S. No.	Date of Inspection	Purpose of inspection	Team of inspector/ Expert	Remarks/ Conclusions	Regulatory actions (certification and/or enforcement)
A.	M/s Serum Inst	itute of India Pvt Ltd, S. N	lo. 212/2, Hadapasr,	Pune 411 028, Maharashtra.	
1.	02.05.2024 to 0.3.05.2024	Joint inspection for addition of new manufacturing facility.	Team comprising of officers from CDSCO and State.	The joint inspection of the firm was carried out from 02.05.2024 to 0.3.05.2024 for the purpose of addition of new manufacturing facility. During inspection certain minor observations were noted with respect to facility, validation, Quality Risk Assessment and documentation.	 Firm was directed to submit compliance with respect to the observations noted during inspection. Post approval change NOC issued for the proposed change.
B.	M/s Serum Inst	itute of India Pvt Ltd, S. N	lo. 105 – 110, Manja	ri Bk., Pune 412 307, India, Ma	aharashtra, India
1.	21.03.2024 to 22.03.2024	Compliance verification for inspection dated 20.05.2023.	Team comprising of officers from CDSCO and State	The compliance of the observations submitted by the firm as CAPA were verified through joint inspection from 21.03.2024 to 22.03.2024.	Firm was directed to submit CAPA plan with respect to the remaining observations.

S. No.	Date of Inspection	Purpose of inspection	Team of inspector/ Expert	Remarks/ Conclusions	Regulatory actions (certification and/or enforcement)
C.	M/s Gennova E Maharashtra, li	•	ed, Block 1, Plot No.	P-1 & P-2, ITBT park, Phase	II, MIDC, Hinjawadi, Pune,
1.	24.04.2024 to 26.04.2024	Routine inspection under CIP 2024.	Team comprising of officers from CDSCO, State and CDL Kasauli.	The joint inspection of the firm was carried out from 24.04.2024 to 26.04.2024 for the purpose of routine inspection as per CIP 2024. During inspection certain observations were noted with respect to procedure and process and adherence to SOPs.	 Firm was directed to submit compliance with respect to the observations noted during inspection. The compliance of the observations submitted by the firm as CAPA were verified through desktop review on.05.06.2024 and was considered as acceptable.
D.	M/s. Panacea E	Biotec Limited, Village Ma	lpur, Baddi, Distt. So	olan (H.P).	
1.	28.02.2024	Compliance verification for inspection dated 28.12.2023 to 30.12.2023 for routine as per CIP 2023 and CPP as per WHO GMP	Team comprising of officers from CDSCO and State	The compliance of the aforesaid observations were verified through joint inspection on 28.02.2024 in which most of the observations were found complied.	Grant of CPP was recommended.

Details of GMP inspection 2024

S. No.	Date of Inspection	Purpose of inspection	Team of inspector/ Expert	Remarks/ Conclusions	Regulatory actions (certification and/or enforcement)
E.	M/s. Biological R.R.Dist, T.S.	E Limited., Plot No. 1,	SP Biotechnology	Park, Phase-II, Kolthur Villa	age, Shameerpet Mandal,
1.	26.03.2024 to 28.03.2024	Joint inspection for grant of license in form 28D for additional product.	Team comprising of officers from CDSCO, State and CDL Kasauli.	The joint inspection of the firm was carried out from 26.03.2024 to 28.03.2024 for the purpose of grant of additional products manufacturing licence in Form 28D. During inspection certain minor observations were noted with respect to facility, validation, Quality Control and documentation.	Firm was directed to submit compliance with respect to the observations noted during inspection.
2.	22.05.2024	Compliance verification for inspection dated 26.03.2024 to 28.03.2024.	Team comprising of officers from CDSCO and State	The compliance of the aforesaid observations were verified through joint inspection on 22.05.2024 in which most of the observations were found complied.	Licence to manufacture the applied vaccine under Form 28D granted.
3.	22.04.2024 to 24.04.2024	Joint inspection for Grant of permission in Form CT 11 for manufacture,	Team comprising of officers from CDSCO, State and	The joint inspection of the firm was carried out from 22.04.2024 to 24.04.2024 for the purpose of for	Firm was directed to submit compliance with respect to the observations noted during

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S. No.	Date of Inspection	Purpose of inspection	Team of inspector/ Expert	Remarks/ Conclusions	Regulatory actions (certification and/or enforcement)
		examination, test and analysis.	CDL Kasauli.	grant of permission in Form CT 11 for manufacture, examination, test and analysis. During inspection minor observation was noted with respect to facility, quality management system and documentation.	inspection.
4.	20.6.2024 to 21.06.2024	Compliance verification for inspection dated 22.04.2024 to 24.04.2024	Team comprising of officers from CDSCO and State	The compliance of the observations submitted by the firm as CAPA were verified through joint inspection from 20.6.2024 to 21.06.2024.	Form CT-11 issued for manufacturing for examination, test and analysis purpose.
F.	M/s. Human B 500019, T.S	iologicals Institute (A D	Division of Immuno	logical Limited) Rakshapura	am Gachibowli Hyderabad
1.	06.06.2024 & 07.06.2024	Joint inspection for grant of CPP as per WHO guidelines.	Team comprising of officers from CDSCO and state.	The joint inspection of the firm was carried out from 06.06.2024 & 07.06.2024 for the purpose of grant of CPP as per WHO guidelines.	Firm was directed to submit compliance with respect to the observations noted during inspection.

