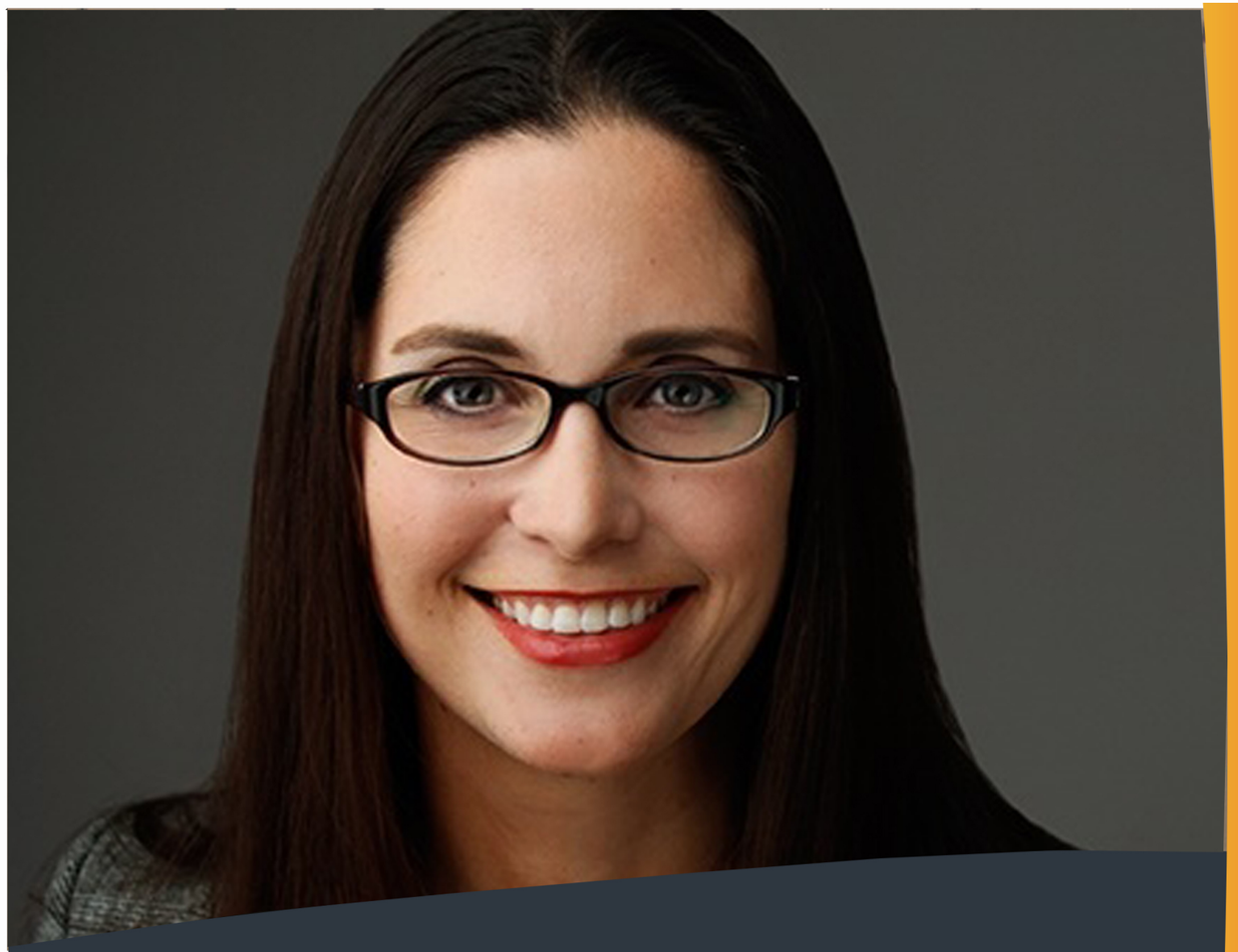
Stemmed Implant™
TECHNOLOGY INC.
Implants that Nature Designed

Ms. Komiyma is a former Food and Drug Administration reviewer who has served as an expert in regulatory submissions. She will develop and manage the required regulatory process - from clinical trial administration to communication with government regulatory agencies, nationally and internationally.

Her involvement will expedite the approval process by avoiding duplication of effort and ensuring procedures undertaken for one regulatory authority will be suitable to meet the standards of other jurisdictions.

Ms Komiyma is the Principal of AcKnowledge Regulatory Strategies, a consultancy for biomedical manufacturer who work with implantable and other patient-contacting medical devices.



Dr. Allison Komiyma