Fact Sheet Regenerative Medicine Consortium (RMC)

Mission	• To accelerate the development and regulatory approval of stem cells and regenerative medicine therapies
Strategic Objectives	• Provide a forum for the discussion of areas of mutual interest to the FDA and the RMC in order to shape and advance clear regulatory pathways for novel technologies • Serve as a technical resource which may be consulted during development of guidelines, standards and best practices • Accelerate stem cell research through the sharing of best practices and resources • Serve as a clearing house for guidances, white papers, standard operating procedures, protocols and other resources
Membership	• Membership reflects leading companies, academic and funding institutions that are engaged in and support stem cell research
Meetings and Agenda	• Periodic roundtable meetings with the FDA; the California Institute for Regenerative Medicine (CIRM) will host and organize the meetings. All meetings are expected to occur at or near FDA. CIRM (with input from members) and FDA will agree on the agenda topics. • Periodic webinars on topics of general interest open to public at large
Fees and Expenses	• No fees requested; members are responsible for their own travel costs.
Communication and Outreach	• See our website at http://www.cirm.ca.gov/RegenerativeMedicineConsortium
Logistics	If you have questions concerning the RMC and or any Webinar or Roundtable content, please contact **Elona Baum, General Counsel** **T: (415) 396-9275** **E: ebaum@cirm.ca.gov** or **Cynthia Schaffer, CIRM Administrator** **T: (415) 396-9241** **E: cschaffer@cirm.ca.gov**