

**David,** a real person with Parkinson's disease

For adults with Parkinson's disease (PD)

## FROM LUNCH TO LEASH TO LAKE

Stay in step with your day with CREXONT: More "Good On" time\* with less frequent dosing compared with immediate-release carbidopa/levodopa (IR CD/LD)<sup>+</sup>

\*"Good On" time is defined as "On" time without troublesome dyskinesia. †As seen in a clinical study.

#### INDICATION

CREXONT® (carbidopa and levodopa) extended-release capsules is a prescription medication for the treatment of Parkinson's disease, Parkinson's disease caused by infection or inflammation of the brain, or Parkinson's disease-like symptoms that may result from carbon monoxide or manganese poisoning in adults.

#### **IMPORTANT SAFETY INFORMATION**

Do not take CREXONT with antidepressant medications known as nonselective monoamine oxidase (MAO) inhibitors.

Please see additional Important Safety Information on other pages and accompanying <u>full Prescribing Information</u>.

# Are you optimizing your "Good On" time\* with your current PD treatment?

# Ups and downs with immediate-release CD/LD treatments are common

IR CD/LD has been the standard of care for many years. It is designed to work quickly but may wear off quickly too As Parkinson's progresses, you might notice that your IR CD/LD doesn't last as long, leading to ups and downs

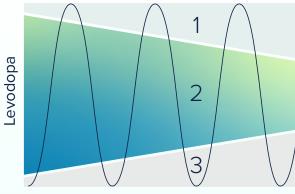
\*"Good On" time is defined as "On" time without troublesome dyskinesia.

#### **IMPORTANT SAFETY INFORMATION (cont'd)**

Do not take CREXONT with other carbidopa-levodopa preparations without consulting your healthcare provider.

CREXONT may cause falling asleep during activities of daily living, somnolence, or dizziness. Avoid activities that require alertness such as driving and operating machinery until you know how CREXONT affects you. The most common side effects that may occur with CREXONT are nausea and anxiety.

It is important to avoid sudden discontinuation or rapid dose reduction of CREXONT. If you are discontinuing CREXONT, work with your healthcare provider to taper the dose over time to reduce the risk of fever or confusion.





Over time with IR CD/LD, the duration of your symptom control may become more unpredictable, and you may experience more "Off" time. In fact, about half of patients taking a PD medication may experience "Off" episodes within 2 to 3 years of being on treatment

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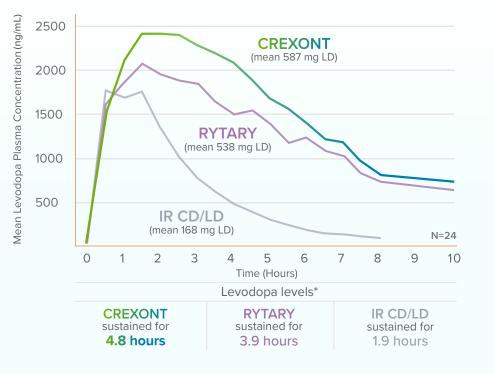


and symptoms return



### IN A CLINICAL PHARMACOLOGY STUDY. **CREXONT** levodopa levels were compared with other forms of oral CD/LD

### Main measurement: Levodopa levels of CREXONT compared with RYTARY<sup>®</sup> and IR CD/LD



(+)In a different analysis of the same study, **CREXONT** levodopa levels outlasted RYTARY and IR CD/LD\*+

\*In a clinical pharmacology study, the duration of levodopa levels was defined by how long the levels of CREXONT, RYTARY, and IR CD/LD were maintained in the blood above half the maximum concentration.

<sup>+</sup>This was a post hoc analysis of a secondary measure in the clinical pharmacology study; CREXONT vs RYTARY P=0.010: CREXONT vs IR CD/LD P<0.0001.

**Please see additional Important** Safety Information on other pages and accompanying full **Prescribing Information.** 

### IN A CLINICAL PHARMACOLOGY STUDY, CREXONT PROVIDED The longest-lasting oral levodopa levels\*\*

### Discover the dual components inside CREXONT:

Immediate-release granules Dissolve quickly, so levodopa levels peak at ~1 hour



Capsule and contents shown for illustrative purposes only.



#### **IMPORTANT SAFETY INFORMATION (cont'd)**

You may take CREXONT with or without food, but taking it with food may decrease or delay its effect. Consider taking the first dose of the day about 1 to 2 hours before eating.

Extended-release pellets Release levodopa slowly so levels last longer\*



#### CREXONT is designed to gradually release levodopa in the part of the gut where it is best absorbed<sup>‡</sup>

\*In a clinical pharmacology study, longest lasting was defined as how long the concentration of CREXONT, RYTARY, and IR CD/LD stayed in the blood above half the maximum concentration.

<sup>+</sup>This was a post hoc analysis of a secondary measure in the clinical pharmacology study; CREXONT vs RYTARY P=0.010; CREXONT vs IR CD/LD P<0.0001

<sup>‡</sup>The exact site and duration of absorption is unknown.



