W. B. PHARMACEUTICAL & PHYTOCHEMICAL DEV. CORPN LTD.

(A Government of West Bengal Undertaking)

ACHARYA P. C. RAY R & D DRUG TESTING & POLLUTION TESTING LABORATORY

Behala Chowrasta, 620 Diamond Harbour Road, Calcutta 34, West Bengal

Report of Analysis [Form 39, See Rule 150 (f)]

License No: DL.NO.T/8-DCH dt: 27.10.80. & T/13-DCH dt: 24.9.90

Dt of Receipt: 15.12.00

3374/07 2

Sample Receiving No:D/2155/00	Dt of Receipt: 15.12.00
I Name, address and license No. of Manufacturer /	Adviser (Health), State Urban Development Agency, llgus
Supplier's from whom sample received	Bhawan, II-C Block, Sec-III, Bidhan Nagar, Cal-91,
2.Reference No. and date of Mapufacturer / Supplier's	SUDA-120/96 (Pt-III)/633 dt. 24.11.00
letter under which the sample was forwarded	
3.Name of Drug / Cosmetics ' Raw material purporting to	Mebendazole tablet I.P. 100mg
be contained in the sample	
4. Detail of Raw material fight product in bulk / final	50 tabs
product (in finished pack) as obtained from the	
manufacturer for analysis	
(a) Original manufacturer's name (in case of raw	M/s. Kansas Laboratories (P) Ltd
materials and drugs repacked)	8/1 , Lal bazar St. Cal-1
(b) Batch Number	H 1427
(c) Batch Size	Nil
(d) Date of Manufacture if any	08/00
(e) Date of Expiry if any	07/03
5. Result of Analysis Recorded Below:	

Method : I.P' 96, VoL-II.

Description :-Buff coloured round shaped tablet having a bisecting mark on one side.

Avg. wt. of a tablet	0.3702gm	
Weight variation	Passes	
Identification	Positive	
Disintegration time	5 minutes	
Related substances	Passes	

Assay:--

Mebendazole I.P.

Found tab 97.05mg <u>Claint/tab</u> 100.00mg

In the opinion of the undersigned, the sample referred to above is of standard quality / is not of standard quality as defined in the Act and Rules made thereunder for the reasons given below ...

for Analyst

Signature of Office? In Charge

Sample Receiving No:D/2155/00

W. B. PHARMACEUTICAL & PHYTOCHEMICAL DEV. CORPN LTD.

(A Government of West Bengal Undertaking)

ACHARYA P. C. RAY R & D DRUG TESTING & POLLUTION TESTING LABORATORY

Behala Chowrasta, 620 Diamond Harbour Road, Caloutta 34, West Bengal

Report of Analysis [Form 39, See Rule 150 (f)]

License No: DL.NO.T/8-DCH dt: 27.10.80. & T/13-DCH dt: 24.9.90

Dt of Receipt: 15.12.00

3374/07 3

<u></u>	Ch of Receipt. 15.12.00
I Name, address and license No. of Manufacturer /	Adviser (Health), State Urban Development Agency, ligus
Supplier's from whom sample received	Bhawan, II-C Block, Sec-III, Bidhan Nagar, Cal-91,
2.Reference No. and date of Manufacturer / Supplier's	SUDA-120/96 (Pt-III)/633 dt. 24.11.00
letter under which the sample was forwarded	
3.Name of Drug / Cosmetics ' Raw material purporting to	Mebendazole tablet I.P. 100mg
be contained in the sample.	e de la construcción de la constru
4. Detail of Raw material final product in bulk / final	50 tabs
product (in finished pack) as obtained from the	
manufacturer for analysis	
(a) Original manufacturer's name (in case of raw	M/s. Kansas Laboratories (P) Ltd
materials and drugs repacked).	8/1 , Lal bazar St. Cal-1
(b) Batch Number	_H 1427
(c) Batch Size	Nil
(d) Date of Manufacture if any	08/00
(e) Date of Expiry if any	07/03
5. Result of Analysis Recorded Below:	

Method : I.P' 96, VoL-II.

Description :- Buff coloured round shaped tablet having a bisecting mark on one side.

Avg. wt. of a tablet	0.3702gm	
Weight variation	Passes	
Identification	Positive	
Disintegration time	5 minutes	
Related substances	Passes	

Assay:-

Mebendazoie I.P.

Found tab 97.05mg

<u>Claim/tab</u> 100.00mg

In the opinion of the undersigned, the sample referred to above is of standard quality / is not of standard quality as defined in the Act and Rules made thereunder for the reasons given below ...

for Analyst

Signature of Officer In-Charge

A. 3375/01.

W. B. PHARMACEUTICAL & PHYTOCHEMICAL DEV. CORPN LTD.

(A Government of West Bengal Undertaking)

ACHARYA P C RAY R & D DRUG TESTING & POLLUTION TESTING LABORATORY

Beliala Chowrasta, o20 Diamond Harbour Road, Calcutta 34, West Bengal

Report of Analysis [Form 39, See Rule 150 (f)]

License No. DL NO.T-8-DCH dr: 27 10 80. & T/13-DCH dr: 24.9.90

Dt of Receipt: 26.12.00

Sample Receiving NotD/2358/00	Dt of Receipt: 26.12.00
i Name address and license No. of Manufacturer	Adviser (Health), State Urban Development Agency, ligus
Supplier's from whom sample received.	Bhawan, H-C Block, Sec-III, Bidhan Nagar, Cal-91.
2.Reference No. and date of Manufacturer / Supplier's	SUDA-120/96 (Pt-III)/ dt. 13.12.00
letter under which the sample was forwarded.	
3 Name of Drug / Cospeties ' Raw meterial purporting to	Aspirin Tablet 1.P.
be contained in the sample	
4. Detail of Raw material final product in bulk / final	2 X 50 tabs
product (in finished pack) as obtained from the	
manufacturer for analysis	
(a) Original manufacturer's name (in case of raw	Mrs. Kansas Labs (P) Ltd
materials and drugs repacked)	8/1, Lal Bazar St. Cal-1
(b) Batch Number	K 1477
(c) Batch Size	Nil
(d) Date of Manufacture if any	11/00
(e) Date of Expiry if any.	04/02

5 Result of Analysis Recorded Below:

Method : I.P' 96, VoL-II.

Description :-White round oval shaped tablet.

Avg. wt. of a tablet	0.3.41 tgm
Weight variation	Passes
Identification	Positive
Disintegration time	1 minutes
Salicylic Acid	Passes

Assav ...

Aspirin I P

Found tab SIDE V SUE Claimab 300 00mg

In the opinion of the undersigned, the sample referred to above is of standard quality Lis not of standard quality as defined in the Act and Rules made thereunder for the reasons given below with respect of the above tests only.

ior Analyst

Signature of

Sample Receiving No:D/2358/00

A1 33 75/01.

W. B. PHARMACEUTICAL & PHYTOCHEMICAL DEV. CORPN LTD.

(A Government of West Bengal Undertaking)

ACHARYA P. C. RAY R & D DRUG TESTING & POLLUTION TESTING LABORATORY

Behala Chowrasta, 620 Diamond Harbour Road, Calcutta 34, West Bengal

Report of Analysis [Form 39, See Rule 150 (f)]

License No: DL.NO.T/8-DCH dt: 27.10.80: & T/13-DCH dt: 24.9.90

Dt of Receipt: 26.12.00

	i Name, address and license No. of Manufacturer /	Adviser (Health), State Urban Development Agency, Ilgu	s
	Supplier's from whom sample received	Bhawan, II-C Block, Sec-III, Bidhan Nagar, Cal-91,	
	2.Reference No. and date of Manufacturer / Supplier's	SUDA-120/96 (Pt-III)/ dt. 13.12.00	
	letter under which the sample was forwarded		
	3. Name of Drug / Cospetics / Raw material purporting to	Aspirin Tablet I.P.	
	be contained in the sample.		
	4. Detail of Raw material/final product in bulk / final	2 X 50 tabs	
	product (in finished pack) as obtained from the		
	manufacturer for analysis.		
	(a) Original manufacturer's name (in case of raw	Mis. Kansas Labs (P) Ltd	
	materials and drugs repacked)	8/1, Lal Bazar St. Cal-1	
	(b) Batch Number	K 1477	
	(c) Batch Size	Nil	
	(d) Date of Manufacture if any	11/00	
••	(e) Date of Expiry if any.	04/02	
	5. Result of Analysis Recorded Below:		

Method : I.P' 96, VoL-II.

Description :-White round oval shaped tablet.

Avg. wt. of a tablet	0.341.4gm	
Weight variation	Passes	
Identification	Positive	
Disintegration time	1 minutes	
Salicylic Acid	Passes	

Assay ---

Aspirin I.P.

Found/tab 308.93mg Claim/tab 300.00mg

In the opinion of the undersigned, the sample referred to above is of standard quality /-is not of standard quality as defined in the Act and Rules made thereunder for the reasons given below with respect of the above tests only.

Junior Analyst /

Signature of Office

W. B. PHARMACEUTICAL & PHYTOCHEMICAL DEV. CORPN LI

(A Government of West Bengal Undertaking)

ACHARYA P. C. RAY R & D DRUG TESTING & POLLUTION TESTING LABORATORY

Behala Chowrasta, 620 Diamond Harbour Road, Calcutta 34, West Bengal

Report of Analysis [Form 39, See Rule 150 (f)]

License No: DL NO T/8-DCH dt: 27.10.80. & T/13-DCH dt: 24.9.90

Sample Receiving No:D/2360/00

be contained in the sample

1.Name, address and license No. of Manufacturer /

Supplier's from whom sample received 2.Reference No. and date of Manufacturer / Supplier's

letter under which the sample was forwarded

3.Name of Drug / Cosmetics / Raw material purporting to

Dt of Receipt: 26.12.00

A1361-

State Urban Development Agency II-C Block, Sector-III, Bidhannagar Cal-700 091 SUDA/120/96 Pt-III

Oxyphenonium Bromide tablet IP 5.00mg

4.Detail of Raw material/final product in bulk / final product (in finished pack) as obtained from the manufacturer for analysis.	10X5 strip
(a) Original manufacturer's name (in case of raw	Kansus Laboratory Pvt Ltd
materials and drugs repacked)	8/1 Lalbazar Street Cal-1 DL 1446/9
(b) Batch Number	K-1476
(c) Batch Size	
(d) Date of Manufacture if any	11/2000
(e) Date of Expiry if any	10/2003
5. Result of Analysis Recorded Below:	

As per IP Suppliments-1975

Description :- A small white round tablet having a bisecting mark on one side

Net avg content of a capsule	0.1163 gm
Wt. variation	passes
Identification	positive
Disintegration time	9 mins

Assay:-

Oxyphenonium Bromide IP

Found/tab 4.81 mg

Claim/tab 5.00 gm

In the opinion of the undersigned, the sample referred to above is of standard quality / is not of standard quality as defined in the Act and Rules made thereunder for the reasons given below with respect to above tests only. However no opinion can be given regarding the quality of the sample as per Drugs & Cosmetics Act & Rules as complete test could not be performed as the Manufacturer / Supplier did not ask for complete test.

Signature of Officer In

A. 3612/09

W. B. PHARMACEUTICAL & PHYTOCHEMICAL DEV. CORPN LTD.

(A Government of West Bengal Undertaking)

ACHARYA P. C. RAY R & D DRUG TESTING & POLLUTION TESTING LABORATORY

Behala Chowrasta, 620 Diamond Harbour Road, Calcutta 34, West Bengal

Report of Analysis [Form 39, See Rule 150 (f)]

License No: DI. NO T/8-DCH dr 27 10 80 & T/13-DCH dr 24 9 90

Sample Receiving No:D/2360/00	Dt of Receipt: 26 12 00
1.Name, address and license No. of Manufacturer	State Urban Development Agency
Supplier's from whom sample received	II-C Block, Sector-III, Bidhannagar Cal-700 09
2.Reference No. and date of Manufacturer / Supplier's	SUDA/120/96 Pt-III
letter under which the sample was forwarded	
3.Name of Drug / Cospeties / Raw material purporting to	Oxyphenonium Bromide tablet IP 5 00mg
be contained in the sample.	
4. Detail of Raw material/final product in bulk / final	10X5 strip
product (in finished pack) as obtained from the	
manufacturer for analysis.	
(a) Original manufacturer's name (in case of raw	Kansus Laboratory Pvt Ltd
materials and drugs repacked).	8/1 Lalbazar Street Cal-1 DL 1446/9
(b) Batch Number	K-1476
(c) Batch Size	
(d) Date of Manufacture if any	11/2000
(e) Date of Expiry if any	10/2003
5.Result of Analysis Recorded Below:	
As per IP Su	appliments-1975

Description :- A small white round tablet having a bisecting mark on one side

Net avg content of a capsule	0.1163 gm	
Wt. variation	passes	
Identification	positive	
Disintegration time	9 mins	
Assay:-		
	Found/tab	Claim tab
Oxyphenonium Bromide II'	4.81 mg	5.00 gm

In the opinion of the undersigned, the sample referred to above is of standard quality / is-not of standard quality as defined in the Act and Rules made thereunder for the reasons given below with respect to above tests only. However no opinion can be given regarding the quality of the sample as per Drugs & Cosmetics Act & Rules as complete test could not be performed as the Manufacturer / Supplier did not ask for complete test

TILVOD Signature of Office

419/M

W. B. PHARMACEUTICAL & PHYTOCHEMICAL DEV. CORPN LTE

(A Government of West Bengal Undertaking)

ACHARYA P. C. RAY R & D DRUG TESTING & POLLUTION TESTING LABORATORY

Behala Chowrasta, 620 Diamond Harbour Road, Calcutta 34, West Bengal

Report of Analysis [Form 39, See Rule 150 (f)]

License No: DL NO.T/8-DCH dt: 27.10.80. & T/13-DCH dt: 24.9.90

Sample Receiving No:D/2359/00	Dt of Receipt: 26.12.00
1.Name, address and license No. of Manufacturer /	State Urban Development Agency
Supplier's from whom sample received	II-C Block.Sector-III,Bidhannagar Cal-700 091
2. Reference No. and date of Manufacturer / Supplier's	SUDA/120/96 Pt-111 dt 13.12.00
Ietter under which the sample was forwarded 3.Name of Drug / Cosmetics / Raw material purporting to be contained in the sample.	Chloramphenicol Bye Ointment IP
4.Detail of Raw material/final product in bulk / final product (in finished pack) as obtained from the manufacturer for analysis.	
(a) Original manufacturer's name (in case of raw materials and drugs repacked).	M/s Jyoti Capsule 123/37 Saresh Bagh, Kanpur-208012
(b) Batch Number	JGC21010
(c) Batch Size	
(d) Date of Manufacture if any	11/2000
(e) Date of Expiry if any	04/2002
5. Result of Analysis Recorded Below:	

As per IP 1996 Vol-2

Description :- Yellowish white coloured applicaps

Net avg content of a capsule 0.2634 gm Wt. variation passes Identification positive

Assay:-

Chloraphenicol IP

Found/daip

0.95%

Claim/tab d 1.00%

0.90% to 1.20%

In the opinion of the undersigned, the sample referred to above is of standard quality / is not of standard quality as defined in the Act and Rules made thereunder for the reasons given below with respect to above tests only. However no opinion can be given regarding the quality of the sample as per Drugs & Cosmetics Act & Rules as complete test could not be performed as the Manufacturer /-Supplier did not ask for complete test.

