# Vaccine Manufacturing Platform With Covid19 Vaccine Case Study

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## **Company Introduction**





#### Founded in 1993

- Headquartered in Shanghai  $\geq$
- Over 100,000 m<sup>2</sup> of professional  $\geq$ production workshop

#### Listed in 2011

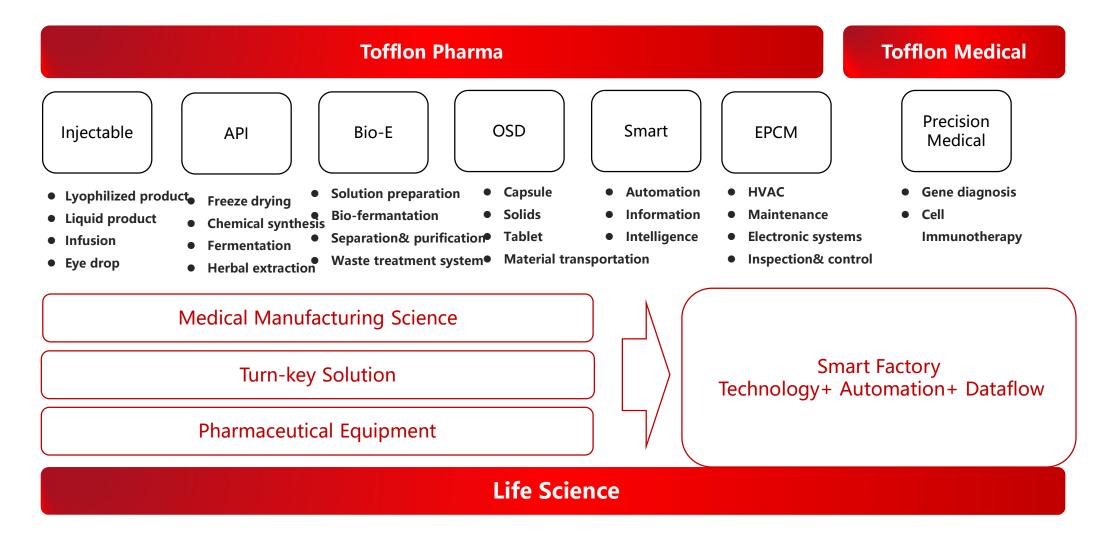
Canada

- GEM Listing Stock Code: 300171
- Over 2,000 professionals

#### **Globalization**

- > China's leading pharmaceutical equipment industry
- > International market share ranked top





Equipment Supplier to Solution provider



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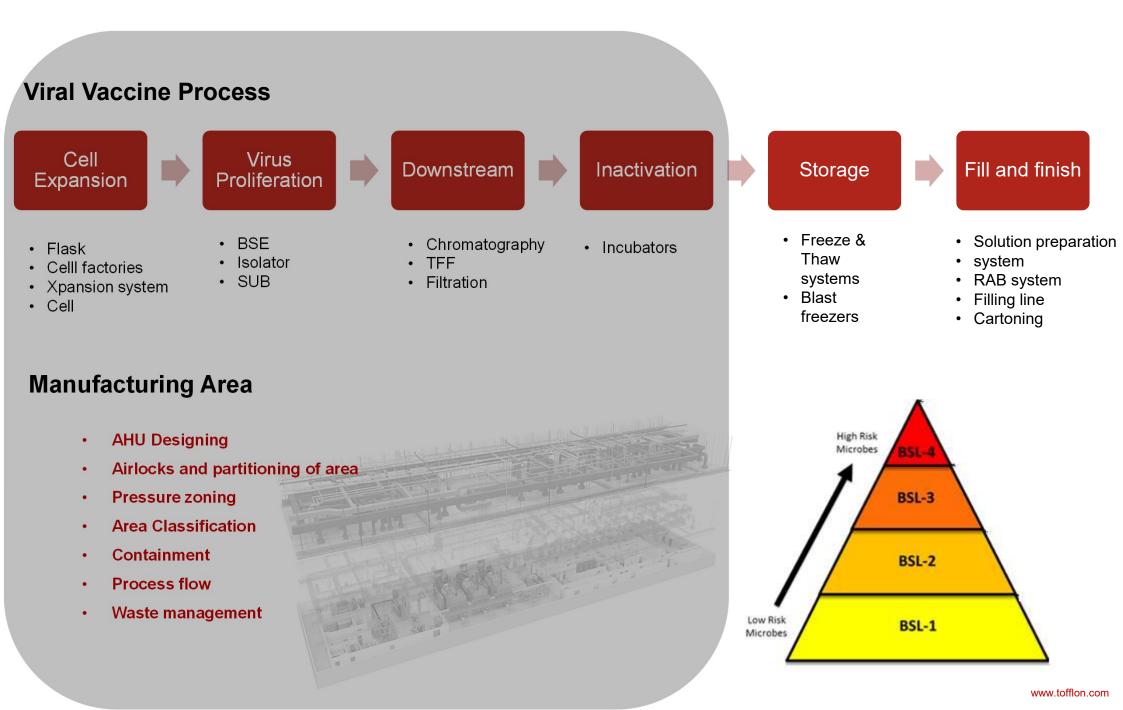
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# Tofflon For Vaccine Platform

### **Vaccine Manufacturing Platform**





### **Upstream Capabilities**







Bioreactors



Vessel System

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#### **Downstream Capabilities**



