अखिल भारतीय आयुर्विज्ञान संस्थान, रायपुर (छत्तीसगढ़)
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AIIMS/R/HS/Pharma/153/Disinfectant/Antiseptic/2018/RC/ Date: 16/05/2019

Corrigendum

Tender No. AIIMS/R/HS/30/Drug & Medicine/2018/RC/

CPPP Tender ID No:- 2019_IMSRP_451568_1

S/ N.	Page no. /Clause no. as per bid documents	Existing Entry	To be Read as
1	Page no. 11, Clause 4 point no. xvii	WHO –GMP as per schedule M (license no. and MFG. unit address highlighted	WHO –GMP/GMP as per revised schedule M (license no. and MFG. unit address highlighted
2	Page no. 11, Clause 4 point no. xviii d	A valid WHO-GMP as per Schedule 'M' certificate as per requirement of the Drugs and cosmetic act and rules of Govt. of India from Competent Authority.	A valid WHO-GMP/GMP as per revised Schedule 'M' certificate as per requirement of the Drugs and cosmetic act and rules of Govt. of India from Competent Authority.
3	Page no. 12, Clause 4 point no. xix e	WHO-GMP as per Schedule 'M' certificate as per requirement of the Drugs and cosmetic act and rules of Govt. of India from Competent Authority.	WHO-GMP/GMP as per revised Schedule 'M' certificate as per requirement of the Drugs and cosmetic act and rules of Govt. of India from Competent Authority.
4	Page no. 13, Clause 5 sub clause no 5.1 point no. a	Manufacturer should having valid License (issued by F.D.A) under the Drugs and Cosmetics Act, 1940 (including its amendments) and should comply with Schedule M (WHO-GMP) of Drugs and Cosmetic Act. The bidder should furnish self-attested photocopy of manufacturing License (Own license/ loan license/ third party license) for the product duly approved by the Licensing authority for each and every product quoted as per specification in	Manufacturer should having valid License (issued by F.D.A) under the Drugs and Cosmetics Act, 1940 (including its amendments) and should comply with Revised Schedule M (WHO-GMP/GMP) of Drugs and Cosmetic Act. The bidder should furnish self-attested photocopy of manufacturing License (Own license/ loan license/ third party license) for the product duly approved by the Licensing authority for each and every product quoted as per specification in the tender.

		the tender. The license must have been duly renewed up to date and the items quoted shall be clearly highlighted in the license. Items for which FDA do not issue manufacturing license but issue repacking license, firm should submit valid repacking license. If revalidation of drug license has been applied, copy of application to State Drug / Licensing authority should be attached.	The license must have been duly renewed up to date and the items quoted shall be clearly highlighted in the license. Items for which FDA do not issue manufacturing license but issue repacking license, firm should submit valid repacking license. If revalidation of drug license has been applied, copy of application to State Drug / Licensing authority should be attached.
5	Page no. 13, Clause 5 sub clause no 5.1 point no. b	The manufacturer should have received a valid WHO-GMP as per revised Schedule 'M' issued by Licensing Authority and WHO-GMP inspection certificate in line with the WHO certification scheme, on pharmaceuticals being quoted. Bidder should submit a valid product wise Certificate of WHO-GMP as per schedule "M" or WHO-GMP issued by Regulatory Authority, for each item offered. License no. and manufacturing unit address clearly highlighted on WHO-GMP certificate.	The manufacturer should have received a The manufacturer should have received a valid WHO-GMP/GMP as per revised Schedule 'M' issued by Licensing Authority and WHO-GMP/GMP inspection certificate in line with the WHO certification scheme, on pharmaceuticals being quoted. Bidder should submit a valid product wise Certificate of WHO-GMP/GMP as per revised schedule "M" or WHO-GMP/GMP issued by Regulatory Authority, for each item offered. License no. and manufacturing unit address clearly highlighted on WHO-GMP/GMP certificate.
6	Page no. 13, Clause 5 sub clause no 5.2 point no. c	Manufacturer must have valid License (issued by F.D.A) under the Drugs and Cosmetics Act, 1940 (including its amendments) and should comply with Revised Schedule M WHO-GMP of Drugs and Cosmetic Act. The bidder should furnish self-attested photocopy of manufacturing License (Own license/loan license/third party license) for the product duly approved by the Licensing authority for each and every product quoted as per specification in the tender. The license must have been duly renewed up to date and the items quoted shall be clearly highlighted in the license. Items for which FDA do not issue manufacturing license but issue repacking license, firm should submit valid repacking license. If revalidation	Manufacturer must have valid License (issued by F.D.A) under the Drugs and Cosmetics Act, 1940 (including its amendments) and should comply with Revised Schedule M WHO-GMP/GMP of Drugs and Cosmetic Act. The bidder should furnish self-attested photocopy of manufacturing License (Own license/loan license/third party license) for the product duly approved by the Licensing authority for each and every product quoted as per specification in the tender. The license must have been duly renewed up to date and the items quoted shall be clearly highlighted in the license. Items for which FDA do not issue manufacturing license but issue repacking license, firm should submit valid repacking license. If revalidation of drug license has been applied, copy of

		of drug license has been applied, copy of application to State Drug / Licensing authority may be attached.	application to State Drug / Licensing authority may be attached.
7	Page no. 14, Clause 5 sub clause no 5.2 point no. d	The manufacturer should have received a valid WHO-GMP as per revised Schedule 'M' issued by Licensing Authority or WHO-GMP inspection certificate in line with the WHO certification scheme, on pharmaceuticals being quoted. Bidder should submit a valid WHO-GMP issued by Regulatory Authority, for each item offered. License no. and manufacturing unit address clearly highlighted on WHO-GMP certificate.	The manufacturer should have received a valid WHO-GMP/GMP as per revised Schedule 'M' issued by Licensing Authority or WHO-GMP/GMP inspection certificate in line with the WHO certification scheme, on pharmaceuticals being quoted. Bidder should submit a valid WHO-GMP/GMP issued by Regulatory Authority, for each item offered. License no. and manufacturing unit address clearly highlighted on WHO-GMP/GMP certificate.

Remarks:-

01 All other terms and condition of tender will remain unchanged.

Issued by

Rishi Gupta
Stores Officer (H)
AIIMS Raipur (C.G.)