User Requirement Specifications
Mobile CIP system

Block Code	Area	Identification #	Qty (Nos)	Capacity(WV)
F1	Viral Vaccine Formulation-Rabies	F1-CIP-01	1	100 L
B4	Rabies bulk block	B4-CIP-01	1	150 L

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URS Annexure List

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1	P&ID
2	List of preferred MAKE of components

1.0 APPROVAL SIGNATURES

This document is prepared by the Process, Validation and GMP Compliance team of "NNE Pharmaplan India" for the project "Integrated Vaccines Complex, Chengalpattu, Chennai" (**project number:** 120310) of HLL BIOTECH LIMITED (Chennai) under the authority of their Project Manager. Hence, this document before being effective should be reviewed by HBL user/s and project/ engineering team, approved by team lead of user department & QA and authorized by the appropriate Project authority.

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2.0 EQUIPMENT DESCRIPTION

The equipment described by this URS is Mobile CIP system for Rabies Formulation block & Rabies Bulk Block.

Mobile CIP system is used for cleaning solution preparation, recirculation and cleaning of the process equipment/vessel. The scope of the Mobile CIP system which includes the preparation of cleaning solution, transfer / once through the CIP solution with the required flow rate & velocity, Re-circulating the CIP solution to the vessel and draining of the CIP solution at the end of the sequence.

Design, function and control of the unit should be cGMP compliant. The Equipment should be designed considering easy accessibility and maintenance. All piping should be designed to achieve 100% drainability as per the guidelines of ASME BPE 2012.

The Skid mounted Mobile CIP system with integrated pump, Lockable castor wheels, Heat exchanger, Process vessel, HMI, valves, instrumentation and control panel with following features:

S No.	Description	Remarks
A.	**The vessel should be equipped with the following requirements:** Sterile vent filter should be provided for venting and pressurizing the vessel. The filter should be of sterilization grade filter cartridge with code 7 type. The filter housing should be constructed of 316L stainless steel with necessary assembly.	
B.	Air Pressure regulator should be provided in the air line for regulating the air pressure.	
C.	Spray ball should be provided in the top dish of the vessel for WFI supply.	
D.	Dosing bottles with necessary accessories like level switch, Non-Return valve, Dosing pumps should be provided for acid & alkali solutions.	
E.	Differential pressure sensor (Hydrostatic Type) should be provided for measurement and control of liquid level in the vessel.	
F.	Port for recirculation line should be provided on the top dish. The recirculation line should be provided with Suitable heat exchanger for heating CIP solution, centrifugal pump for recirculation and transfer of CIP solution.	
G.	Two numbers of dosing metering pumps for addition of acid and alkali, all necessary automation and instrumentation should be provided for fully automatic functionality of the system.	
H.	Centrifugal pump with Suitable flow rate should be provided for re circulation and transfer of the CIP solution, all necessary automation and instrumentation should be provided for functionality of the system. Pump should be provided with a drain line.	
I.	A flow switch should be provided to prevent the centrifugal pump from dry running.	
J.	Conductivity sensor (0-200mS/cm) should be provided on the recirculation line and drain line for the measurement and control during CIP solution preparation & CIP cycle.	
K.	Temperature sensor duplex Pt 100 should be provided on the discharge line of the centrifugal pump for the measurement and control of CIP solution temperature during heating.	
L.	All the condensate drain lines should be provided with necessary steam traps.	

The equipment should consist of following parts for operation:

TABLE 1

S. No.	Description	Purpose	MOC	Remarks
1	Shell	Cylindrical	SS316L	
2	Top closure	Tori spherical dish	SS316L	
3	Bottom closure	Tori spherical dish	SS316L	
4	Insulation	To avoid heat loss	Mineral wool	
5	Cladding	Outer cover to Insulation	SS304	
6	Heat exchanger [Shell & Tube]	Maintain the CIP solution temperature	SS 316L	

