



## Regulatory Affairs Committee

**CHAIR:** Scott Webber

**PURPOSE:** Serve as a resource to AABB on Transfusion Medicine and Cellular Therapy related regulatory issues.

- CHARGES:**
1. Provide input for the development of position statements for the Food and Drug Administration (FDA) and Centers for Medicare and Medicaid (CMS).
  2. Review FDA/CMS documents and notices and provide input on the operational and technical implications.
  3. Develop, revise and review documents and toolkits for alignment with FDA regulations and recommendations.
  4. Review AABB Standards to identify areas where the addition of regulatory references would be relevant and useful.
  5. Assist with developing content for the Ask the FDA and CMS/CLIA session at Annual Meeting.

### Current Personnel as of May 20, 2024

Name	Roles
<b><u>Chair</u></b>	
Scott Webber	
<b><u>Consultant</u></b>	
Anne Chenoweth	
<b><u>Junior Committee Member</u></b>	
Richard Bewley	
Jennifer Vrieze	
<b><u>Liaison</u></b>	
Nina Sen, SBB(ASCP)	BB/TS AC Liaison
Dr. Aasawari Bapat, CABP(H),MD,PHD	Quality, Regulatory, and Management Subsection Liaison
<b><u>Member</u></b>	
Janice Habel	
Roxanne Tata, MT(ASCP)SBB	
Jean Reece	
Valery Freeman-Allen	
Jessica Lantz	
Mrs Susan Sullivan	
<b><u>Staff Liaison</u></b>	
Karen Palmer	

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