



CAPRELSA (VANDETANIB)
DOSING AND MONITORING
GUIDE FOR PATIENTS AND
PATIENTS' CAREGIVERS
(PAEDIATRIC USE)

For full information, please read the Patient Information Leaflet

This document is approved by The Executive Directorate of Pharmacovigilance, at SFDA

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Table of Contents

- **What is Caprelsa® and what does it treat?**
- **How is the dose of Caprelsa ® calculated?**
- **How is Caprelsa® used?**
- **What are the side effects associated with Caprelsa® ?**
- **Which monitoring is requested?**
- **How to use the daily tracker table ?**
- **GENERAL DAILY TRACKER FOR 14 DAYS**
- **EXAMPLE OF STARTING DOSE FOR A CHILD WITH BSA FROM 0.7 m² to <0.9 m²**
- **EXAMPLE OF INCREASED DOSE FOR A CHILD WITH BSA FROM 0.9m² to <1.2m²**
- **EXAMPLE OF INCREASED DOSE FOR A CHILD WITH BSA > 1.6 m²**

What is Caprelsa and what does it treat?

Caprelsa is a medicine that contains the active substance vandetanib. It is available in film-coated tablets (100 mg and 300 mg). It is used to treat medullary thyroid cancer that is called Rearranged during Transfection (RET) mutant and which cannot be removed by surgery or has spread to other parts of the body.

Caprelsa works by slowing down the growth of new blood vessels in tumours (cancers). This cuts off the supply of food and oxygen to the tumour. Caprelsa may also act directly on cancer cells to kill them or slow down their growth.

How is the dose of Caprelsa calculated?

The calculation of the dose of vandetanib is made by the treating physician, based on the child/adolescent's body surface area (BSA), depending of body height and weight of the patient.

Depending on the calculated BSA, the physician will prescribe to your child a **starting dose**, which can be changed (dose adjustments):

- for an **increased dose**, if vandetanib is well tolerated after 8 weeks at starting dose
- for a **reduced dose** in case of undesirable side effects, after a suspension of treatment (at least a week)

The dose can also change if the BSA changes during the treatment.

The treatment schedule will correspond to one of the 3 following schemes:

“daily” schedule (same dose every day)*

“every other day” schedule (same dose every other day)*

“7 day” schedule (treatment every day but two different doses alternately)*

Please note that the dosing scheme can change during treatment. For example, you can follow a daily schedule for the starting dose period and switch to a 7 day schedule after a dose adjustment.

You will have to report each dose taken on a daily tracker table (see below).

How is Caprelsa used?

The prescribed calculated dose should be taken:

- at about the same time
- with or without food.

The total daily dose in children must not exceed 300 mg.

If the child has trouble swallowing the tablet, you can mix it with water as follows:

- Take half a glass of still (non carbonated) water. Only use water, do not use any other liquids.
- Put the tablet into the water.
- Stir the tablet until it has dispersed into the water. This may take about 10 minutes.
- Then make the child drink it straight away.
- To ensure no medicine remains, refill the glass halfway with water and ask your child to finish it.

What are the side effects associated with Vandetanib ? Which monitoring is requested?

Your doctor will inform you of the main risks of vandetanib. Please also read carefully the package leaflet for more information about the drug. The most commonly reported side effects with vandetanib are :

- diarrhoea
- rash or other skin reaction
- nausea (feeling sick)
- hypertension (high blood pressure) and headache.

Monitoring of blood and heart will be necessary BEFORE and regularly DURING the treatment by vandetanib, especially:

- blood potassium, calcium, magnesium, and thyroid stimulating hormone (TSH)
- the electrical activity of the heart with a test called an electrocardiogram (ECG)

A good skin protection is requested (wearing clothes, sunscreen), especially if you are sensitive to sun.

Tell your doctor or pharmacist if you are taking, have recently taken or might take any other medicines, including medicines that you buy without a prescription and herbal medicines. These might interact with vandetanib and provoke lack of efficacy or increase side effects.

Report any side effect to your doctor. He/she may prescribe other medicines to help control the patient's side effects. A suspension of treatment and dose reduction may also be necessary.

