no longer in use. These measures will
help to protect the environment.

6. FURTHER INFORMATION

**In addition to the active ingredient, the
medicine also contains:**

Microcrystalline cellulose, croscarmellose
sodium, stearic acid, magnesium stearate,
colloidal silicon dioxide, hypromellose,
titanium dioxide, polyethylene glycol.

**What the medicine looks like and the
contents of the package:**

Escitalopram Teva 10 mg: White,
film-coated, round convex tablet, scored
on one side and debossed with "10" on the
other side.

Escitalopram Teva 20 mg: White,
film-coated, round convex tablet, scored on
one side and debossed with "20" on the left
side of the score and "3" on the right side
of the score. The other side of the tablet is
debossed with "7463".

There are packages of 28 or 30 tablets.
Not all package sizes may be marketed.

**Name of Manufacturer and License
Holder and Address:**

Teva Israel Ltd.,
124 Dvora HaNevi'a St., Tel Aviv 6944020.

**The leaflet was revised in November
2022 according to MOH guidelines.**

**Registration number of the medicine
in the National Drug Registry of the
Ministry of Health:**

Escitalopram Teva 10 mg:
137.84.31381

Escitalopram Teva 20 mg:
137.85.31382

teva

ESCITAB PL SH 031222