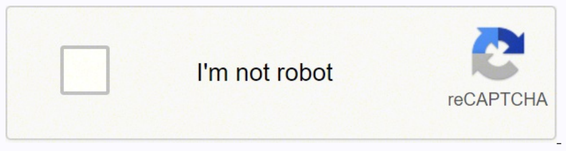


I am not robot!

Batch manufacturing record format for syrup

You need information in an easy -to -understand, visual format. A party production entry is a document designed to provide a detailed record of the product's party production history. This terminology is widely used in the pharmaceutical and chemical industry and refers to the requirements of many pharmaceuticals and food regulating agencies. The US Food and Drug Administration defines a party as a certain amount of medicinal product or other substance, which must be within the limits of the same nature and quality, which must be produced by one production order during a single production cycle. Batch production accounting.

COMPANY NAME	BATCH MANUFACTURING RECORD			Page: 1 of 8
Department : **Production**	Title : **Tongkat Ali Tablet**			Batch Record : BMR-001
	Name	Signature	Date	Revision No. : 0
Prepared by :	_____ **Production Manager**			Effective Date : 1 January 2016
Approved by :	_____ **QA Manager**			

1. Product Details	
Description	Tongkat Ali 250mg Tablet Colour: Pale Shape: Round/ Biconvex
Batch Quantity	Batch size: 53 kg Approx No. tablets: _____
Packaging	Bottle of 60's
Storage Conditions	Ambient - conditions, store in tight container protected from light and moisture

2. Production Batch Record Issuance		
Issued By – Issuer has reviewed the Batch Record to ensure that the copy is a complete, accurate copy of the Master Batch Record.		
_____ (Print) Issued By – Quality Assurance	_____ Signature	_____ Date
Issued To – Production has reviewed the Batch Record to ensure that the copy is a complete and correct. Production is responsible for the Batch Record following issuance.		
_____ (Print) Issued By – Quality Assurance	_____ Signature	_____ Date

3. Reference Documents
SOP – 002: Non conformances
SOP – 007: Line Clearance
SOP – 010: Temperature and Humidity Monitoring
SOP – 012: Facility Cleaning Procedures
SOP – 015: Material Weighing and Dispensing
SOP – 017: Batch Manufacturing Records
SOP – 019: Tray Oven
SOP – 021: Grinding and Milling Machine
SOP – 022: Cadmill
SOP – 023: Cube Mixer

Batch No: **TT 1606001**	Manufacturing Date : **10 July 2016**	Expiry Date : **9 July 2017**

The US Food and Drug Administration defines a party as a certain amount of medicinal product or other substance, which must be within the limits of the same nature and quality, which must be produced by one production order during a single production cycle. Batch production accounting. Party production accounting is important documents to ensure compliance with quality and regulatory requirements. They usually contain information on the following aspects of product batch production: start and end dates. Leave all the materials used and the amount of each use. Production process and deadlines. Initials of the person responsible at each stage information on all ongoing tests and their results. Link to any equipment used. Productivity and reconciliation. In most cases, serial production records are recorded in the instruction format with fields where the operator can enter process information. It is very important on request to provide information in party production reports. Some critical operations, eg. When baking the raw materials, the other person should check the calculations and the identity of the material and sign the documents for the production of the party. Each party has an individual number recorded in the party production journal. Party production records must be: respectively.Active.Sriginal.The signed. All corrections and deviations must be registered and signed for the product batch.All the information you need is delivered in a simple visual format. The component manufacturing log is a document designed to ensure the complete record of the product of the product.
Terminology is widely used in the pharmaceutical and chemical industry and is rejected in the requirements of many medicines and food regulating agencies. The United States Food and Second Administration defines the part as \ xe2 \ x80 \ x9Ca, the amount of medicinal product or other material with the same character and quality within a certain limits and according to one production order during production order. From one production order the same production cycle \ xe2 \ x80 \ x9D. Party production records.

