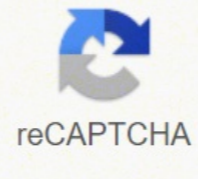


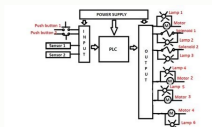
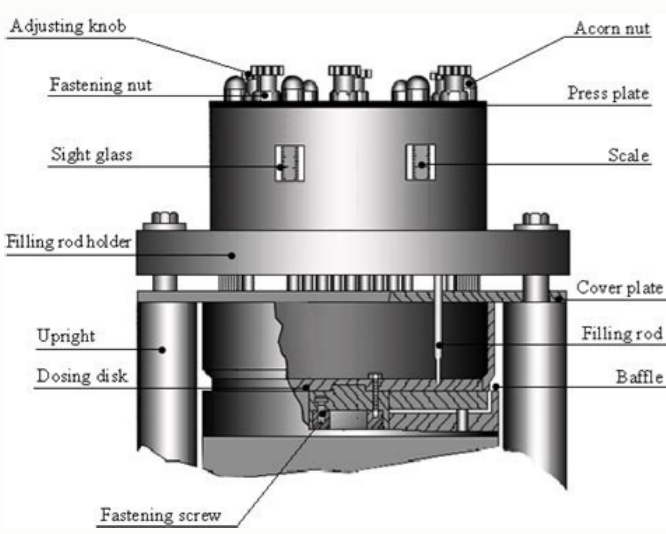


I'm not robot



Next

Ampoule filling machine pdf



Quality..... 5. Check weight mechanism of the machine helps to maintain consistency in each batch. Automatic Tablet Coater is designed for the application of: Film Coating with organic solvent, Enteric Coating and aqueous coating Sugar Coating 42. The use of the following labeling instructions is recommended: On the label Means "Do not store over 30°C" from +2°C to +30°C "Do not store over 25°C" from +2°C to +25°C "Do not store over 15°C" from +2°C to +15°C "Do not store below 8°C" from +2°C to +8°C "Do not store below 8°C" from +8°C to +25°C "Protect from moisture" no more than 60% relative humidity in normal storage conditions; to be provided to the patient in a moisture-resistant container. Foil is generally used as a component of the backing lamination if barrier protection is a critical requirement; however, metallized polyester is replacing foil for some barrier applications. 3. Our operations are driven by best-in-class technology and processes, abiding by all major stringent regulatory approvals. Report On Industrial Visit For Partial Fulfillment Of The Requirement For The Award Of Degree Of Bachelor of Pharmacy For Dr. A.P.J. Abdul Kalam University Through Under the supervision of Miss. The shrink wrap concept has a diversity of uses in packaging, one of which is its use as an over wrap An L-shaped sealer seals the remainder of the over wrap and trims off the excess film. DRY-POWDER INJECTABLE SECTION (VIAL) - CAPACITY - 12 MILLION VIALS PER ANNUM. These products are manufactured in accordance with highest international standards in facilities as per norms of WHO-GMP & ISO 9000-2008. > To see Brooks as an Institution for ethical & respectable world class Pharmaceutical company. 6.1.3-Colored Glass—Light Protection: Glass containers for drugs are generally available in clear flint or amber color. 5. Various inner seal compositions may be used, but the structures most frequently encountered are glassine and foil laminations. C) GENERAL SECTION: In general section we have injectable both Dry and Liquid. Assuring quality from the root upwards... If there is one attribute to which we attach the utmost importance it is quality. However, the desire to reduce the time needed for the sugar coating process has led to the trend towards side-vented pans. Soft and Hard gelatin capsules The basic difference between the hard and soft gelatin encapsulation processes is that in the hard gelatin capsule process, the capsule is prefabricated and supplied empty, whereas in the soft gelatin capsule process the encapsulation and filling take place simultaneously. No Composition Pack Size Indication ANTI-PSYCHOTICS 1 Amoxapine 50mg, 100mg 10's Endogenous depression 2 Buspirone HCl 5mg, 10mg 10's Anxiety disorder 3 Carbamazepine 200mg, 400mg 10's Epilepsy, Trigeminal Neuralgia 4 Chlorpromazine HCl 50mg, 100mg, 200mg, Trihexyphenidyl HCl 2mg. Research & Development (R&D) Helios's focused towards developing new products, improving existing products as well as drug delivery systems and expanding product applications. This lid is sealed to the base layer by heat, pressure or both. It's a journey, which is a wonderful experience. . ND RR240.pdf 5. 34. To achieve this, we follow a four-step process: Adopt Quality by Design (QbD) approach in Manufacturing and clearly identify sources of variability and minimize them on an ongoing basis. The following package configuration have been identified by the FDA as examples of packaging systems that are capable of meeting the requirements of tamper-resistant packaging as defined by FDA regulation 1. Film wrappers 2. 10. Packaging should assist in patient compliance. DRY-POWDER INJECTABLE SECTION CAPACITY - 12 Million Vials per Annum. 