

Clinical Trial Agreement Checklist

Sl. No	Description	Yes	No
1.	Protocol Number and Title		
2.	Effective date		
3.	Parties Involved - (Sponsor / CRO, Principal Investigator, Institution and or SMO) Bipartite		
	Tripartite		
	Quadra parted		
4.	Agreed terms - Definition, Conduct of the study, Responsibility of the company, Principal investigator, Institution		
5.	Study drug and Materials		
6.	Study and Protocol		
7.	The Study Schedules		
8.	Monitoring and audit by the company		
9.	Inspection by the regulatory authorities		
10.	Payment Details- Budget and Payment scheduled, Payment of cost outside budget and payment schedule, Payment terms, payment recipient and address, Reimbursement, Payment for screen failure, payment for study coordinator.		
11.	Obligations of the institution and Principal Investigator - EC Approval, Performance of the study, Key personnel, sponsor Visit, Supplies		
12.	Study Records , reports and Data - Study records , Case report form, Annual reports, Final Reports , (In case of PI is no longer associated with the institute, Institute head or authorized designee will be responsible for maintenance and retention of study records) , Reporting of SAE(Sponsor, EC,DCGI and head of institution), 14th day PI analysis Report (Sponsor, EC,DCGI and head of institution).		
13.	Confidentiality		
14.	Publications		
15.	Ownership of materials, data, inventions and discoveries.		
16.	Representations, warranties and covenant.- Of the PI, Of the Sponsor, No other Representations or warranties, Of the Institutions		

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17.	Governing Law -This agreement and any dispute or claim out of or in connections with it or its subject matter (including non- contractual disputes or claims) shall be governed by and constructed in accordance with the laws of India without regard to the conflict of law principles thereof. The parties irrevocably agree that the courts of India shall have exclusive jurisdiction to settle any dispute or claim that arises out of or in connection with this agreement or its subject matter (including non-contractual disputes or claims).		
18.	Indemnification - Sponsor Indemnification, Institution Indemnification, Notification, Claims, Representation, subject injury.		
19.	Insurance - Sponsor insurance, Institution Insurance		
20.	Compliance, Transparency, Anti - bribery, Anti- corruption and Conflict of Interest.		
21.	Term and Termination		
22.	Miscellaneous		
23.	Agreed by the parties - Sponsor/ CRO, PI, Institution, SMO(if involved)		
24.	Witness details		
25.	Payee Details of the Hospital: Head of the Institution: Cheque in the Name of Registrar-KAHER for clinical trial -Institutional overhead charges: 25%		
26.	Payee Details of the Ethics Committee of KAHER Cheque: Registrar, KAHER, Belagavi-10		
27.	Research Pharmacy funds to be paid in the name of Registrar KLE University For the Non- Global study Global study		
Name and sign of the reviewer			