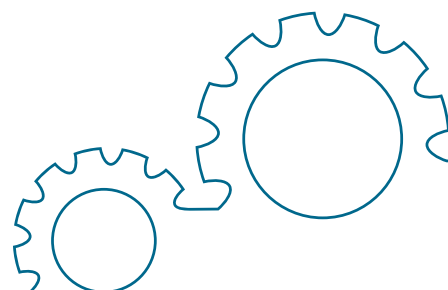
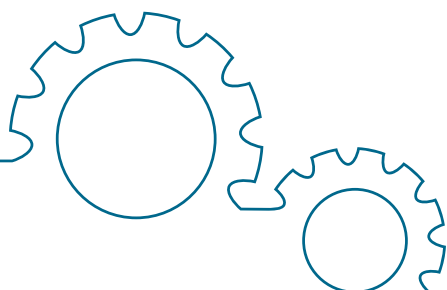


A Complete Solution For Pharma Industries®

NAV SARJAN
TURNKEY SOLUTIONS
♦ NOVELTY ♦ TRUST ♦ SATISFACTION
ISO 9001:2015 & CE Certified Organisation

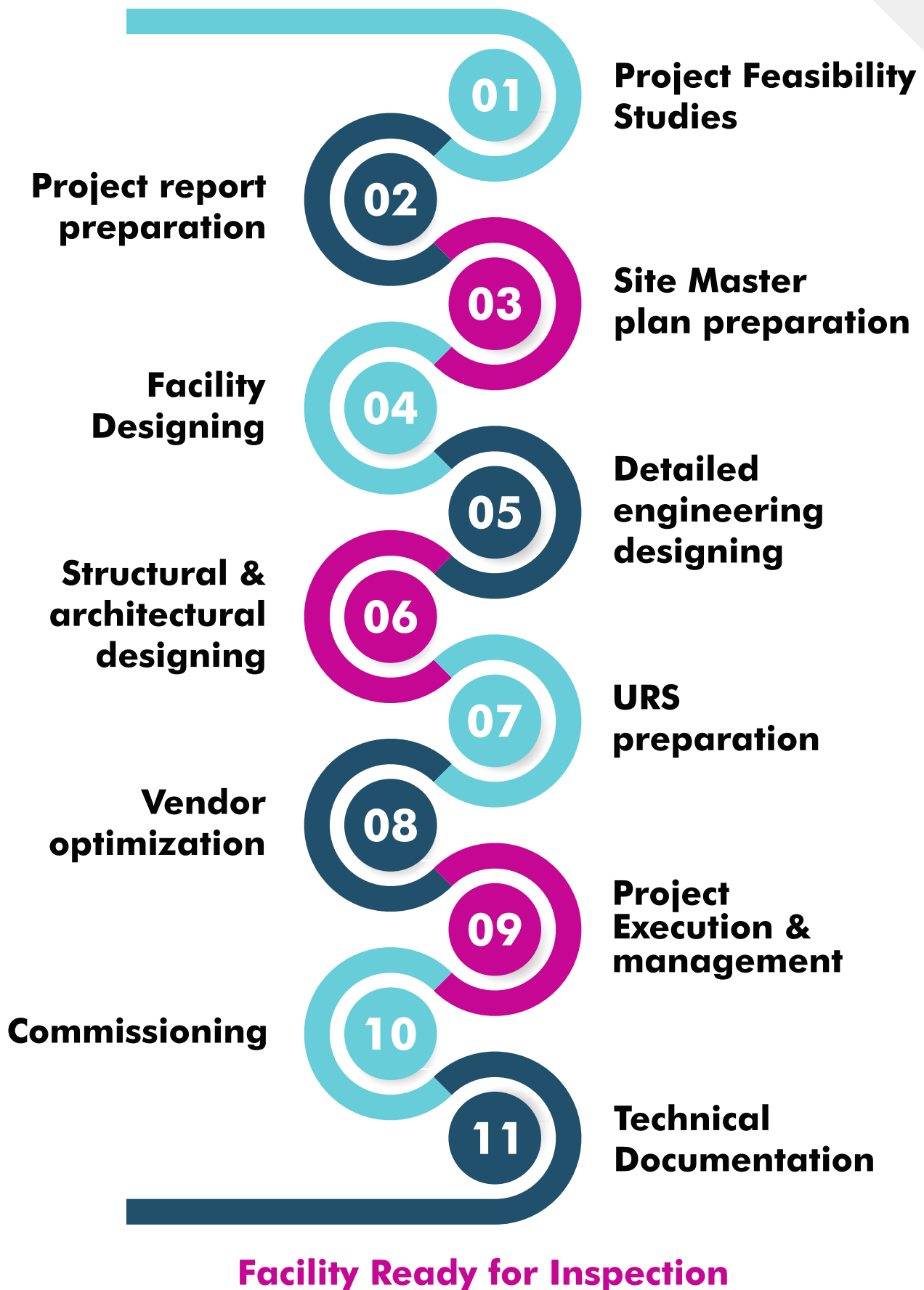
Turnkey Solutions



WHO WE ARE

- We are one stop solution for Pharmaceutical & Allied industries.
- We perform Turnkey project execution to render facilities compliant to Schedule- M, cGMP, WHO-GMP, PICS, EU-GMP, TGA, ANVISA, USFDA





WE PERFORM

PROJECT FEASIBILITY STUDIES

- Economic Feasibility
- Technical Feasibility
- Operational Feasibility
- Schedule Feasibility

PROJECT REPORT PREPARATION

Comprises of:

- Building construction details
- Supporting utilities details
- Plant equipments details
- Quality control instruments details
- Plant accessories details
- Technical requirement details
- Financial details

SITE MASTER PLAN PREPARATION

Comprises of:

- Key Plan
- Site Plan
- Building Orientation
- Utility Block Placement
- Administrative Arrangements

FACILITY DESIGNING

- Basic Technical Plan Layout
- Section and Elevation Drawings
- Area Table with detailed Containments
- Nomenclature with necessary approval

DETAILED ENGINEERING DESIGNING

Comprises of:

- Basic Design
- Technical Datasheet
- Bought Out Quantity list

STRUCTURAL & ARCHITECTURAL DESIGNING

Comprises of:

- Soil Evaluation
- Load Requirements
- Structural details
- Load Bearing Calculations
- Interior and Exterior 3D imposements
- Overall Elegance Management





URS PREPARATION

Comprises of:

- Required Standards of cGMP
- Output requirements
- Load Calculation
- Automation requirements
- Cost Effectiveness

