



**MEMO: PIDTC REQUIREMENTS FOR PERSONNEL
AUTHORIZED TO OBTAIN INFORMED CONSENT**

DATE: 25 FEBRUARY 2026
TO: ALL PIDTC SITE PRINCIPAL INVESTIGATORS AND STUDY TEAMS
FROM: PIDTC PRINCIPAL INVESTIGATORS
RE: REQUIREMENTS FOR PERSONNEL AUTHORIZED TO OBTAIN INFORMED
CONSENT

Dear Colleagues,

After review of prior consortium documentation, historical site practices, and applicable regulatory standards, the Principal Investigators of the Pediatric Immune Deficiency Treatment Consortium (PIDTC)—Drs. Chris Dvorak, Elie Haddad, Troy Torgerson, and Rebecca Marsh—confirm the following:

There is no requirement that informed consent for PIDTC studies be obtained exclusively by the physician Site Principal Investigator. In accordance with standard regulatory expectations, informed consent may be obtained by any study team member who:

1. Is appropriately qualified and trained to conduct the consent process,
2. Is listed on the study delegation of authority log for obtaining consent, and
3. Conducts the discussion in a manner consistent with regulatory, ethical, and protocol-specific standards.

The physician Site PI should be available for questions, clarification, or further discussion if needed.

This clarification is intended to support operational efficiency while maintaining full compliance with ethical, regulatory, and institutional requirements for informed consent.

Sincerely,

Drs. Chris Dvorak, Elie Haddad, Rebecca Marsh, and Troy Torgerson on behalf of the PIDTC.

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