3374/01 2

Computerised File Name: D&Creport1880

A/

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/2155/00 Dt of Receipt: 15.12.00

1.Name, address and license No. of Manufacturer / Supplier's from whom sample received.....	Adviser (Health),State Urban Development Agency, Ilgus Bhawan, II-C Block, Sec-III, Bidhan Nagar, Cal-91,
2.Reference No. and date of Manufacturer / Supplier's letter under which the sample was forwarded	SUDA-120/96 (Pt-III)/633 dt. 24.11.00
3.Name of Drug / Cosmetics / Raw material purporting to be contained in the sample.....	Mebendazole tablet I.P. 100mg
4.Detail of Raw material final product in bulk / final product (in finished pack) as obtained from the manufacturer for analysis.....	50 tabs
(a) Original manufacturer's name (in case of raw materials and drugs repacked).....	M/s. Kansas Laboratories (P) Ltd 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:--

	Found/tab	Claim/tab
Mebendazole I.P.	97.05mg	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~~ .

(Junior Analyst)

Signature of Officer In-Charge

3374/01 2

Computerised File Name: D&Creport1880

A.

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/2155/00 Dt of Receipt: 15.12.00

1.Name, address and license No of Manufacturer / Supplier's from whom sample received.....	Adviser (Health),State Urban Development Agency, Ilgus Bhawan, II-C Block, Sec-III, Bidhan Nagar, Cal-91,
2.Reference No. and date of Manufacturer / Supplier's letter under which the sample was forwarded	SUDA-120/96 (Pt-III)/633 dt. 24.11.00
3.Name of Drug / Cosmetics / Raw material purporting to be contained in the sample.....	Mebendazole tablet I.P. 100mg
4.Detail of Raw material final product in bulk / final product (in finished pack) as obtained from the manufacturer for analysis.....	50 tabs
(a) Original manufacturer's name (in case of raw materials and drugs repacked).....	M/s. Kansas Laboratories (P) Ltd
	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:--

	Found/tab	Claim/tab
Mebendazole I.P.	97.05mg	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~~ .

Signature of Junior Analyst

8/1/2001

Signature of Officer In-Charge

Computerised File Name: D&Creport1907

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

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

Report of Analysis [Form 39, See Rule 150 (1)]

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

Sample Receiving No:D/2358/00 Dt of Receipt: 26.12.00

1.Name, address and license No. of Manufacturer / Supplier's from whom sample received	Adviser (Health),State Urban Development Agency, Nigas Bhawan, 11-C Block, Sec-III, Bidhan Nagar, Cal-91.
2.Reference No. and date of Manufacturer / Supplier's letter under which the sample was forwarded	SUDA-120/96 (Pt-III)/ dt. 13.12.00
3.Name of Drug / Cosmetics / Raw material purporting to be contained in the sample	Aspirin Tablet I.P.
4.Detail of Raw material/final product in bulk / final product (in finished pack) as obtained from the manufacturer for analysis	2 X 50 tabs
(a) Original manufacturer's name (in case of raw materials and drugs repacked)	M/s. Kansas Labs (P) Ltd 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 lgm
Weight variation	Passes
Identification	Positive
Disintegration time	1 minutes
Salicylic Acid	Passes

Assay:-

	Found tab	Claim tab
Aspirin I P	308.93mg	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.

(Miss) Junior Analyst /

Signature of Officer In-Charge

5

C. Computerised File Name: D&Creport1907

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

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/2358/00 Dt of Receipt: 26.12.00

1.Name, address and license No. of Manufacturer / Supplier's from whom sample received.....	Adviser (Health),State Urban Development Agency, Ilgus Bhawan, 11-C Block, Sec-III, Bidhan Nagar, Cal-91,
2.Reference No. and date of Manufacturer / Supplier's letter under which the sample was forwarded	SUDA-120/96 (Pt-III)/ dt. 13.12.00
3.Name of Drug / Cosmetics / Raw material purporting to be contained in the sample.....	Aspirin Tablet I.P.
4.Detail of Raw material/final product in bulk / final product (in finished pack) as obtained from the manufacturer for analysis.....	2 X 50 tabs
(a) Original manufacturer's name (in case of raw materials and drugs repacked).....	M/s. Kansas Labs (P) Ltd 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	0.3414gm
Weight variation	Passes
Identification	Positive
Disintegration time	1 minutes
Salicylic Acid	Passes

Assay:-

	Found/tab	Claim/tab
Aspirin I.P.	308.93mg	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.

