BMSIC/40020/11-2017/1493 Date:- 02/8/17

BMSICL
(A Govt. of Bihar undertaking)

निविदा से संबंधित तकनीकी मूल्यांकन समिति की बैठक की कार्यवाही

निविदा संख्याः BMSIC/DRUGS/17-05
दिनांक 31-07-2017
समय 15:30 बजे अपराह्न

उपस्थिति–पंजी के अनुसार

बिहार चिकित्सा सेवाएँ एवं आधारभूत संरचना निगम लिमिटेड द्वारा Snake Venom Anti Serum औषधि का दर अनुबन्ध करने हेतु दिनांक 23/06/2017 को निविदा संख्या BMSIC/DRUGS/17-05 का प्रकाशन विभिन्न समाचार पत्रों में किया गया। निविदा सूचना में उल्लेखित कार्यक्रम के अनुसार ई–निविदा के माध्यम से निविदा जमा करने की अंतिम तिथि 26/07/2017 तक कुल 2 निविदादाताओं की निविदाएँ ई–निविदा के माध्यम से प्राप्त हुई। सभी प्राप्त तकनीकी निविदाओं का मूल्यांकन प्रपत्र तकनीकी मूल्यांकन समिति की उपसमिति द्वारा सत्यापित कराया गया। उपसमिति द्वारा सत्यापित मूल्यांकन प्रपत्र के अनुसार कतिपय बिन्दुओं पर सबंधित निविदादाता से स्पष्टीकरण की आवश्यकता के मद्धेनजर उक्त मूल्यांकन प्रपत्रों को निगम के वेबसाईट पर एक सप्ताह के लिए प्रकाशित कर दावा/आपत्ति आमंत्रिंत करने की अनुशंसा करने का सर्व सम्मति से निर्णय लिया गया । अन्त में धन्यवाद ज्ञापन के साथ बैठक की कारर्वाई समाप्त की गई।

31/07/2017
(रवीन्द्र कुमार सिन्हा)
राज्य औषधि नियंत्रक
बिहार

31.7.17
(सुधीर कुमार)
महाप्रबंधक (अधिप्राप्ति)
बी.एम.एस.आई.सी.एल
बिहार

31/7/17
(खालिद अरशद)
प्रषासी पदाधिकारी–सह–प्रभारी प्रोक्योरमेंट
राज्य स्वास्थ्य समिति, बिहार

31/7/17
(सच्चिदानन्द चौधरी)
विशेष सचिव, स्वास्थ्य विभाग
बिहार

31.7.17
(डॉ0 शशि रानी)
निदेशक प्रमुख
स्वास्थ्य सेवाएँ, बिहार

5th floor, Biscomaun Bhavan, Gandhi Maidan, Patna 800 001
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Tender Reference no. BMSIC/DRUGS/17-05

