Abstract for IV Bag Production Contract Manufacturing

This abstract refers to contract manufacturing of intravenous
Our partner company is an industrial and commercial company that is “revolutionary” in its markets of references. It is innovative regarding products, careful about using quality materials and unique technological production process. The company is provided with sophisticated equipment and instrumentation. Their key feature process is an integrated combined production-filling cycle: the production of infusion bags and the filling of themselves with solution, then the final sterilization in autoclave. This innovation offers protection against a wide range of contamination and environmental risk. Moreover, all processes are performed by ISO standards 9001-2000 and 14001-2004.

Manufacturing
Due to a long manufacture experience and availability of technical data, such as stability studies, manufacturing process and batch validation, the company is involved in offering assistance to the clients in preparing of registration files in CTD format for submission to the authorities. The quality control laboratories are the core of the production line. Qualified and performed staffs (in conformity to the EU-GMP) check throughout the manufacturing process the product meets the quality specification by regulation, through physical, chemical and microbiological tests.

For packaging material our partner chooses the Polypropylene “Cryovac” that has good reputation for its quality and the primary packaging characteristics are successful in preparing sterile fluid. The Food and Drug Administration (FDA) approved this material suited perfectly for sterile pharmaceutical solutions. But the use of PVC for the infusion bags and of the glass bottles is also possible. Besides, it is possible to manufacture different size of the infusion bags, up to 5000 ml. After production and filling of the bags, it could be put in an over-bag that should exhibit an under-vacuum or not under-vacuum.
PHARMACEUTICAL CONTRACT MANUFACTURING SERVICES FOR IV BAGS AND IV BOTTLES

As a cGMP API manufacturer, our pharmaceutical contract manufacturing can support the development process of your IV business regarding the procurement of empty IV bags. All projects are supported by our chemistry expertise, GMP compliance and extensive quality control. Our partner currently manufactures sterile medical devices and sterile commercial products under GMP conditions. They have manufactured many products for clinical trials (Phase I, II and III).

Validation
In a regulated industry, special care must be taken to ensure all equipment and processes are performing to specifications. Deviations from the norm could have unpredictable and sometimes devastating results. Therefore, special attention is payed to equipment qualification and process validation to provide full regulatory compliance to the clients. Strictest guidelines are followed to perform the most rigorous testing and execute studies in a timely fashion to ensure your product is of the highest quality when it reaches its destination, has capabilities to develop, create and execute validation protocols and studies required for your products and equipment that are required for your projects in accordance with current Guidelines (HPFBI, FDA, EP, ICH, WHO) and acceptable formats (prospective, retrospective and concurrent). These include process, cleaning, aseptic and terminal sterilization processes, environmental control, IQ/OQ/PQ/PV and utility qualifications. Statistical and decision making analysis is also part of the service. We stand by our products which are manufactured to the highest quality standards. Please review our given product list.
# PRODUCT LIST

<table>
<thead>
<tr>
<th>For Human Use (PP/PVC)</th>
<th>For Human Use (Glass)</th>
<th>Medical Devices</th>
</tr>
</thead>
<tbody>
<tr>
<td>Water for Injection</td>
<td>Water for Injection</td>
<td>Destillate Water</td>
</tr>
<tr>
<td>Glucose 10%</td>
<td>Glucose and Sodium Chloride</td>
<td>Sodium Chloride Solution</td>
</tr>
<tr>
<td>Sodium Chloride 0,9%</td>
<td>Electrolytic Enteric Equilibrate</td>
<td>Sodium Bicarbonate Solution</td>
</tr>
<tr>
<td>Mannitol 5%, 10%, 18%, 20%</td>
<td>Electrolytic Gastric Equilibrate</td>
<td></td>
</tr>
<tr>
<td>Glucose</td>
<td></td>
<td>Hydratant Solution</td>
</tr>
<tr>
<td>Electrolytic Enteric Equilibrate</td>
<td>Maintenance Equilibrate Solution with Glucose 5%</td>
<td></td>
</tr>
<tr>
<td>Dextrose 5%, 10%, 25%, 33%</td>
<td></td>
<td>Glucose</td>
</tr>
</tbody>
</table>

We also contract manufacture every kind of SPECIAL IV solutions on demand and order.
ensymmm is a German based premier project consulting company for Life Sciences, serving biotech companies, pharmaceutical industry and food ingredient companies. We provide clients with a variety of business and technology consulting services as well as with specialized teams in various areas of our competence.

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