TABLE 2

S. No.	Description	Viral Vaccine Formulation - Rabies	Rabies Bulk	Remarks
1	No. of system	1	1	
2	Max. working Volume	100 L	150 L	
3	Geometric Volume	Vendor to specify	Vendor to specify	
4	Min Operating Volume	Vendor to specify	Vendor to specify	
5	Acid dosage container	Vendor to specify	Vendor to specify	
6	Alkali dosage container	Vendor to specify	Vendor to specify	
7	H/D	Vendor to specify	Vendor to specify	

2.1 General Requirement for vessel:

S No.	Nozzle schedule :	Remarks
2.1.1	**Top Dish:**	
	Light and sight port	
	Port for Spray ball	
	Spare port –TC end	
	Acid addition port	
	Alkali addition port	
	Port for vent filter	
2.1.2	**Bottom dish:**	
	Transfer line port/vessel outlet port	

NOTE:

1.	This Technical Specification is the basis for an inquiry to a vendor and therefore the basis for the vendor's proposal.
2.	The vendor is asked to state in "REMARKS" column with "yes" if the described requirement will be completely fulfilled and with "no" in case the requirement will not or cannot be fulfilled with the proposed equipment. I n case of an deviation a comment must be inserted or enclosed as a separate annexure by referring to the respective URS specification number.
3.	The vendor must clearly comment each item of the Technical Specification. The comments must be in English language. If extra cost for necessary options becomes necessary the item must be clearly stated.
4.	In case that the requirement includes a question or request or information from the vendor, the answer / information should be stated in the "REMARKS" column.
5.	The final version of this document including the vendor's comments will become basis of a potential purchase order or contract.
6.	The Technical Specification serves to define a summary of all vendors' requirements concerning scope of delivery and services.
7.	The vendor is responsible for technically unobjectionable function of the equipment. This TS is not intended to dictate a technical design to the vendor. If agreed upon with the vendor, the vendor can apply his practically proven design.
8.	Special Instruction a. If no comments against any specification should be considered as "NO" and b. If there is no reply / comments against the complete URS by the vendor then it should be treated as unresponsive / technically non-compliant and rejected.
9.	All the instruments and controls mentioned in the URS(s) are expected to be standard supply and part of your standard equipment model. In case of any deviation or redundancy or additional scope of supply is noticed, vendor is required to obtain clarification from HBL before submitting the quotes.
10.	The makes requested are standard international makes. In case of any deviation, vendor to seek clarification from HBL before submitting the offers.
11.	Refer document Installation Requirement Specification and Specific Instructions with URS; NPI_120310_IRS_S1_01
12.	Refer Tender document with URS; NPI_120310_EQP_S1_TD_08

Note: The following points which are there in the IRS (Installation Requirement Specifications) are NOT APPLICABLE for this equipment:

a. Sl.NO 5 CE Conformity,
b. Sl.NO 7 ANSI/NSF 49-2008, ISO 14664, ISO 8362
c. Sl.NO 8 ISO 14664
d. Sl.NO 9 ISO 8362

Specifications	Remarks
3.0 PROCESS DESCRIPTION	
3.1 Input & Charging method	
The alkali and acid solutions are prepared in Mobile CIP system with the help of Dosing and metering pump and transferred to the equipment for cleaning. Cleaning solution can be charged continuously to the CIP tank so that tank should be provided with 'HIGH' and 'LOW' level functions to control the addition and discharge from the tank.	
Vendor to provide the necessary valve assembly with automation to integrate the above mentioned sequence	
3.2 Brief Process Steps	
Mobile CIP system will perform all the sequence i.e., Preparation, Transfer, Once through, Recirculation etc. from the system itself by taking the IO from victim Vessel PLC. The system should be capable of the following sequence of cleaning cycle: 1) Wash Phases: Once through with PW rinse- 55-60°C 2) Alkali addition through metering/dosing pump: PW addition through totalizer set point flow meter Re-circulation with Alkali solution- 55-60 °C 3) Intermediate wash: Once-through with PW 4) Acid addition through metering /Dosing pump: PW addition through totalizer set point flow meter Re-circulation with Acid solution Once-through with PW 4) Rinse Phases: Recirculation/Once-through with WFI 5) Drying in place: Air blow to the equipment **Note:** CIP solution preparation in the respective vessel is based on conductivity set point, a feedback signal from conductivity sensor in the recirculation line of the pump is employed for controlled acid/alkali dosing.	
3.3 Output & Discharging method	
The prepared CIP solution will be transferred to the process vessel which will be recirculated with the help of Mobile CIP system itself connected to the vessel by using flexible hoses.	
After achieving required conductivity the solution will be drained.	

