Patient ID __ __ - __ __ __ __ - __ __ __ ID

NAC Trial Criteria

DOEM/DOED/DOEY Date of Evaluation __ __- __ __ -__ __
mm-dd-yy

SECTION I: INCLUSION CRITERIA

1.	Patient enrolled in PALF registry? EREG	☐ Yes	☐ No
2.	Patient/guardian provided informed consent for participation in the NAC trial? CONN If No, reason: ☐ Refused ☐ Unobtained ☐ Other, specify CONNS _____ CONNR	☐ Yes	☐ No

SECTION II: EXCLUSION CRITERIA

3.	Acute Acetaminophen toxicity? AACT	☐ Yes	☐ No
4.	Patient already on N-acetylcysteine? ANAC	☐ Yes	☐ No
5.	Patient is pregnant? PREG	☐ Yes	☐ No
6.	Patient has a known malignancy? MALG	☐ Yes	☐ No
7.	Patient is on a liver support device? LVSD	☐ Yes	☐ No
8.	Sepsis? SEP	☐ Yes	☐ No
9.	Signs of cerebral herniation? CHERN	☐ Yes	☐ No
10.	Intractable hypotension: systolic BP < 85 mm Hg or requires ionotropic drugs other than renal dosing of dopamine? HYPO	☐ Yes	☐ No
11.	Does the patient have a history or other evidence of severe illness or any other condition that would make the patient, in the opinion of the investigator, unsuitable for the study? If Yes, specify _____HXEL HXELS	☐ Yes	☐ No

If the responses to all inclusion criteria are YES and all exclusion criteria are NO, the patient is eligible to participate in the NAC Trial.