S.N.	*Technical Eligibility Criteria as per NIT*	*Firm Name - Bharat Serum and Vaccines limited* *Corporate Address: - 17th Floor, Hoechst House, Nariman Point, Mumbai 400021* *Manufacturer Address:- Plot No. K 27 Additional MIDC Anandnagar Ambernathe East Dist.,Thane 421501 Maharastra*
1	Constitution of the Bidding Company/Firm such as Memorandum of Association and Article of Association with complete address. As per Clause 3(c).	(a) Scanned copy of Articles of Assciation of Bharat Serum & Vaccines limited is submitted. (pg. no-192-79) (b) Memorandum of Association of Bharat Serums and Vaccines limited is submitted. (pg.no-78-73) **Note- I. Articles of Assciation is not self attested by Authorised signatory with official seal of the firm which is not in accordance with clause-5 (j) of NIT rateher it is submitted in general documents.** **Note:- II. Bidder information/Bidder Details as per Annexure-V is not submitted by bidder in mandatory documents which is not in accordance with clause 3 (d) of NIT.** (c) Self attested copy of certificate of incorporation of Gautam Laboratories Private Limited dt 29.04.1971 submitted (page no.205) (d) Self attested copy of fresh certificate of incorporation dt 14.03.1983 reagarding the change of name of company from Gautam Laboratories private limited to Bharat Serum and Vaccines private limited is submitted (page no.204) **Note: Certificate of incorporation of Bharat Serum and Vaccines Limited not submitted.**
2	Power of Attorney or Resolution of Board by which the authorised signatory has been authorised by bidder firm to sign the documents. As per Clause 3 (e) of the NIT.	(a) Scanned copy of certified true copy of the resolution regarding power of attorney to be issued to Mr. Adeet Ghosh Vice president passed by the board of directors of the company at its meeting held on January 31, 2014 is submitted. [Pg. no.-63] (b) Scanned copy of Power of Attorney dated 14th day of feburary, 2014 is submitted wherein it is stated that Bharat Serums And Vaccines Limited, a Company do hereby nominate, constitute authorise and appoint Mr. Adeet Ghosh, Group Vice President, to be true and lawful Attorney for and on behalf of the company [Pg. no. 62 to 60]
3	List of item Quoted in prescribed format as Annexure III as per Clause 3 (o)	(a)Scanned copy of list of items quoted as per Annexure-III is submitted. [Pg. no.-56] No of items quoted 01 (one) nit s. no name of Drug specification pack size 1 Anti Snake Venom Serum 10 ml (Lyophilized) 10 ml vial **(With sterile water for injecton)** **Note:-(1) In column of Name of Drug in submitted list of items quoted the words "with sterile water for Injection" is not mentioned.** **(2) As per IP 2014 - Name of the said product is mentioned in IP as (i) Snake Venom AntiSerum -** **(ii) Snake Antivenin** **(iii) Snake Antivenom Serum** **(iv) Snake Venom Antitoxin.** **whereas the name of the product in NIT is mentioned as Anti Snake Venom Serum.**