Signature of Officer In-Charge

•	31.	
Computerised File Name: D&Creport2102	A 3619/01	
W. B. PHARMACEUTICAL & PI	HYTOCHEMICAL DEV CORPN LTD > 01 11	
(A Government of V	Vest Bengal Undertaking)	
ACHARYA P. C. RAY R & D DRUG TEST	ING & POLLUTION TESTING LABORATORY	
Behala Chowrasta, 620 Diamond H	larbour Road, Calcutta 34, West Bengal	
Report of Analysis [F	orm 39. See Rule 150 (f)]	
License No: DL NO T/8-DCH of	lt 27.10 80 & T/13-DCH dt 24.9 90	
Sample Receiving No:D/2359/00	Dt of Receipt: 26.12.00	
1.Name, address and license No of Manufacturer /	State Urban Development Agency	
Supplier's from whom sample received	II-C Block.Sector-III,Bidhannagar Cal-700 091	
2.Reference No. and date of Manufacturer / Supplier's	SUDA 120/96 Pt-III dt 13.12.00	
letter under which the sample was forwarded	andlen	
3.Name of Drug / Cosmétics / Ray material purporting to	Chloramphenicol Eye Omment IP	
be contained in the sample		
4.Detail of Raw material final product in bulk / final		
product (in finished pack) as obtained from the		
manufacturer for analysis		
(a) Original manufacturer's name (in case of raw	M/s Jyon Capsule	
materials and drugs repacked)	123/37 Saresh Bagh, Kanpur-208012	
(b) Batch Number	JGC21010	
(c) Batch Size		
(d) Date of Manufacture if any	11/2000	
(e) Date of Expiry if any	04/2002	
5. Result of Analysis Recorded Below:		
As per I	P 1996 Vol-2	
Description :- Yellowish white coloured applicaps		
Net avg content of a capsule 0.2634 gm		
Wt. variation passes		
Identification positive		
Assay:-		
Found Cap	Claim tap d	
	Clamitap	
Chloraphenicol IP 0.95%	1.00% 0.90% to 1.20%	
In the opinion of the undersioned the same is the start		

In the opinion of the undersigned, the sample referred to above is of standard quality is not of standard quality as defined in the Act and Rules made thereunder for the reasons given below with respect to above tests only. However no opinion can be given regarding the quality of the sample as per Drugs & Cosmetics Act & Rules as complete test could not be performed as the Manufacturer / Supplier did not ask for complete test.

W. B. PHARMACEUTICAL & PHYTOCHEMICAL DEV. CORPN LTD.

(A Government of West Bengal Undertaking)

ACHARYA P. C. RAY R & D DRUG TESTING & POLLUTION TESTING LABORATORY

Behala Chowrasta, 620 Diamond Harbour Road, Calcutta 34, West Bengal

Report of Analysis [Form 39, See Rule 150 (f)]

License No: DL.NO.T/8-DCH dt: 27.10.80. & T/13-DCH dt: 24.9.90

Sample Receiving No:D/2142/00

Dt of Receipt: 15.12.00

A/35

I.Name, address and license No. of Manufacturer /	Adviser (Health), State Urban Development Agency, Ilgus
Supplier's from whom sample received.	Bhawan, II-C Block, Sec-III, Bidhan Nagar, Cal-91,
2. Reference No. and date of Manufacturer / Supplier's	SUDA-15/98 (Pt-II)/720 dt. 14.12.00
letter under which the sample was forwarded	
3.Name of Drug / Cosmetics / Raw material purporting to	O.R.S.
be contained in the sample	
4.Detail of Raw material/final product in bulk / final	16 sachets
product (in finished pack) as obtained from the	
manufacturer for analysis	
(a) Original manufacturer's name (in case of raw	Pure Pharma
materials and drugs repacked)	41, 42 & 44, Industrial Area. Pologround (Indore)
(b) Batch Number	0021
(c) Batch Size	Nil
(d) Date of Manufacture if any	11/00
(e) Date of Expiry if any	10/02
5. Result of Analysis Recorded Below:	

As per I.P.'96

Description :- A white fine crystalline powder in PVC sachet ...

Identification	Positive for Sodium, Potassium, Chloride, Dextrose
Uniformity of weight	Passes
Weight variation	Passes
Net content of a sachet	28.45136gm

Assay:--

	Found/27.9gm	Claim/27.9gm	Limit/27.9gm
Anhydrous Dextrose I.P.	19.46gm	20.00gm	18.0gm to 22.0gm
Chloride Content	2.735gm	2.8386gm	2.554gm to 3.122gm
Sodium Content	2.172gm	2.056gm	1.85gm to 2.26gm
Potassium content	0.747gm	0.786gm	0.707 to 0.864gm
Sodium Citrate I.P.	2.807gm	2.9gm	2.61gm to 3.19gm

In the opinion of the undersigned, the sample referred to above is of standard quality \perp is not of standard quality as defined in the Act and Rules made thereunder for the reasons given below.

N Analyst / Junior Analyst

Signature of Officer In-Charge

W. B. PHARMACEUTICAL & PHYTOCHEMICAL DEV. CORPN LTD.

(A Government of West Bengal Undertaking)

ACHARYA P. C. RAY R & D DRUG TESTING & POLLUTION TESTING LABORATORY

Behala Chowrasta, 620 Diamond Harbour Road, Calcutta 34, West Bengal

Report of Analysis [Form 39, See Rule 150 (f)]

License No: DI, NO T/8-DCH dt: 27.10.80 & T/13-DCH dt: 24 9 90

Sample Receiving No:D/2142/00

Dt of Receipt: 15.12.00

1.Name, address and license No. of Manufacturer /	Adviser (Health), State Urban Development Agency, Ilgus
Supplier's from whom sample received.	Bhawan, II-C Block, Sec-III, Bidhan Nagar, Cal-91,
2.Reference No. and date of Manufacturer / Supplier's	SUDA-15/98 (Pt-II)/720 dt. 14.12.00
letter under which the sample was forwarded	
3.Name of Drug ' Cosmetics / Raw material purporting to	O.R.S.
be contained in the sample.	
4. Detail of Raw material/final product in bulk / final	16 sachets
product (in finished pack) as obtained from the	
manufacturer for analysis	
(a) Original manufacturer's name (in case of raw	Pure Pharma
materials and drugs repacked)	41, 42 & 44, Industrial Area. Pologround (Indore)
(b) Batch Number	0021
(c) Batch Size	Nil
(d) Date of Manufacture if any	11/00
(e) Date of Expiry if any	10/02
5. Result of Analysis Recorded Below:	

As per I.P.'96

Description :- A white fine crystalline powder in PVC sachet ...

Identification	Positive for Sodium, Potassium, Chloride, Dextrose
Uniformity of weight	Passes
Weight variation	Passes
Net content of a sachet	28.45136gm

Assay:--

	Found/27.9gm	Claim/27.9gm	Limit/27.9gm
Anhydrous Dextrose IP.	19.46gm	20.00gm	18.0gm to 22.0gm
Chloride Content	2.735gm	2.8385gm	2.554gm to 3.122gm
Sodium Content	2.172gm	2.056gm	1.85gm to 2.26gm
Polassium content	0.747gm	0.786gтн	0.707 to 0.864gm
Sodium Citrate I.P.	2.807gm	2.9gm	2.61gm to 3.19gm

In the opinion of the undersigned, the sample referred to above is of standard quality / is not of standard quality as defined in the Act and Rules made thereunder for the reasons given below.

Signature of Officer In-Charge

Sample Receiving No:D/2143/00

W. B. PHARMACEUTICAL & PHYTOCHEMICAL DEV. CORPN LTD.

(A Government of West Bengal Undertaking)

ACHARYA P. C. RAY R & D DRUG TESTING & POLLUTION TESTING LABORATORY

Behala Chowrasta, 620 Diamond Harbour Road, Calcutta 34, West Bengal

Report of Analysis [Form 39, See Rule 150 (f)]

License No: DL.NO.T/8-DCH dt: 27.10.80. & T/13-DCH dt: 24.9.90

Dt of Receipt: 15.12.00

ARE

A	
1.Name, address and license No. of Manufacturer /	Adviser (Health), State Urban Development Agency, Ilgus
Supplier's from whom sample received	Bhawan, II-C Block, Sec-III, Bidhan Nagar, Cal-91,
2. Reference No. and date of Manufacturer / Supplier's	SUDA-15/98 (Pt-II)/720 dt. 14.12.00
letter under which the sample was forwarded	
3.Name of Drug / Cospectics / Raw material purporting to	O.R.S.
be contained in the sample	
4.Detail of Raw material/final product in bulk / final	16 sachets
product (in finished pack) as obtained from the	
manufacturer for analysis	
(a) Original manufacturer's name (in case of raw	Pure Pharma
materials and drugs repacked)	
	41, 42 & 44, Industrial Area. Pologround (Indore)
(b) Batch Number	0022
(c) Batch Size	Nil
(d) Date of Manufacture if any	11/00
(e) Date of Expiry if any	10/02
5. Result of Analysis Recorded Below:	

As per I.P.'96

Description :- A white fine crystalline powder in PVC sachet.

Identification	Positive for Sodium, Potassium, Chloride, Dextrose
Uniformity of weight	Passes
Weight variation	Passes
Net content of a sachet	28.45773gm

Assay:--

	Found/27.9gm	Claim/27.9gm	Limit/27.9gm
Anhydrous Dextrose IP.	19.29gm	20.00gm	18.0gm to 22.0gm
Chloride Content	2.71gm	2.8386gm	2.554gm to 3.122gm
Sodium Content	2.055gm	2.056gm	1.85gm to 2.26gm
Potassium content	0.736gm	0.786gm	0.707 to 0.864gm
Sodium Citrate I.P.	2.798gm	2.9gm	2.61gm to 3.19gm

In the opinion of the undersigned, the sample referred to above is of standard quality / is not of standard quality as defined in the Act and Rules made thereunder for the reasons given below.

Analyst / Junior Analyst

Signature of Off

A1357

W. B. PHARMACEUTICAL & PHYTOCHEMICAL DEV. CORPN LTD. (A Government of West Bengal Undertaking)

ACHARYA P. C. RAY R & D DRUG TESTING & POLLUTION TESTING LABORATORY Behala Chowrasta, 620 Diamond Harbour Road, Calcutta 34, West Bengal

Report of Analysis [Form 39. See Rule 150 (f)]

License No: DL.NO.T/8-DCH dt: 27.10.80. & T/13-DCH dt: 24.9.90

Sample Receiving No:D/2143/00	r: 27.10.80. & T/13-DCH dt: 24.9.90
 Name, address and license No. of Manufacturer / Supplier's from whom sample received	Dt of Receipt: 15.12.00 Adviser (Health), State Urban Development Agency, Ilgus Bhawan, II-C Block, Sec-III, Bidhan Nagar, Cal-91, SUDA-15/98 (Pt-II)/720 dt. 14.12.00
3. Name of Drug ' Cospiletics / Raw material purporting to be contained in the sample	O.R.S.
4.Detail of Raw material/final product in bulk / final product (in finished pack) as obtained from the manufacturer for analysis	16 sachets
 (a) Original manufacturer's name (in case of raw materials and drugs repacked). 	Pure Pharma
(D) Batch Number	41, 42 & 44, Industrial Area. Pologround (Indore) 0022
	Nil
(d) Date of Manufacture if any(e) Date of Expire if any	11/00
(e) Date of Expiry if any.5. Result of Analysis Recorded Below:	10/02

As per I.P.'96

Description :- A white fine crystalline powder in PVC sachet ...

Identification	Positive for Sodium, Potassium, Chloride, Dextrose
Uniformity of weight	Passes
Weight variation	Passes .
Net content of a sachet	28.45773gm

Assay:--

Analyst / Junior Analyst

	Found/27.9gm	Claim/27.9gm	Limit/27.9gm
Anhydrous Dextrose IP.	19.29gm	20.00gm	18.0gm to 22.0gm
Chloride Content	2.71gm	2.8386gm	2.554gm to 3.122gm
Sodium Content	2.055gm	2.056gm	
Potassium content	0.736gm	0.786gm	1.85gm to 2.26gm 0.707 to 0.864gm
Sodium Citrate I.P.	2.798gm	2.9gm	2.61gm to 3.19gm

N

In the opinion of the undersigned, the sample referred to above is of standard quality / is not of standard quality as defined in the Act and Rules made thereunder for the reasons given below.

TILN Signature of Officer In-Charge

AT 75

W. B. PHARMACEUTICAL & PHYTOCHEMICAL DEV. CORPN LTD.

(A Government of West Bengal Undertaking)

ACHARYA P. C. RAY R & D DRUG TESTING & POLLUTION TESTING LABORATORY

Behala Chowrasta, 620 Diamond Harbour Road, Calcutta 34, West Bengal

Report of Analysis [Form 39, See Rule 150 (f)]

License No: DL NO. T/8-DCH dt: 27.10 80. & T/13-DCH dt: 24.9.90

Sample Receiving No:D/2144/00

Dt of Receipt: 15.12.00

Adviser (Health), State Urban Development Agency, Ilgus 1.Name, address and license No. of Manufacturer / Bhawan, II-C Block, Sec-III, Bidhan Nagar, Cal-91, Supplier's from whom sample received 2. Reference No. and date of Manufacturer / Supplier's SUDA-15/98 (Pt-II)/720 dt. 14.12.00 letter under which the sample was forwarded 3. Name of Drug / Cospetics / Raw material purporting to O.R.S. be contained in the sample. 4. Detail of Raw material final product in bulk / final 16 sachets product (in finished pack) as obtained from the manufacturer for analysis. (a) Original manufacturer's name (in case of raw Pure Pharma 41, 42 &44, Industrial Area. Pologround (Indore) materials and drugs repacked) (b) Batch Number 0023 Nil (c) Batch Size (d) Date of Manufacture if any 11/00 (e) Date of Expiry if any..... 10/02 5. Result of Analysis Recorded Below:

As per I.P.'96

Description :- A white fine crystalline powder in PVC sachet ..

