



Legal and Regulatory Affairs Committee: Australia and New Zealand

ESP Leadership Development Program – Role Description

Committee's scope and research interests

The ANZ LRA considers legal and regulatory matters in relation to cell and gene therapy from an Australian and New Zealand or international perspective. It contributes to public consultations, the ISCT Watch Dog report and the members share their experience and knowledge in implementing related clinical trials and GMP.

Projects and responsibilities for the ESP member

Co-authoring the Watch Dog report, contributing to submissions to relevant public consultations and assisting with a webinar being developed by the LRA on a relevant topic.

ESP member's opportunities within the Committee

Contributing to the above but also the opportunity to speak with members in relation to their sphere of interest. We are a small committee (approximately 10 members) with diverse roles covering cord blood banking, cell therapy, industry, public hospital clinical services, commercial cell therapy manufacturing, legal affairs, gene technology and administration.

[Committee Webpage](#)