Onset and duration of effect of extended-release carbidopa-levodopa in advanced Parkinson’s disease

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Background: In patients with Parkinson’s disease (PD), oral dosing of extended-release carbidopa-levodopa (Rytary, IPX066 [ER CD-LD]) achieves peak levodopa plasma concentrations within 1 hour and maintains them for 4–6 hours.

Aims: To compare the onset and duration of ER CD-LD benefit with those of immediate-release carbidopa-levodopa (IR CD-LD) in PD patients with motor fluctuations, using crossover data, and to evaluate which threshold values of improvement in finger-tapping and Unified Parkinson’s Disease Rating Scale (UPDRS) motor scores yield results most similar to those for trained raters’ “on”/“off” assessments.

Methods: Patients underwent serial “on”/“off” rating and provided serial finger-tapping and UPDRS motor scores after receiving, in an “off” state, their usual morning IR dose or an ER dose designed to produce a similar levodopa peak concentration. Predefined improvement thresholds for analysis were 10%, 15%, and 20% increases in finger-tapping score and 2.5, 5, 7, and 11-point decreases in UPDRS motor score. Serial plasma samples were assayed for levodopa.

Results: Among 27 patients, mean time to onset of an “on” state was similar for ER compared with IR CD-LD (0.83 vs 0.81 hour), but mean duration was significantly longer for ER CD-LD than for IR CD-LD (5.56 vs 2.69 hours; P<0.0001). Duration was best matched by a ≥20% improvement in finger-tapping, a ≥11-point improvement in UPDRS motor score, and a levodopa plasma concentration ≥1,000 ng/mL.

Conclusion: For ER CD-LD, observer assessments of “on” state were corroborated by sustained treatment effects. Correlations among “on”-state duration, finger-tapping score, and UPDRS motor score may suggest clinically relevant thresholds for acute assessment of treatment benefit.

Keywords: Rytary, carbidopa-levodopa, Parkinson’s disease, treatment, duration of effect, motor fluctuations

Introduction
Carbidopa and levodopa extended-release capsules1 (ER CD-LD, Rytary, IPX066) are a multiparticulate oral formulation of carbidopa-levodopa (CD-LD) consisting of immediate-release (IR) and extended-release (ER) components, as well as the functional excipient tartaric acid.2 In patients with Parkinson’s disease (PD), the formulation achieves peak levodopa plasma concentrations within 1 hour after dosing and maintains them for 4–6 hours.3–6 Because of its distinct immediate and extended levodopa-releasing properties in the gastrointestinal tract, the plasma levodopa profile produced by an ER CD-LD dose differs markedly from those of other oral levodopa products.7 In a study of patients with advanced PD, single doses resulted in ~70% of the levodopa exposure seen after a similar milligram dose of IR CD-LD (as measured...
by area under the plasma concentration curve) and ~30% of
the peak levodopa plasma levels ($C_{\text{max}}$). For doses provid-
ing similar peak levodopa levels, the onset of improvement
of motor function observed for ER CD-LD resembled that
observed for IR CD-LD, as judged by change in score on
Part III of the Unified Parkinson’s Disease Rating Scale
(UPDRS) and by an investigator rating of motor state, and
the duration of effect was longer by both measures. For doses
providing similar peak levodopa levels, the onset of improve-
ment was taken in an “off” state. Patients experiencing three
levodopa keys 20 cm apart in 1 minute, using the index finger of the arm
more affected by PD, as determined at screening. A UPDRS
Part III score was obtained hourly by qualified study-site
personnel, twice pre-dose and for 8 hours post-dose.

For each pharmacokinetic/pharmacodynamic day, each
patient was assessed by a trained rater at approximate half-
hour intervals, three times pre-dose and for 8 hours post-dose,
as being “on” with no dyskinesia, “on” with non-troublesome
dyskinesia, “on” with troublesome dyskinesia, or “off”. For
each patient, the same rater performed all assessments.
In addition, finger-tapping was measured three times pre-dose
and half-hourly for 8 hours post-dose. Each test was scored
as the number of times the patient could alternately tap two
keys 20 cm apart in 1 minute, using the index finger of the arm
more affected by PD, as determined at screening. A UPDRS
Part III score was obtained hourly by qualified study-site
personnel, twice pre-dose and for 8 hours post-dose.