Identification	Positive for Sodium, Potassium, Chloride, Dextrose
Uniformity of weight	Passes
Weight variation	Passes
Net content of a sachet	28.45512gm

Assay:--

	Found/27.9gm	Claim/27.9gm	Limit/27.9gm
Anhydrous Dextrose IP.	19.65gm	20.00gm	18.0gm to 22.0gm
Chloride Content	2.742gm	2.8386gm	2.554gm to 3.122gm
Sodium Content	2.093gm	2.056gm	1.85gm to 2.26gm
Potassium content	0.750gm	0.786gm	0.707 to 0.864gm
Sodium Citrate I.P.	2.849gm	2.9gm	2.61gm to 3.19gm

In the opinion of the undersigned, the sample referred to above is of standard quality / is not of standard quality as defined in the Act and Rules made thereunder for the reasons given below.

Analyst / Junior Anal

Signature of Officer In-Cl

W. B. PHARMACEUTICAL & PHYTOCHEMICAL DEV. CORPN LTD.

(A Government of West Bengal Undertaking)

ACHARYA P. C. RAY R & D DRUG TESTING & POLLUTION TESTING LABORATORY

Behala Chowrasta, 620 Diamond Harbour Road, Calcutta 34, West Bengal

Report of Analysis [Form 39. See Rule 150 (f)]

License No: DL.NO T/8-DCH dt: 27.10.80 & T/13-DCH dt: 24.9.90

Sample Receiving No:D/2144/00

Dt of Receipt: 15.12.00

Adviser (Health), State Urban Development Agency, Ilgus Bhawan, II-C Block, Sec-III, Bidhan Nagar, Cal-91, SUDA-15/98 (Pt-II)/720 dt. 14.12.00

2.Reference No. and date of Manufacturer / Supplier's letter under which the sample was forwarded

3 Name of Drug / Cospetics / Raw material purporting to

be contained in the sample.

1.Name, address and license No. of Manufacturer /

Supplier's from whom sample received

O.R.S.

4 Detail of Raw material final product in bulk / final 16 sachets

• • • • • • • • • • • • • • • • • • •
Pure Pharma
41, 42 &44, Industrial Area. Pologround (Indore)
0023
Nil
11/00
10/02

5. Result of Analysis Recorded Below:

As per I.P.'96

Description :- A white fine crystalline powder in PVC sachet ...

Identification	Positive for Sodium, Potassium, Chloride, Dextrose
Uniformity of weight	Passes
Weight variation	Passes
Net content of a sachet	28.45512gm

Assay:--

	Found/27.9gm	Claim/27.9gm	Limit/27.9gm
Anhydrous Dextrose IP.	19.65gm	20.00gm	18.0gm to 22.0gm
Chloride Content	2.742gm	2.8386gm	2.554gm to 3.122gm
Sodium Content	2.093gm	2.056gm	1.85gm to 2.26gm
Polassium content	0.750gm	0.786gm	0.707 to 0.864gm
Sodium Citrate I.P.	2.849gm	2.9gm	2.61gm to 3.19gm

In the opinion of the undersigned, the sample referred to above is of standard quality / is not of standard quality as defined in the Act and Rules made thereunder for the reasons given below.

Analyst / Junior Analyst

Signature of Officer In

Sample Receiving No:D/2145/00

A/ 35

W. B. PHARMACEUTICAL & PHYTOCHEMICAL DEV. CORPN LTD.

(A Government of West Bengal Undertaking)

ACHARYA P. C. RAY R & D DRUG TESTING & POLLUTION TESTING LABORATORY

Behala Chowrasta, 620 Diamond Harbour Road, Calcutta 34, West Bengal

Report of Analysis [Form 39, See Rule 150 (f)]

License No: DL.NO.T/8-DCH dt: 27.10.80. & T/13-DCH dt: 24.9.90

Dt of Receipt: 15.12.00

1.Name, address and license No. of Manufacturer /	Adviser (Health), State Urban Development Agency, Ilgus
Supplier's from whom sample received	Bhawan, II-C Block, Sec-III, Bidhan Nagar, Cal-91,
2.Reference No. and date of Manufacturer / Supplier's	SUDA-15/98 (Pt-II)/720 dt. 14.12.00
letter under which the sample was forwarded	
3.Name of Drug / Cospetics / Raw material purporting to	O.R.S.
be contained in the sample	
4.Detail of Raw material/final product in bulk / final	16 sachets
product (in finished pack) as obtained from the	
manufacturer for analysis	
(a) Original manufacturer's name (in case of raw	Pure Pharma
materials and drugs repacked)	41, 42 &44, Industrial Area. Pologround (Indore)
(b) Batch Number	0024
(c) Batch Size	Nil
(d) Date of Manufacture if any	11/00
(e) Date of Expiry if any	10/02
5. Result of Analysis Recorded Below:	

As per I.P.'96

Description :- A white fine crystalline powder in PVC sachet ...

Identification	Positive for Sodium, Potassium, Chloride, Dextrose
Uniformity of weight	Passes
Weight variation	Passes
Net content of a sachet	28.47158gm

Assay:--

	Found/27.9gm	Claim/27.9gm	Limit/27.9gm
Anhydrous Dextrose IP.	19.49gm	20.00gm	18.0gm to 22.0gm
Chloride Content	2.708gm	2.8386gm	2.554gm to 3.122gm
Sodium Content	2.165gm	2.056gm	1.85gm to 2.26gm
Potassium content	0.748gm	0.786gm	0.707 to 0.864gm
Sodium Citrate I.P.	2.798gm	2.9gm	2.61gm to 3.19gm

In the opinion of the undersigned, the sample referred to above is of standard quality / is not of standard quality as defined in the Act and Rules made thereunder for the reasons given below.

Analyst / Junior Ana

Signature of Office

3 AI 3578/17 W. B. PHARMACEUTICAL & PHYTOCHEMICAL DEV. CORPN LTD.

(A Government of West Bengal Undertaking)

ACHARYA P. C. RAY R & D DRUG TESTING & POLLUTION TESTING LABORATORY

Behala Chowrasta, 620 Diamond Harbour Road, Calcutta 34, West Bengal

Report of Analysis [Form 39, See Rule 150 (f)]

License No: DL.NO.T/8-DCH dt: 27 10 80 & T/13-DCH dt: 24 9 90

Sample Receiving No:D/2145/00

Dt of Receipt: 15.12.00

Sumple recently reader and	
1.Name, address and license No. of Manufacturer /	Adviser (Health), State Urban Development Agency, Ilgus
Supplier's from whom sample received	Bhawan, II-C Block, Sec-III, Bidhan Nagar, Cal-91,
2.Reference No. and date of Manufacturer / Supplier's	SUDA-15/98 (Pt-II)/720 dt. 14.12.00
letter under which the sample was forwarded	
3.Name of Drug / Cospletics / Raw material purporting to	O.R.S.
be contained in the sample.	
4. Detail of Raw material/fingl product in bulk / final	to sachets
product (in finished pack) as obtained from the	
manufacturer for analysis	
(a) Original manufacturer's name (in case of raw	Pure Pharma
materials and drugs repacked)	41, 42 & 44, Industrial Area. Pologround (Indore)
(b) Batch Number	0024
(c) Batch Size	Nil
(d) Date of Manufacture if any	11/00
(e) Date of Expiry if any	10/02
5. Result of Analysis Recorded Below:	

As per I.P.'96

Description :- A white fine crystalline powder in PVC sachet ...

Identification	Positive for Sodium, Potassium, Chloride, Dextrose
Uniformity of weight	Passes
Weight variation	Passes
Net content of a sachet	28.47158gm

Assay:--

	Found/27.9gm	Claim/27.9gm	Limit/27.9gm
Anhydrous Dextrose IP.	19.49gm	20.00gm	18.0gm to 22.0gm
Chloride Content	2.708gm	2.8386gm	2.554gm to 3.122gm
Sodium Content	2.165gm	2.056gm	1.85gm to 2.26gm
Polassium content	0.748gm	0.786gm	0.707 to 0.864gm
Sodium Citrate I.P.	2.798gm	2.9gm	2.61gm to 3.19gm

In the opinion of the undersigned, the sample referred to above is of standard quality / is not of standard quality as defined in the Act and Rules made thereunder for the reasons given below.

Analyst/Junior Analyst/

Signature of Officer In-Charge

A: 356 PM

W. B. PHARMACEUTICAL & PHYTOCHEMICAL DEV. CORPN LTD.

(A Government of West Bengal Undertaking)

ACHARYA P. C. RAY R & D DRUG TESTING & POLLUTION TESTING LABORATORY

Behala Chowrasta, 620 Diamond Harbour Road, Calcutta 34, West Bengal

Report of Analysis [Form 39, See Rule 150 (f)]

License No: DL NO T/8-DCH df: 27 10.80 & T/13-DCH df: 24 9.90

Sample Receiving No:D/2363/00

Dt of Receipt: 26.12.00

Adviser (Health), State Urban Development Agency, Ilgus Bhawan, II-C Block, Sec-III, Bidhan Nagar, Cal-91, SUDA/120/96 (Pt-III) dt. Nil

3.Name of Drug / Cosmetics / Ray material purporting to Cotrimoxazole tablet 1.P.

be contained in the sample. 4.Detail of Raw material/final product in bulk / final product (in finished pack) as obtained from the manufacturer for analysis.

I.Name, address and license No. of Manufacturer /

2.Reference No. and date of Manufacturer / Supplier's letter under which the sample was forwarded

Supplier's from whom sample received

5X10's in strip

n	nanufacturer for analysis.	
(a)	Original manufacturer's name (in case of raw	Kansas Laboratones Pvt. Ltd
	materials and drugs repacked).	8/1, Lal Bazar Street, Cal-1, DL No- 1446M
(b)	Batch Number	K 1462
(c)	Batch Size	Nil
(d)	Date of Manufacture if any	11/00
(e)	Date of Expiry if any	10/03

5. Result of Analysis Recorded Below:

As per I.P.'96

Description :- A white round tablet having a bisecting mark on one side .

Identification	Positive
Avg. wt. of a tablet	0.5742gm
Weight variation	Passes
Time of disintegration	9 minutes

Assay:--

Trimethoprim I.P. 79.90mg Sulphamethoxazole I.P. 406.15mg

 Found/ml
 b
 Claim/ml
 b
 L

 79.90mg
 80.00mg
 400.00mg
 400.00mg

In the opinion of the undersigned, the sample referred to above is of standard quality *is not of standard quality* as defined in the Act and Rules made thereunder for the reasons given below.

Signature of Officer In-Charge

Sample Receiving No:D/2363/00

A! 3566 01

W. B. PHARMACEUTICAL & PHYTOCHEMICAL DEV. CORPN LTD.

(A Government of West Bengal Undertaking)

ACHARYA P. C. RAY R & D DRUG TESTING & POLLUTION TESTING LABORATORY

Behala Chowrasta, 620 Diamond Harbour Road, Calcutta 34, West Bengal

Report of Analysis [Form 39, See Rule 150 (f)]

License No: DL.NO. T/8-DCH dt: 27.10.80. & T/13-DCH dt: 24.9.90

Dt of Receipt: 26.12.00

Duripas researcing researcing and	
1.Name, address and license No. of Manufacturer /	Adviser (Health), State Urban Development Agency, Ilgus
Supplier's from whom sample received	Bhawan, II-C Block, Sec-III, Bidhan Nagar, Cal-91,
2.Reference No. and date of Manufacturer / Supplier's	SUDA/120/96 (Pt-III) dt. Nil
letter under which the sample was forwarded	
3.Name of Drug / Cosmetics / Ray material purporting to	Cotrimoxazole tablet I.P.
be contained in the sample.	
4.Detail of Raw material/final product in bulk / final	5X10's in strip
product (in finished pack) as obtained from the	
manufacturer for analysis	
(a) Original manufacturer's name (in case of raw	Kansas Laboratories Pvt. Ltd
materials and drugs repacked)	8/1, Lal Bazar Street, Cal-1, DL No- 1446M
(b) Batch Number	K 1462
(c) Batch Size	Nil
(d) Date of Manufacture if any	11/00
(e) Date of Expiry if any	10/03

5. Result of Analysis Recorded Below:

As per I.P.'96

Description :- A white round tablet having a bisecting mark on one side .

Identification	Positive
Avg. wt. of a tablet	0.5742gm
Weight variation	Passes
Time of disintegration	9 minutes

Assay:--

Trimethoprim I.P. Sulphamethoxazole I.P. Found/ml back 79.90mg 406.15mg <u>Claim/ml</u> het \$ 80.00mg 400.00mg

In the opinion of the undersigned, the sample referred to above is of standard quality / is not of standard quality as defined in the Act and Rules made thereunder for the reasons given below.

Signature of Officer

W. B. PHARMACEUTICAL & PHYTOCHEMICAL DEV. CORPN LTD.

(A Government of West Bengal Undertaking)

ACHARYA P. C. RAY R & D DRUG TESTING & POLLUTION TESTING LABORATORY

Behala Chowrasta, 620 Diamond Harbour Road, Calcutta 34, West Bengal

Report of Analysis [Form 39, See Rule 150 (f)]

License No: DL NO. T/8-DCH dt: 27.10.80. & T/13-DCH dt: 24.9.90

Sample Receiving No:D/2161/00

be contained in the sample.

1.Name, address and license No. of Manufacturer /

2. Reference No. and date of Manufacturer / Supplier's

A Datail of Down and interfer and in 1 11 1 5 1

letter under which the sample was forwarded 3. Name of Drug / Cosmetics / Raw material purporting to

Supplier's from whom sample received.....