43. We tap every possible lead and source to add to our human capital. Improve efficiency. A strip package is formed by feeding two webs of a heat-sealable flexible film through either a heated crimping roller or a heated reciprocating plate. Packaging must also function as a means of drug administration. Storage areas should be designed or adapted to ensure good storage conditions. LIQUID INJECTABLE SECTION (VIALS) - CAPACITY - 5 MILLION VIALS PER ANNUM. Geneva, World Health Organization, 2002 (unpublished document QAS/01.014, available on request from Essential Drugs and Medicines Policy, World Health Organization, 1211 Geneva 27, Switzerland). Advantages: • Economical • Readily available container of variety of sizes and shapes • Impermeability • Strength and rigidity • Has FDA clearance • Does not deteriorate with age • Easy to clean • Effective closure and resolves are applicable. However, they may be adapted to meet individual needs where necessary, provided that the desired standards of quality are still achieved. It is also borne in mind that this best quality is provided at affordable costs to our valued customers... giving them products which adhere to the specifications and also meet the aspirations of the society. 24. With almost 5-6% of the company turnover being invested towards R&D each year, our strategy focuses on: Developing new drug formulations for existing and newer drug substances Improving processes for existing API and formulation products Developing new drug delivery systems for existing and newer active drug substances, as well as newer medical devices, mainly in the area of respiratory medicine Tie-ups with independent research teams to develop new products Strengthening our intellectual property, including the patenting of new products, drug delivery systems and medical devices, mainly in the area of respiratory medicine. Bubble pack 5. So process materials (product) are coated due to rotating pan. Some of them are: • Type B straight -stem • Type C funnel -tip • Type D closed 6.1.6.Bottles, vials and syringes: These are more or less thick walled containers with closures of glass or of material other than glass such as plastic materials or elastomers. Each site has well trained personnel for quality control along with a regulatory affairs department ensuring strict adherence to quality systems and procedures. T he risk is greatest with cellulosic substance and if the use of such materials is unavoidable, the attack may be minimized by impregnation. Prof. J. Miss Jyoti Srivastava (Asst. The tube holders are fitted on the circular turret which intermittently takes the tube to all the stations. The center is supported by high-tech instruments like HPLC, UV-spectrophotometer, stability chambers, Lyophilizers, osmolality meter and GC etc. - It bears appropriate label(s) providing content and usage information. Film coating techniques reduce process times, offer greater control over coating parameters and provide more opportunity for innovation. Student's Signature Date: 4. The contents may be removed in several proportions on one or more occasions. Pack Types Multidose/Reclosables Unit Dose/Non-reclosables Bulk Bottles Aerosol packs Tubes Ampoules Blister packs Prefilled syringes Vials Sachets Form/Blow-Fill-Seal (FFS, BFS) - various pack formats Bottles Drums/Kegs Sacks/Bags 20. Storage and labeling conditions Normal storage conditions Storage in dry, well-ventilated premises at temperatures of 15-25°C or, depending on climatic conditions, up to 30°C. We believe that we all together can drive pharma machinery market to next generation So. This is generally accomplished by heat-softening the plastic film and vacuum-drawing a pocket into the film in a manner similar to the formation of a blister in a blister package. For decorative purposes, special colors such as blue, emerald green, and opal may be obtained from the glass manufacturer. Quality is considered paramount at all locations where we conduct regulated research, development, manufacture, testing and distribution of pharmaceutical products. We are currently manufacturing for drug majors Dr Reddys, Sandoz, Nicholas Piramal, Cipla, Alembic, Glenmark, Unichem, and Cadila among others. Labeling and containers All materials and pharmaceutical products should be stored in containers which do not adversely affect the quality of the materials or products concerned, and which offer adequate protection from external influences. They are more rigid than strip packages and are not used for powders or semi-solids.. 