Duopa® Medication and Device Information

Patient Name: _____

Date of Birth: _____

MRN: _____

I use carbidopa/levodopa enteral suspension (Duopa®)? ☐ Yes ☐ No
If yes, please complete the information below.

Surgery & Device Information

Surgery and Device Details: _____

Movement Disorder Neurologist: _____

Movement Disorder Neurologist Phone: _____

Surgeon: _____

Date of Implantation: _____

Daily Dosage Schedule

Remove the refrigerated cassette 20 minutes before connecting.

Start Time: _____

Morning Dose: _____

Stop Time: _____

Continuous Dose Rate: _____

Extra Dose: _____

Time(s): _____

Titrated Dose(s): _____

Time(s): _____

Tubing Information

Type of Tubing: ☐ AbbVie ☐ ENFit

Date of Last Tube Replacement: _____

Due to a fixed, interior bumper plate, AbbVie tubing should be removed endoscopically if replacement is indicated.

Duopa® Support Contacts

DuoConnect RN Name: _____

Phone: _____

Device Representative Name: _____

Device Representative Phone: _____

PEG-J Tube Information

My PEG-J tube for this device was implanted by: ☐ Gastroenterologist ☐ Interventional Radiologist ☐ General Surgeon

Name: _____

Phone: _____

(For tubing system complications)

Important Duopa® Safety Information

- **Duopa® cassettes must be refrigerated** at 36°F–46°F and should not be frozen. The cassette should be brought to room temperature before use.
- Remove a cassette from the refrigerator **20 minutes before use.** An opened cassette should not be reused.
- **Do not use the cassette if its contents appear yellowed.**
- Avoid sudden discontinuation or rapid reduction of the Duopa® dose.
- If the pump is disconnected for l**ess than 2 hours, do not supplement with oral medications.**
- If imaging is required, including ultrasound, **ECG, CT scan, X-ray, or MRI,** turn off the pump, detach it from the patient, and remove the pump from the procedure room.