Dt of Receipt: 15.12.00

A356/M

Adviser (Health), State Urban Development Agency, Ilgus Bhawan, II-C Block, Sec-III, Bidhan Nagar, Cal-91, SUDA/120/96 (Pt-III) dt. Nil

Cotrimoxazole tablet I.P.

	product (in finished pack) as obtained from the manufacturer for analysis	OX10's in strip
(a) Original manufacturer's name (in case of raw	Kansas Laboratories Pvt. Ltd
	materials and drugs repacked)	8/1, Lal Bazar Street, Cal-1, DL No- 1446M
(t) Batch Number	K 1462
(0) Batch Size	Nil
(d	Date of Manufacture if any	11/00
(e) Date of Expiry if any.	10/03
E	Demle flash to bar t	

5. Result of Analysis Recorded Below:

As per I.P.'96

Description :- A white round tablet having a bisecting mark on one side .

Identification	Positive
Avg. wt. of a tablet	0.5828gm
Weight variation	Passes
Time of disintegration	7 minutes

Assay:--

	Found/mil AL	Claimini for V
Trimethoprim I.P.	79.65mg	80.00mg
Sulphamethoxazole I.P.	385.73mg	400.00mg

In the opinion of the undersigned, the sample referred to above is of standard quality / is not of standard quality as defined in the Act and Rules made thereunder for the reasons given below.

Signature of Officer In-Charge

A: 3563/04

Computerised File Name: D&Creport2059

W. B. PHARMACEUTICAL & PHYTOCHEMICAL DEV. CORPN LTD.

(A Government of West Bengal Undertaking)

ACHARYA P. C. RAY R & D DRUG TESTING & POLLUTION TESTING LABORATORY

Behala Chowrasta, 620 Diamond Harbour Road, Calcutta 34, West Bengal

Report of Analysis [Form 39, See Rule 150 (f)]

License No: DL.NO T/8-DCH dt: 27 10 80. & T/13-DCH dt: 24.9.90

Sample Receiving No:D/2363/00	Dt of Receipt: 26.12.00
I.Name, address and license No. of Manufacturer /	Adviser (Health), State Urban Development Agency, Ilgus
Supplier's from whom sample received	Bhawan, II-C Block, Sec-III, Bidhan Nagar, Cal-91,
2.Reference No. and date of Manufacturer / Supplier's	SUDA/120/96 (Pt-III) dt. Nil
letter under which the sample was forwarded	
3.Name of Drug / Cosmetics / Raw material purporting to	Cotrimoxazole tablet I.P.
be contained in the sample.	
4. Detail of Raw material/final product in bulk / final	5X10's in strip
product (in finished pack) as obtained from the	
manufacturer for analysis	
(a) Original manufacturer's name (in case of raw	Kansas Laboratories Pvt. Ltd
materials and drugs repacked).	8/1, Lal Bazar Street, Cal-1, DL No- 1446M
(b) Batch Number	K 1462
(c) Batch Size	Nil
(d) Date of Manufacture if any	11/00
(e) Date of Expiry if any	10/03
5. Result of Analysis Recorded Below:	

As per 1.P.'96

Description :- A white round tablet having a bisecting mark on one side

Identification	Positive
Avg. wt. of a tablet	0.5742gm
Weight variation	Passes
Time of disintegration	9 minutes

Assay:--

Trimethoprim I.P. Sulphamethoxazole I.P. Found/ml bot 79.90mg 406.15mg

Claimint he 1 80.00mg

400.00mg

In the opinion of the undersigned, the sample referred to above is of standard quality is not of standard quality as defined in the Act and Rules made thereunder for the reasons given below

Signature of Officer In-

letter under which the sample was forwarded 3.Name of Drug / Cosmetics / Raw material purporting to

be contained in the sample ...

W. B. PHARMACEUTICAL & PHYTOCHEMICAL DEV. CORPN LTD

(A Government of West Bengal Undertaking)

ACHARYA P. C. RAY R & D DRUG TESTING & POLLUTION TESTING LABORATORY

Behala Chowrasta, 620 Diamond Harbour Road, Calcutta 34, West Bengal

Report of Analysis [Form 39, See Rule 150 (f)]

License No: DL.NO.T/8-DCH dt: 27.10.80 & T/13-DCH dt: 24.9.90

Sample Receiving No:D/2162/00 Dt of Receipt: 15.12.00 1.Name, address and license No. of Manufacturer / Adviser (Health), State Urban Development Agency, Ilgus Supplier's from whom sample received Bhawan, II-C Block, Sec-III, Bidhan Nagar, Cal-91, 2. Reference No. and date of Manufacturer / Supplier's SUDA/120/96 (Pt-III) dt. Nil

Cotrimoxazole tablet I.P.

4	Detail of Raw material/final product in bulk / final product (in finished pack) as obtained from the manufacturer for analysis.	5X10's in strip
(a)	Original manufacturer's name (in case of raw	Kansas Laboratones Pvt. Ltd
	materials and drugs repacked)	8/1, Lal Bazar Street, Cal-1, DL No- 1446M
(b)	Batch Number	K 1463
(c)	Batch Size	Nil
(d)	Date of Manufacture if any	11/00
(e)	Date of Expiry if any	10/03
5.	Result of Analysis Recorded Below:	

As per I.P.'96

Description :- A white round tablet having a bisecting mark on one side .

Identification	Positive
Avg. wt. of a tablet	0.5714gm
Weight variation	Passes
Time of disintegration	6 minutes

Assay:--

	Found/ml 26	Claim/ml he
Trimethoprim I.P.	79.80mg	80.00mg
Sulphamethoxazole I.P.	391.37mg	400.00mg

in the Act and Rules made thereunder for the reasons given below.

In the opinion of the undersigned, the sample referred to above is of standard quality / is not of standard quality as defined

Signature of Officer In-Charge

A 9568

W. B. PHARMACEUTICAL & PHYTOCHEMICAL DEV. CORPN LID

(A Government of West Bengal Undertaking)

ACHARYA P. C. RAY R & D DRUG TESTING & POLLUTION TESTING LABORATORY

Behala Chowrasta, 620 Diamond Harbour Road, Calcutta 34, West Bengal

Report of Analysis [Form 39, See Rule 150 (f)]

License No: DL NO T/8-DCH dt: 27.10.80. & T/13-DCH dt: 24.9.90

Sample Receiving No:D/2162/00

be contained in the sample ...

Supplier's from whom sample received.

1.Name, address and license No. of Manufacturer /

2. Reference No. and date of Manufacturer / Supplier's

letter under which the sample was forwarded 3. Name of Drug / Cospnetics / Raw material purporting to

Dt of Receipt: 15.12.00

Adviser (Health), State Urban Development Agency, Ilgus Bhawan, II-C Block, Sec-III, Bidhan Nagar, Cal-91, SUDA/120/96 (Pt-III) dt. Nil

Commoxazole tablet 1.P

4. Detail of Raw material final product in bulk / final product (in finished pack) as obtained from the	5X10's in strip
manufacturer for analysis	
(a) Original manufacturer's name (in case of raw	Kansas Laboratories Pvt. Ltd
materials and drugs repacked)	8/1, Lal Bazar Street, Cal-1, DL No- 1446M
(b) Batch Number	K 1463
(c) Batch Size	Nil
(d) Date of Manufacture if any	11/00
(e) Date of Expiry if any	10/03

5. Result of Analysis Recorded Below:

As per I.P.'96

Description :- A white round tablet having a bisecting mark on one side

Identification	Positive
Avg. wt. of a tablet	0.5714gm
Weight variation	Passes
Time of disintegration	ó minutes

Assay:--

Trimethoprim I.P. Sulphamethoxazole I.P.

Found mil-6 79.80mg 391.37mg

Claim mi ho/ 80.00mg 400.00mg

In the opinion of the undersigned, the sample referred to above is of standard quality his not of standard quality as defined in the Act and Rules made thereunder for the reasons given below

Signature of Officer In-Charge

Sample Receiving No:D/214/00

A: 3532/01

W. B. PHARMACEUTICAL & PHYTOCHEMICAL DEV. CORPN LTD.

(A Government of West Bengal Undertaking)

ACHARYA P. C. RAY R & D DRUG TESTING & FOLLOTION TESTING LADORATORY

Behala Chowrasta, 620 Diamond Harbour Road, Calcutta 34, West Bengal

Report of Analysis [Form 39, See Rule 150 (f)]

License No: DL.NO.T/8-DCH dt: 27.10.80. & T/13-DCH dt: 24.9.90

Dt of Receipt: 15.12.00

PROPERTY F A DEMD A TENNS

1.Name, address and license No. of Manufacturer / Adviser (Health), State Urban Development Agency, Ilgus Bhawan, II-C Block, Sec-III, Bidhan Nagar, Cal-91, Supplier's from whom sample received. 2.Reference No. and date of Manufacturer / Supplier's SUDA/120/96 (Pt-III) dt. Nil letter under which the sample was forwarded 3.Name of Drug / Cosmetics / Raw material purporting to Merborin be contained in the sample /... 2X20gm app in polythene container 4.Detail of Raw material/final product in bulk / final product (in finished pack) as obtained from the manufacturer for analysis..... C. D. Pharmaceuticals (a) Original manufacturer's name (in case of raw 352/3/1, G.T.Road, (S), Howrah-3, Mfg Lics No.-44RP3 materials and drugs repacked)..... 377 (b) Batch Number Nil (c) Batch Size (d) Date of Manufacture if any..... 11/00 Not mentioned (e) Date of Expiry if any

5. Result of Analysis Recorded Below:

As per N.F

Description :- A green scales.

Identification	Positive
Solubility	Passes
Identification	Positive
Loss on drying	2.12%
Bromine Ion	Passes
Mercury Ion	Passes

Assav:--

Bromine Content

Mercury Content

Foundant 20.82% 24 95%

Limit 18% to 21.30% 24.00% to 26.70%

In the opinion of the undersigned, the sample referred to above is of standard quality / is not of standard quality as defined in the Act and Rules made thereunder for the reasons given below-.

Signature of Offic

Analyst / Junior Analyst /

UNTERVENT A DEAD ATTAD V

W. B. PHARMACEUTICAL & PHYTOCHEMICAL DEV. CORPN LTD.

(A Government of West Bengal Undertaking)

ACHARYA P. C. RAY R & D DRUG TESTING & FOLLOTION TESTING LADORATORY

Behala Chowrasta, 620 Diamond Harbour Road, Calcutta 34, West Bengal

Report of Analysis [Form 39, See Rule 150 (f)]

License No: DL NO T/8-DCH dt: 27 10 80 & T/13-DCH dt: 24 9 90

Sample Receiving No:D/21 4/00 Dt of Receipt: 15.12.00 1.Name, address and license No. of Manufacturer / Adviser (Health), State Urban Development Agency, Ilgus Supplier's from whom sample received. Bhawan, II-C Block, Sec-III, Bidhan Nagar, Cal-91, 2. Reference No. and date of Manufacturer / Supplier's SUDA/120/96 (Pt-III) dt. Nil letter under which the sample was forwarded 3. Name of Drug / Cospicetics / Raw material purporting to Merborin be contained in the sample / 4.Detail of Raw material/final product in bulk / final 2X20gm app in polythene container product (in finished pack) as obtained from the manufacturer for analysis..... (a) Original manufacturer's name (in case of raw C D Pharmaceuticals materials and drugs repacked) 352/3/1, G.T.Road, (S), Howrah-3, Mfg Lics No -44RP3 (b) Batch Number 277 (c) Batch Size Nil (d) Date of Manufacture if any..... 11.00 (e) Date of Expiry if any Not mentioned

5. Result of Analysis Recorded Below:

As per NF

Description :- A green scales.

Identification	Positive
Solubility	Passes
Identification	Positive
Loss on drying	2.12%
Bromine Ion	Passes
Mercury Ion	Passes

Assay:--

Bromine Content Mercury Content Foundard 47 20.82% 24.95%

Limit 18% to 21.30% 24.00% to 26.70%

In the opinion of the undersigned, the sample referred to above is of standard quality *is not of standard quality* as defined in the Act and Rules made thereunder ter the reasons given below.

Signature of Officer In

Sample Receiving No:D/2140/00

A/ 3574/01

W. B. PHARMACEUTICAL & PHYTOCHEMICAL DEV. CORPN LTD.

(A Government of West Bengal Undertaking)

ACHARYA P. C. RAY R & D DRUG TESTING & POLLUTION TESTING LABORATORY

Behala Chowrasta, 620 Diamond Harbour Road, Calcutta 34, West Bengal

Report of Analysis [Form 39, See Rule 150 (f)]

License No: DL.NO.T/8-DCH dt: 27.10.80. & T/13-DCH dt: 24.9.90

Dt of Receipt: 15.12.00

1.Name, address and license No. of Manufacturer /	Adviser (Health), State Urban Development Agency, Ilgus
Supplier's from whom sample received	Bhawan, II-C Block, Sec-III, Bidhan Nagar, Cal-91,
2.Reference No. and date of Manufacturer / Supplier's	SUDA/120/96 (Pt-III) dt. Nil
letter under which the sample was forwarded	
3.Name of Drug / Cosmetics / Raw material purporting to	Fesolic- L
be contained in the sample	
4. Detail of Raw material/final product in bulk / final product (in finished pack) as obtained from the	50 tabs
manufacturer for analysis	
(a) Original manufacturer's name (in case of raw	Pure Pharma Ltd, 41,42 &44, Industrial Estate,
materials and drugs repacked)	(Pologround), Indore-452015, Mfg Lics No30/64
(b) Batch Number	0017
(c) Batch Size	Nil
(d) Date of Manufacture if any	10/00
(e) Date of Expiry if any	09/02

5. Result of Analysis Recorded Below:

As per I.P.'96, Vol-I

Description :-Brick red shaped blunt edge film coated tablets in blister pack.

Identification	Positive
Avg. wt. of a tablet	0.1102777gm
Weight variation	Passes
Time of disintegration	21 minutes

Assay:--

Folic Acid

Dried Ferrous Sulphate I.P.

0.0573gm (85.61%) 0.45mg (90%)

Found/tab

<u>Claim/tab</u> 0.067gm

0.5mg

Limit

80% to 90%

Not less than 90%

In the opinion of the undersigned, the sample referred to above is of standard quality / is not of standard quality as defined in the Act and Rules made thereunder for the reasons given below.