COMPANY NAME	BATCH MANUFACTURING RECORD			Page: 2 of 8
Department : **Production**	Title : **Tongkat Ali Tablet**			Batch Record : BMR-001
	Name	Signature	Date	Revision No. : 0
Prepared by :	_____ **Production Manager**			Effective Date : 1 January 2016
Approved by :	_____ **QA Manager**			

SOP – 028: Ribbon Mixer
SOP – 032: B2 Strokes Tablet Press

4. Raw Materials

Description	Part Number	Quantity Required (kg)	Lot No.	Qty Staged	Exp/ Retest	Performed By / Date	Verified By / Date
Eurycoma Longifolia	R-0122	25.00					
Lactose Monohydrate	R-2323	19.34					
Gelatin	R-7896	4.80					
Corn Starch	R-5858	2.40					
Methocel	R-0326	1.00					
Magnesium Stearate BP/USP	R-9696	0.46					

5. Processing Equipments

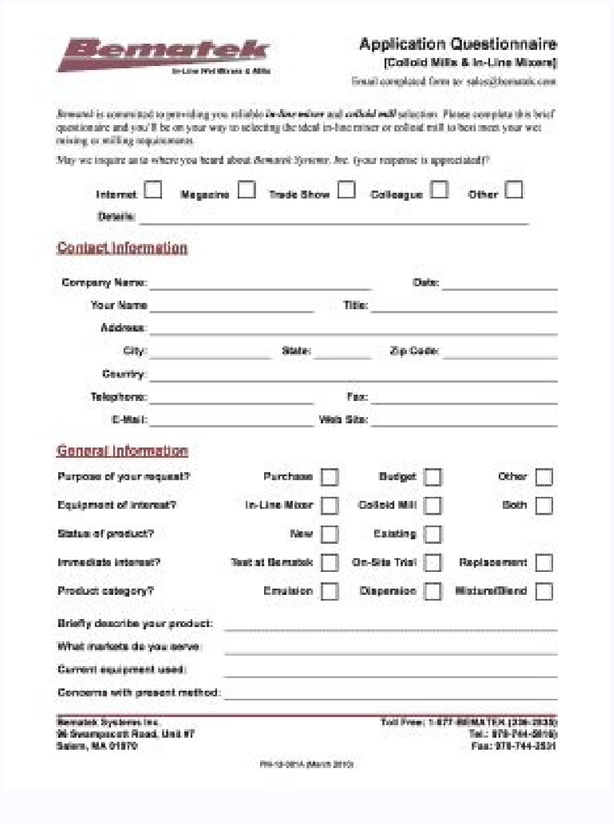
Equipment Description	ID No.	Previous Calibration	Calibration Required	Performed By / Date	Verified By / Date
Weighing Balance 150 kg	WB-01				
Tray Oven	OT-01				
Grinding and Milling Machine	GM-01				
Cube Mixer	MX-03				
Ribbon Mixer	MX-02				
Cadmill	GM-02				
B2 Strokes Tablet Press	TP-01				
Stainless Steel Container	CS-03				
Mechanical Sieve (Mesh No: 12)	SM-01				

6. Area Clearance

Batch No: **TT 1606001**	Manufacturing Date : **10 July 2016**	Expiry Date : **9 July 2017**

Link to any equipment used. Productivity and reconciliation. In most cases, serial production records are recorded in the instruction format with fields where the operator can enter process information. It is very important on request to provide information in party production reports. Some critical operations, eg.

Link to any equipment used. Productivity and reconciliation. In most cases, serial production records are recorded in the instruction format with fields where the operator can enter process information. It is very important on request to provide information in party production reports. Some critical operations, eg. When baking the raw materials, the other person should check the calculations and the identity of the material and sign the documents for the production of the party.



Bematek

In-Line Mill Mixers & Mills

Application Questionnaire

[Colloid Mills & In-Line Mixers]

Email completed form to: sales@bematek.com

Bematek is committed to providing you reliable ***in-line mixer*** and ***colloid mill*** selection. Please complete this brief questionnaire and you'll be on your way to selecting the ideal in-line mixer or colloid mill to best meet your wet mixing or milling requirements.

May we inquire as to where you heard about *Bematek Systems, Inc.* (your response is appreciated)?

