SYN-ONE TEST™ REQUISITION FORM

CND Internal Use Only

Case #: ____________________

Date Received: ____________________

☐ Syn-One Test™ for Synucleinopathy

Syn-One is an anatomical pathology test to detect, visualize, and quantify the presence of abnormal, phosphorylated alpha-synuclein in cutaneous nerve fibers to support a diagnosis of a synucleinopathy: Parkinson's disease (PD), dementia with Lewy bodies (DLB), multiple system atrophy (MSA), pure autonomic failure (PAF), or REM sleep behavior disorder (RBD). Syn-One is also able to confirm and quantify the density of intra-epidermal nerve fibers to support a diagnosis of small fiber neuropathy and includes modified Congo red staining to identify amyloid proteins to support a diagnosis of amyloidosis.

Complete all the information below. **Sign and fax with the documents listed to (480) 569-2910:**

☐ Copy of patient's insurance card (front and back)
☐ Copy of patient's drivers license or state ID (front and back)
☐ Relevant medical records
☐ Pathology report
☐ DaTscan
☐ Other

PATIENT INFORMATION

First Name: ____________________
Middle Initial: ____
Last Name / Surname: ____________________

Date of Birth (Month/Day/Year): ____________________
☐ Male
☐ Female

ICD-10
☐ G60.3 Idiopathic Neuropathy
☐ G20 Parkinsonism
☐ Other: ____________________

PRACTICE INFORMATION

Ordering Physician/Clinician: ____________________
Physician NPI (US) or Clinician ID Number (International): ____________________

Practice Name: ____________________

Street Address: ____________________
City: ____________________
State: ____________________

Zip or Postal Code: ____________________
Country (International Only): ____________________

Phone Number: ____________________
Fax Number: ____________________
Email Address: ____________________

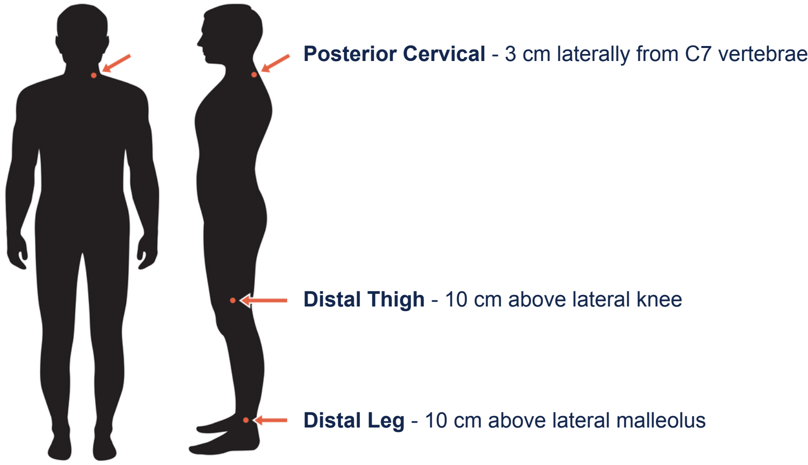
SYNUCLEINOPATHY CLINICAL INFORMATION

Family History of Parkinson's Disease	☐ Yes	☐ No	☐ Unknown
Response to L-Dopa	☐ Yes	☐ No	☐ Unknown
Response to Dopaminergic Agonists	☐ Yes	☐ No	☐ Unknown
Dementia	☐ Yes	☐ No	☐ Unknown
Irregular Autonomic Function	☐ Yes	☐ No	☐ Unknown
Resting Tremors	☐ Yes	☐ No	☐ Unknown
Orthostatic Hypotension	☐ Yes	☐ No	☐ Unknown
REM Sleep Behavioral Disorder	☐ Yes	☐ No	☐ Unknown
Loss of Smell (Anosmia)	☐ Yes	☐ No	☐ Unknown
Periods of Confusion/Hallucinations	☐ Yes	☐ No	☐ Unknown
Constipation	☐ Yes	☐ No	☐ Unknown
Bladder Dysfunction	☐ Yes	☐ No	☐ Unknown
DaTScan Result	☐ Normal	☐ Abnormal	☐ Unknown

PLEASE COMPLETE AND SIGN THE OTHER SIDE OF THIS DOCUMENT

CND Life Sciences™

3MM SKIN BIOPSY SPECIMEN INFORMATION



Guidelines:

- When performing the biopsy, the metal head of the punch tool should be used with gentle pressure and rotation until **fully into the skin**
- **Gently** handle the biopsy with the forceps
- Make sure the biopsy is **free floating** in the vial

Clinician Performing Biopsy

Physician NPI (US) or Clinician ID Number (International)

Date of Specimen (Month/Day/Year)

Time of Specimen ☐ AM ☐ PM

	Side (Choose One)		Location (Choose One)			
Specimen 1	☐ Right	☐ Left	☐ Posterior Cervical	☐ Distal Thigh	☐ Distal Leg	☐ Other: ______
Specimen 2	☐ Right	☐ Left	☐ Posterior Cervical	☐ Distal Thigh	☐ Distal Leg	☐ Other: ______
Specimen 3	☐ Right	☐ Left	☐ Posterior Cervical	☐ Distal Thigh	☐ Distal Leg	☐ Other: ______

The undersigned certifies that he/she is licensed to order the test(s) selected and that such test(s) are medically necessary for the care/treatment of this patient

Authorized Signature

Date

SHIPPING INSTRUCTIONS

1. Label each vial with patient name, date, time, and body site.

2. Place this Requisition Form with supporting documentation and specimens into the insulated mailer and seal.

3. Place the sealed mailer into the FedEx UN 3373 Pak and seal.

4. Place the return label onto the return package in the designated area.

5. Contact **FedEx at 1 (800) 463-3339** to arrange same-day pickup with Next Day Delivery

CND Life Sciences | 5070 N 40th St., Suite 220 Phoenix, AZ 85018-2135 | cndlifesciences.com | Phone: 480.569.2900 | Fax: 480.569.2910 | CLIA# 03D2151444

Immunohistochemistry tests were developed and their performance characteristics were determined by CND Life Sciences, Phoenix, AZ. They have not been cleared or approved by the U.S. Food and Drug Administration. The FDA has determined that such clearance or approval is not necessary. These tests are used for clinical purposes. They should not be regarded as investigational or for research. CND Life Sciences is certified under the Clinical Laboratory Improvement Amendments of 1988 (CLIA-88) as qualified to perform high complexity clinical laboratory testing. Unless otherwise specified, all histochemical and immunohistochemical controls are adequate.