Analyst / Junior Analyst /

Signature of Officer Ih-Charg

A: 3574/01

W. B. PHARMACEUTICAL & PHYTOCHEMICAL DEV. CORPN LTD.

(A Government of West Bengal Undertaking)

ACHARYA P. C. RAY R & D DRUG TESTING & POLLUTION TESTING LABORATORY

Behala Chowrasta, 620 Diamond Harbour Road, Calcutta 34, West Bengal

Report of Analysis [Form 39, See Rule 150 (f)]

License No: DL NO.T/8-DCH dt: 27.10.80. & T/13-DCH dt: 24.9.90

Sample Receiving No:D/2140/00	Dt of Receipt: 15.12.00
1.Name, address and license No. of Manufacturer /	Adviser (Health), State Urban Development Agency, Ilgus
Supplier's from whom sample received	Bhawan, II-C Block, Sec-III, Bidhan Nagar, Cal-91,
2.Reference No. and date of Manufacturer / Supplier's	SUDA/120/96 (Pt-111) dt. Nil
letter under which the sample was forwarded	
3.Name of Drug / Cosmetics / Raw material purporting to	Fesolic- L
be contained in the sample	
4.Detail of Raw material/final product in bulk / final	50 tabs
product (in finished pack) as obtained from the	
manufacturer for analysis	
(a) Original manufacturer's name (in case of raw	Pure Pharma Ltd, 41,42 &44, Industrial Estate.
materials and drugs repacked)	(Pologround), Indore-452015, Mfg Lics No30/64
(b) Batch Number	0017
(c) Batch Size	Nil
(d) Date of Manufacture if any	10/00
(e) Date of Expiry if any	09/02
5. Result of Analysis Recorded Below:	

As per I.P.'96, Vol-I

Description :-Brick red shaped blunt edge film coated tablets in blister pack.

Identification	Positive
Avg. wt. of a tablet	0.1102777gm
Weight variation	Passes
Time of disintegration	21 minutes

Assay:--

Dried Ferrous Sulphate I.P.

0.067gm

Limit

80% to 90%

Not less than 90%

Folic Acid

0.45mg (90%) 0.5mg

In the opinion of the undersigned, the sample referred to above is of standard quality / is not of standard quality as defined in the Act and Rules made thereunder for the reasons given below.

Signature of Officer In

Analyst / Junior Analyst /

Found/tab Claim/tab 0.0573gm (85.61%)

AI 257

W. B. PHARMACEUTICAL & PHYTOCHEMICAL DEV. CORPN LTD.

(A Government of West Bengal Undertaking)

ACHARYA P. C. RAY R & D DRUG TESTING & POLLUTION TESTING LABORATORY

Behala Chowrasta, 620 Diamond Harbour Road, Calcutta 34, West Bengal

Report of Analysis [Form 39, See Rule 150 (f)]

License No: DL NO. T/8-DCH dt: 27.10.80. & T/13-DCH dt: 24.9.90

Sample Receiving No:D/2141/00	Dt of Receipt: 15.12.00
1. Name, address and license No. of Manufacturer /	Adviser (Health), State Urban Development Agency, Ilgus
Supplier's from whom sample received	Bhawan, II-C Block, Sec-III, Bidhan Nagar, Cal-91,
2. Reference No. and date of Manufacturer / Supplier's	SUDA/120/96 (Pt-III) dt. Nil
letter under which the sample was forwarded	
3.Name of Drug / Cosmetics / Raw material purporting to	Fesohic- S
be contained in the sample	
4.Detail of Raw material/final product in bulk / final	50 tabs
product (in finished pack) as obtained from the	
manufacturer for analysis	
(a) Original manufacturer's name (in case of raw	Pure Pharma Ltd, 41,42 &44, Industrial Estate,
materials and drugs repacked)	(Pologround), Indore-452015, Mfg Lics No30/64
(b) Batch Number	0009
(c) Batch Size	Nil
(d) Date of Manufacture if any	10/00
(e) Date of Expiry if any.	09/02
5.Result of Analysis Recorded Below:	

As per I.P.'96, Vol-I

Description :-Brick red shaped blunt edge film coated tablets in blister pack.

Identification	Positive
Avg. wt. of a tablet	0.53705gm
Weight variation	Passes
Time of disintegration	18 minutes

Assay:--

	Found/tab	Claim/tab	Limit
Dried Ferrous Sulphate I.P.	0.27336gm	0.335gm	80% to 90%
Folic Acid	0.091mg (91%)	0.1mg	Not less than 90%

In the opinion of the undersigned, the sample referred to above is of standard quality / is not of standard quality as defined in the Act and Rules made thereunder for the reasons given below.

Anayst Junior Analyst /

Signature of Officer In-Charge

Sample Receiving No:D/2141/00

A. 35

W. B. PHARMACEUTICAL & PHYTOCHEMICAL DEV. CORPN LTD.

(A Government of West Bengal Undertaking)

ACHARYA P. C. RAY R & D DRUG TESTING & POLLUTION TESTING LABORATORY

Behala Chowrasta, 020 Diamond Harbour Road, Calcutta 34, West Bengal

Report of Analysis [Form 39, See Rule 150 (f)]

License No: DL NO T/8-DCH dt: 27 10 80. & T/13-DCH dt: 24 9 90

Dt of Receipt: 15.12.00

I.Name, address and license No. of Manufacturer / Supplier's from whom sample received.....

2. Reference No. and date of Manufacturer / Supplier's

Adviser (Health), State Urban Development Agency, Ilgus Bhawan, II-C Block, Sec-III, Bidhan Nagar, Cal-91, SUDA/120/96 (Pt-III) dt. Nil

letter under which the sample was forwarded 3.Name of Drug / Cosmetics / Raw material purporting to be contained in the sample.....

4. Detail of Raw material/final product in bulk / final product (in finished pack) as obtained from the manufacturer for analysis.

- (a) Original manufacturer's name (in case of raw materials and drugs repacked).(b) Batch Number
- (c) Batch Size
- (d) Date of Manufacture if any.....

(e) Date of Expiry if any.....

5. Result of Analysis Recorded Below:

Fesolic-S

50 tabs

Pure Pharma Ltd, 41,42 &44, Industrial Estate, (Pologround), Indore-452015, Mfg Lics No.-30/64 0009 Nil 10/00 09/02

As per I.P.'96, Vol-I

Description :-Brick red shaped blunt edge film coated tablets in blister pack.

IdentificationPositiveAvg. wt. of a tablet0.53705gmWeight variationPassesTime of disintegration18 minutes

Assay:--

Found/tabClaim/tabDried Ferrous Sulphate I.P.0.27336gm0.335gmFolic Acid0.091mg (91%)0.1mg

Limit 80% to 90% Not less than 90%

In the opinion of the undersigned, the sample referred to above is of standard quality - is not of standard quality as defined in the Act and Rules made thereunder for the reasons given below.

mior Analyst

Signature of Officer In

Sample Receiving No:D/2146/00

35-JS/

W. B. PHARMACEUTICAL & PHYTOCHEMICAL DEV. CORPN LTD.

(A Government of West Bengal Undertaking)

ACHARYA P. C. RAY R & D DRUG TESTING & POLLUTION TESTING LABORATORY

Behala Chowrasta, 620 Diamond Harbour Road, Calcutta 34, West Bengal

Report of Analysis [Form 39, See Rule 150 (f)]

License No: DL.NO.T/8-DCH dt: 27.10.80. & T/13-DCH dt: 24.9.90

Dt of Receipt: 15.12.00

1.Name, address and license No. of Manufacturer	/ Adviser (Health), State Urban Development Agency, Ilgus
Supplier's from whom sample received	Bhawan, II-C Block, Sec-III, Bidhan Nagar, Cal-91,
2.Reference No. and date of Manufacturer / Supplier'	s SUDA-120/96 (Pt-II)/633 dt. 24.12.00
letter under which the sample was forwarded	
3.Name of Drug / Cosmetics / Raw material purporting to	D Antiseptic Lotion
be contained in the sample	
4.Detail of Raw material/final product in bulk / fina	2X200ml
product (in finished pack) as obtained from th	e
manufacturer for analysis	
(a) Original manufacturer's name (in case of raw	W Hindusthan Drugs
materials and drugs repacked)	129, Shyam Nagar Road, Cal- 700 055
(b) Batch Number	LE 06
(c) Batch Size	Nil
(d) Date of Manufacture if any	. 04/99
(e) Date of Expiry if any	03/04
5. Result of Analysis Recorded Below:	

As per I.P.'96, Page- 87

Description :- A light brown coloured liquid.

Identification

Positive

Found

2.10%

Assay:--

Benzal Konium Chloride

<u>Claim</u> 2.00%

In the opinion of the undersigned, the sample referred to above is of standard quality / is not of standard quality as defined in the Act and Rules made thereunder for the reasons given below. with respect of the above tests only.

Signature of Officer In-

A135750

Computerised File Name: D&Creport2078

W. B. PHARMACEUTICAL & PHYTOCHEMICAL DEV. CORPN LTD.

(A Government of West Bengal Undertaking)

ACHARYA P. C. RAY R & D DRUG TESTING & POLLUTION TESTING LABORATORY

Behala Chowrasta, 620 Diamond Harbour Road, Calcutta 34, West Bengal

Report of Analysis [Form 39, See Rule 150 (f)]

License No: DL.NO.T/8-DCH dt: 27.10.80. & T/13-DCH dt: 24.9.90

Sample Receiving No:D/2146/00	Dt of Receipt: 15.12.00
1.Name, address and license No. of Manufacturer /	Adviser (Health), State Urban Development Agency, Ilgus
Supplier's from whom sample received.	Bhawan, II-C Block, Sec-III, Bidhan Nagar, Cal-91,
2. Reference No. and date of Manufacturer / Supplier's	SUDA-120/96 (Pt-II)/633 dt. 24.12.00
letter under which the sample was forwarded	
3.Name of Drug / Cospetics / Ray material purporting to	Antiseptic Lotion
be contained in the sample	
4. Detail of Raw material/final product in bulk / final	2X200ml
product (in finished pack) as obtained from the	
manufacturer for analysis	
(a) Original manufacturer's name (in case of raw	Hindusthan Drugs
materials and drugs repacked)	129,Shyam Nagar Road, Cal- 700 055
(b) Batch Number	LE 06
(c) Batch Size	Nil .
(d) Date of Manufacture if any	04/99
(e) Date of Expiry if any	03/04
5. Result of Analysis Recorded Below:	
As per I.P.'96, Page- 87	

Description :- A light brown coloured liquid.

Identification

Positive

Assay:--

Benzal Konium Chloride

Found 2.10% <u>Claim</u> 2.00%

In the opinion of the undersigned, the sample referred to above is of standard quality *is not of standard quality as defined* in the Act and Rules made thereunder for the reasons given below .with respect of the above tests only.

Signature of Officer In-Charge

Benzyl Benzoate Application 1P.

· B.m - 106974

Ni tro furazone Cream B. mo - NF 1234

Rel no. pp&c/kab/1012 de 21-12-2000

A/ 32-15
HYTOCHEMICAL DEV. CORFN LTD.
West Bengal Undertaking)
ING & POLLUTION TESTING LABORATORY
Jarbour Road, Calcutta 34, West Bengal
orm 39, See Rule 150 (f)]
lt: 27.10.80. & T/13-DCH dt: 24.9.90
Dt of Receipt: 15.12.00
Adviser (Health), State Urban Development Agency, Ilgu
Bhawan, II-C Block, Sec-III, Bidhan Nagar, Cal-91,
SUDA-15/98 (Pt-II)/633 dt. 24.11.00
Benzyl Benzoate Application I.P.
2X100ml
Hindusthan Drugs, 139, Shyarn Nagar Road, Cal-700 056
DL No1069M
11
Nil
11/00
10/02

Description :- White viscous emulsion in glass bottle

Identification

Positive

Assay:--

Benzyl Benzoate I.P.

Found 25.89% Limit 22.5% to 27.5%

In the opinion of the undersigned, the sample referred to above is of standard quality / is not of standard quality as defined in the Act and Rules made thereunder for the reasons given below.

Claim

25.0%

Augustutte

Signature of Officer I

A. 945/n

W. B. PHARMACEUTICAL & PHYTOCHEMICAL DEV. CORPN LTD.

(A Government of West Bengal Undertaking)

ACHARYA P. C. RAY R & D DRUG TESTING & POLLUTION TESTING LABORATORY

Behala Chowrasta, 620 Diamond Harbour Road, Calcutta 34, West Bengal

Report of Analysis [Form 39, See Rule 150 (f)]

License No: DL.NO.T/8-DCH dt: 27.10.80. & T/13-DCH dt: 24.9.90

Sample Receiving No:D/2153/00	Dt of Receipt: 15.12.00
1.Name, address and license No. of Manufacturer /	Adviser (Health), State Urban Development Agency, Ilgus
Supplier's from whom sample received	Bhawan, II-C Block, Sec-III, Bidhan Nagar, Cal-91,
2.Reference No. and date of Manufacturer / Supplier's	SUDA-15/98 (Pt-II)/633 dt. 24.11.00
letter under which the sample was forwarded	
3. Name of Drug / Cosmetics / Raw praterial purporting to	Benzyl Benzoate Application I.P.
be contained in the sample	Butes i Delizone Application I.P.
4.Detail of Raw material/final/product in bulk / final	2X100ml
product (in finished pack) as obtained from the	2210011
manufacturer for any here	
(a) Original manufacturer's name (in case of raw	
	Hindusthan Drugs, 139, Shyam Nagar Road, Cal-700 056
materials and drugs repacked)	DL No1069M
(b) Batch Number	11
(c) Batch Size	Nil
(d) Date of Manufacture if any	11/00
(e) Date of Expiry if any	10/02
5. Result of Analysis Recorded Below:	
Meth	od : I.P.
Description :- White viscous emulsion in glass bottle	
Identification Positive	

Assay ---

Benzyl Benzoate I.P.

Found 25.89% Claim 25.0%

Limit

22.5% to 27.5%

In the opinion of the undersigned, the sample referred to above is of standard quality / is not of standard quality as defined in the Act and Rules made thereunder for the reasons given below .