**Efficacy analyses**

For ER CD-LD and IR CD-LD, the half-hourly “on/off”
classifications made by trained raters were utilized to cal-
culate the mean time to onset and the mean duration of an
“on” state. Differences between ER CD-LD and IR CD-LD
were tested for statistical significance, set at $P<0.05$, using a mixed-model analysis of covariance (ANCOVA) with baseline value as a covariate.

Details on the finger-tapping scores have been reported previously. In the present post hoc analysis, to assess the proportions of patients attaining various levels of motor-function improvement, thresholds of improvement from a patient’s pre-dose value (obtained in an “off” state) were defined as increases of 10%, 15%, and 20% in finger-tapping score and decreases of 2.5, 5, 7, and 11 points in UPDRS Part III score. For each such threshold, the time at which each patient achieved the threshold was defined as the time to onset of effect, and the time at which the response subsequently fell below the threshold for the first time during the 8-hour post-dose period was defined as the loss of effect. Duration of effect was defined as time from onset of effect to time of loss of effect. For finger-tapping, each patient’s baseline was the average of the patient’s three pre-dose values. For UPDRS Part III score, it was the average of the two pre-dose values. Differences between ER CD-LD and IR CD-LD were tested for statistical significance by the ANCOVA model. Only the patients who reached each threshold were included in each analysis. No adjustments were made for multiple testing. For patients who were redosed during the 8 hours after their pre-study morning dose, any measurement obtained 10 minutes or more after taking a second dose was assigned a value of the pre-dose average of the treatment period and treated as if there was no benefit.

Pharmacokinetic analyses
Venous blood was collected pre-dose and for 8 hours post-dose, half-hourly for 6 hours and at hours 7 and 8. The resulting plasma samples were stored at $-70^\circ\text{C}$ or below until analysis for levodopa concentration by liquid chromatography coupled to tandem mass spectrometry, using a validated bioanalytical method with a minimum quantifiable levodopa concentration of 10.0 ng/mL. If patients were redosed, their pharmacokinetic data were censored after the redosing time point.

Results
Study participants
In total, 27 patients were randomized at six study sites, all in the US. Their baseline characteristics are summarized in Table 1. All patients completed the study.

Dosing
Each patient received individualized regimens of ER CD-LD and IR CD-LD. On day 1 of each treatment, the mean (SD) levodopa morning doses were 663.9 (185.2) mg for ER CD-LD and 164.8 (61.7) mg for IR CD-LD. The medians were 585.0 mg for ER CD-LD and 200.0 mg for IR CD-LD. In total, six patients on ER CD-LD and ten patients on IR CD-LD were redosed before the end of the 8-hour pharmacokinetic/pharmacodynamic assessment period. One ER CD-LD patient and two IR CD-LD patients were redosed before hour 4, three IR CD-LD patients were redosed between hours 4 and 6, and five patients in each group were redosed between hours 6 and 8.

Table 1 Patients’ baseline characteristics (safety population)