Analyst / Junior Analyst /

Signature of Officer In-Charge

3

Computerised File Name: D&Creport2097 A/3617/07

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/2360/00 Dt of Receipt: 26.12.00

1.Name, address and license No. of Manufacturer / Supplier's from whom sample received.....	State Urban Development Agency II-C Block,Sector-III,Bidhannagar Cal-700 091
2.Reference No. and date of Manufacturer / Supplier's letter under which the sample was forwarded	SUDA/120/96 Pt-III
3.Name of Drug / Cosmetics / Raw material purporting to be contained in the sample.....	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 materials and drugs repacked).....	Kansus Laboratory Pvt Ltd 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:-

	Found/tab	Claim/tab
Oxyphenonium Bromide IP	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.

Analyst / Junior Analyst / Signature of Officer In-Charge

24/11/2007

3.

A. 3617/01

Computerised File Name: D&Creport2097

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/2360/00 Dt of Receipt: 26.12.00

1.Name, address and license No. of Manufacturer / Supplier's from whom sample received	State Urban Development Agency II-C Block,Sector-III,Bidhannagar Cal-700 091
2.Reference No. and date of Manufacturer / Supplier's letter under which the sample was forwarded	SUDA/120/96 Pt-III
3.Name of Drug / Cosmetics / Raw material purporting to be contained in the sample	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 materials and drugs repacked)	Kansus Laboratory Pvt Ltd 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 Supplments-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 IP	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.

Analyst / Junior Analyst /

Signature of Officer In-Charge

24/11/2001

Computerised File Name: D&Creport2102

A/ 3619/07

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/2359/00 Dt of Receipt: 26.12.00

1.Name, address and license No. of Manufacturer / Supplier's from whom sample received.....	State Urban Development Agency II-C Block,Sector-III,Bidhannagar Cal-700 091
2.Reference No. and date of Manufacturer / Supplier's letter under which the sample was forwarded	SUDA/120/96 Pt-III dt 13.12.00
3.Name of Drug / Cosmetics / Raw material purporting to be contained in the sample.....	Chloramphenicol Eye ~~Ointment~~ IP applicap
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:-

	Found/tab	Claim/tab	
Chloraphenicol IP	0.95%	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.

Analyst / Junior Analyst /

24/1/2007

Signature of Officer In-Charge

Computerised File Name: D&Creport2102

A 3619/01

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

(A Government of West Bengal Undertaking)

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

Belvala 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 / Supplier's from whom sample received	State Urban Development Agency II-C Block,Sector-III,Bidhannagar Cal-700 091
2.Reference No. and date of Manufacturer / Supplier's letter under which the sample was forwarded	SUDA/120/96 Pt-III dt 13.12.00
3.Name of Drug / Cosmetics / Raw material purporting to be contained in the sample	Chloramphenicol Eye ~~Ointment~~ applicap 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:-

	Found/cap	Claim/cap	
Chloraphenicol IP	0.95%	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.

Analyst / Junior Analyst /

24/11/2000

Signature of Officer In-Charge

Computerised File Name: D&Creport2067

A/3570/07

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/2142/00 Dt of Receipt: 15.12.00

1.Name, address and license No. of Manufacturer / Supplier's from whom sample received.....	Adviser (Health),State Urban Development Agency, Ilgus Bhawan, II-C Block, Sec-III, Bidhan Nagar, Cal-91,
2.Reference No. and date of Manufacturer / Supplier's letter under which the sample was forwarded	SUDA-15/98 (Pt-II)/720 dt. 14.12.00
3.Name of Drug / Cosmetics / 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 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 ~~/ is not of standard quality~~ as defined in the Act and Rules made thereunder ~~for the reasons given below~~.

18/1/2007

Analyst / Junior Analyst /

18/1/2007

Signature of Officer In-Charge

Computerised File Name: D&Creport2067

A.3570/04

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

1.Name, address and license No. of Manufacturer / Supplier's from whom sample received.....	Adviser (Health),State Urban Development Agency, Ilgus Bhawan, II-C Block, Sec-III, Bidhan Nagar, Cal-91,
2.Reference No. and date of Manufacturer / Supplier's letter under which the sample was forwarded	SUDA-15/98 (Pt-II)/720 dt. 14.12.00
3.Name of Drug / Cosmetics / 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 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 / is not ~~of standard quality~~ as defined in the Act and Rules made thereunder ~~for the reasons given below~~.

18/1/05

Analyst / Junior Analyst /

18/1/2005

Signature of Officer In-Charge

Computerised File Name: D&Creport2068

A/ 3577/07

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/2143/00 Dt of Receipt: 15.12.00

1.Name, address and license No. of Manufacturer / Supplier's from whom sample received.....	Adviser (Health),State Urban Development Agency, Ilgus Bhawan, II-C Block, Sec-III, Bidhan Nagar, Cal-91,
2.Reference No. and date of Manufacturer / Supplier's letter under which the sample was forwarded	SUDA-15/98 (Pt-II)/720 dt. 14.12.00
3.Name of Drug / Cosmetics / 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 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 I..P.	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~~.