Autorfabil

land atin for Signature of Officer In-Charg

326/N

W. B. PHARMACEUTICAL & PHYTOCHEMICAL DEV. CORPN LTD.

(A Government of West Bengal Undertaking)

ACHARYA P. C. RAY R & D DRUG TESTING & POLLUTION TESTING LABORATORY

Behala Chowrasta, 620 Diamond Harbour Road, Calcutta 34, West Bengal

Report of Analysis [Form 39, See Rule 150 (f)]

License No: DL.NO.T/8-DCH dt: 27.10.80. & T/13-DCH dt: 24.9.90

Dt of Receipt: 15.12.00 Sample Receiving No:D/2141/00 1.Name, address and license No. of Manufacturer / Adviser (Health), State Urban Development Agency, Ilgus Bhawan, II-C Block, Sec-III, Bidhan Nagar, Cal-91, Supplier's from whom sample received SUDA-15/98 (Pt-II)/633 dt. 24.11.00 2. Reference No. and date of Manufacturer / Supplier's letter under which the sample was forwarded 3. Name of Drug / Cosmetics / Raw material purporting to Nitrofurazone cream U.S.P. be contained in the sample 10X15gm 4.Detail of Raw material/final product in bulk / final product (in finished pack) as obtained from the manufacturer for analysis (a) Original manufacturer's name (in case of raw Pilco Pharma Pvt. Ltd. 123/37, Suresh Bagh, Kanpur- 208 072, DL No - 21 of 88 materials and drugs repacked)..... (b) Batch Number NF 1234 (c) Batch Size Nil 11/00 (d) Date of Manufacture if any. 10/02 (e) Date of Expiry if any 5. Result of Analysis Recorded Below:

Method : U.S.P., XXIII

Description :- White soft creamy base in metal tube.

Identification

Positive

Assay:--

Nitrofurazone

Found 0.22% Claim

0.2%

Limit

90% to 110%

In the opinion of the undersigned, the sample referred to above is of standard quality is not of standard quality as defined in the Act and Rules made thereunder for the reasons given below. With a lura my

Inalyst/Junior Analyst/ Photochul

Signature of Officer In

W. B. PHARMACEUTICAL & PHYTOCHEMICAL DEV. CORPN LTD. (A Government of West Bengal Undertaking)

ACHARYA P. C. RAY R & D DRUG TESTING & POLLUTION TESTING LABORATORY

Behala Chowrasta, 620 Diamond Harbour Road, Calcutta 34, West Bengal

Report of Analysis [Form 39, See Rule 150 (f)]

License No: DL.NO.T/8-DCH dt: 27.10.80. & T/13-DCH dt: 24.9.90

Sample Receiving No:D/2141/00	Dt of Receipt: 15.12.00
1.Name, address and license No. of Manufacturer /	Adviser (Health), State Urban Development Agency, Ilgus
Supplier's from whom sample received	Bhawan, II-C Block, Sec-III, Bidhan Nagar, Cal-91,
2.Reference No. and date of Manufacturer / Supplier's	SUDA-15/98 (Pt-II)/633 dt. 24.11.00
letter under which the sample was forwarded	
3.Name of Drug / Cosmetics / Raw material purporting to	Nitrofurazone cream U.S.P.
be contained in the sample	
4.Detail of Raw material/Anal product in bulk / final	10X15gm
product (in finished pack) as obtained from the	
manufacturer for analysis.	
(a) Original manufacturer's name (in case of raw	Pilco Phanna Pvt. Ltd
materials and drugs repacked)	123/37, Suresh Bagh, Kanpur- 208 072, DL No - 21 of 88
(b) Batch Number	NF 1234
(c) Batch Size	Nil
(d) Date of Manufacture if any	11/00
(e) Date of Expiry if any	10/02

5.Result of Analysis Recorded Below:

Method : U.S.P., XXIII

Description :- White soft creamy base in metal tube.

Identification

Positive

Assay:--

Nitrofurazone

Found 0.22% Limit 90% to 110%

In the opinion of the undersigned, the sample referred to above is of standard quality / is not of standard quality as defined in the Act and Rules made thereunder for the reasons given below. With a lura his

Claim

0.2%

Tab. Mehonidazole-200 2009 K. -1472 Tab. Bromhexine 8mg. G 1414 Tab. Paracetamil-K-1459 K-1459 K-1460 k = 1468
	mi PPSe /Lab / 1056 dt 4-1-20 Receiver on Receiver on 2001 Computerised File Name: D&CrepOrtisco 1 2001	0 1
	opge and som	3 *
. ;	mit. Receiver - 101/2001	~ /
p.º	Computerised File Name: D&CrepOrt1805	A. 3326/01
	W. B. PHARMACEUTICAL & PH	YTOCHEMICAL DEV. CORPN LTD.
-	(A Government of W	est Bengal Undertaking)
-)	ACHARYA P. C. RAY R & D DRUG TESTI	NG & POLLUTION TESTING LABORATORY
7/	Behala Chowrasta, 620 Diamond Ha	arbour Road, Calcutta 34, West Bengal
	Report of Analysis [Fo	rm 39, See Rule 150 (f)]
	License No: DL NO T/8-DCH dt	27.10.80. & T/13-DCH dt: 24.9.90
	Sample Receiving No:D/2362/00	Dt of Receipt: 26.12.00
	1.Name, address and license No. of Manufacturer /	Adviser (Health), State Urban Development Agency, Ilgue
	Supplier's from whom sample received	Bhawan, II-C Block, Sec-III, Bidhan Nagar, Cal-91,
	2.Reference No. and date of Manufacturer / Supplier's	SUDA/120/96 (Pt-III) dt. Nil
	letter under which the sample was forwarded	
	3.Name of Drug / Cosmetics / Ray material purporting to	Metronidazole tab I.P. 200mg
	be contained in the sample	
	4.Detail of Raw material/fings product in bulk / final	50X2
	product (in finished pack) as obtained from the	
	manufacturer for analysis	
	(a) Original manufacturer's name (in case of raw	Kansas Laboratories Pvt. Ltd
	materials and drugs repacked).	8/1, Lal Bazar Street, Cal-700 001, DL No DL 1446M
	(b) Batch Number	K-1472
	(c) Batch Size	Nil
	(d) Date of Manufacture if any	11/00
	(e) Date of Expiry if any	10/03
	5 Result of Analysis Recorded Below:	

Method :As per I.P' 96.

Description :- A small size orange yellow colour circular film coated tablet, with a break mark on one side of each tablet.

Identification	Positive
Avg. wt. of a tablet	0.3551gm
Uniformity of weight	Passes
Related substances	Passes
Dissolution	Passes

Assay:--

Metronidazole I.P.

Found/tab 206.5mg Claim/tab 200.00mg

190mg to 210mg

Limit

In the opinion of the undersigned, the sample referred to above is of standard quality / is not of standard quality as defined in the Act and Rules made thereunder for the reasons given below

Anal

Sig nature of Offi

N3326/17

W. B. PHARMACEUTICAL & PHYTOCHEMICAL DEV. CORPN LTD.

(A Government of West Bengal Undertaking)

ACHARYA P. C. RAY R & D DRUG TESTING & POLLUTION TESTING LABORATORY

Behala Chowrasta, 620 Diamond Harbour Road, Calcutta 34, West Bengal

Report of Analysis [Form 39, See Rule 150 (f)]

License No: DL NO. T/8-DCH dt: 27.10.80. & T/13-DCH dt: 24.9.90

Dt of Receipt: 26.12.00

Sample Receiving No:D/2362/00	Dt of Receipt: 26.12.00
1.Name, address and license No. of Manufacturer /	Adviser (Health), State Urban Development Agency, Ilgu
Supplier's from whom sample received	Bhawan, II-C Block, Sec-III, Bidhan Nagar, Cal-91,
2.Reference No. and date of Manufacturer / Supplier's	SUDA/120/96 (Pt-III) dt. Nil
letter under which the sample was forwarded	
3.Name of Drug / Cosmetics / Raw material purporting to	Metronidazole tab I.P. 200mg
be contained in the sample	
4.Detail of Raw material/final product in bulk / final	50X2
product (in finished pack) as obtained from the	
manufacturer for analysis.	
(a) Original manufacturer's name (in case of raw	Kansas Laboratories Pvt. Ltd
materials and drugs repacked).	8/1, Lal Bazar Street, Cal-700 001, DL No DL 1446M
(b) Batch Number	K-1472
(c) Batch Size	Nil
(d) Date of Manufacture if any	11/00
(e) Date of Expiry if any	10/03
5. Result of Analysis Recorded Below:	

Method :As per I.P' 96.

Description :- A small size orange yellow colour circular film coated tablet, with a break mark on one side of each tablet.

Identification	Positive
Avg. wt. of a tablet	0.3551gm
Uniformity of weight	Passes
Related substances	Passes
Dissolution	Passes

Assay:--

Metronidazole 1 P.

Found/tab 206.5mg

Claim/tab 200.00mg

190mg to 210mg

Limit

In the opinion of the undersigned, the sample referred to above is of standard quality Lis not of standard quality as defined in the Act and Rules made thereunder for the reasons given below-

Signature of

A 3327/01

ligus

W. B. PHARMACEUTICAL & PHYTOCHEMICAL DEV. CORPN LTD

(A Government of West Bengal Undertaking)

ACHARYA P. C. RAY R & D DRUG TESTING & POLLUTION TESTING LABORATORY

Behala Chowrasta, 620 Diamond Harbour Road. Calcutta 34, West Bengal

Report of Analysis [Form 39, See Rule 150 (f)]

License No: DI. NO T/8-DCH dt: 27.10.80. & T/13-DCH dt: 24.9.90

Dt of Receipt: 15.12.00

Sample Receiving No:D/2156/00	Dt of Receipt: 15.12.00
I.Name, address and license No. of Manufacturer /	Adviser (Health), State Urban Development Agency, I
Supplier's from whom sample received.	Bhawan, II-C Block, Sec-III, Bidhan Nagar, Cal-91,
2.Reference No. and date of Manufacturer / Supplier's	SUDA/120/96 (Pt-II)/631 dt. 24.11.00
letter under which the sample was forwarded	
3.Name of Drug / Cosmetics / Raw material purporting to	Bromhexine Tablet B.P. 8mg
be contained in the sample	
4.Detail of Raw material/final product in bulk / final	10 tabs \times 5 strips
product (in finished pack) as obtained from the	
manufacturer for analysis.	
(a) Original manufacturer's name (in case of raw	Kansas Laboratories Pvt. Ltd
materials and drugs repacked)	8/1, Lal Bazar Street, Cal-700 001, DL No DL 1446M
(b) Batch Number	G1414
(c) Batch Size	Nil
(d) Date of Manufacture if any	07/00
(e) Date of Expiry if any	06/03
5. Result of Analysis Recorded Below:	

Method :As per I.P' 96, Vol-I

Description:-Small white round tablet having a bisecting mark on one side.

Identification	Positive
Avg. wt. of a tablet	0.1715gm
Weight variation	Passes
Disintegration time	1 min
Uniformity of content	Passes
Related substances	Passes

Assay:--

Found/tab

Claim/tab

Bromhexine HCL I.P.

Junior Analyst /

7.88mg

8.00mg

In the opinion of the undersigned, the sample referred to above is of standard quality / is not of standard quality as defined

in the Act and Rules made thereunder for the reasons given below .

Signature of C harge

Sample Receiving No:D/2156/00

A1 732/19 W. B. PHARMACEUTICAL & PHYTOCHEMICAL DEV. CORPN LTD.

(A Government of West Bengal Undertaking)

ACHARYA P. C. RAY R & D DRUG TESTING & POLLUTION TESTING LABORATORY

Behala Chowrasta, 620 Diamond Harbour Road, Calcutta 34, West Bengal

Report of Analysis [Form 39, See Rule 150 (f)]

License No: DL NO T/8-DCH dt: 27.10.80 & T/13-DCH dt: 24.9.90

Dt of Receipt: 15.12.00

	Lit of acoust. 10.12.00
1.Name, address and license No. of Manufacturer /	Adviser (Health), State Urban Development Agency, Ilgus
Supplier's from whom sample received	Bhawan, II-C Block, Sec-III, Bidhan Nagar, Cal-91,
2.Reference No. and date of Manufacturer / Supplier's	SUDA/120/96 (Pt-II)/631 dt. 24.11.00
letter under which the sample was forwarded	
3.Name of Drug / Cosmetics / Raw material purporting to	Bromhexine Tablet B.P. 8mg
be contained in the sample	
4.Detail of Raw material/final product in bulk / final	10 tabs X 5 strips
product (in finished pack) as obtained from the	
manufacturer for analysis	
(a) Original manufacturer's name (in case of raw	Kansas Laboratories Pvt. Ltd.
materials and drugs repacked)	8/1, Lal Bazar Street, Cal-700 001, DL No DL 1446M
(b) Batch Number	G1414
(c) Batch Size	Nil
(d) Date of Manufacture if any	07/00
(e) Date of Expiry if any	06/03
5. Result of Analysis Recorded Below:	

Method :As per I.P' 96, Vol-I

Description:-Small white round tablet having a bisecting mark on one side.

Identification	Positive
Avg. wt. of a tablet	0.1715gm
Weight variation	Passes
Disintegration time	l min
Uniformity of content	Passes
Related substances	Passes

Assay:--

Bromhexine HCL I.P.

Analyst/ Junior Analyst /

Found/tab 7.88mg

Claim/tab 8.00mg

In the opinion of the undersigned, the sample referred to above is of standard quality A is not of standard quality as defined in the Act and Rules made thereunder for the reasons given below ...

ignature of Offi r In-Charge

S 1

2

3

4

iame: D&Creport1869 W. B. PHARMACEUTICAL & PHYTOCHEMICAL DEV. CORPN LTD. (A Government of West Report Linderty)

ACHARYA P. C. RAY R & D DRUG TESTING & POLLUTION TESTING LABORATORY

Behala Chowrasta, 620 Diamond Harbour Road, Calcutta 34, West Bengal

Report of Analysis [Form 39, See Rule 150 (f)]

License No: DL NO. T/8-DCH dt: 27.10.80. & T/13-DCH dt: 24.9.90

Sample Receiving No:D/2160/00	Dt of Receipt: 15.12.00
Name, address and license No. of Manufacturer /	Adviser (Health), State Urban Development Agency, llgus
Supplier's from whom sample received	Bhawan, II-C Block, Sec-III, Bidhan Nagar, Cal-91,
2.Reference No. and date of Manufacturer / Supplier's	SUDA/120/96 (Pt-III)/633 dt. 24.12.00
letter under which the sample was forwarded	
3. Name of Drug / Cospeties / Ray material purporting to	Paracetamol tablets I.P.
be contained in the sample	
4.Detail of Raw material/final product in bulk / final	10 tabs X 5 strips
product (in finished pack) as obtained from the	
manufacturer for analysis	
(a) Original manufacturer's name (in case of raw	Kansas Laboratories Pvt. Ltd
materials and drugs repacked).	8/1, Lal Bazar Street, Cal-700 001, DL No DL 1446M
(b) Batch Number	K-1468
(c) Batch Size	Nil
(d) Date of Manufacture if any	11/00 .
(e) Date of Expiry if any	10/03
5 Result of Analysis Recorded Below:	
Method :/	As per I.P' 96.
Description . White almular second tablet in blister neals	

Description :-White circular scored tablet in blister pack.

