Technical Specification of Operating Microscope

S.N.	Purchaser's Specifications		
	Portable Operating	Microscope (Ophthalmic)	
	Manufacturer	Should be Europe, USA and Australia	
	Country of Origin	Europe, USA and Australia	
1	Technical Specific	AND ADDRESS OF THE PARTY OF THE	
1.1	Microscope should have4.2x to 25x continuous zoom.		
1.2			
AND SHALL	Should have binacular streoscopic convergent angle 10 degree for viewing comfort.		
1.3	Should have focus range of +-25 mm		
1.4	Binocular tube: swivel tube 45°		
1.5	Objective Lens: F= max 200mm		
1.6	Should have swivel tube 45°, Binocular tube		
1.7	Should havemax. 200mm from Aux Objective Lens for working/focal distance		
1.8	Should have +-5D refractive error correction anisometropai.		
1.9	Should have anti fungus pellets (Three year Protection)		
1.10	Should have inclined binocular tube Inclination 45 Degree, high field number 22 for wide field of view. High eye relief 24.6mm		
1.11	Should have UV and Blue Light Hazard Protection, (Filtered to 435nm) for Patient Safety.		
1.12			
1.13	0.10		
1.14	- Microccope head for easy movement for doctor during ourgon		
1.15	Tablemounting left or right side of table		
1.16	Clamp opening 120mm, throat 160mm		
1.17	Weight should be not more then 40 Kg		
2	A securior spares and consumables		
2.1	Must provide dust cover, operational manual, Aluminium Transit Case		
2.2			
2.3 All standard accessories, consu		ssories, consumables and parts required to operate the equipment	
	offer. Bidders mus	ard tools and cleaning and distriction with the state of	

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S.N.	Purchaser's Specifications		
	not specified above).		
3	Operating Environment		
3.1	The product offered shall be designed to be stored and to operate normally under the		
	conditions of the purchaser's country. The conditions include Power Supply, Climate,		
	Temperature, and Humidity.		
3.2	Power supply: AC 100-240VAC; Frequency: 50-60Hz fitted with appropriate plug, 2.0A Max @100VA and The power cable must be at least 3 metre in length.		
4	Standards and Safety Requirements		
4.1	Must submit ISO certificate and AC 20016 for Medical Devices		
4.2	CE (93/42 EEC Directives) or USFDA approved product certificate.		
4.3	Electrical safety conforms to standards for electrical safety IEC 60601-1 General		
	requirement for Electrical safety of Medical Equipment.		
5	User Training User Training		
5.1	Must provide user training (how to use the equipment).		
5.2	Must provide On-site Service training (how to maintain the equipment).		
6	Warranty		
6.1	Comprehensive warranty for 2 years after acceptance.		
7	Maintenance Service During Warranty Period		
7.1	During warranty period supplier must ensure corrective/breakdown maintenance whenever required.		
8	Installation and Commissioning		
8.1	The bidder must arrange for the equipment to be installed and commissioned by certified or		
1	qualified personnel; any prerequisites for installation to be communicated to the purchaser		
1976	in advance, in detail.		
9	Documentation		
9.1	User (Operating) manual in English.		
9.2	in English.		
9.3	List of important spare parts and accessories with their part numbers and decessories		
9.4	Certificate of calibration and inspection from factory.		

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