Europe Legal and Regulatory Affairs Committee
ESP Leadership Development Program

Committee's scope and research interests

The European Legal and Regulatory Affairs Committee (EU LRA) of the ISCT serves the membership and others working in cellular therapy by providing information regarding European regulations, standards, and guidance documents and presenting the regulatory concerns of the membership back to the regulatory agencies.

Projects and responsibilities for the ESP member

- The EU LRA is recruiting an ESP to be the key liaison for monitoring the current regulatory landscape including new consultation/guidelines from the EMA, ICH, CAT and WHO.
- This ESP will author the WatchDog report for the ISCT community while supporting educational initiatives such as webinars.
- The ESP will support the consultation efforts, such as providing and consolidating comments for draft guidance documents.

ESP member's opportunities within the Committee

Contributing to the committee's deliverables along with the opportunity to work with experienced members in relation to their sphere of interest.

EU LRA Committee Webpage