Identification	Positive
Avg. wt. of a tablet	0.5704gm
Weight variation	Passes
Dissolution test	Passes

Assay:--

Paracetamol I.P.

Found/tab 479.0mg

Claim/tab 500.0mg

Limit 95% to 105%

In the opinion of the undersigned, the sample referred to above is of standard quality / is not of standard quality as defined in the Act and Rules made thereunder for the reasons given below...

Analyst/Junior Analyst/ PUSerful

Sign

A13324/01

W. B. PHARMACEUTICAL & PHYTOCHEMICAL DEV. CORPN LTD.

(A Government of West Bengal Undertaking)

ACHARYA P. C. RAY R & D DRUG TESTING & POLLUTION TESTING LABORATORY

Behala Chowrasta, 620 Diamond Harbour Road, Calcutta 34, West Bengal

Report of Analysis [Form 39, See Rule 150 (f)]

License No: DL.NO.T/8-DCH dt: 27.10.80. & T/13-DCH dt: 24.9.90

Sample	Receiving	No:D/2160/00	0
--------	-----------	--------------	---

Dt of Receipt: 15.12.00 Adviser (Health), State Urban Development Agency, Ilgus wan, II-C Block, Sec-III, Bidhan Nagar, Cal-91,

A A A A A A A A A A A A A A A A A A A	
1.Name, address and license No. of Manufacturer /	Advi
Supplier's from whom sample received	Bhav
2. Reference No. and date of Manufacturer / Supplier's	SUD
letter under which the sample was forwarded	
3.Name of Drug / Cosmetics / Raw material purporting to	Parad
be contained in the sample	
4.Detail of Raw material/final product in bulk / final	10 ta
product (in finished pack) as obtained from the	
manufacturer for analysis	
(a) Original manufacturer's name (in case of raw	Kan
materials and drugs repacked).	8/1,
(b) Batch Number	K-14
(c) Batch Size	Nil
(d) Date of Manufacture if any	11/0
(e) Date of Expiry if any	10/0
5. Result of Analysis Recorded Below:	

DA/120/96 (Pt-III)/633 dt. 24.12.00

cetamol tablets I.P.

abs X 5 strips

isas Laboratories Pvt. Ltd Lal Bazar Street, Cal-700 001, DL No. - DL 1446M 468 00 03

Method :As per I.P' 96.

Description :-White circular scored tablet in blister pack.

Identification	Positive
Avg. wt. of a tablet	0.5704gm
Weight variation	Passes
Dissolution test	Passes

Assay .--

Paracetamol I.P.

Found/tab 479.0mg

plasenfutur.

Claim/tab 500.0mg

Limit 95% to 105%

In the opinion of the undersigned, the sample referred to above is of standard quality / is not of standard quality as defined in the Act and Rules made thereunder for the reasons given below.

Signature of Off

Analyst / Junior Analyst /

W. B. PHARMACEUTICAL & PHYTOCHEMICAL DEV. CORPN LTD. 3323/0/ (A Government of West Bergel Lindert

ACHARYA P. C. RAY R & D DRUG TESTING & POLLUTION TESTING LABORATORY

Behala Chowrasta, 620 Diamond Harbour Road, Calcutta 34, West Bengal

Report of Analysis [Form 39, See Rule 150 (f)]

License No: DL NO. T/8-DCH dt: 27.10.80 & T/13-DCH dt: 24.9.90

Dt of Receipt: 15.12.00

Sample Receiving No:D/2159/00 Adviser (Health), State Urban Development Agency, Ilgus I.Name, address and license No. of Manufacturer / Bhawan, II-C Block, Sec-III, Bidhan Nagar, Cal-91, Supplier's from whom sample received SUDA/120/96 (Pt-III)/633 dt. 24.12.00 2. Reference No. and date of Manufacturer / Supplier's letter under which the sample was forwarded 3.Name of Drug / Cospletics / Raw material purporting to Paracetamol tablets I.P. be contained in the sample 10 tabs X 5 strips 4. Detail of Raw material/and product in bulk / final product (in finished pack) as obtained from the manufacturer for analysis..... (a) Original manufacturer's name (in case of raw materials and drugs repacked) (b) Batch Number (c) Batch Size (d) Date of Manufacture if any (e) Date of Expiry if any

Kansas Laboratories Pvt. Ltd 8/1, Lal Bazar Street, Cal-700 001, DL No. - DL 1446M K-1460 Nil 11/00 10/03

5. Result of Analysis Recorded Below:

Method :As per I.P' 96.

Description :- White circular scored tablet in blister pack.

Identification	Positive
Avg. wt. of a tablet	0.5735gm
Weight variation	Passes
Dissolution test	Passes

Assay .--

Paracetamol I P.

Found/tab 509.79mg Claim/tab 500.0mg

Limit 95% to 105%

In the opinion of the undersigned, the sample referred to above is of standard quality / is not of standard quality as defined in the Act and Rules made thereunder for the reasons given below-

Analyst/Junior Analyst/ Phile-fubl-

A1 3323/07

W. B. PHARMACEUTICAL & PHYTOCHEMICAL DEV. CORPN LTD.

(A Government of West Bengal Undertaking)

ACHARYA P. C. RAY R & D DRUG TESTING & POLLUTION TESTING LABORATORY

Behala Chowrasta, 620 Diamond Harbour Road, Calcutta 34, West Bengal

Report of Analysis [Form 39, See Rule 150 (f)]

License No: DL NO. T/8-DCH dt: 27.10.80. & T/13-DCH dt: 24.9.90

	Dt of	Receip	t: 15.12.00		
Adviser	(Health),State	Urban	Development	Agency,	llgus

Bhawan, H-C Block, Sec-III, Bidhan Nagar, Cal-91,

SUDA/120/96 (Pt-III)/633 dt. 24.12.00

Paracetamol tablets I.P.

10 tabs X 5 strips

Sample Receiving No:D/2159/00
1.Name, address and license No. of Manufacturer /
Supplier's from whom sample received
2.Reference No. and date of Manufacturer / Supplier's
letter under which the sample was forwarded
3.Name of Drug / Cospietics / Ray material purporting to
be contained in the sample
4.Detail of Raw material/final product in bulk / final
product (in finished pack) as obtained from the
manufacturer for analysis.
(a) Original manufacturer's name (in case of raw
materials and drugs repacked)
(b) Batch Number
(c) Batch Size
(d) Date of Manufacture if any
(e) Date of Expiry if any
5.Result of Analysis Recorded Below:

Kansas Laboratories Pvt. Ltd 8/1, Lal Bazar Street, Cal-700 001, DL No. – DL 1446M K-1460 Nil 11/00 10/03

Description :-White circular scored tablet in blister pack.

Identification	Positive
Avg. wt. of a tablet	0.5735gm
Weight variation	Passes
Dissolution test	Passes

Assay:--

Paracetamol I P.

Found/tab

Claim/tab

Method : As per I.P' 96.

Limit 95% to 105%

In the opinion of the undersigned, the sample referred to above is of standard quality / is not of standard quality as defined in the Act and Rules made thereunder for the reasons given below.

Sidne

Analyst / Junior Analyst /

plagengube.

A 3322/01 W. B. PHARMACEUTICAL & PHYTOCHEMICAL DEV. CORPN LTD.

(A Government of West Bengal Undertaking)

ACHARYA P. C. RAY R & D DRUG TESTING & POLLUTION TESTING LABORATORY

Behala Chowrasta, 620 Diamond Harbour Road, Calcutta 34, West Bengal

Report of Analysis [Form 39, See Rule 150 (f)]

License No: DL NO.T/8-DCH dt: 27.10.80. & T/13-DCH dt: 24.9.90

Sample Receiving No:D/2158/00
1.Name, address and license No. of Manufacturer /
Supplier's from whom sample received.
2.Reference No. and date of Manufacturer / Supplier's

letter under which the sample was forwarded 3.Name of Drug / Cosmetics / Ray material purporting to

be contained in the sample. 4.Detail of Raw material/final product in bulk / final Dt of Receipt: 15.12.00

Adviser (Health), State Urban Development Agency, Ilgus Bhawan, II-C Block, Sec-III, Bidhan Nagar, Cal-91, SUDA-120/96 (Pt-III)/633 dt. 24.11.00

1446M

Paracetamol tablets I.P.

10 tabs X 5 strips

	product (in finished pack) as obtained from the	
1	nanufacturer for analysis	
(a)	Original manufacturer's name (in case of raw	Kansas Laboratories Pvt. Ltd
	materials and drugs repacked)	8/1, Lal Bazar Street, Cal-1, DL NoDL
(b)	Batch Number	1459
(c)	Batch Size	Nil
(d)	Date of Manufacture if any	11/00
	Date of Expiry if any	10/03
	esult of Analysis Recorded Below:	

Method :1.P' 96.

Description :- White circular scored tablet in blister pack.

Identification	Positive
Avg. wt. of a tablet	0.5738gm
Weight variation	Passes
Dissolution test	Passes

Assay:-

Paracetamol I.P.

Found/tab 481.8mg

Claim/tab 500 00mg

Lunit

95% to 105%

In the opinion of the undersigned, the sample referred to above is of standard quality anderd quality as defined in the Act and Rules made thereunder for the reasons given below-...

Does not comply with respect to Cresol Content.

Analyst / Junior Analyst /

Stab. Furazolidine K-1470

Tab Chlorphenisamine malate

J- 1458

	Prove Hab 1045 - 200 Front	2 ~	6
•	pp dt 5T 300	So en:	~
Dan	Computerised File Name: D&Creport1826	A/ 5 66/0)
TY	W. B. PHARMACEUTICAL & PH	YTOCHEMICAL DEV. CORPN LTD.	
_	(A Government of W	est Bengal Undertaking)	
2	ACHARYA P. C. RAY R & D DRUG TESTI	NG & POLLUTION TESTING LABORATORY	
\mathcal{V}	Behala Chowrasta, 620 Diamond Harbour Road, Calcutta 34, West Bengal		
	Report of Analysis [Fo	m 39, See Rule 150 (f)]	
	License No: DL NO. T/8-DCH dt	: 27.10.80. & T/13-DCH dt: 24.9.90	
	Sample Receiving No:D/2361/00	Dt of Receipt: 26.12.00	
	I.Name, address and license No. of Manufacturer /	Adviser (Health), State Urban Development Agency, Ilgue	5
	Supplier's from whom sample received.	Bhawan, II-C Block, Sec-III, Bidhan Nagar, Cal-91,	
	2.Reference No. and date of Manufacturer / Supplier's	SUDA-120/96 (Pt-III)/512 dt. 13.12.00	
	letter under which the sample was forwarded		
	3.Name of Drug / Cosmetics / Raw material purporting to be contained in the sample.	Furazolidone Tablet I.P.	
	4. Detail of Raw material/final product in bulk / final	50 tablets	
	product (in finished pack) as obtained from the		
	manufacturer for analysis		
	(a) Original manufacturer's name (in case of raw	M/s. Kansas Labs (P) Ltd	
	materials and drugs repacked)	8/1,Lalbazar St., Cal-2	
	(b) Batch Number	K 1470	
	(c) Batch Size	Nil	
	(d) Date of Manufacture if any	11/00	
	(e) Date of Expiry if any	10/03	
	5. Result of Analysis Recorded Below:		

Method : I..P' 96, Vol-II.

Description :- Yellow coloured round shaped tablet having a bisecting mark on one side.

Avg. wt. of a tablet	0.1703gm
Weight variation	Passes
Identification	Positive
Disintegration time	7 minutes

Assay:---

Furazolidone I.P.

Claim/tab

Found/tab

94.83mg

100.00mg

Limit

90.00mg to 110.00mg

In the opinion of the undersigned, the sample referred to above is of standard quality / is not of standard quality as defined in the Act and Rules made thereunder for the reasons given below- in respect of the above tests only.

Signature of Oth harge

80/07

W. B. PHARMACEUTICAL & PHYTOCHEMICAL DEV. CORPN LTD.

(A Government of West Bengal Undertaking)

ACHARYA P. C. RAY R & D DRUG TESTING & POLLUTION TESTING LABORATORY

Behala Chowrasta, 620 Diamond Harbour Road, Calcutta 34, West Bengal

Report of Analysis [Form 39, See Rule 150 (f)]

License No: DL.NO.T/8-DCH dt: 27.10.80. & T/13-DCH dt: 24.9.90

Sample Receiving No:D/2361/00	Dt of Receipt: 26.12.00
LName, address and license No. of Manufacturer /	Adviser (Health), State Urban Development Agency, Igus
Supplier's from whom sample received.	Bhawan, II-C Block, Sec-III, Bidhan Nagar, Cal-91,
2.Reference No. and date of Manufacturer / Supplier's	SUDA-120/96 (Pt-III)/512 dt. 13.12.00
letter under which the sample was forwarded	(· ···, · ·· · ·· ·· ·· ·· ·· ·· ·· ·· ·
3.Name of Drug / Cosmetics / Raw material purporting to	Furazolidone Tablet I.P.
be contained in the sample.	
4. Detail of Raw material/final product in bulk / final	50 tablets
product (in finished pack) as obtained from the	
manufacturer for analysis	
(a) Original manufacturer's name (in case of raw	M/s. Kansas Labs (P) Ltd
materials and drugs repacked)	8/1,Lalbazar St., Cal-2
(b) Batch Number	K 1470
(c) Batch Size	Nil
(d) Date of Manufacture if any	
(e) Date of Expiry if any	11/00
5. Result of Analysis Recorded Below:	10/03
" TELAN WE'L DERIW.	

Method : I..P' 96, Vol-II.

Description :- Yellow coloured round shaped tablet having a bisecting mark on one side.

Avg. wt. of a tablet	0.1703gm
Weight variation	Passes
Identification	Positive
Disintegration time	7 minutes

Assay:--

Furazolidone I.P.

Found/tab 94.83mg Claim/tab 100.00mg

Limit

90.00mg to 110.00mg

In the opinion of the undersigned, the sample referred to above is of standard quality / is not of standard quality as defined in the Act and Rules made thereunder for the reasons given below in respect of the above tests only.