<table>
<thead>
<tr>
<th>Variables</th>
<th>Values</th>
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<tbody>
<tr>
<td>N</td>
<td>27</td>
</tr>
<tr>
<td>Age (years)</td>
<td>Mean (SD) 62.7 (8.6)</td>
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<tr>
<td></td>
<td>Median (range) 62 (48–81)</td>
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<tr>
<td>Sex, n (%)</td>
<td>Male 21 (78)</td>
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<tr>
<td></td>
<td>Female 6 (22)</td>
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<tr>
<td>Race, n (%)</td>
<td>Caucasian 24 (89)</td>
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<tr>
<td></td>
<td>Asian 3 (11)</td>
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<td>Weight (kg)</td>
<td>Mean (SD) 84.5 (14.8)</td>
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<td>Median (range) 85 (51–120)</td>
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<td>Age at PD onset (years)</td>
<td>Mean (SD) 52.3 (8.7)</td>
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<tr>
<td></td>
<td>Median (range) 50 (36–74)</td>
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<td>&quot;Off&quot; time (h/day)</td>
<td>Mean (SD) 5.9 (2.2)</td>
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<tr>
<td></td>
<td>Median (range) 5.7 (2.0–11.0)</td>
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<tr>
<td>H&amp;Y stage, n (%)</td>
<td>2 13 (48)</td>
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<tr>
<td></td>
<td>3 10 (37)</td>
</tr>
<tr>
<td></td>
<td>4 4 (15)</td>
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<tr>
<td>Levodopa IR dosage (mg/day)</td>
<td>Mean (SD) 815.7 (249.7)</td>
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<tr>
<td></td>
<td>Median (range) 800 (500–1,250)</td>
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<td>Levodopa IR dosing frequency (doses/day)</td>
<td>Mean (SD) 5.4 (1.6)</td>
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<td></td>
<td>Median (range) 5 (4–10)</td>
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<td>Finger-tapping score (taps/min)</td>
<td>&quot;On&quot; state Mean (SD) 170.4 (64.6)</td>
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<tr>
<td></td>
<td>Median (range) 163 (73–327)</td>
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<tr>
<td></td>
<td>&quot;Off&quot; state Mean (SD) 117.9 (44.1)</td>
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<tr>
<td></td>
<td>Median (range) 116 (44–229)</td>
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<tr>
<td>UPDRS Part III score</td>
<td>&quot;On&quot; state Mean (SD) 17.6 (8.5)</td>
</tr>
<tr>
<td></td>
<td>Median (range) 17 (1–32)</td>
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<tr>
<td></td>
<td>&quot;Off&quot; state Mean (SD) 35.9 (9.4)</td>
</tr>
<tr>
<td></td>
<td>Median (range) 36 (20–51)</td>
</tr>
</tbody>
</table>

Note: During "on" time. 
Abbreviations: IR, immediate release; PD, Parkinson’s disease; SD, standard deviation; UPDRS, Unified Parkinson’s Disease Rating Scale.
Time to “on” and duration of “on”
For patients rated as achieving an “on” state, the mean time to onset and the mean duration are displayed in Figure 1 (leftmost charts). The mean (SD) time to onset showed no significant difference between ER CD-LD and IR CD-LD (0.83 [0.52] vs 0.81 [0.45] hours; \( P=0.89 \)), but the mean duration was significantly longer for ER CD-LD than for IR CD-LD (5.56 [1.78] vs 2.69 [0.92] hours; \( P<0.0001 \)).

Time to onset of effect and duration of effect
For patients achieving each predefined threshold level of improvement in finger-tapping score or in UPDRS Part III score, the mean time to onset of effect and the mean duration of effect are displayed in Figure 1 (center and rightmost charts). For all improvement levels, the mean time to onset of effect showed no significant difference between ER CD-LD and IR CD-LD. However, the mean duration of effect was significantly longer for ER CD-LD than for IR CD-LD.

Pharmacokinetics
Mean plasma levodopa concentrations following single doses of ER CD-LD and IR CD-LD are presented in Figure 2. The initial concentrations increased rapidly after both treatments, but were sustained longer by ER CD-LD than by IR CD-LD. The fluctuation index is a measure of the magnitude of rise and fall of levodopa plasma concentrations relative to the average concentration and may be of particular interest for an ER formulation. The lower the fluctuation index, the more likely the \( C_{\text{max}} \) is blunted relative to the trough, thus improving the pharmacodynamic profile and minimizing \( C_{\text{max}} \)-related adverse effects. The fluctuation index measured

![Image of Figure 1](https://www.dovepress.com/)

**Figure 1** Mean time to onset (top graphs) and mean duration (bottom graphs) for an “on” state, finger-tapping score improvement, and UPDRS Part III score improvement after single doses of ER CD-LD vs IR CD-LD.

**Notes:** *\( P<0.05 \), **\( P<0.01 \), ***\( P<0.001 \) (ANCOVA). Each analysis includes only the patients who reached each outcome.

**Abbreviations:** ANCOVA, analysis of covariance; CD-LD, carbidopa-levodopa; ER, extended release; IR, immediate release; SE, standard error; UPDRS, Unified Parkinson’s Disease Rating Scale.
of mean durations of motor-function improvement, the thresholds that yielded the most similar set of durations were a 20% increase from baseline for finger-tapping and a 11-point decrease from baseline for UPDRS Part III score (Figure 1). In the study’s pharmacokinetic data, a levodopa plasma level ≥1,000 ng/mL provided a similar set of durations (Figure 2).