4. During the rotation motion, coating material is sprayed by spraying system according to the technological process and rational technological process. Career..... Pellegrino Coating Pan Machine Capsule Manufacturing Process In pharmaceutical industry, there are various pharma company manufactures types of machineries for the different process and application as per there specialty. 12. We correlate their theory with practical application in pharma manufacturing field. This team is working in development of latest applications, a paper/polyethylene/foil/polyethylene lamination is commonly used. We insist that our business partners comply with national and international regulatory and business standards which are in alignment with those of our own. Finishing operations take place in clean rooms equipped with clean air supply, air curtains and dust handling/collection systems. The main functions of the primary package are to contain and to restrict any chemical, climatic or biological or occasionally mechanical hazards that may cause or lead to product deterioration. - A pack component with no product contact but may add protection to that provided by - The sum of packaging components that together contain and protect the dosage form. We are committed to being an employer of choice; offering the best performers a career path beating the normal line of the industry. PHOTO 5.2: END FOLDED WRAPPER 5.3.2-Shrink Wrapper: Film over wrapping can also be accomplished with the use of a shrink wrapper. The shrink wrap concept involves the packaging of a product in a thermoplastic film that has been stretched and oriented during its manufacture and that has the property of reverting back to its un-stretched dimension once the molecular structure is "unfrozen " by the application of heat. Manufacturing Helios leverages state-of-art manufacturing facilities at Baddi, conforming to WHO-GMP standards, to ensure quality products at all times. WHO Expert Committee on Specific Capions for Pharmaceutical Preparations. PHOTO 5.4: STRIP PACKAGING MACHINE 5.3.5- Bubble Pack: The bubble pack can be made in several ways but is usually formed by sandwiching the product between a thermoformable, extensible, or heat-shrinkable plastic film and a rigid backing material. Product Quality Over the years we have endeavored to set the standards for providing the highest quality products through the utilization of state of the art production techniques with an ultimate focus of satisfying the requirements and need of our customers. Loading and transport functions: Packaging has a crucial impact on the efficiency of transport, handling and storage of goods. 6.1.1.-Composition of Glass: Glass is composed principally of silica with varying amount of metal oxides, soda-ash, limestone, and cullet. 1.1.2-For solid dosage forms: Tamper - evident containers are closed containers fitted with a device that irreversibly breaks on opening. - Package should preferably have an aesthetically acceptable design. The contents are withdrawn after rupture of the glass, or a single occasion only. Thrust on improving healthcare delivery as well as pharmaceuticals manufacturing infrastructure by many states government. 2. Career Working with HELIOS PHARMACEUTICALS is not a job. Ex-Date Purpose 21-Sep-2015 : ANNUAL GENERAL MEETING 31-Jul-2014 : ANNUAL GENERAL MEETING 29-Aug-2013 : ANNUAL GENERAL MEETING 14-Sep-2012 : ANNUAL GENERAL MEETING / DIVIDEND - RS 9/- PER SHARE 19-Sep-2013 : ANNUAL GENERAL MEETING 11-Jun-2013 : INTERIM DIVIDEND RS 7.50/- PER EQUITY SHARE - PressRelease Oct23, 2015, 08:19 - TradingWindow - Oct 06, 2015, 14:24 - RecordDate Sep 23, 2015, 13:43 - Disclosure of VotingPattern - Clause 35A Aug 14, 2015, 09:56 - StockSplit Aug 12, 2015, 14:41 57. Bottle seals 8.Tape seals 9. About Company Brooks Laboratories Ltd. Market Secretary/Compliance Officer May 29, 2015, 18:33 - Code of Conduct under SEBI(PIT) Reg., 2015 May 20, 2015, 18:10 - TradingWindow May 19, 2015, 17:11 26. OUTPUT Model - A 40 40,000 capsules per hour for powder & 30,000 capsules per hour for pellets. Special scrapers are provided for transfer of