4	• Minimum three years old valid manufacturing licence of the product quoted with latest license renewal certificate. As per clause 3(f). • Approved product list as per the license issued for quoted drugs for minimum three years. As per clause 3(f). • Manufacturing License along with approved product list must be valid till the last date of the submission of tender. As per clause 3(f). • In Case of those drugs which are notified first time in IP 2014 then Manufacturing and Marketing license should be in any official compendium (USP/BP/IP) along with current valid License in IP in continuation. • Market standing certificate & Manufacturing certificate issued by the Licensing Authority as a Manufacturer for each drug quoted for the last 3 years (Certificate should be enclosed with list of items) except for the drugs falling under the category of 'New Drug' as defined by CDSCO (Central Drugs Standard Control Organization). If permission in Form 46 from DCGI has been obtained, then the 3 years Manufacturing & Market standing clause will be relaxed.The provisions of Rule 122 E of Drugs and Cosmetics Act rule 1945 shall apply. • For all regulated products, the bidder should have at least two years of manufacturing and marketing experience of the particular items as a manufacturer for each regulated product quoted in the tender. However, this would not apply to regulated products which have been licensed by DCGI less than two years ago. A certificate from DCG (I) shall be required for all new regulated products to this effect. • FFS (Flow Fill & Seal Process) Technology will be accepted wherever applicable. Bidders shall submit self attested copies of required manufacturing license and approved product list in support of above mentioned condition and they are required to specify the quoted product in their approved product list by highlighting it. As per Clause 3(f)	(a)Self attested copy of licence to manufacture for sale or distribution of large volume parenterals/Sera and vaccine in Form 28 D [Licence No.-KD/4 date of issue 27-06-2003] issued by licensing Authority Joint commissioner (Konkan Division) FDA (M.S.) thane and central licence ,approving authority, New Delhi along with a attached list of items to be manufactured is submitted. wherein it is stated that the licence shall be in force from 27-06-2003 to 26-06-2008. The quoted items is mentioned at S. No. 1 of the attached list. [Pg. no.50 to 49] (b) Self attested copy of approved list of product dt 27.06.2003 under license No.-KD-4 is submitted in which quoted product is mentioned at S. No.-01 as Snake Venom AntiSerum IP (lyophilized). (page no.51) (c) Self attested copy of certificate of renewal of licence in Form-26-H bearing licence No.-KD-4 dated 27-06-2003 issued by FDA, Govt of Maharastra for period 27-06-2008 to 26-06-2013 is submitted. [Pg. no.-52] (d) Self attested copy of list of product under licence NO.-KD-4 is submitted in which quoted product is mentioned at S. No.-01as Snake Venom AntiSerum IP (lyophillized). [Pg. no.-51] (e) Self attested copy of certificate of renewal of licence in Form-26-H bearing licence No.-KD-4 dated 27-06-2003 issued by FDA, Govt of Maharastra for period 27-06-2013 to 26-06-2018 is submitted. [Pg. no.-55] (f) Self attested copy of list of product under licence NO.-KD-4 is submitted in which quoted product is mentioned at S. No.-01 as Snake Venom AntiSerum IP (lyophillized). [Pg. no.-53] **Note:- Pack Size not mentioned in approved list.** (g) Self attested copy of Manufacturing & Marketing certificate [No.-6076420 dated 01/07/2017] issued by Licensing Authority, FDA, Konkan Division, valid upto 30/06/2018 is submitted wherein it is stated that the above products(quoted product-mentioned at sl.no.12) are being ,marketed by the firm since last 3 years.(page no.47-46) (h) Self attested copy of Licence to sell, stock or exhibit (or offer) for sale in Form 20B and 21B is also submitted. wherein it is stated that the licence shall be in force from 03 dec. 2013 to 02 dec 2018. (Pg. no.- 71 to70) (i) self attested copy of Permormance certificate (no. 6072887 dated 15/12/2016) issued by Licensing Authority, FDA, Maharastra is submitted. wherein it is stated that this certificate is valid for a period 15/12/2016 to 14/12/2017. (Pg. no. - 69 to 68) **Note - Signature of Authorized signatory on most of the submitted documents does not match with the specimen signature.**
5	In case of Importer, the bidder (importer) firm must have minimum three years old valid import License of the quoted product. All Quoted products should be accompanied by their invoices, statement and import License showing that the quoted product are being imported and sold in India by the bidder (Importer) firm minimum for Last three years. Import license must be valid on the last date of submission of tender.As per Clause 3(g)	(a) scanned copy of letter regarding IMPORT LICENCE NOT APPLICABLE on letter head is submitted. (Pg.no.- 48)
6	Self attested copies of Market standing certificate (in India) of minimum three years, issued by the concerned Licensing Authority from Drugs Control Department for the quoted product. As per Clause 3(h)	N/A
7	Self attested copy of Non Conviction Certificate (NCC), issued by the concerned Licensing Authority from Drug Control Administration of the state for last three years. It should be not more than one year old. As per Clause 3(i).	(a) Self attested copy of Non conviction certificate (no. 6076151) dt. 15-06-2017 for manufacturing licence 28-KD/4, 28E-KD/5 & 28-KD/360) issued Licensing Authority, FDA, Maharastra valid upto 14.06.2018 is submitted. where in it is stated that Bharat Serums & vaccines limited or any office bearer does not stand convicted at the instance of this administration under D & C Act 1940 and rules thereunder so far (pg. no. 45-44)