Tell your doctor straight away if you notice any of the following side effects – you may need urgent medical treatment:

- Fainting, dizziness or heart rhythm changes. These may be signs of a change in the electrical activity of the heart. They are seen in 8% of people taking Caprelsa for medullary thyroid cancer. Your doctor may recommend you take Caprelsa at a lower dose or stop taking Caprelsa. Caprelsa has uncommonly been associated with life threatening changes in heart rhythm.
- Severe skin reactions affecting large areas of your body. The signs may include redness, pain, ulcers, blisters and shedding of the skin. The lips, nose, eyes and genitals may also be affected. These may be common (affecting less than 1 in 10 people) or uncommon (affects less than 1 in 100 people) depending on the type of skin reaction.
- Severe diarrhoea.
- Serious breathlessness, or sudden worsening breathlessness, possibly with a cough or a high temperature (fever). This may mean that you have an inflammation of the lungs called 'interstitial lung disease'. This is uncommon (affects less than 1 in 100 people) but can be life-threatening.
- Seizures, headache, confusion or finding it difficult to concentrate. These may be signs of a condition called RPLS (Reversible Posterior Leukoencephalopathy Syndrome). These usually go away when Caprelsa is stopped. RPLS is uncommon (affects less than 1 in 100 people).

How to use the daily tracker table ?

While prescribing the starting dose, your doctor will complete the “prescriber part” of the daily tracker and explain how to use it. The daily tracker is made to help you:

- to remember when to take a new dose and which dose. It has to be completed by you after each dose intake.
- to report side effects and follow dose adjustments.

The daily tracker is **adapted to all dose regimens**. In the event that a dose change is made, a new daily tracker sheet should be provided by your doctor to the patients and/or patient's caregivers.

Please find blank copies of the daily tracker after the examples of completed daily trackers.

GENERAL DAILY TRACKER FOR 14 DAYS







Space reserved prescriber		Daily tracker for patient			
Weight : _____ Height : _____ BSA: _____ m ² Date of prescription : _____ <input type="checkbox"/> starting dose <input type="checkbox"/> increased dose <input type="checkbox"/> reduced dose (see posology recommendations)		Name of patient : _____ Date of Birth : _____ If you forget to take Caprelsa : - If it is 12 hours or more until your next dose: take the missed tablet as soon as you remember. Then take the next dose at the normal time. - If it is less than 12 hours until your next dose: skip the missed dose. Then take the next dose at the normal time. DO NOT TAKE a double dose (two doses at the same time) to make up for a forgotten tablet.			
Day of week	Dose prescribed	Week 1-2 Start:	Week 3-4 Start:	Week 5-6 Start:	Week 7-8 Start:
Monday D1					
Tuesday D2					
Wednesday D3					
Thursday D4					
Friday D5					
Saturday D6					
Sunday D7					

Day of week	Dose prescribed	Week 1-2 Start:	Week 3-4 Start:	Week 5-6 Start:	Week 7-8 Start:
Monday D8					
Tuesday D9					
Wednesday D10					
Thursday D11					
Friday D12					
Saturday D13					
Sunday D14					

Doses available	Comments for patient and/or patient's caregiver (side effects, other treatment, or important information)			
 100 mg =				
  2 tablets of 100 mg =				
 300 mg =				

**EXAMPLE OF STARTING DOSE FOR A CHILD WITH BSA FROM
0.7 m² to <0.9 m² ("EVERY OTHER DAY" SCHEDULE : D1 ≠ D8)**




Space reserved prescriber		Daily tracker for patient				
Weight : _____ Height : _____ BSA : <u>0,8</u> m ² Date of prescription : <u>12/09/16</u> <input checked="" type="checkbox"/> starting dose <input type="checkbox"/> increased dose <input type="checkbox"/> reduced dose (see posology recommendations)		Name of patient : _____ Date of Birth : _____ If you forget to take Caprelsa : - If it is 12 hours or more until your next dose: take the missed tablet as soon as you remember. Then take the next dose at the normal time. - If it is less than 12 hours until your next dose: skip the missed dose. Then take the next dose at the normal time. DO NOT TAKE a double dose (two doses at the same time) to make up for a forgotten tablet.				
Day of week	Dose prescribed	Week 1-2 Start: <u>12/09/16</u>	Week 3-4 Start: <u>26/09/16</u>	Week 5-6 Start: <u>10/10/16</u>	Week 7-8 Start: <u>24/10/16</u>	
Monday D1	-	0	0	0	0	
Tuesday D2	100 mg	1 x 100 mg	1 x 100 mg	1 x 100 mg	1 x 100 mg	
Wednesday D3	-	0	0	0	0	
Thursday D4	100 mg	1 x 100 mg	1 x 100 mg	1 x 100 mg	1 x 100 mg	
Friday D5	-	0	0	0	0	
Saturday D6	100 mg	1 x 100 mg	1 x 100 mg	1 x 100 mg	1 x 100 mg	
Sunday D7	-	0	0	0	0	