4.0 PRODUCTIVITY REQUIREMENT

4.1. Change Over Time

Not Applicable	

4.2. Others (if any)

Not Applicable	

5.0 CONTAINMENT

Not Applicable	

6.0 GMP REQUIREMENTS

6.1 Process control

	The equipment must operate and control the following process cycle:	
6.1.1	Duration of each cycle.	
6.1.2	Number of cycles.	
6.1.3	Quantities of wash liquid in each cycle.	
6.1.4	Temperature of washing liquid.	
6.1.5	Cycle sequence	
6.1.6	Conductivity	
6.1.7	pH(0-14)	

6.2 Failure mode detection

6.2.1	Equipment should be capable to detect the following failure, notify the operator with alarm and shutdown the process:	
a)	Emergency stop activated	
b)	Power	
6.2.2	Following condition need only notification to operator for procedural control	
a)	End of any/all process sequence.	
b)	Low compressed air	

6.3 In – Process control

NA	

6.4 Level of instrumentation

Sufficient and suitable instrumentation for the process, safety and productivity control as indicated in the following table:

Type of control	Purpose	Instrumentation
Temperature	To monitor , control and record the temperature	Temperature sensor and transmitter
DP Sensor (Hydrostatic Type)	To monitor , control and record the level in Vessel	DP sensor and transmitter
pH & Conductivity	To monitor , control and record the Conductivity	pH and Conductivity sensor and transmitter
Flow	To check flow	Flow switch
Speed	To control the speed of Pump	VFD

6.5 Batch data display and record printing

The system should be provided with all necessary automation and instrumentation for establishing interface (Handshake b/w the system) with other systems i.e., Blending vessel, media prep. Vessel, Buffer Prep. Vessel etc.

Controller: - PLC Based Controller (Non-editable data format to be obtainable) with minimum of 10" size HMI (Displaying data trends as Graphs, synoptic view of running parameters etc)

The HMI should be of touch screen type (Provision for manual operation to be provided). All settings should be user adjustable. HMI and Control Panel should be mounted on skid.

- Human machine interface must be used to enter the process details, which should appear in the print out.
- All critical alarms, Critical parameters and interlocks
- All Recipes/ sequences
- P&ID of the vessel along with instrumentation details
- Login details
- HMI screen showing simulation of valves

6.6 GMP requirements (Others)

6.6.1 All valve and joints should be sanitary type (preferably tri-clover connection).

6.6.2 Equipment should be completely drainable.

6.7 Specific requirements

6.7.1 All attachments required for fixing nozzles, supply pipes and return pipes should be provided by vendor only.

6.7.2 All the operations should be automatic through PLC, HMI and provision to be provided for

	manual interventions.	
6.7.3	All the flexible piping used for cleaning services should be of SS re-enforced and PTFE lined to withstand pressure, temperature and very high and low pH.	
6.7.4	Vendor should provide the following details in the quotes apart from these mentioned in the URS (a) Makes of pumps (Supply and return), Conductivity meter, Valves, PLC etc. (b) Schematics of the Mobile CIP system (c) Vessel Capacity	
6.7.5	The Vendor should ensure maintenance parts availability for a minimum of 15 months from delivery.	
6.7.6	Cables, air tubes, etc required from the point (single utility point) to equipment is in scope of vendor.	
6.7.7	Vendor to perform a criticality assessment to assess the applicability of the system to Part 11 regulation. Software, if used to generate, process, store the quality critical data must be validated and must comply 21 CFR Part 11 requirements	
6.7.8	Vendor should provide the FRL (Filter, regulator, lubricator), automatic valve assembly and air pressure switch for instrument air. Connections to automatic diaphragm valve should be in vendor scope.	