### Explore the benefits of CREXONT

#### Main measurement

### 30 mins more "Good On" time per day with less frequent dosing\*\*

In a clinical study, people who took CREXONT an average of 3 times a day achieved 30 minutes more "Good On" time per day than those taking IR CD/LD an average of 5 times a day

\*"Good On" time is defined as "On" time without troublesome dyskinesia. Change in "Good On" time was measured by comparing values at the end of study to baseline; P=0.019 vs IR CD/LD.

<sup>+</sup>This was the average dose frequency during the double-blind maintenance period.

The most common side effects (incidence  $\geq$ 3% and greater than IR CD/LD) were nausea and anxiety.

Secondary measurement

30 mins less "Off" time per day<sup>±§</sup>

People who took CREXONT had 30 minutes less "Off" time per day compared with those who took IR CD/LD

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<sup>‡</sup>This was a secondary endpoint of the study. <sup>§</sup>Study end=Week 20 or early termination: P=0.025 vs IR CD/LD

### IN A DIFFERENT ANALYSIS OF THE SAME STUDY. CREXONT provided more "Good On" time" per dose

A different analysis of the same study

### >1 hr 30 mins more "Good On" time" per dose<sup>1</sup>

In a clinical study of a dose-to-dose comparison, people took CREXONT fewer times per day and experienced over an hour and a half more "Good On" time per dose than people taking IR CD/LD

"Good On" time is defined as "On" time without troublesome dyskinesia. <sup>1</sup>This was a post hoc analysis of the study; P<0.0001 vs IR CD/LD.

#### **IMPORTANT SAFETY INFORMATION (cont'd)**

Swallow CREXONT whole. Do not chew, divide, or crush the capsules. Do not take CREXONT with alcohol.

Please see additional Important Safety Information on other pages and accompanying full Prescribing Information.







### Your dose of CREXONT will be personalized

**CREXONT** dosina may start at 2 to 3 times per day



A typical dose of **CREXONT** is 1 or 2 capsules

After 1 to 3 days on treatment, your doctor may adjust the frequency and dosage of your medicine. Your frequency may go up to 4 times per day

Do not take CREXONT with antidepressant medications known as nonselective monoamine oxidase (MAO) inhibitors.

#### **IMPORTANT SAFETY INFORMATION (cont'd)**

Tell your healthcare provider if you:

3 times

per day for

most patients

- Have any heart conditions, especially if you have had a heart attack or irregular heartbeats
- Experience hallucinations or abnormal thoughts and behaviors
- Have an inability to control urges to gamble, have increased sexual urges, or experience other intense urges

- Have thoughts of suicide or have attempted suicide
- Have abnormal involuntary movements that appear or get worse during treatment
- Have ever had a peptic ulcer or glaucoma
- Become or intend to become pregnant. Based on animal data, CREXONT may cause fetal harm
- Are breastfeeding during therapy

## Multiple dose strengths for dosing flexibility

CREXONT comes in 4 dose strengths

Capsules shown are not actual size





35 mg carbidopa 140 mg levodopa

52.5 mg carbidopa 210 mg levodopa

You can work with your doctor to find which dose will help you achieve the "Good On" time\* you need

\*"Good On" time is defined as "On" time without troublesome dyskinesia.

Please see additional Important Safety Information on other pages and accompanying full Prescribing Information.





### CREXONT was generally well-tolerated

Side effects that occurred in ≥2% of participants while switching from IR CD/LD to CREXONT and also occurred at a higher rate than IR CD/LD in the double-blind part of the study

	Switching from IR CD/LD to CREXONT	Double-blind phase	
	CREXONT (n=589)	CREXONT (n=256)	IR CD/LD (n=250)
Nausea	5%	4%	1%
Anxiety	2%	3%	0%
Dizziness	3%	2%	1%
Dyskinesia	7%	2%	0.4%
Constipation	2%	2%	0.4%
Headache	2%	1%	0%
Vomiting	2%	1%	0%
Insomnia	2%	1%	0.4%

### Established safety profile of CREXONT

YEAR

To report SUSPECTED ADVERSE REACTIONS, contact Amneal Specialty, a division of Amneal Pharmaceuticals, LLC at 1-877-835-5472 or the FDA at 1-800-FDA-1088 or www.fda.gov/medwatch.

Please see additional Important Safety Information on other pages and accompanying full Prescribing Information.

CREXONT was studied for over a year in adults with Parkinson's disease, which established its well-understood safety profile



## Could CREXONT be right for you?



More "Good On" time per day with less frequent dosing compared with IR CD/LD\*





- CREXONT was generally well-tolerated
- Your healthcare provider will personalize your CREXONT dosing

\*\*Good On" time is defined as "On" time without troublesome dyskinesia. Change in "Good On" time was measured by comparing values at the end of study to baseline; P=0.019 vs IR CD/LD. People who took CREXONT an average of 3 times per day saw significant improvements in "Good On" time vs people who took IR CD/LD an average of 5 times per day. This was the average dose frequency during the double-blind maintenance period. "Good On" time is defined as "On" time without troublesome dyskinesia. \*This was a post hoc analysis of the study; P<0.0001 vs IR CD/LD.

Take your next step Ask your doctor about more "Good On" time with CREXONT



To access support and resources, scan this QR code



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