Day of week	Dose prescribed	Week 1-2 Start:	Week 3-4 Start:	Week 5-6 Start:	Week 7-8 Start:
Monday D8	100 mg	12/09/16 1 x 100 mg	26/09/16 1 x 100 mg	10/10/16 1 x 100 mg	24/10/16 1 x 100 mg
Tuesday D9	-	0	0	0	0
Wednesday D10	100 mg	1 x 100 mg	1 x 100 mg	1 x 100 mg	1 x 100 mg
Thursday D11	-	0	0	0	0
Friday D12	100 mg	1 x 100 mg	1 x 100 mg	1 x 100 mg	1 x 100 mg
Saturday D13	-	0	0	0	0
Sunday D14	100 mg	1 x 100 mg	1 x 100 mg	1 x 100 mg	1 x 100 mg
Doses available	Comments for patient and/or patient's caregiver (side effects, other treatment, or important information)				
 100 mg =					
 200 mg =					
 300 mg =					
	<p>side effect : small skin reaction. called doctor, no interruption. recovered using stronger sun protection.</p>				
	<p>well tolerated after 8 week, new prescription with increased dose (100 mg daily) -- new daily tracker sheet.</p>				

EXAMPLE OF INCREASED DOSE FOR A CHILD WITH BSA FROM 0.9m² to <1.2m² ("7 DAYS" SCHEDULE: D1 = D8)

Space reserved prescriber		Daily tracker for patient				
Weight : <u>35 kg</u> Height : <u>125 cm</u> BSA : <u>1.1</u> m ² Date of prescription : <u>12/09/16</u> <input type="checkbox"/> starting dose <input checked="" type="checkbox"/> increased dose <input type="checkbox"/> reduced dose (see posology recommendations)		Name of patient : _____ Date of Birth : _____ If you forget to take Caprelsa : - If it is 12 hours or more until your next dose: take the missed tablet as soon as you remember. Then take the next dose at the normal time. - If it is less than 12 hours until your next dose: skip the missed dose. Then take the next dose at the normal time. DO NOT TAKE a double dose (two doses at the same time) to make up for a forgotten tablet.				
Day of week	Dose prescribed	Week 1-2 Start: <u>12/09/16</u>	Week 3-4 Start: <u>26/09/16</u>	Week 5-6 Start: <u>10/10/16</u>	Week 7-8 Start: <u>24/10/16</u>	
Monday D1	<u>100 mg</u>	<u>1 x 100 mg</u>	<u>1 x 100 mg</u>	<u>1 x 100 mg</u>		
Tuesday D2	<u>200 mg</u>	<u>2 x 100 mg</u>	<u>2 x 100 mg</u>	<u>2 x 100 mg</u>		
Wednesday D3	<u>100 mg</u>	<u>1 x 100 mg</u>	<u>1 x 100 mg</u>	<u>1 x 100 mg</u>		
Thursday D4	<u>200 mg</u>	<u>2 x 100 mg</u>	<u>2 x 100 mg</u>	<u>2 x 100 mg</u>		
Friday D5	<u>100 mg</u>	<u>1 x 100 mg</u>	<u>1 x 100 mg</u>	<u>suspended</u>		
Saturday D6	<u>200 mg</u>	<u>2 x 100 mg</u>	<u>2 x 100 mg</u>	<u>suspended</u>		
Sunday D7	<u>100 mg</u>	<u>1 x 100 mg</u>	<u>1 x 100 mg</u>	<u>suspended</u>		