18/1/07

Analyst / Junior Analyst / Signature of Officer In-Charge

Computerised File Name: D&Creport2068

A/ 3571/07

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/2143/00 Dt of Receipt: 15.12.00

1.Name, address and license No. of Manufacturer / Supplier's from whom sample received.....	Adviser (Health),State Urban Development Agency, Ilgus Bhawan, II-C Block, Sec-III, Bidhan Nagar, Cal-91,
2.Reference No. and date of Manufacturer / Supplier's letter under which the sample was forwarded	SUDA-15/98 (Pt-II)/720 dt. 14.12.00
3.Name of Drug / Cosmetics / 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
(b) Batch Number	41, 42 &44, Industrial Area. Pologround (Indore)
(c) Batch Size	0022
(d) Date of Manufacture if any.....	Nil
(e) Date of Expiry if any.....	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	28.45773gm

Assay:--

	Found/27.9gm	Claim/27.9gm	Limit/27.9gm
Anhydrous Dextrose I..P.	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~~.

18/1/2007

Analyst / Junior Analyst /

18/1/2007

Signature of Officer In-Charge

3

Computerised File Name: D&Creport2069

A/3572/07

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/2144/00 Dt of Receipt: 15.12.00

1.Name, address and license No. of Manufacturer / Supplier's from whom sample received.....	Adviser (Health),State Urban Development Agency, Ilgus Bhawan, II-C Block, Sec-III, Bidhan Nagar, Cal-91,
2.Reference No. and date of Manufacturer / Supplier's letter under which the sample was forwarded	SUDA-15/98 (Pt-II)/720 dt. 14.12.00
3.Name of Drug / Cosmetics / 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 41, 42 &44, Industrial Area. Pologround (Indore)
(b) Batch Number	0023
(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	28.45512gm

Assay:--

	Found/27.9gm	Claim/27.9gm	Limit/27.9gm
Anhydrous Dextrose I..P.	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 Analyst / 18/1/07

Signature of Officer In-Charge 18/1/07

3

Computerised File Name: D&Creport2069 A. 3572/07

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/2144/00 Dt of Receipt: 15.12.00

1.Name, address and license No. of Manufacturer / Supplier's from whom sample received.....	Adviser (Health),State Urban Development Agency, Ilgus Bhawan, II-C Block, Sec-III, Bidhan Nagar, Cal-91,
2.Reference No. and date of Manufacturer / Supplier's letter under which the sample was forwarded	SUDA-15/98 (Pt-II)/720 dt. 14.12.00
3 Name of Drug / Cosmetics / 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 41, 42 &44, Industrial Area. Pologround (Indore)
(b) Batch Number	0023
(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	28.45512gm

Assay:--

	Found/27.9gm	Claim/27.9gm	Limit/27.9gm
Anhydrous Dextrose I..P.	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~~.

28/11/07

Analyst / Junior Analyst /

28/11/2007

Signature of Officer in-Charge

3

Computerised File Name: D&Creport2070 A/ 3573/07

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/2145/00 Dt of Receipt: 15.12.00

1.Name, address and license No. of Manufacturer / Supplier's from whom sample received.....	Adviser (Health),State Urban Development Agency, Ilgus Bhawan, II-C Block, Sec-III, Bidhan Nagar, Cal-91,
2.Reference No. and date of Manufacturer / Supplier's letter under which the sample was forwarded	SUDA-15/98 (Pt-II)/720 dt. 14.12.00
3.Name of Drug / Cosmetics / 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 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 I..P.	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 Analyst / 18/1/2007 Signature of Officer In-Charge 18/1/2007

3

Computerised File Name: D&Creport2070

A/ 3578/07

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/2145/00 Dt of Receipt: 15.12.00

1.Name, address and license No. of Manufacturer / Supplier's from whom sample received.....	Adviser (Health),State Urban Development Agency, Ilgus Bhawan, II-C Block, Sec-III, Bidhan Nagar, Cal-91,
2.Reference No. and date of Manufacturer / Supplier's letter under which the sample was forwarded	SUDA-15/98 (Pt-II)/720 dt. 14.12.00
3.Name of Drug / Cosmetics / 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 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 I..P.	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~~.