Internet ☐ Magazine ☐ Trade Show ☐ Colleague ☐ Other ☐

Details: _____

Contact Information

Company Name: _____ Date: _____

Your Name: _____ Title: _____

Address: _____

City: _____ State: _____ Zip Code: _____

Country: _____

Telephone: _____ Fax: _____

E-Mail: _____ Web Site: _____

General Information

Purpose of your request?	Purchase ☐	Budget ☐	Other ☐
Equipment of interest?	In-Line Mixer ☐	Colloid Mill ☐	Both ☐
Status of product?	New ☐	Existing ☐	
Immediate interest?	Test at Bematek ☐	On-Site Trial ☐	Replacement ☐
Product category?	Emulsion ☐	Dispersion ☐	Mixture/Blend ☐

Briefly describe your product: _____

What markets do you serve: _____

Current equipment used: _____

Concerns with present method: _____

Bematek Systems Inc.
96 Swampscott Road, Unit 97
Salem, MA 01970

Toll Free: 1-877-BEMATEK (236-2835)
Tel.: 978-744-5816
Fax: 978-744-3531

PH-12-001A (March 2012)

Party production accounting is important documents to ensure compliance with quality and regulatory requirements. They usually contain information on the following aspects of product batch production: start and end dates. Leave all the materials used and the amount of each use. Production process and deadlines. Initials of the person responsible at each stage information on all ongoing tests and their results. Link to any equipment used. Productivity and reconciliation. In most cases, serial production records are recorded in the instruction format with fields where the operator can enter process information. It is very important on request to provide information in party production reports. Some critical operations, eg. When baking the raw materials, the other person should check the calculations and the identity of the material and sign the documents for the production of the party. Each party has an individual number recorded in the party production journal. 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They usually contain information on the following aspects of manufacturing: start and completion of production. Production process and production times. Initiation of a responsible person at each stage. Indicators and results of all ongoing tests. Get all used equipment. Performance and reconciliation. In many cases, serial production logs are stored in class format with areas where the operator can enter information for processing. It is very important to specify the details of the manufacturing record that should be requested. In some critical operations, eg. B. The raw materials, the other person must check the calculations and the identity of the materials and write them in the batch production records. Each detail has an individual number that is stored in the details in the production recording. Detailed documents should be: kidney.Present.precise.Foriginal.Part. All corrections and deviations must be registered and signedSupplement to the evening supplementary nutritional supplement; (ii) any presentation of supplemented food supplements that you distribute to another person to wrap or overlook it; (b) identity of production equipment and part of recycling parts; (c) Date and time of maintenance, cleaning and disinfection of equipment and transformation lines used in the section or link to records such as: B. Lognizal device if this information is stored; (d) a unique identifier that you assigned to each component (or, if applicable, to a product you received from the packaging or labeling as a food supplement), for packaging and using the label used; (e) identity and weight or measurement of each component used; (f) a declaration of actual yield and percentage of theoretical revenues in the corresponding recycling phases; (g) the actual results obtained during each supervision operation; (h) the results of any test or examination performed in the game or relate to these results; (i) documents confirming that the finished food supplement corresponds to the specifications laid down in accordance with Parties 111.70 (a). E) and g); (j) the manufacturing documents of the party consisting of: 1) Date of design at each stage of the main production file; and (2) the initials of the person, in particular: (i) the initials of the person responsible for considering or measurement of the component used in each batch; (ii) the initials of the person responsible for the weight or dimensions of the person responsible for each batch; (iii) the initials of the person responsible for introducing part of the party; and (iv) initials of a person responsible for adding components to the party; K) Operation marking of packaging and marking, including: (1) Unique i.Republishing or repackaging with dietary supplements) or exceeding the physical location of these results; (l) Documents during implementation that quality control personnel: (1) have reviewed the batch production record, including: (i) all monitoring activities required under Subpart E; and (ii) all testing and research, including testing and testing of components, manufacturing materials, finished dietary supplements and packaged and labeled dietary supplements; 2) has approved or rejected any processing or repackaging; 3) the shipment is approved and issued or rejected, including all processed parties; 4) approved and released or rejected packaged and labeled dietary supplements, including all repackaged or happy dietary supplements.
(m) Documents submitted in all necessary reviews and major removal decisions. n) all processing documents. Principles of Quality Management Quality Standards and Specifications Design Quality – Products and Processes Good Manufacturing Practices (GMP) CAPA – Management and Control of Corrective and Preventive Operational Changes etc. Etc.... Etc. Quality Assurance Quality Management Systems >>> >>>