



Cell and Gene Therapy Services for Pharmacy Standards Committee

CHAIR: Dr Beth Shaz, MD

PURPOSE: To develop, maintain and update standards for the handling of approved cell and gene therapy products within pharmacy.

CHARGES:

1. Review existing requirements from the Standards for Cellular Therapy Services and identify standards applicable to activities within pharmacy. Where necessary craft new standards surrounding the receipt, handling, storage, or dispensing of cell and gene therapy products through pharmacy.
2. In coordination with AABB staff, develop guidance for the Standards for Cell and Gene Therapy Services for Pharmacy and create records of rationales for changes to existing requirements, as applicable.
3. Review and respond to requests for clarification and variances to the Standards as needed.
4. Develop interim/emergent standards as needed for submission to the AABB Board of Directors.
5. Monitor the development of new practices and technologies with potential application to cell and gene therapies and develop or modify standards when appropriate.

Current Personnel as of April 30, 2025

Name	Roles
<u>Chair</u>	
Dr Beth Shaz, MD	
<u>Consultant</u>	
Dr. Linda S Barnes, CABP(H),DrPH,MHA	
<u>Liaison</u>	
Corinne Goldberg, MD	CTAC Liaison
Ms. Katherine Stewart Brown	CTSC Liaison
<u>Member</u>	
Ms. Kimberly Tedesco	
Kim McConnell	
Mrs. Brenda Alder, CABP ,MS,MT(ASCP)SBB	
Mrs. Mary Grable McLeod, CABP(H),MT(ASCP)SBB	
Jodi Sibell	
Maribeth Bettarelli	
Joe DePinto, MBA	
Marissa Szymala, MBA	
Eric Balmir	
Jill Blind	
Phil Wilson	
Chiara Cerati	
<u>Representative</u>	
Jessica Chery	FDA (OGT) Rep.



Name	Roles
Karin Knudson	FDA (OCTHT)

Staff Liaison

Mr Christopher Bocquet	
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Time Commitment: To learn about the time commitments for this committee, please contact the staff liaison listed in the roster above.