Speakers Profile: ISCR Workshop (Clinical Trial Transparency Track, 14 Feb 2019)

	**Sonali Parmar** Sonali Parmar, M.Pharm. in Pharmaceutics, presently Senior Manager in Kinapse in Clinical Trial Disclosures, has more than 15 years of experience of working in pharmaceutical industry across different domains. She is WIPO certified scientist and holds a certificate in Regulatory Affairs and CDM from IOCB, Bangaluru
	**Sonam Vijan** Sonam Vijan, Ph.D. in Nanoscience, presently Team Lead in TCS, has experience in Document Publishing and Regulatory Medical Writing. She has 2 publications in peer reviewed journals and has won several awards for presentation in many international conferences
	**Deepa Trivedi** Deepa Trivedi, Ph.D. in Biochemistry and PDF in Pharmacology, presently Delivery Lead at TCS, has more than 15 years of experience in pharmaceutical and IT outsourcing industries in medical writing for pharma and medical devices. She has been closely involved with data transparency initiatives at TCS and preparation of CTR summaries when Lilly became the first company to voluntarily post CT study results in a registry in 2004
	**Neha Tickoo** Neha Tickoo, M.Pharm from BITS, Pilani, presently Manager in Kinapse, has more than 6.5 years of experience in Clinical Trial Disclosures. Neha provides advisory support and guidance to pharma companies for conducting User Acceptance Testing for client specific platforms for disclosure
	**Hitendra Pandey** Hitendra Pandey, M.Sc. in Total Quality Management, M.Sc.& B.Sc. Hons in Statistics from BHU, presently Subject Matter Expert (SME) with TCS, has more than 13 years of experience as a Statistician in CT domain across all phases of CTs, with extensive experience pharmacokinetic (PK) Phase I studies
	**Ajay Yeola** Ajay Yeola, M.Sc. in Statistics, has over 17 years of industry experience in Clinical Research, Analytics etc. Ajay is presently a Subject Matter Expert on Data Transparency and Disclosures initiatives at TCS

	**Shalini Dwivedi** Shalini Dwivedi, M.Pharm from Jamia Hamdard, presently Director in Kinapse, has about 14 years of professional experience in academic research and regulatory medical and publication writing. Presently she manages clinical trial disclosures including EMA Policy 0070 and 0043 projects at Kinapse