Europe Legal and Regulatory Affairs Committee
ESP Leadership Development Program

Committee's scope and research interests

The EU LRA strives to:

- Provide input on key regulatory issues within Europe that impact cell and gene therapy researchers and related industries through building and promoting regulatory partnerships with CAT-EMA, the European Commission, and other relevant authorities (i.e. stakeholder workshops, regulatory consultations)
- Provide education opportunities to our members about regulatory issues within Europe that impact cell and gene therapy researchers and related industries (i.e. webinars, publications, workshops)
- Drive committee and society membership
- Build and promote the visibility of ISCT's regulatory partnerships

ESP member's opportunities within the Committee

Membership on the EU LRA provides the unique opportunity to access, network and learn from numerous regulatory leaders from industry and academia spanning Europe. Participation in the meetings and various committee-led initiatives allows for collaboration with these leaders to impact regulations throughout the region.

Projects and responsibilities for the ESP member

- Serve as the key liaison for monitoring the current regulatory landscape, specifically CAT-EMA, European Commission, ICH, WHO, etc.
- Author at least one WatchDog report per year for the ISCT community while supporting educational initiatives such as webinars.
- Support the consultation efforts, such as providing and consolidating comments for draft guidance documents.
- Support the development of at least one workshop/roundtable at the ISCT Annual or Regional Meeting

EU LRA Committee Webpage