**Similarities among outcomes**

By observers’ assessments, the mean duration of an “on” state was ~5.6 hours following ER CD-LD dosing and ~2.7 hours following IR CD-LD dosing (Figure 1). By inspection of mean durations of motor-function improvement, the thresholds that yielded the most similar set of durations were a 20% increase from baseline for finger-tapping and a 11-point decrease from baseline for UPDRS Part III score (Figure 1). In the study’s pharmacokinetic data, a levodopa plasma level ≥1,000 ng/mL provided a similar set of durations (Figure 2).

**Discussion**

In this post hoc analysis of ER CD-LD and IR CD-LD in patients with advanced PD, the onset of clinical effect following a single dose was similar for both treatments, but the duration of effect was significantly longer for ER CD-LD than for IR CD-LD, as documented by investigator assessments of motor state, finger-tapping, and UPDRS Part III scores. Consistent with these clinical endpoints, plasma levodopa concentrations showed a similar time to reach a threshold level ≥1,000 ng/mL for both treatments, but were maintained for a longer duration by ER CD-LD.

Pharmacokinetic data in healthy subjects had indicated that a dose-adjusted basis (mg ER CD-LD:mg IR CD-LD), the ER CD-LD treatment resulted in a mean $C_{\text{max}}$ ratio of 0.34. Thus, an approximately threefold higher dose of ER CD-LD compared to IR CD-LD would result in similar $C_{\text{max}}$ values. Recognizing that there may be differences in the pharmacokinetics of ER CD-LD in healthy subjects and patients, and the importance of achieving an “on” state in both treatments to allow for pharmacodynamic comparisons, a conservative approach was adopted for the initial conversion, allowing physicians to subsequently titrate the ER CD-LD dose for the first 3 days as needed to optimize the therapeutic effect. Similarly, physicians could also adjust the IR CD-LD regimen for the first 3 days as needed. Subsequent pharmacokinetic data from patients in this study confirmed the dose conversion. Following a single IR CD-LD dose of 50–200 mg, the mean levodopa peak concentration was 2,331 ng/mL compared to 2,424 ng/mL following 122.5–490 mg ER CD-LD. In addition, multiple dose pharmacokinetic data indicated that the $C_{\text{max}}$ and average levodopa concentration following the IR 50–200 mg CD-LD were 3,057 ng/mL and 969 ng/mL, respectively. In comparison, following multiple doses of 490 mg ER CD-LD, the $C_{\text{max}}$ and average levodopa concentration were 3,227 ng/mL and 1,623 ng/mL, respectively. These data indicate that similar to observations in healthy subjects, a 2.45-fold higher levodopa dose with ER CD-LD compared to IR CD-LD resulted in very comparable (ie, within 4%–6%) mean peak levodopa concentrations following single and multiple doses. The higher average levodopa concentration with ER CD-LD is due to its pharmacokinetic profile with more sustained levodopa concentration and contributes to the longer duration of effect for this treatment.

Of the three efficacy measures utilized in the present analyses – observers’ “on”/“off” assessments, finger-tapping scores, and UPDRS Part III scores – the “on”/“off” assessments may come closest to being considered a “gold standard” for judging a patient’s clinical status. The observation of an “on” state within 1 hour after levodopa dosing is consistent with that reported by others. For IR CD-LD, Lewitt et al reported that the mean (SD) time to “on”, as assessed via patient diary, was 0.95 (0.55) hours. Similarly, Merims et al reported that the mean (SD) time to beginning of an “on” state after levodopa dosing was 46 (21) minutes. The original study did not use a stopwatch method to measure time to “on” and its duration. Investigator assessments of motor state were done every 30 minutes, and measurements of the time to “on” and the duration of “on” may be expected to be influenced by the frequency of assessment.