the complete product and to avoid wastage. We establish uniform standards for all products, regardless of geography. Manufacturing facilities: The stability of the manufacturing facilities should be considered due to new package, increased sale, improvements in Good Manufacturing Practice, revised product, new product etc. Manufacturing Brooks manufactures standardized herbal ingredients for many countries around the world. Jyoti Srivastava (Asst. 6.1.2-Manufacture of Glass: Four basic processes are used in the production of glass: blowing, drawing, pressing, and casting. Optional Pellets filling station, Dozing format for pellet station. Jacketed bowls available for heating or cooling of products during mixing. The break system OPC(one -point cut) or the color break ring offer consistent breaking force. Mr. Manoj Kumar Prajapati , Mr. Alok giri, Miss Jyoti Srivastava. 16. With the support of our distinguished clientele we have made a mark as one of the leading contract manufacturing facilities in the country. The package is made of two layers of film or laminate material. 5)Packaging materials used in different formulations: 5.1-Paper and board: The use of paper/board materials (cellulose fiber) remains a significant part of pharmaceutical packaging in spite of the facts that paper is rarely used on its own for a primary package. 8. * : 1455.64(Cr) Impact Cost:0.09 as on Oct- 2015 52 weekhigh/low price : 453.30/271.02 Ex-Date Purpose 06-Oct-2015 : FACI VALUE SPLIT (SUB-DIVISION) - FROMRS 5/- PER SHARE TO RE 1/- PER SHARE 30-Jul-2015 : ANNUAL GENERAL MEETING/DIVIDEND - RS 12/- PER SHARE 17-Jul-2014 : ANNUAL GENERAL MEETING / DIVIDEND - RS 9/- PER SHARE 19-Sep-2013 : ANNUAL GENERAL MEETING 11-Jun-2013 : INTERIM DIVIDEND RS 7.50/- PER EQUITY SHARE - PressRelease Oct23, 2015, 08:19 - TradingWindow - Oct 06, 2015, 14:24 - RecordDate Sep 23, 2015, 13:43 - Disclosure of VotingPattern - Clause 35A Aug 14, 2015, 09:56 - StockSplit Aug 12, 2015, 14:41 57. Bottle seals 8.Tape seals 9. About Company Brooks Laboratories Ltd. Market Tracker..... HELIOS PHARMACEUTICALS 1. Be right the first time. It conforms to relevant Schedule M and W.H.O. GMP standards and the facility can be upgraded to US FDA norms in the future. 7. Sealing is done either by laser sealing system or conventional gas flame, are also implemented. LIQUID INJECTABLE SECTION (AMPOULES) - CAPACITY - 12 MILLION VIALS PER ANNUM. Introduction..... 2. Storage areas should be of sufficient capacity to allow the orderly storage of the various categories of materials and products, namely starting and packaging materials, intermediates, bulk and finished products, products in quarantine, and released, rejected, returned or recalled products. 5.3.7- Foil, Paper, or Plastic Pouches: The flexible pouch is a packaging concept capable of providing not only a package that is tamper-resistant, but also, by the proper selection of material, a package with a high degree of environmental protection. You have helped me in all possible ways to make this industrial visit report a reality. The shape, size and color are all visual characteristics reflected by the core and the coating such as sugar coat, film coat. Every aspect of manufacturing is performed in accordance with current Good Manufacturing Practices (cGMP) guidelines by experienced and trained personnel and is validated by Standard Operating Procedures (SOPs). QC, as a Centre of excellence, ensures compliance and follows systematic interventions like streamlining SOPs around critical quality parameters, bullet-proofing complex procedures and targeted capacity building. > To establish world class Research & Development infrastructure for new products & Drug delivery systems. It is provided with fully automatic lami tube filling line inbuilt with tube cleaning, filling, sealing and embossing stations integrated with online carbonator, online batch coding and check weighing system. Contract Manufacturing: Our focus on delivering quality products and customer service has attracted several notable Pharma giants for whom we are doing contract manufacturing. Work-life balance is our winning formula, ensuring effectiveness across all spheres of