Chromatography column & Systems



ILC, ILD, Integrated Systems





Hollow Fiber systems



Vessel systems

### **Bioinactivation & Decontamination capabilities**







**VHP** Pass Box

VHP

- BSL3/BSL4 manufacturing area
- Sterile time:2 Hr for the passbox 6 logarithmic reduction killing effects of
- microorganism
- Efficient decomposition system to shorten the ventilation time to reduce H2O2conent to less than 1 ppm
- Robotic VHP

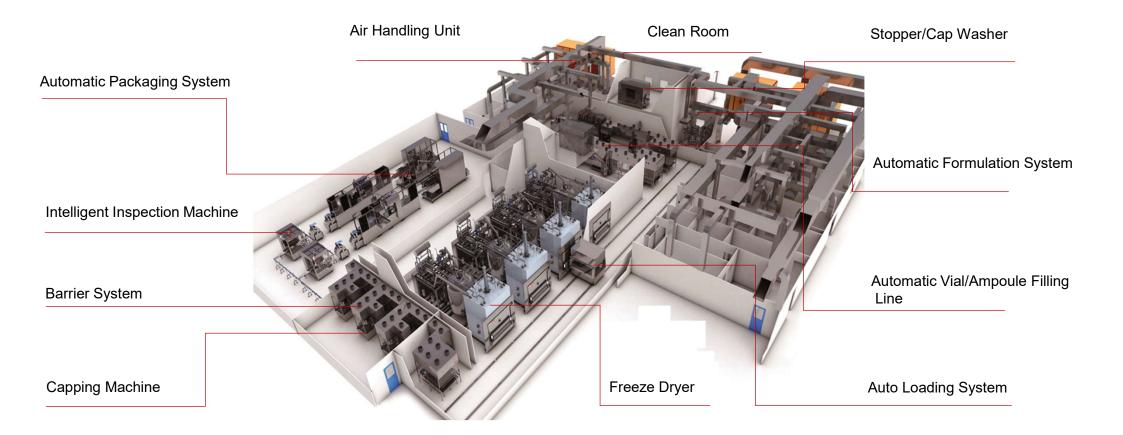


Continuous Bioinactivation system

- Efficient heat energy utilization
- The inactivation effect is the same as that of steam sterilization at 121.1 °Cfor 30 minutes
- Continuous operation
- Scada platform controlling and data acquisition
- Economic compared to batch

### **Injectable Capabilities**







Washing & Drying

Rubber Cleaning

Filling

Capping

Inspection

Packaging www.tofflon.com





## Case study 1 : Beijing National Institute of Biological Sciences Tofflon

Product : Inactivated Vius vaccine

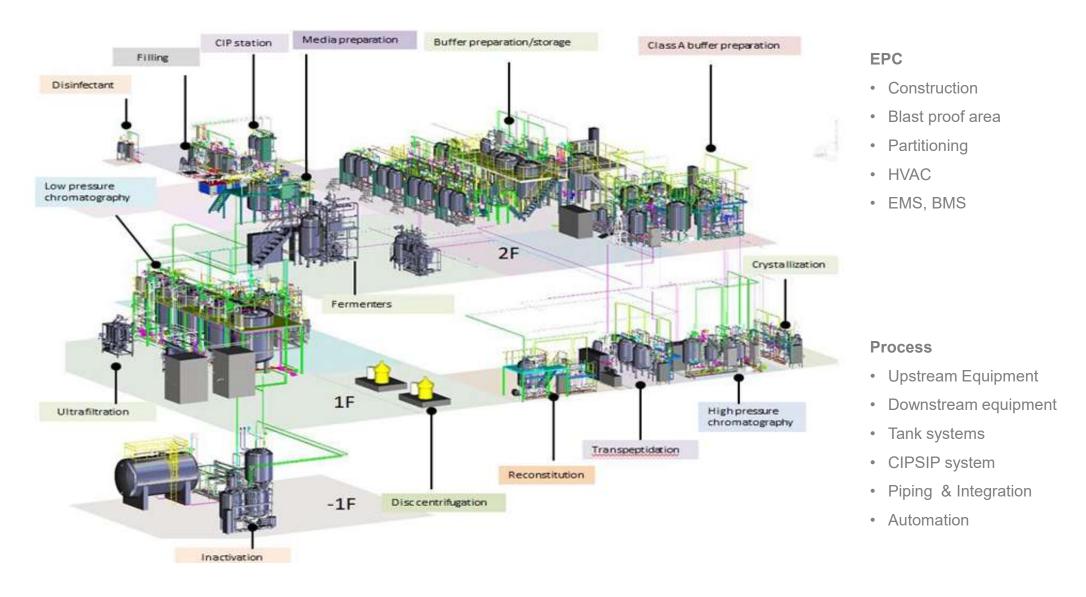
#### **Problem statement :**

Closed and contained operation for handling of virus



- In production for virus inoculation, virus distribution, harvesting fluid handling, in QC for QC purpose viz virus sterility inspection, mycoplasma inspection, titer and other experiments
- Isolator designed with incubator cart to transfer material from isolator to incubator through RPT port.
- Considered sterility requirements and toxicity protection requirements for isolator.
- The internal dimensions of the isolator chamber are reasonably customized according to the different equipment, which are involved in different process.
- Separate air supply and separate air exhaust for isolator to ensure that isolator has no effect on the control of the differential pressure in the laboratory.

#### Case study 2 : Jilin Jinsheng Pharmaceutical Co., Ltd. China



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# **Thank You**