7.0 CONSTRAINTS

7.1 Equipment location and available space

These equipment will be installed in the Bacterial Vaccine Formulation and Viral Vaccine Formulation block:

a. **Block:** Viral Vaccine Formulation- Rabies
Floor Ground floor
False ceiling height: 4000 mm
Physical condition of the room:
1. Class: EU Class "C"
2. Differential Pressure:50 Pa
3. Temperature maintained: 22±2 ºC
4. Relative Humidity: < 55%RH

b. **Block:** Rabies Bulk Block
Floor: Ground floor
False ceiling height: 2700 mm
Physical condition of the room:
1. Class: EU Class "C"
2. Differential Pressure: 30 Pa
3. Temperature maintained: 22±2 ºC
4. Relative Humidity: < 55 % RH

7.2 Available Utility

7.2.1	Plant steam – 130 °C – 150°C at 3 – 3.5 bar (g) ----------(Report requirement)	
7.2.2	WFI (Hot loop) – 80-85°C at 2 bar (g) ----------(Report requirement)	

7.2.3 Purified Water- 28 – 30°C at 2.5 bar (g) ----------(Report requirement)	
7.2.4 Electricity – Vendor to specify----------(Report requirement)	
7.2.5 Compressed air- 6.0– 8.0 bar (g) ----------(Report requirement)	
Note: Utility consumption to be specified by the vendor, in case if there is a deviation in the values mentioned above.	

8.0 ABBREVIATION

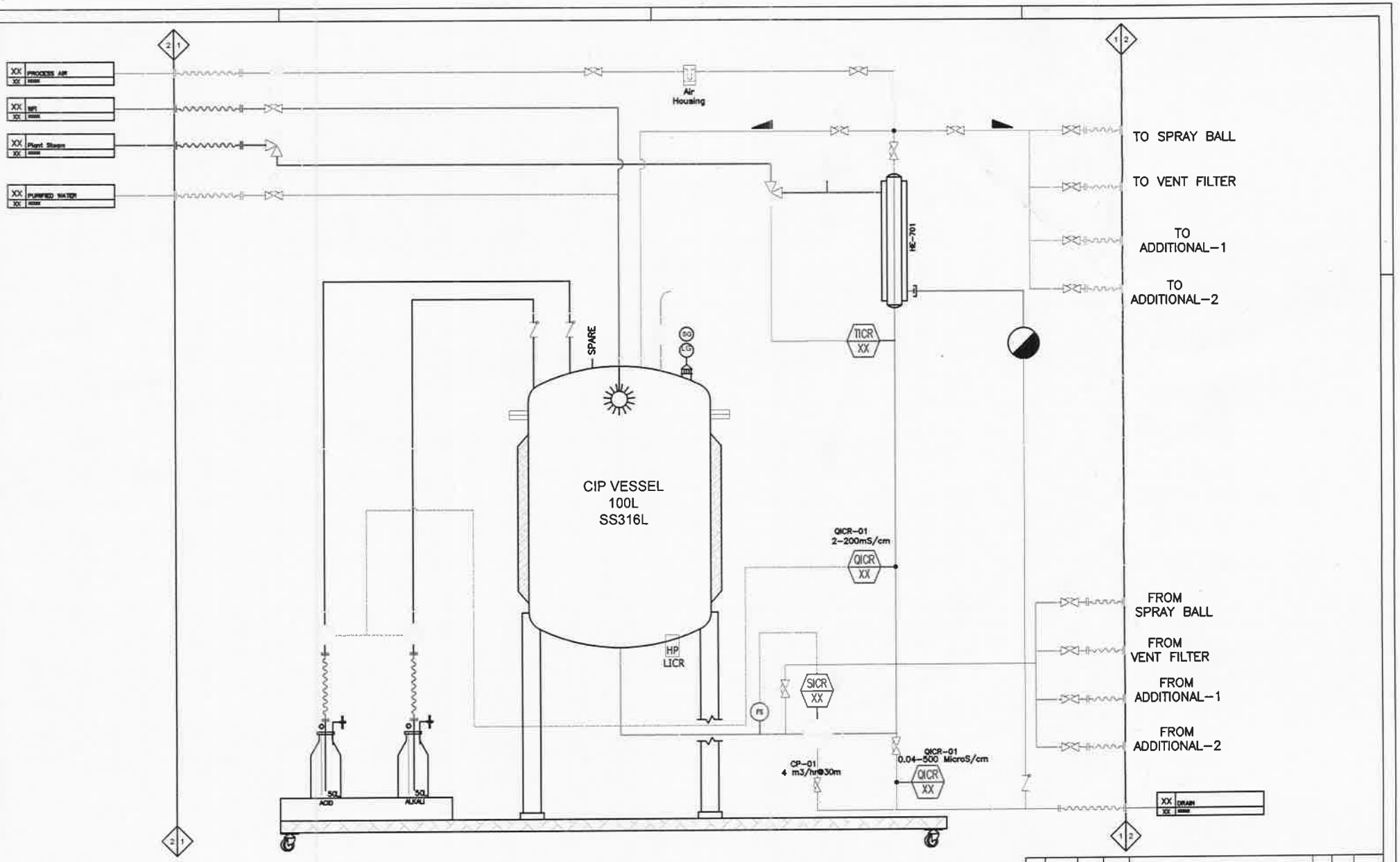
Abbreviation	Definition
IVC	Integrated Vaccines Complex
HBL	HLL Biotech limited
CIP	Clean In Place
SIP	Sterilization In Place
GMP	Good Manufacturing Practices
NPI	NNE Pharmaplan India Ltd
ISO	International Standards Organization
HMI	Human Machine Interface
PLC	Programmable Logic Controller
SCADA	Supervisory control and data acquisition system
NMT	Not More Than
TBD	To be discussed
PTFE	Polytetrafluoroethylene
ASME BPE	American Society of Mechanical Engineers Bioprocessing Equipment Standards
RH	Relative Humidity
Class EU	Class European Union