S. No.	Date of Inspection	Purpose of inspection	Team of inspector/ Expert	Remarks/ Conclusions	Regulatory actions (certification and/or enforcement)
				inspection certain minor observations were noted with respect to material management, manufacturing process and control, and documentation.	
2.	25.07.2024	Compliance verification for inspection dated 06.06.2024 & 07.06.2024	Team comprising of officers from CDSCO and State	The compliance of the observations submitted by the firm as CAPA were verified through joint inspection 25.07.2024	Grant of CPP was recommended.
G.		ologicals Institute (A div Karkapatla (V), Markook		nunologicals Limited), Sy.No angana	o.281-284 & 321, Biotech
1.	17.06.2024 to 19.06.2024	Joint inspection for grant of CPP as per WHO guidelines.	Team comprising of officers from CDSCO and state.	The joint inspection of the firm was carried out from 17.06.2024 to 19.06.2024 for the purpose of grant of CPP as per WHO guidelines. During inspection certain minor observations were noted with respect to quality risk	Firm was directed to submit compliance with respect to the observations noted during inspection.

S. No.	Date of Inspection	Purpose of inspection	Team of inspector/ Expert	Remarks/ Conclusions	Regulatory actions (certification and/or enforcement)
				assessment, material management, manufacturing process and control and documentation.	
Н.		iotech International Ltd. al- Malkajgiri Dist- 500 07		⊔ 235 Genome Valley, Turkar India.	pally village, Shameerpet
1.	20.03.2024 to 21.03.2024	Compliance verification for inspection dated 13.10.2023 to 14.10.2023 for routine inspection under CIP 2023.	Team comprising of officers from CDSCO and State	The compliance of the aforesaid observations were verified through joint inspection from 20.03.2024 to 21.03.2024.	Firm was directed to submit CAPA plan with respect to the remaining observations.
2.	18.04.2024	Compliance verification for inspection dated 27.12.2023 to 29.12.2023 for grant of CPP as per WHO guidelines.	Team comprising of officers from CDSCO and State	The compliance of the aforesaid observations were verified through joint inspection on 18.04.2024.	Firm was directed to submit CAPA plan with respect to the remaining observations.

S. No.	Date of Inspection	Purpose of inspection	Team of inspector/ Expert	Remarks/ Conclusions	Regulatory actions (certification and/or enforcement)
3.	27.05.2024 to 29.05.2024	Joint inspection for grant of license in form 28D for additional product.	Team comprising of officers from CDSCO, State and CDL Kasauli.	The joint inspection of the firm was carried out from 27.05.2024 to 29.05.2024 for the purpose of grant of additional products manufacturing licence in Form 28D. During inspection certain minor observations were noted with respect to validation, material management, Quality management system and documentation.	Firm was directed to submit compliance with respect to the observations noted during inspection.
4.	11.06.2023 to 12.06.2023	Compliance verification for inspection dated 13.10.2023 to 14.10.2023 for routine inspection under CIP 2023	Team comprising of officers from CDSCO and State	The compliance of the aforesaid observations were verified through joint inspection from 11.06.2023 to 12.06.2023.	Firm was directed to submit CAPA plan with respect to the remaining observations.
5.	13.06.2024	Compliance verification for inspection dated	Team comprising of officers from	The compliance of the aforesaid observations were verified through joint	Firm was directed to submit CAPA plan with respect to the remaining

S. No.	Date of Inspection	Purpose of inspection	Team of inspector/ Expert	Remarks/ Conclusions	Regulatory actions (certification and/or enforcement)
		27.12.2023 to 29.12.2023 for grant of CPP as per WHO guidelines.	CDSCO and State	inspection on 13.06.2024.	observations.
I.	_	•	•	ila Healthcare Limited), Sur ist. Ahmedabad - 382 210, Gu	
1.	26.02.2024	Joint inspection for Grant of permission in Form CT 11 for manufacture, examination, test and analysis.	Team comprising of officers from CDSCO, State and CDL Kasauli.	The joint inspection of the firm was carried out 26.02.2024 for the purpose of for grant of permission in Form CT 11 for manufacture, examination, test and analysis. During inspection observation was noted with respect to validation, personnel and training and documentation.	Firm was directed to submit compliance with respect to the observations noted during inspection.
2.	16.04.2024	Compliance verification for inspection dated 26.02.2024	Team comprising of officers from CDSCO and State.	The compliance of the observations submitted by the firm as CAPA were verified through joint	Form CT-11 issued for manufacturing for examination, test and analysis purpose

