

**Bid Evaluation For Drugs/ Medicines (Injectables, Inhalations & IV Fluids) Financial year 2017-18
(Sole Agents of Not Prequalified Items)**

BID EVALUATION CRITERIA

Tender Sr. No. 91

Name of Item . Inj. metoprolol 5mg/ 5ml ,amp. Of 5ml, packed in carton with leaflet.

Name of Bidder	Name of firm/ Manufacturer	Brand Name	Compulsory Parameters										Marking Criteria										Obt. Marks	Approval of Sample by TSC/ End User (A/R)	Technical Eligibility of Product	Remarks			
			Prequalification, (Preq.) / Sole Agency Agreement (SAA) / Drug Sale License ("DSL") / Valid Drug Registration Certificate of the Product (DRC) / Sample Specifications (SS) / Undertaking, Regarding non declarations of Spurious Batch by DTL's of Punjab/ any other competent lab. within last three years (Und.T of DTL).	Preq. Status by the Govt. (Y/N)	S.A.A. (Y/N)	DSL (Y/N)	DRC (Y/N)	SS (Y/N)	Und. T. of DTL (Y/N)	cGMP (Y/N)	Bidder and Manufacturer Relationship (20)				Local Market Business (How many years the quoted product is being marketed in Pakistan) (25)					Compliance of Quality Standards (20)		International Testing (Reports From WHO ACC. Int. Labs. Performed on the product by any procuring agency) (15)					Export of Quoted Product (20)		
Novartis Pharma Pak.	Cibabergy Switzerland	Lopressor 5mg/ 5ml	NA	Y	Y	AFR	No Sample	Y	Y	Less Than 01 Year (0)	Upto 02 Years (05)	Upto 05 Years (10)	More Than 05 Years (20)	Less Than 01 Year (0)	Upto 02 Years (05)	Upto 05 Years (10)	Upto 09 Years (20)	More than 09 Years (25)	FDA/ WHO (20)	Others (10)	1-2 Labs (07)	3-4 Labs (15)	Developed countries (3-5=10 Above 05= 20)	Other Countries (01 Mark for Each Country)	0	62	Subject to the provision of sample	Subject to the provision of sample	

Total Marks = 100
Qualifying Marks = 60%

(Handwritten signatures and initials in blue ink)

**Bid Evaluation For Drugs/ Medicines (Injectables, Inhalations & IV Fluids) Financial year 2017-18
(Sole Agents of Not Prequalified Items)**

BID EVALUATION CRITERIA

Tender Sr. No. 102

Name of Item . Inj. Fondaparinux Sodium 2.5mg/0.5ml, Prefilled syringe, packed in carton with leaflet.

Name of Bidder	Name of firm/ Manufacturer	Brand Name	Compulsory Parameters							Marking Criteria										Obt. Marks	Approval of Sample by TSC/ End User (A/R)	Technical Eligibility of Product	Remarks						
			Prequalification, (Preq.) / Sole Agency Agreement (SAA) / Drug Sale License ("DSL") / Valid Drug Registration Certificate of the Product (DRC) / Sample Specifications (SS) / Undertaking, Regarding non declaration of Spurious Batch by DTL's of Punjab/ any other competent lab. within last three years (Und.T of DTL).	Preq. Status by the Govt. (Y/N)	S.A.A. (Y/N)	DSL (Y/N)	DRC (Y/N)	SS (Y/N)	Und. T. of DTL (Y/N)	cGMP (Y/N)	Bidder and Manufacturer Relationship (20)				Local Market Business (How many years the quoted product is being marketed in Pakistan) (25)				Compliance of Quality Standards (20)					International Testing (Reports From WHO ACC. Int. Labs. Performed on the product by any procuring agency) (15)		Export of Quoted Product (20)			
GSK Pak.	Glaxo Wellcome Production France (Aspen Notre Dame De Bonderville France)	Arixtra 2.5mg/ 0.5ml	NA	N	N	Y Not Valid	Y	Y	Y	Y	Less Than 01 Year (0)				Less Than 01 Year (0)				FDA/ WHO (20)		1-2 Labs (07)		0	0	0	45	A	Non Responsive, due to short documents.	Compulsory parameter not fulfilled
											Upto 02 Years (05)				Upto 02 Years (05)				Others (10)		3-4 Labs (15)								
											Upto 05 Years (10)				Upto 05 Years (10)														
											More Than 05 Years (20)				More than 09 Years (25)														

Total Marks = 100
Qualifying Marks = 60%

(Handwritten signatures and initials in blue ink)