VENDOR OPTIMIZATION

Comprises of:

- Vendor assessment with respect to technical capabilities, timelines, delivery schedules, maintenance, etc.
- URS compatibility
- Dynamic balance between quality and economy

PROJECT EXECUTION & MANAGEMENT

Comprises of:

- Project review based on design drawings
- Phase wise target completion
- Project flow direction
- Troubleshooting and streamline execution

COMMISSIONING

Comprises of:

- Facility Qualification
- Report preparation and review
- Finishing and elegance

TECHNICAL DOCUMENTATION

We perform documentation set up as to comply with Schedule M, cGMP, WHO-GMP, MHRA, PIC/S & USFDA standards.

OUR CORE TEAM



Mr. Jitendra Prajapati

Designation: Partner
Qualification : M.Pharm
(Quality Assurance)
Professional Experience : 15 years



Mr. Dharmesh Prajapati

Designation: Partner
Qualification : B.E
(Instrumentation & Control)
Professional Experience : 11 years



Mr. Jigar Patel

Designation: Partner
Qualification : B.E
(Instrumentation & Control)
Professional Experience : 10 years



SUPPORTING TEAM

DEPARTMENT	NO. OF PERSONNEL
Project Co – Ordinator	1
Civil draftsman	1
Site Engineer	1
Architectural Design	1
Structural Design	1
Electrical Design	1
HVAC Design	1
Utility Design	1
Technical Documentation	3
Computer Operator and Office Administration	3

OUR CLIENTELE

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 + NOVELTY + TRUST + SATISFACTION
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Pulse
 PHARMA MFG. CO.

asmed
 sharing excellence

Many More



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Corporate office & Manufacturing Workshop

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