life. Disadvantages: • Fragility • Heavy weight 53. Good trade and distribution practice (TGD) of pharmaceutical's starting materials. This facility is having well garnished & well maintained lawns and greenery to add to the healthy eco-system of Himachal Pradesh. N.K MANNA Director Internal examiner External examiner 3. The product is to be filled inside the pan. The entire approach here is based on quality. Quality Quality has always been at the epicenter of the growth of an industry and the success of a company in a competitive world. Such as Pharma machines for Tablet filling, Machines for Tablet Counting, Machines for Tablet Pressing, and Machines for Tablet Labeling etc. The final extracts are then spray dried or vacuum dried. Storage and Packaging Packaging 1) Introduction: Packaging can be defined as an economical means of providing presentation, protection, identification information, containment, convenience and compliance for a product during storage, carriage, display and until the product is consumed. The packaging external to the primary package is known as the secondary packaging, whether it be to improve the life of the human being, extending the benefits to the members of the organisation or to the society at large, we believe in imparting the best possible. Product built mounted on castor wheels for easy portability, washing & transporting mixed materials. Film wrapping can be accomplished in several ways and varies in configuration with packaging equipment. PHOTO 5.1: FILM WRAPPER MACHINE 50. After aging in stainless steel receiving tanks, the gelatin solution is transferred to stainless steel feed tanks. A challenging and enjoyable work environment is provided which will help you take a leap in your professional, technical and personal growth. To maintain quality standards, every plant has well defined procedures and systems in place in compliance with the requirements of the current Good Manufacturing Practices (GMP). WHO, PIC's and EU GMP in order to ensure that our operating procedures meet the very exacting standards of regulators like the US FDA, EMA, HC, WHO and TGA, among others. Defined storage instructions Drug products that must be stored under defined conditions require appropriate storage instructions. 3.4- Biological properties: The material of the container must be able to withstand attack by insects if this hazard is likely to be encountered. 6.1.4-Glass for Drugs: The USP and NF describe the various types of glass and provide the powdered glass and water attack tests for evaluating the chemical resistance of glass. * : 40.7(Cr) Impact Cost: 1.21 as on Oct-2015 52 weekhigh/low price : 106.80/36.05 * Free-float market capitalization as on the previous tradingday. Transfer from storage vessel to the filling hoppers is achieved by means of reciprocating metering pumps at the required rate. of Brooks is professionally managed. Capsule fillers are used to fill hard gelatin and non gelatin capsules with pre determined quantity of liquids, powders, pellets, tablets. Entire Dept. This can occur with certain combination of dissimilar materials. We have focused ourselves in developing a world class team to develop new molecules in injectable and clavulanic acid based products supported by Sophisticated Infrastructure for Research & Development. Thus, our passion for quality goes beyond business and statutory requirements. Blister packages are composed of a base layer, with cavities called blisters which contain the pharmaceutical product, and a lid. No. Name of the drugs/items Strength 1 Hydrocotrisone Sodium Succinate Injection 100 mg 2 Methylprednisolone Injection 40 mg 3 Methylprednisolone Injection 500 mg GENERAL INJECTABLE (LIQUID) S. If the container is to be opened on more than one occasion it must remain airtight after re closure. Today, Helios Pharmaceutical's is a mid-sized company, but it's you who will be helping it grow by leaps and bounds. The packaging is thus intended to protect the goods from loss, damage and theft. Storage conditions for pharmaceutical products and materials should be in compliance with the labelling, which is based on the results of stability testing (see Appendix). 