9.0 REVISION INDEX

Revision	Date	Reason for Revision
00	05-05-2014	First Draft for Client's Review
01	04-10-2014	Philosophy changed as per the mail dated 04-10-2014

URS Annexure 2: List of Preferred Make of components

SL.NO	DESCRIPTION	MAKE
A	**INSTRUMENTATION**	
1	PLC	Allen Bradley/ Siemens
2	Operator Interface/HMI	Allen Bradley/ Siemens
3	Temperature transmitter	Radix/ Yokogawa/Emerson
4	Temperature sensor	Negele/ Radix/Negale
5	Conductivity sensor	Metler Toledo/E+H
6	DP transmitter	E+H/Negele
7	DP sensor	E+H/Rosemount /Emerson
8	Flow switch	Negele/ Danfoss/E+H
9	VFD	Allen Bradley/ Siemens /Danfoss
B	**MECHANICAL**	
10	Centrifugal pump	Alfa Laval/ Grundfos
11	Vent filter cartridge	Sartorius/Pall/ Millipore
12	Spray ball	Hake/Lechler/Alfa laval
13	Diaphragm valve(Manual)	Gemu / SED
14	Ball valve(Manual)	Modentic/Saunders/Alfa laval
15	Flexible hose	BBS/ AMI Polymer / Venair
16	Dosing metering pump	Prominent/ Masterflex
17	Steam trap	Steriflow/spirax marshall
C	**PNEUMATIC**	
18	Diaphragm valve(Automatic)	Gemu / SED
19	Angle seat valve(Automatic)	Gemu / SED



NOTE : THIS P&ID IS ONLY FOR INFORMATION. ACTUAL P&ID HAVE TO BE PROVIDED BY THE VENDOR

Rev.	Date	Changed	Checked	Kind of revision	No. Of Prints	Date	Issued To

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Originated From Drg. No.:

Project:
INTEGRATED VACCINES COMPLEX,
HLL BIOTECH LIMITED

Project No.: 120310
Location: CHENGALPATTU

Description:
P&ID
MOBILE CIP SYSTEM

Sheet:- 1 of 1

nne pharmaplan
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	Date	Name
Drawn	2014.10.10	SATR
Checked	2014.10.10	BCJP
Approved	2014.10.10	VKRA

Scale-NTS | Units : mm | Size : A3

Drawing no: | Rev. 000