8	Self attested copies of WHO-GMP/GMP as per revised Schedule- 'M'/COPP Certificate of the manufacturing unit issued by the Licensing Authority / Drugs Control Department. The GMP certificate must not be older than one year form the last date of submission of tender. As per clause 3(j).	(a) Self attested copy of certificate of good manufacturing practices (No. NEW-WHO-GMP/CERT/KD/36205/2016/11/16049) issued on 28-07-2016 is submitted wherein it is stated that this certificate remains valid until 26 jul. 2018. (Page no. 7 to 3) (b) self attested copy of GMP certificate no. 6076149 dt- 15.-06-2017 valid upto date 14-06-2018 for Mfg. licence 28-KD/4,28E-KD/5 & 28-KD/360 is submitted. (pg. no. - 2) (c) self attested copy of Good Laboratory Practices Certificate (6076148 dated 15/06/17) is also submitted. (Pg. no.- 1)
9	Self attested copies of Maximum Production Capacity Certificate (section wise) issued by concerned Licensing Authority form Drugs Control Department highlighting the quoted product section.In case of Importer An affidavit (With Stamp) sworn before first class magistrate/Notary stating the batch production capacity of the firm and also that said production (Importing)capacity shall be adequate for requirement laid in NIT. Importers will have also to submit Invoices/Evidence of import in items of said product with quantity details. As per Clause 3(k).	Self attested copy of capacity and quality certification issued on 05-07-2016 by licensing authority , FDA, Konkan division Mahrastra state is submitted wherein the quoted product is mentioned at s.no.92. (pg. no. - 43 to 32) **Note: Capacity and quality certification(product wise) is submitted. As per 3(k) of NIT maximum production capacity certificate (section wise) is required**
10	An affidavit (with stamp)sworn before first class magistrate/Notary stating that the firm & its quoted product is not black listed currently (as on the last date of submission of the tender) by Central Government/ Central Government Agencies/any state government/any of the state government agencies/any Drug Procurement Agencies or by BMSICL as per Annexure II (clause 3(n) of NIT).	self attested copy of notarised affidavit (with stamp of Rs-100/-) for non blacklisting dt 20.07.2017 is submitted as per Annexure- II of NIT. (Pg. no. - 31 to 29)
11	EMD details (DD number/BG number and date with issuing bank) as per Clause 3(b) .	D.D. No.002311, Dt : 19-07-2017 Rs. 1,00,000/- Page No.- 13.
12	Tender Fee Rs 10,000/- in form of DD as per Clause 3(a).	D.D. No.002312, Dt : 19-07-2017 Rs. 10,000/- Page No.- 27.
13	Self attested copies of the Audited Annual Balance Sheet and Profit and Loss statement showing details of their annual average turnover not less than 25 crores for any three of the last four consecutive financial years. (Auditor/ C.A. Certificate of turnover will not be accepted). As per Clause 3(l)	F.Y - 2013-14 Rs.364.39/- Crore Page No.- 22-23, F.Y - 2014-15 Rs.493.16/- Crore Page No. - 23-24, F.Y -2015-16 Rs.560.36/- Crore Page No.-25-26.
14	Self attested copy of Income Tax Return for any three of last four consecutive Assessment years. As per Clause 3(m).	A.Y - 2014-15 Page No.- 18, A.Y - 2015-16 Page No.- 19, A.Y - 2016-17 Page No.-20.
15	Self attested copy of PAN Card of the Bidder Company. As per Clause 3(p)	PAN No. - AAACB2431M, Page No. - 17.
16	Self attested copy of Certificate of valid Sales tax/VAT registration of the bidder company. As per Clause 3(q)	GSTIN No. - 27AAACB2431M1ZT, Page No. - 72
17	Affidavit (with stamp)declaration regarding acceptance of tender conditions to be submitted by the bidding firm as per Annexure IV (Clause 5(k) of NIT).	A notarised affidavit (with stamp of Rs-100) regarding acceptance of tender conditions dt 18.07.2017 is submitted as per Annexure-IV of NIT. (pg. no. - 16 to 15)

The facts in clause No-1 to 10 and 17 of the provided sheet has been compiled with due deligence and care, on the basis of document provided by BMSICL, in compliance of letter No 1146(15) Dt-05.11.2016 of Health Department, Govt of Bihar . Inspite, some inadvertent discrepencies could have been crept in. Humble request to all concerned to bring to notice in due time if any discrepencies is observed for rectification.