5.2- Rubber based components: Rubber components may be made from either natural or synthetic sources. We believe that our people are our most valuable assets for us. References 1. Helios is a fully integrated, global healthcare provider, with strengths all along the pharmaceutical value chain. In this type of glass a substantial part of the alkali and earth cations are replaced by boron and/or aluminum and zinc. Child Resistant Containers, commonly referred to as CRC's, are designed to prevent the child accessing the potentially hazardous product.2.1.1.3-Containers for semi solid and pressurized products: Semi solid dosage forms like ointments, creams etc are packed in metallic collapsible tubes. Jyoti Srivastava 79 (Asst. Pharmaceutical ampoules are most commonly used to contain pharmaceutical hypodermic solutions. ResourcesandInfrastructure Advanced, sophisticated machineries and instruments State-of-the-art manufacturing facilities and laboratories Competent reporting team to support and focus on key functions Comprehensive quality policies and procedures Competent, enthusiastic and dedicated staff Continual monitoring by dedicated quality assurance and self-inspection team Controlled computerized systems Quality Control o Actual manufacturing process o Written definition or policy o Corporate pathways o Authority o Product standards 35. B) CEPHALOSPORINS: In cephalosporin's section we have dry injectable powder facility. Documentation such as SOPs, calibration and validation records are maintained assiduously and all documented procedures are strictly followed at all levels. The surface must be capable of clear labeling, often difficult, for example, with plastics. Contact for Complete range of Tableting Pharmaceutical machineries such as Rotary Tableting Machine, Granulator, Blister Packing Machine, Drying Oven, Coating Machine, Mult Mill, Mass Mixer, Capsule Filling Machine, Sieving, Grading & Straining, Rotary Piston Surface & Sealing Machine etc. Marketing Helios Pharmaceuticals is a division of PKTP Pvt.Ltd. Growth in opportunities for medical tourism Low-cost production and R&D Highly skilled workforce with significant expertise in chemical synthesis World-class facilities at national laboratories specializing in process and cost-effective technology development Increasing international trade in the pharmaceuticals sector Cost-effective source for generic drugs, especially, for those going off patent A separate QA team handles all in-process quality controls. Capsule quality is monitored throughout the production process including size, moisture content, single wall thickness, and color. MARKET PREPARATION PRODUCTS We have wide range of products catering to critical care segment in Parental Section like Beta Lactam, Cephalosporin & General dry powder Injectable, Ampoules and Liquid vial. The packing should not support mould growth. Prof. T. 5.3.8- Bottle Seals: A bottle may be made tamper-resister by bonding an inner seal to the rim of the bottle in such a way that access to the product can only be attained by irreparably destroying the seal. The capacities are as under: We have a well-equipped Quality Control / Quality Assurance department with sophisticated analytical instruments. 47. It is the totality of the arrangements made with the object of ensuring that pharmaceutical products are of the quality required for their intended use. Natural rubber has got good resealing (multi-dose injection), fragmentation and coring(description for the means by which particles are created when a needle is passed through a rubber) 5.3-Tamper resistant packaging: The requirement for tamper -resistant packaging is now one of the major consideration in the development of packaging for pharmaceutical products As defined by the FDA "a tamper -resistant package is one having an indicator or barrier to entry which, if breached or trussing ,can reasonably be expected to provide visible evidence to consumers that tampering has occurred Tamper -container (carton systems or any combination thereof intended to provide a visual indication of package integrity when handled in a reasonable manner during manufacture , distribution , and retail display " . Rather, it is the result of a well-conceived, rigorously implemented Quality Monitoring System (QMS). HELIOS PHARMACEUTICALS About Us HELIOS, is one of the well-reputed manufacturer and marketer of Ethical & Generic pharmaceutical products and also a leading contract manufacturer. BETA LACTUM S. Prof.) By PRAVEEN KUMAR SINGH (Roll no. We remain committed to the highest levels of quality and will ensure that all our facilities, offices and legal entities continue to meet the exemplary standards that are expected of a global pharmaceutical company. 