Day of week	Dose prescribed	Week 1-2 Start:	Week 3-4 Start:	Week 5-6 Start:	Week 7-8 Start:
Monday D8	100 mg	12/09/16 1 x 100 mg	26/09/16 1 x 100 mg	10/10/16 suspended	24/10/16
Tuesday D9	200 mg	2 x 100 mg	2 x 100 mg	suspended	
Wednesday D10	100 mg	1 x 100 mg	1 x 100 mg	suspended	
Thursday D11	200 mg	2 x 100 mg	2 x 100 mg	suspended	
Friday D12	100 mg	1 x 100 mg	1 x 100 mg	suspended	
Saturday D13	200 mg	2 x 100 mg	2 x 100 mg	Restart with reduced dose	
Sunday D14	100 mg	1 x 100 mg	1 x 100 mg		

Doses available	Comments for patient and/or patient's caregiver (side effects, other treatment, or important information)			
 100 mg =	 200 mg =	 300 mg =	<p>increased dose after 8 w 100 mg daily tired (weakness)</p> <p>friday - w1 : diarrhoea (1 episode), called doctor-- no change</p> <p>tonsillitis: amoxicillin</p> <p>thursday - w 4 : small skin reaction, called doctor-- survey and no sun exposition.</p> <p>thursday - w 5 : skin reaction worsening, treatment suspended</p> <p>friday - w 6 : recovered from skin reaction.</p> <p>new prescription starting a reduced dose-- new daily tracker sheet.</p>	

**EXAMPLE OF INCREASED DOSE FOR A CHILD WITH BSA
> 1.6 m² ("DAILY" SCHEDULE: D1 = DX)**

Space reserved prescriber		Daily tracker for patient			
Weight : _____ Height : _____ BSA : <u>1.68</u> m ² Date of prescription : <u>12/09/16</u> <input type="checkbox"/> starting dose <input checked="" type="checkbox"/> increased dose <input type="checkbox"/> reduced dose (see posology recommendations)		Name of patient : _____ Date of Birth : _____ If you forget to take Caprelsa : - If it is 12 hours or more until your next dose: take the missed tablet as soon as you remember. Then take the next dose at the normal time. - If it is less than 12 hours until your next dose: skip the missed dose. Then take the next dose at the normal time. DO NOT TAKE a double dose (two doses at the same time) to make up for a forgotten tablet.			
Day of week	Dose prescribed	Week 1-2 Start: <u>12/09/16</u>	Week 3-4 Start: <u>26/09/16</u>	Week 5-6 Start: <u>10/10/16</u>	Week 7-8 Start: <u>24/10/16</u>
Monday D1	<u>300 mg</u>	<u>1 x 300 mg</u>	<u>1 x 300 mg</u>		
Tuesday D2	<u>300 mg</u>	<u>1 x 300 mg</u>	<u>1 x 300 mg</u>		
Wednesday D3	<u>300 mg</u>	<u>1 x 300 mg</u>	<u>1 x 300 mg</u>		
Thursday D4	<u>300 mg</u>	<u>1 x 300 mg</u>	<u>1 x 300 mg</u>		
Friday D5	<u>300 mg</u>	<u>1 x 300 mg</u>	<u>1 x 300 mg</u>		
Saturday D6	<u>300 mg</u>	<u>1 x 300 mg</u>	<u>1 x 300 mg</u>		
Sunday D7	<u>300 mg</u>	<u>1 x 300 mg</u>	<u>etc.</u>		

Day of week	Dose prescribed	Week 1-2 Start:	Week 3-4 Start:	Week 5-6 Start:	Week 7-8 Start:
Monday D8	300 mg	12/09/16 1 x 300 mg	26/09/16	10/10/16	24/10/16
Tuesday D9	300 mg	1 x 300 mg			
Wednesday D10	300 mg	1 x 300 mg			
Thursday D11	300 mg	1 x 300 mg			
Friday D12	300 mg	1 x 300 mg			
Saturday D13	300 mg	1 x 300 mg			
Sunday D14	300 mg	1 x 300 mg			

Doses available	Comments for patient and/or patient's caregiver (side effects, other treatment, or important information)			
<div>100 mg = </div>	<div>increased dose after 8 w 200 mg daily</div>			
<div>200 mg = </div>				
<div>300 mg = </div>				

Blank copies
Package leaflet has to be provided with dosing guide.

In case of any drug related adverse events, please
contact: The National Pharmacovigilance Center (NPC):
Call Center: 19999
E-mail: npc.drug@sfd.gov.sa
Website: <https://ade.sfd.gov.sa>



For SANOFI Pharmacovigilance center, please contact:
+966-544-284-797
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