Tender Reference no. BMSIC/DRUGS/17-05		
S.N.	*Technical Eligibility Criteria as per NIT*	*Firm Name - VINS BIOPRODUCTS LIMITED* *Corporate Address: - 806,ESSJAY HOUSE,ROAD NO.3,BANJARA HILLS ,HYDERABAD-500034,TELANGANA STATE* *Manufacturer Address:- SY 117 THIMMAPUR VILLAGE KOTHUR MANDAL,RANGA REDDY DISTRICT-509325 TELANGANA STATE*
1	Constitution of the Bidding Company/Firm such as Memorandum of Association and Article of Association with complete address. As per Clause 3(c).	(1) Self attested copy of Memorandum of association and Article of Association submitted (page no.1 to 47) (2) Self attested copy of certificate of incorporation dt 17.02.1997 in the name of Vins Bioproducts Private Limited submitted (page no.55) (3) Another self attested copy of Fresh certificate of incorporation dt 27.11.2000 regarding change of name of company **from Vins Bioproducts private limited to Vins Bioproducts Limited submitted** (page no.54) **A copy of Ammalgamation of SPL with VBPL sanctioned/Confirmed by the Honerable High court Hyderabad submitted (page no.48-53)** **(4) Bidder Information as per Annexure-V is submitted wherein at Sl No. 06 only enclosed list of director is mentioned. (page no.150)** **Note: List of directors is not submitted**
2	Power of Attorney or Resolution of Board by which the authorised signatory has been authorised by bidder firm to sign the documents. As per Clause 3 (e) of the NIT.	1) Self attested copy of Resolution of Board dt 19.06.2017 issued by the Whole time director Ajit Nair without issued date by which Mr. Vinai Babu vice president (Domestic sales & Marketing of the company is authorised to sign, Execute, file forms, make corrections, modification, give undertakings, declarations, participate in all tenders for supply of the goods & Services on behalf of the company. (page no.62) 2) A copy of Power of Atorny on the Non Judicial Stamp Paper of Rs. 100 addressed to The Managing Director BMSICL, Patna issued by Ajit Nair, Director without issue date submitted in which authorises Mr. P. Vinay Babu, Vive President Sales and Marketing is authorised to sign, submit all tender documents and execute the order, aggrement with BMSICL Patna. (page no.61) **Note:- In Resolution passed by the board, the name of Authorised person is written as MR. Vinai Babu whereas in the Power of Atorny on the Non Judicial Stamp Paper of Rs. 100, the name of Authorised Person is written as MR. P. Vinai Babu.**

