North American Consortium for Histiocytosis

# HLHRUXO STUDY INFORMATION SHEET

## PATIENTS ELIGIBLE FOR TREATMENT

All patients must have active disease based on presence of >5 of 8 HLH-2004 diagnostic criteria for those with secondary HLH or HLH of unknown etiology at diagnosis, or >4 of 8 HLH-2004 diagnostic criteria for those with primary HLH. Patients must be and be ≥6 weeks to ≤ 22 years of age.

To be eligible for for the ***Frontline Arm***, patients must not have received any other HLH-directed therapy except steroids or anakinra at any dose.

To be eligible for the ***Salvage Arm***, HLH must have failed to adequately respond or reactivated after previous standard of care therapy (etoposide, ATG, dexamethasone). Patients must not have received JAK inhibitors or other anti-cyokine therapies in the 3 months prior to enrolling.

For more information about inclusion/ exclusion criteria, please go to ClinicalTrials.gov (NCT04551131).

## HOW TO ENROLL PATIENTS

Please contact the HLHRUXO team at: HLHRUXO@stjude.org

Study website: nacho-consortium.org

## PRINCIPAL INVESTIGATORS

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## COLLABORATING SITES

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