S. No.	Date of Inspection	Purpose of inspection	Team of inspector/ Expert	Remarks/ Conclusions	Regulatory actions (certification and/or enforcement)
				inspection on 16.04.2024.	
J.	M/s. Cadila Pha	armaceuticals Limited, 13	89, Trasad Road, Dh	olka , Dist: Ahmedabad -382	225, Gujarat
1.	16.02.2024	Joint inspection for Grant of permission in Form CT 11 for manufacture, examination, test and analysis.	Team comprising of officers from CDSCO, State and CDL Kasauli.	The joint inspection of the firm was carried out 16.02.2024 for the purpose of for grant of permission in Form CT 11 for manufacture, examination, test and analysis. During inspection observation was noted with respect to validation, facility and equipment, and documentation.	Firm was directed to submit compliance with respect to the observations noted during inspection.
K.	M/s Chiron Bel 393002,Gujarat	•	Plot No. 3502, G.I.D.	C. Estate, P. B. No. 136, Ank	leshwar, Dist :- Bharuch-
1.	15.03.2024	Compliance verification for inspection dated 22.11.2023	Team comprising of officers from CDSCO and State.	The compliance of the observations submitted by the firm as CAPA were verified through joint inspection on 15.03.2024.	Form CT-11 issued for manufacturing for examination, test and analysis purpose

S. No.	Date of Inspection	Purpose of inspection	Team of inspector/ Expert	Remarks/ Conclusions	Regulatory actions (certification and/or enforcement)
L.	M/s. Bio-Med P	vt, C-96, Site No.1, Bulan	dshahr, Road Indl. A	rea, Ghaziabad	
1.	06.05.2024 to 07.05.2024	Compliance verification for inspection dated 26.12.2023 to 28.12.2023 as routine inspection under CIP 2023.	Team comprising of officers from CDSCO, State and CDL Kasauli.	The compliance of the aforesaid observations were verified through another joint inspection team from 06.05.2024 to 07.05.2024, in which most of the observations were found complied.	 Firm has complied most of the deficiencies of routine inspection as per Central Inspection Plan 2023. Firm was directed to submit CAPA plan with respect to the remaining observations.
M.	M/s. Bio Vaccir	nes (India) Pvt.Ltd., 46- A,	Sanwar Road, Indus	strial Area, Indore (M.P.) 4520)15.
1.	18.03.2024	Compliance verification for inspection dated 01.03.2023 to 03.03.2023 and 28.06.2023 for routine inspection under CIP 2023	Team comprising of officers from CDSCO and State	The compliance of the observations submitted by the firm as CAPA were verified through joint inspection on 18.03.2024.	The compliance of the observations submitted by the firm as CAPA were considered as acceptable.
N.		ologicals Institute (A div Iam643007, Tamilnadu.	ision of Indian Imm	unologi cals Limited) Kozhi	pannai, Pudumund Post,

S. No.	Date of Inspection	Purpose of inspection	Team of inspector/ Expert	Remarks/ Conclusions	Regulatory actions (certification and/or enforcement)
1.	04.01.2024	For cause inspection	Team comprising of officers from CDSCO, State and CDL Kasauli.	The joint investigation of the firm was carried out on 04.01.2024 for the purpose of investigation for AEFI reported on 15.12.2023.	Firm was directed to submit detailed investigation report to CDSCO. The compliance of the observations submitted by the firm as detailed investigation report were verified through desktop review on 08.02.2024 and was considered as acceptable.
Ο.	Ms. Green Sign	nal Bio Pharma Ltd., No. 4	9. Pappankuppam V	illage, Gummidipoondi, Cher	nai-601201.
1.	09.08.2024	Compliance verification for inspection dated 19.09.2022 to 21.09.2022 for post approval change.	Team comprising of officers from CDSCO, State and CDL Kasauli.	The compliance of the observations submitted by the firm as CAPA were verified through joint inspection on 09.08.2024.	PAC NOC issued for the proposed change.
2.	04.06.2024 to 06.06.2024	Routine inspection as per Central Inspection Plan-2024 and additional manufacturing line.	Team comprising of officers from CDSCO, State and CDL Kasauli.	The joint inspection of the firm was carried out from 04.06.2024 to 06.06.2024 for Routine inspection as per Central Inspection Plan-2024 and additional	Firm was directed to submit compliance with respect to the observations noted during inspection.

S. No.	Date of Inspection	Purpose of inspection	Team of inspector/ Expert	Remarks/ Conclusions	Regulatory actions (certification and/or enforcement)
				manufacturing line. During inspection observation was noted with respect to validation, facility and equipment, and documentation.	

^{*}Details are available in inspection report along with applicable regulatory action as internal record.