3	List of item Quoted in prescribed format as Annexure III as per Clause 3 (o)	Self attested copy of List of Item quoted as per Annexure III is submitted. (Page No. 145) NIT S. No. -01-No. of item quoted-01 (one) - Anti Snake Venom Serum (with sterile water of injection) 10 ml vial Lyophillesed. **Note: As per IP 2014 - Name of the said product is mentioned in IP as (i) Snake Venom AntiSerum -** **(ii) Snake Antivenin** **(iii) Snake Antivenom Serum** **(iv) Snake Venom Antitoxin.** **whereas the name of the product in NIT is mentioned as Anti Snake Venom Serum.**
4	• Minimum three years old valid manufacturing licence of the product quoted with latest license renewal certificate. As per clause 3(f). • Approved product list as per the license issued for quoted drugs for minimum three years. As per clause 3(f). • Manufacturing License along with approved product list must be valid till the last date of the submission of tender. As per clause 3(f). • In Case of those drugs which are notified first time in IP 2014 then Manufacturing and Marketing license should be in any official compendium (USP/BP/IP) along with current valid License in IP in continuation. • Market standing certificate & Manufacturing certificate issued by the Licensing Authority as a Manufacturer for each drug quoted for the last 3 years (Certificate should be enclosed with list of items) except for the drugs falling under the category of 'New Drug' as defined by CDSCO (Central Drugs Standard Control Organization). If permission in Form 46 from DCGI has been obtained, then the 3 years Manufacturing & Market standing clause will be relaxed.The provisions of Rule 122 E of Drugs and Cosmetics Act rule 1945 shall apply. • For all regulated products, the bidder should have at least two years of manufacturing and marketing experience of the particular items as a manufacturer for each regulated product quoted in the tender. However, this would not apply to regulated products which have been licensed by DCGI less than two years ago. A certificate from DCG (I) shall be required for all new regulated products to this effect. • FFS (Flow Fill & Seal Process) Technology will be accepted wherever applicable. Bidders shall submit self attested copies of required manufacturing license and approved product list in support of above mentioned condition and they are required to specify the quoted product in their approved product list by highlighting it. As per Clause 3(f)	(1) Self attested scanned copy of manufacturing licence in Form 28D having licence no. 01/MN/AP/2003/SERA/G issud on 17.12.2003 is submited in which name of drug is writtrn as SNAKE VENOM ANTI SERUM (BULK) [pg. no.-90] (2) Self attested scanned copy of renewal certificate in FORM 26H for the lic. no. 01/MN/AP/03/SERA/G issued on 25-07-2009 for the period 17.12.2008 to 16-12-2013 and dt 26-07-2014 for removal period 17-12-2013 to 16-12-2018 submitted (page no.105 & 122) (3) Self attested letter issued by deputy director (enforcement)drug control administration Hydrabad is submitted in which renewal of drug licence in FORM28D bearing no.01/MN/AP/2003/SERA/G approved by central licence approving authority New Delhi is mentioned (page no 124) (4) Self attested scaned copy of letter issud by DCGI to deputy director , licensing and controlling authority Hydrabad regarding amendment in the renewal of manufacturing license in FORM26H with respect to nomenclature of the product for the period of 17.12.2013 to 16.12.2018 is submitted (page no 123) (5) Self attested copy of Additional product approval in FORM28D for the product snake venom antiserum IP lyophilised issued on 26.09.2007 by director drug control administration Hydrabad is submited (page no.92-93) (6) Self attested copy of List of additional product to be approved to M/S VINS BIOPRODUCT LIMITED Sy no.117 Thimmapur village, Kothur Mandal Mahaboob Nagar District under Drug licence bearing no. 01/MN/AP/03/SERA/G approved by DCGI New Delhi on 02.03.2015 is submitted the quoted product being highlighted is mentioned at Sl. No.-9.(page no.108 to 121). (7.) Self attested copy of Market Standing Certificate L.Dis. No. 8623/E(k)/TS/2017, dated 29/06/2017 is submitted in which it is stated that the firm has standing of more than three years (2014-15, 2015-16, 2016-2017) in manufacturing and marketing of the products in which at S. No. 01 Snake Venom Antiserum I.P. (Lyophilliosed) is mentioned. This certificate is issued by Joint Director & Licensing Authority, Telangana. Page No. 85-86) (8) Self attested copy of Market Standing Certificate L.Dis. No. DI/SAN/KTR/MSC/04/07/2016-6, dated 16/07/2016 is submitted in which it is stated that the firm has standing of more than three years (2013-14, 2014-15, 2015-2016) in manufacturing and marketing of the products in which at S. No. 01 Snake Venom Antiserum I.P. (Lyophilliosed) is mentioned. This certificate is issued by Deputy Director & Certifying Authority, Telangana. (Page No. 83-84)

5	In case of Importer, the bidder (importer) firm must have minimum three years old valid import License of the quoted product. All Quoted products should be accompanied by their invoices, statement and import License showing that the quoted product are being imported and sold in India by the bidder (Importer) firm minimum for Last three years. Import license must be valid on the last date of submission of tender.As per Clause 3(g)	N/A
6	Self attested copies of Market standing certificate (in India) of minimum three years, issued by the concerned Licensing Authority from Drugs Control Department for the quoted product. As per Clause 3(h)	N/A
7	Self attested copy of Non Conviction Certificate (NCC), issued by the concerned Licensing Authority from Drug Control Administration of the state for last three years. It should be not more than one year old. As per Clause 3(i).	Self attested copy of Non conviction Certificate L.Dis. No.462/E(k)/TS/2017, dated 12-01-2017 issued by Joint Director & Licensing Authority,Telengana is submitted in which it is mentioned that the said licence manufacturer has not been convicted by any government or Central government organization during the preceeding for the last three (3) years. (page no.75-76)
8	Self attested copies of WHO-GMP/GMP as per revised Schedule- 'M'/COPP Certificate of the manufacturing unit issued by the Licensing Authority / Drugs Control Department. The GMP certificate must not be older than one year form the last date of submission of tender. As per clause 3(j).	(1) Self attested scanned copy of GMP certificate issued on 28.10.2016 by deputy derector and certifying authority of drug control administration Telangana is submited(page no.140) (2) Self attested scanned copy of GLP certificate issued on 28.10.2016 by deputy derector and certifying authority of drug control administration Telangana is submited(page no.139) (3) Self attested scanned copy of letter issued on 10.02.2017 by joint derector and licencing authority of drug control administration Telangana along with list of products approved under WHO-GMP certificate Scheme for country and export purpose submitted (the quoted product is mentioned at Sl. No.-01) and stated that the manufacturer confirms to requirements for Good Manufacturing proctices in the manufacturing Practices in the manufacturing Practices in the manufacturer and the quality control (as recommended by the WHO) (page no.129-137)