Of the other two efficacy measures, finger-tapping scores have not been extensively validated, and various derivations...
of minimal clinically relevant changes in UPDRS scores have not resolved the issue of what clearly constitutes a meaningful change in a given patient or population. 14,15,17-19 By observers’ assessments, the mean duration of an “on” state (~5.6 hours after ER CD-LD dosing compared with ~2.7 hours after IR CD-LD dosing) was matched most closely by a ≥20% improvement in finger-tapping, a ≥11-point improvement in UPDRS Part III score, and a levodopa plasma concentration ≥1,000 ng/mL, suggesting that these values may be the most clinically relevant thresholds for acute assessment of benefit, at least in this population of PD patients. Nonetheless, we note that none of the threshold values evaluated provided a perfect match to the “on”/”off” assessments. To some extent, this could be due to methodologic issues including the timing and frequency of the various evaluations and due to the fact that we only evaluated a few thresholds. It is also possible that there would always be some differences in duration of benefit as assessed by the various measures, no matter how carefully such a study were conducted. This could be because the transition from “on” to “off” and “off” to “on” is often gradual, so the designation of “on” or “off” during these transition periods is subjective. It is also possible that benefit occurs in the various signs and body parts at different times, such that finger-tapping could improve before, or after, walking and speech. These issues deserve further study. The concentration threshold of 1,000 ng/mL identified in the present study is comparable to the value of 900 ng/mL identified by Nelson et al23 as the median effective concentration typically needed to maintain control of Parkinsonian features.24

Limitations of the present analyses include the fact that the study was an open-label study and relatively small. In addition, although the doses administered were designed to have a similar $C_{\text{max}}$ according to pharmacokinetic information known at the time, results indicated that $C_{\text{max}}$ was higher in the ER CD-LD group than in the IR CD-LD group. Duration of benefit may depend on treatment dose, which was not tested, and it is possible that higher doses of IR CD-LD would yield a longer duration of benefit. However, it should be noted that the doses of IR CD-LD administered in the original study were the doses patients were typically taking prior to entering the study, and troublesome dyskinesia was not significantly increased with ER CD-LD compared to IR CD-LD. The identified thresholds for benefit are most likely to be applicable to patients with motor fluctuations of a severity similar to that required for original study entry.

The post hoc analyses presented here provide additional evidence that in a population of patients with motor fluctuations on levodopa, time to onset of benefit from ER CD-LD is similar to that for IR CD-LD, and duration of benefit is significantly longer. In PD patients such as those in the present study, for whom IR dosing provides benefit for only about 2.5 hours, ER dosing may provide benefit for about 5.6 hours. Nonetheless, clinicians should be aware that timing of administration may require adjustment based on the individual patient’s duration of benefit.

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Author contributions
RAH: design and execution of the clinical study, conduct of data analyses, writing of the first draft, and review and critique of the manuscript; AE: design and execution of the clinical study, conduct of data analyses, and review and critique of the manuscript; SK: design of the clinical study, conduct of statistical and data analyses, and review and critique of the manuscript; SG: design and execution of the clinical study, conduct of data analyses, and review and critique of the manuscript; NBM: design and execution of the clinical study, conduct of data analyses, and review and critique of the manuscript. All authors contributed toward data analysis and drafting and revising the paper, and agree to be accountable for all aspects of the work.

Disclosure
Robert A Hauser has received honoraria or payments for consulting, advisory services, or speaking services over the past 12 months from AbbVie, Allergan, AstraZeneca, Biote Therapeutics, Ceregene, Chelsea Therapeutics, Cleveland Clinic, Eli Lilly, GE Healthcare, Impax Laboratories, Inc., Neurocrine, Indus, Ipsen Biopharmaceuticals, Lundbeck, Merck/MSD, Noven Pharmaceuticals, Pfizer, Straken Pharmaceuticals, Targacept, Teva Pharmaceutical Industries, Ltd., Teva Neuroscience, Upsher-Smith Laboratories, UCB, UCB Pharma SA, University of Houston, US WorldMeds, XenoPort, and Zambon Company SpA. Dr Hauser’s institution has received research support over the past 12 months from Abbott Laboratories, Addex Therapeutics, Allergan, AstraZeneca, Biote Therapeutics, Chelsea Therapeutics, Civitas, GE Healthcare, Impax Laboratories, Inc., Ipsen Biopharmaceuticals, Merck/MSD, Merz, the Michael J. Fox Foundation for Parkinson’s Research, NINDS, the Parkinson
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References