Signature of Officer In-Charge

Analyst/Junior Analyst/

Sample Receiving No:D/2157/00

be contained in the sample

1.Name, address and license No. of Manufacturer /

Supplier's from whom sample received

2. Reference No. and date of Manufacturer / Supplier's letter under which the sample was forwarded

3.Name of Drug / Cospetics / Raw material purporting to

W. B. PHARMACEUTICAL & PHYTOCHEMICAL DEV. CORPN LTD

(A Government of West Bengal Undertaking)

ACHARYA P. C. RAY R & D DRUG TESTING & POLLUTION TESTING LABORATORY

Behala Chowrasta, 620 Diamond Harbour Road, Calcutta 34, West Bengal

Report of Analysis [Form 39, See Rule 150 (f)]

License No: DL.NO.T/8-DCH dt: 27.10.80. & T/13-DCH dt: 24.9.90

Dt of Receipt: 15.12.00

3329

Adviser (Health), State Urban Development Agency, Ilgus Bhawan, II-C Block, Sec-III, Bidhan Nagar, Cal-91, SUDA-15/98 (Pt-II)/720 dt. 14.12.00

Chlorpheniramine Maleate tabs I.P.400mg

4.Detail of Raw material/final product in bulk / final product (in finished pack) as obtained from the manufacturer for analysis	10X5 strips
(a) Original manufacturer's name (in case of raw	Kansan Laboratories Pvt. Ltd
materials and drugs repacked)	8/1, Lai Bazar Street, Cal-700 001, DL 1464M
(b) Batch Number	J-1458
(c) Batch Size	Nil
(d) Date of Manufacture if any	10/00
(e) Date of Expiry if any	09/03
5. Result of Analysis Recorded Below:	

Method : I.P* 95.

Description :- A small size white circular uncoated tablet with a bisecting mark on one side of each tablet.

Identification	Positive
Related substances	Passes
Uniformity of content	Passes
Time of disintegration	2 minutes
Avg. wt. of a tablet	0.17400gm
Uniformity of weight	Passes

Assay:--

C.P. Maleate I.P.

Found/tab 4.12mg

4.0mg

3.8mg to 4.2mg

In the opinion of the undersigned, the sample referred to above is of standard quality / is not of standard quality as defined in the Act and Rules made thereunder for the reasons given below ...

nature of Offi er In-Charge

Sample Receiving No:D/2157/00

be contained in the sample

Supplier's from whom sample received.

I.Name, address and license No. of Manufacturer /

2. Reference No. and date of Manufacturer / Supplier's

letter under which the sample was forwarded

3.Name of Drug / Cospetics / Raw material purporting to

A/ 3329/0/

(A Government of West Bengal Undertaking)

ACHARYA P. C. RAY R & D DRUG TESTING & POLLUTION TESTING LABORATORY

Behala Chowrasta, 620 Diamond Harbour Road, Calcutta 34, West Bengal

Report of Analysis [Form 39, See Rule 150 (f)]

License No: DL NO T/8-DCH dt 27 10.80. & T/13-DCH dt: 24.9.90

Dt of Receipt: 15.12.00

Adviser (Health), State Urban Development Agency, ligus Bhawan, II-C Block, Sec-III, Bidhan Nagar, Cal-91, SUDA-15/98 (Pt-11)/720 dt. 14.12.00

Chlorpheniramine Maleate tabs I.P. 400mg

4. Detail of Raw material/inal product in bulk / final	10X5 strips
product (in finished pack) as obtained from the	
manufacturer for analysis	
(a) Original manufacturer's name (in case of raw	Kansan Laboratones Pvt. Ltd
materials and drugs repacked)	8/1, Lal Bazar Street, Cal-700 001, DL 1464M
(b) Batch Number	J-1458
(c) Batch Size	Nil
(d) Date of Manufacture if any.	10/00
(e) Date of Expiry if any	09/03
S Dornit of Analyzis Deparded Balans	

Method : I.P' 96.

Description .- A small size white circular uncoated tablet with a bisecting mark on one side of each tablet.

Identification	Positive
Related substances	Passes
Uniformity of content	Passes
Time of disintegration	2 minutes
Avg. wt. of a tablet	0 17400gm
Uniformity of weight	Passes

Assay:--

C.P. Mileate I.P

Claun tab

Found tab

412mg

4 ümg

Limit

3.8mg to 4.2mg

In the opinion of the undersigned, the sample referred to above is of standard quality / is not of standard quality as defined, in the Vet and Dal.

2000 files



R.C.H

STATE URBAN DEVELOPMENT AGENCY

"ILGUS BHAVAN" H-C BLOCK, SECTOR-III, BIDHANNAGAR, CALCUTTA-700 091 West Bengal

Ref No.....

Date

2000

SUDA-120/96(Pt.III)/

From : Adviser (Health), SUDA

To : Sri. Amitava Basak, West Bengal Pharmaceuticals Ltd., (A Govt. of West Bengal undertaking Lab) 620, Diamond Harborur Road, Chowrasta, Behala, Cal.- 700 034

> Sub : Analytical Testing of the samples of Drugs and MSR for use at the Health Units under RCH- Sub Project Asansol

Sir.

I like to forward the samples of Drugs and MSR to you for Analytical Test and Report. A lis of products to be tested is attached herewith.

You are requested to send the testing reports and the bill for payment of charges of testing to the undersigned mentioning "RCH- Sub Project Asansol", as early as possible.

Thanking You,

ours faithfully Glavent dviser (Health) SUDA

LH/0-7



SI. No.	Name of Items	Mfg. By	Qty. of Sample	Batches Of items	Mfg. Date	Exp. Date
V.	Aspirin Tab. IP- 300 mg.	Kansas Lab	3 x 50 tab	K-1477	11/2000	04/2002
2/	Chloramphenicol Eye Ont. I.P 1% w/w	Jyoti Capsules	50 x 2 applicap	JGC-21010	11/2000	04/2002
3.	Oxyphenonium Bromide IP 5 mg. tab	Kansas Lab	10 x 10 tab	K-1476	11/2000	10/2003
4.	Furazolidone 100 mg tab.	Kansas Lab	50 x 2 tab	K-1470	11/2000	10/2003
5.	Metronidazole IP 200 mg tab	Kansas Lab	50 x 2 tab	K-1472	11/2000	10/2003
6.	Cotrimoxazole IP S/S	Kansas Lab	50 x 2 tab	K-1462	11/2000	10/2003

List of samples to be tested

Adviser (Health) SUDA

LH/p-79



STATE URBAN DEVELOPMENT AGENCY

HEALTH WING "ILGUS BHAVAN" H-C BLOCK, SECTOR-III, BIDHANNAGAR, CALCUTTA-700 091 West Bengal

From: Adviser(Health) SUDA

To : Srl. Amitava Basak, West Bengal Pharmaceuticals Ltd., (A Govt. of West Bengal undertaking Lab) 620, Diamond Harbour Road, Chowrasta, Behala, Calcutta-700 034

Sub: Analytical Testing of the samples of Drugs and MSR for use at the Health Units under R.C.H. Sub-Project, Asansol.

Sir.

JLH/017

I like to forward the samples of Drugs and MSR to you for Analytical Test and Report.

Alist of products to be tested is attached herewith. You are requested to send the testing reports and the bill for payment of charges of testing to the undersigned mentioning "R.C.H. Sub-Project, Asansol," as early as possible.

Thanking you.

Yours faithfully, Adviser (health)

Tel/Fax No.: 359-3184



SI. No.	Name of Items	Mfg. By	Qty. of Sample	Batches Of items	Mfg. Date	Exp. Date
y	Mebendazole IP-100 mg	Kansas Lab	1 x 50 tab	H-1427	08/2000	06/2003
/2.	Bromhexine Hcl.BP. 8 mg	Kansas Lab	1 x 50 tab	G-1414	07/2000	09/2003
/3.	Chlorpheniramine Maleate IP 4 mg.	Kansas Lab	1 x 50 tab	J-1458	10/2000	10/2003
J4.	Paracitamol IP 500 mg.	Kansas Lab	3 x 50 tab	/K-1459 /K-1460 /K-1468	11/2000	10/2003
8.	Cotrimoxazole IP S/S	Kansas Lab	2 x 50 tab	K-1462 73 K-1463	11/2000	10/2003

List of samples to be tested

Adviser (Health) SUDA

4

in

LH/p-72



STATE URBAN DEVELOPMENT AGENCY

HEALTH WING "ILGUS BHAVAN" H-C BLOCK, SECTOR-III, BIDHANNAGAR, CALCUTTA-700 091 West Bengal

Ref No.SUDA 120/96(Pt-III)/633

From: Adviser(Health) SUDA

To : Sri. Amitava Basak, West Bengal Pharmaceuticals Ltd., (A Govt. of West Bengal undertaking Lab) 620, Diamond Harbour Road, Chowrasta, Behala, Calcutta-700 034

Sub: Analytical Testing of the samples of Drugs and MSR for use at the Health Units under R.C.H. Sub-Project, Asansol.

Sir,

Labor LH/p17

I like to forward the samples of Drugs and MSR to you for Analytical Test and Report.

Alist of products to be tested is attached herewith. You are requested to send the testing reports and the bill for payment of charges of testing to the undersigned mentioning "R.C.H. Sub-Project, Asansol" as early as possible.

Thanking you.

Yours faithfully,

Adviser (health) 24/11/200



List of samples to be tested

SI. No.	Name of Items	Mfg. By	Qty. of Sample	Batches Of items	Mfg. Date	Exp. Date
M.	Compound Magnesium Trisilicate tab	Avron Lab	10 x 100 tab	AS-(4140-* to 4144)	Nov.2000	Oct.2003
A.	Fesolic -L	Pure Pharma Ltd	1 x 50 tab	0017	Oct.2000	Sep.2002
3.	Fesolic-S	Pure Pharma Ltd	1 x 50 tab	0009	Oct.2000	Sep.2002
14.	Oral Rehidration Salt	Pure Pharma Ltd	16 x	0021 to 0024	Nov.2000	Oct.2002
15.	Antiseptic Lotion	Hisdustan Drugs	2 x 200 bottles	LE/06	Apr.1999	Mar.2004
6.	Merbromin 20 gm	C.D. Pharmaceu- tical Works.	2 x 20 gm phial	377	-	
7.	Nitrofourantoin Skin Cream 15 gm	Pilco Pharma Pvt.Ltd.	10 tubes ~	NF-1234	Nov.2000	Oct.2002
8.	Adhesive Plaster (10 cm x 5 mm)	Precision Coatings Ltd.	3 x 1 reals	010779_ 010776 - 010780_	Oct.2000	3 years
9.	Benzyl Banzoate Application	Hindustan Drugs	2 x 100 ml bottles	n.	Nov.2000	Oct.2002
10.	Absorbent Cotton 50 gm	Jaya & Co.	2(4 x 50) gm	43,46	Nov.2000	3 years

1 Lal Adviser (Health) 24 11 200

LH/p-73

FOR KANSAS LABS. P. LTD.



2 3 NOV	orr	J IND A

		2			2	3		ζ.	<	<			
	1.0.	Nº.	8. 1	~ 7.	6.7	5.	/	4.	3.	2.	1	1.	-Sl.No.
	Absorbent Cotton 50gm.	Benzyl Banzoate Application	Adhesive Plaster (10cmx5mm:	Nitrofurantoin Skin Cream 15gm. Jude	Merbromin 20 gm.	Antiseptic Lotion		Oral Rehidration Salt	Fesolic-S	Fesolic-L	May. Ton'silicate . 250 my Don'ed Alleroninium Hydron idegel-25004	Compound Magnesium Trisilicate Tab.	Name of Item
	Jaya & Co.	Hindustan Drugs	Precision Coatings LTD.	Pilco Pharma Pvt.Ltd.	C.D.Pharmeutical Works	Hindustan Drugs		-do-	-do-	Pure Pharma Ltd.	idezel-25039	Avron Labaratory	Mfg. Name.
	43 46	11 10	010779 010776 010780	NF-1234	1 377 ~	LE/CK	0023	0021	6000	0017	AS4143- AS4144-	AS4140- AS4141-	Batch No.
Jarker M	Var read	NEV-2000	OCT.2000	11/2000	•	Apr.99	-do-	11/2000 -do-	10/2000	10/2000		11/2000	Mfg.Date
1	3 years	ect. wor	3 Years	00/2002		MAR-2004	-do-	10/2002 -do-	9/2002	9/2002		10/2003	Expy.Date
5.11	4x50gm. 4x50gm.	(2) []x100ml.	1 Reel "	10 Tubes /	2x20gm.	2x200		472 (410)	1x50	1x50		2x100 Tabs	Quantity

111	1	
12:4	1. /	• S1.NO.
Bromhexine Hcl.BP.8mg	Mebendazole IP-1.00mg	Name of the items.
-do-	Kansas Lab.	Mfg. by :
,G 1414	H 1427	Batch No.
07/2000	08/2000	Mfg.Date
06/2003	07/2003	Exp.Date
1x50 Tabs.	1x50 Tabs.	Quantity
	Bromhexine Hcl.BP.8mg -do- /G 1414 07/2000 06/2003	./ Mebendazole IP-100mg Kansas Lab. H 1427 08/2000 07/2003 Bromhexine Hcl.BP.8mg -do- iG 1414 07/2000 06/2003

2 3 NON 2000

FOR KANSAS LABS. P. LTD.

5	5	3.	z.	M.	SL.ND.
CHLORAMPHENICOL EYE OINTMENT I.P 1% U/U (EYE Applicaps)	OXYPHENONIUM BROMIDE TABLETS I.P 5mg.	ASPIRIN TABLETS I.P 300mg.	METRONIDAZOLE TABLETS I.P. 200mg.(Film Coated	ABLETS	NAME OF ITEM
Jyoti Capsules	1 R -	ł	-1-	Kensas Labora- tories Pvt.Ltd.	MFG .NAME
JGC 21010	K 1476	K 1477	K 1472	K 1470	BATCH NO .
NOV. 2009	11/2000	11/2000	11/2000	11/2000	MFG.DATE
APRIL 2002	10/2903	4/2002	10/2003	10/2003	EXPY. DATE
2 x 50 Caps.	10×10 Tabs.	3 x 50 Tabs.	2 x 50 Tabs.	2 x 50 Tabs.	QUANTITY



Ter KANDA ABORTING

Tab Fersolie - 4

B.no- 0014

Tab - Fersolie - S Bro - 0009

Antiseptie 200 -LE-OB

Mebendazole tab. 100 mg

H-1427

Aspisin tab K-1477

Oxyphenonium Bromde - 500 %.

K-1476 Chloramphenicol Ex. Applicap. Jeg. Joe 21010