9	Self attested copies of Maximum Production Capacity Certificate (section wise) issued by concerned Licensing Authority form Drugs Control Department highlighting the quoted product section.In case of Importer An affidavit (With Stamp) sworn before first class magistrate/Notary stating the batch production capacity of the firm and also that said production (Importing)capacity shall be adequate for requirement laid in NIT. Importers will have also to submit Invoices/Evidence of import in items of said product with quantity details. As per Clause 3(k).	Self attested scanned copy of production capacity certificate bearing number L. Dis No. DI/SAN/KTR/PCC/18/10/2016-6 dated 24-10-2016 issued by Joint Director, Licensing and controlling authority department of Drug Control Administration Telangan is submitted. **Note:- In this certificate the annual production capacity of M/s-Vins bioproduct ltd. is 4100000 Vials. mentioned. In this certificate there are seven different items are mentioned [pg. no.-65-66]**
10	An affidavit (with stamp)sworn before first class magistrate/Notary stating that the firm & its quoted product is not black listed currently (as on the last date of submission of the tender) by Central Government/ Central Government Agencies/any state government/any of the state government agencies/any Drug Procurement Agencies or by BMSICL as per Annexure II (clause 3(n) of NIT).	Scanned copy of Notarised affidavit for non blacklisting as per Annexure-II dated 20-07-2017 on Indian non Judicial stamp paper of Rs 100 is submitted. In which the name of quoted item Anti Snake Venom Serum (with sterile water for Injection) is mentioned [Pg. no. -146]
11	EMD details (DD number/BG number and date with issuing bank) as per Clause 3(b) .	B.G. No.- 790G0PG171920002, Dt : 11-07-2017 Rs. 1,00,000/- Page No.- 141-143
12	Tender Fee Rs 10,000/- in form of DD as per Clause 3(a).	D.D. No. - 064402, Dt : 03-07-2017 Rs. 10,000/- Page No.- 63
13	Self attested copies of the Audited Annual Balance Sheet and Profit and Loss statement showing details of their annual average turnover not less than 25 crores for any three of the last four consecutive financial years. (Auditor/ C.A. Certificate of turnover will not be accepted). As per Clause 3(l)	F.Y - 2013-14 Rs.51.07/- Crore Page No.- 71-72, F.Y - 2014-15 Rs.54.81/- Crore Page No. - 70-69, and F.Y - 2015-16 Rs.61.24/- Crore Page No. - 67-68.
14	Self attested copy of Income Tax Return for any three of last four consecutive Assessment years. As per Clause 3(m).	A.Y - 2013-14 Page No.- 125, A.Y - 2014-15 Page No.- 126, A.Y - 2015-16 Page No.- 127, A.Y - 2016-17 Page No.-128.
15	Self attested copy of PAN Card of the Bidder Company. As per Clause 3(p)	PAN No. - AABCV0226F, Page No.- 73.
16	Self attested copy of Certificate of valid Sales tax/VAT registration of the bidder company. As per Clause 3(q)	GSTIN No. - 36AABCV0226FIZR, Page No. - 58
17	Affidavit (with stamp)declaration regarding acceptance of tender conditions to be submitted by the bidding firm as per Annexure IV (Clause 5(k) of NIT).	Scanned copy of Notarised affidavit for (acceptance of tender conditions) as per Annexure -IV on Indian non Judicial stamp paper of Rs 100 on dated 17-07-2017 is submitted [Pg. no.-144]

The facts in clause No-1 to 10 and 17 of the provided sheet has been compiled with due deligence and care, on the basis of document provided by BMSICL, in compliance of letter No 1146(15) Dt-05.11.2016 of Health Department, Govt of Bihar . Inspite, some inadvertent discrepencies could have been crept in. Humble request to all concerned to bring to notice in due time if any discrepencies is observed for rectification.