3 Doripenem Injection 250 mg/500 mg 18. Brooks Laboratories LtdDivision believes in the spirit of "growing together". Breakable caps 10. Date: (Praveen Kumar Singh) Place: Varanasi 5. ACKNOWLEDGEMENT As I begin to reflect on the magnitude of this project, I am reminded of the kindness, support and affection offered to me by people to whom I am overwhelmed. Ointment Manufacturing Plant: Operations All waxes and oils are dissolved in the wax phase vessel separately. All aqueous phase materials are added in the water phase vessel and processed separately Both phase vessels are jacketed and are provided with motor driven propeller type of agitators which facilitate thorough mixing Once the phases are ready, they are transferred to the main Ointment manufacturing vessel by opening respective valves. 6.1.4.2-Type II—Treated Soda-Lime Glass: When glassware is stored for several months, especially in a damp atmosphere or with extreme temperature variations, the wetting of the surface by condensed moisture (condensation) results in salts being dissolved out of the glass. Stock rotation and control Periodic stock reconciliation should be performed by comparing the actual and recorded stocks. Cartons are used for a high 49. Our commitment to implementing a robust global quality management system is based on our determination to sustain a culture of operational excellence, meeting and exceeding the expectations of all stakeholders, including patients, customers and regulators. 6.1.5-Ampoules: Ampoules are thin-walled glass containers, which after filling, are sealed by either tip sealing or pull sealing. PHOTO 5.6: SHRINK TUBING 5.3.6- Shrink Banding: The shrink band concept makes use of the heat-shrinking characteristics of a stretch-oriented polymer, usually PVC. Capsules are sorted and visually inspected. Receipt of incoming materials and pharmaceutical containers On receipt, each incoming delivery should be checked against the relevant purchase order and each container physically verified, e.g. by the label description, batch number, type of material or pharmaceutical product and quantity. Packaging also serves as a mean to identify the manufacturer of the product. No. Name of the drugs/items Strength 1 Amoxicillin & Potassium Clavulanate Tablets 228.5 mg /375 mg /625 mg / 1 gm. - Combination of primary and secondary packaging, whether or not the latter has any overt stability maintenance function. DECLARATION I hereby declare that this report entitled "Project report on Industrial Visit" is a work carried out by me under the guidance of Assistant Professor, Miss Jyoti Srivastava Kashi Institute of Pharmacy, Mirzamurad, Varanasi. This is called "blooming" or "weathering," and in its early stages, it gives the appearance of fine crystals on the glass. We look the people who are result oriented and take ownership for the work. Commitment focusing on Learning, Delivering and Growing and ensure speed in action to have timely results. In the Automatic section, capsule halves are individually stripped from the Pins. "Protect from light" to be provided to the patient in a light-resistant container. Self-reliance displayed by the production of 70% of bulk drugs required and almost the entire Formulations requirement within the country. Our unwavering commitment to quality goes beyond ourselves. The cap and body lengths are precisely trimmed to a ±0.15 mm tolerance. 1. 2.1-Function of packaging: The various functions of packaging are: • Protective function • Storage function • Loading & Transport functions • Identification 48. We have our own

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