18/1/2001

Analyst / Junior Analyst /

18/1/2001

Signature of Officer In-Charge

Computerised File Name: D&Creport2059 A/ 3566/99

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

1.Name, address and license No. of Manufacturer / Supplier's from whom sample received.....	Adviser (Health),State Urban Development Agency, Ilgus Bhawan, II-C Block, Sec-III, Bidhan Nagar, Cal-91,
2.Reference No. and date of Manufacturer / Supplier's letter under which the sample was forwarded	SUDA/120/96 (Pt-III) dt. Nil
3.Name of Drug / Cosmetics / Raw material purporting to be contained in the sample.....	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 materials and drugs repacked).....	Kansas Laboratories Pvt. Ltd 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:--

	Found/ml	Claim/~~ml~~
Trimethoprim I.P.	79.90mg	80.00mg
Sulphamethoxazole I.P.	406.15mg	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~~

Analyst / Junior Analyst /

Signature of Officer In-Charge

Computerised File Name: D&Creport2059

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

Sample Receiving No:D/2363/00 Dt of Receipt: 26.12.00

1.Name, address and license No. of Manufacturer / Supplier's from whom sample received.....	Adviser (Health),State Urban Development Agency, Ilgus Bhawan, II-C Block, Sec-III, Bidhan Nagar, Cal-91,
2.Reference No. and date of Manufacturer / Supplier's letter under which the sample was forwarded	SUDA/120/96 (Pt-III) dt. Nil
3.Name of Drug / Cosmetics / Raw material purporting to be contained in the sample.....	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 materials and drugs repacked).....	Kansas Laboratories Pvt. Ltd 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:--

	Found/ml	Claim/ml
Trimethoprim I.P.	79.90mg	80.00mg
Sulphamethoxazole I.P.	406.15mg	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~~.

Analyst / Junior Analyst /

Signature of Officer In-Charge

Computerised File Name: D&Creport2060

A 3561/01

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/2161/00 Dt of Receipt: 15.12.00

1.Name, address and license No. of Manufacturer / Supplier's from whom sample received.....	Adviser (Health),State Urban Development Agency, Ilgus Bhawan, II-C Block, Sec-III, Bidhan Nagar, Cal-91,
2.Reference No. and date of Manufacturer / Supplier's letter under which the sample was forwarded	SUDA/120/96 (Pt-III) dt. Nil
3.Name of Drug / Cosmetics / Raw material purporting to be contained in the sample.....	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 materials and drugs repacked).....	Kansas Laboratories Pvt. Ltd 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	0.5828gm
Weight variation	Passes
Time of disintegration	7 minutes

Assay:--

	Found/~~ml~~	Claim/ml
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~~.

Analyst / Junior Analyst /

8/1/2001

Signature of Officer In-Charge

Computerised File Name: D&Creport2059 A/ 3563/04

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

1.Name, address and license No. of Manufacturer / Supplier's from whom sample received.....	Adviser (Health),State Urban Development Agency, Ilgus Bhawan, II-C Block, Sec-III, Bidhan Nagar, Cal-91.
2.Reference No. and date of Manufacturer / Supplier's letter under which the sample was forwarded	SUDA/120/96 (Pt-III) dt. Nil
3.Name of Drug / Cosmetics / Raw material purporting to be contained in the sample.....	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 materials and drugs repacked).....	Kansas Laboratories Pvt. Ltd 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:--

	Found/ml	Claim/ml
Trimethoprim I.P.	79.90mg	80.00mg
Sulphamethoxazole I.P.	406.15mg	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~~.

Analyst / Junior Analyst / Signature of Officer In-Charge

Computerised File Name: D&Creport2061

A/3568/07

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/2162/00 Dt of Receipt: 15.12.00

1.Name, address and license No. of Manufacturer / Supplier's from whom sample received.....	Adviser (Health),State Urban Development Agency, Ilgus Bhawan, II-C Block, Sec-III, Bidhan Nagar, Cal-91,
2.Reference No. and date of Manufacturer / Supplier's letter under which the sample was forwarded	SUDA/120/96 (Pt-III) dt. Nil
3.Name of Drug / Cosmetics / Raw material purporting to be contained in the sample.....	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 materials and drugs repacked).....	Kansas Laboratories Pvt. Ltd 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	Claim/ml
Trimethoprim I.P.	79.80mg	80.00mg
Sulphamethoxazole I.P.	391.37mg	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~~.

Analyst / Junior Analyst /

Signature of Officer In-Charge

Computerised File Name: D&Creport2061

A 3568/3/07

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/2162/00 Dt of Receipt: 15.12.00

1.Name, address and license No. of Manufacturer / Supplier's from whom sample received.....	Adviser (Health),State Urban Development Agency, Ilgus Bhawan, II-C Block, Sec-III, Bidhan Nagar, Cal-91,
2.Reference No. and date of Manufacturer / Supplier's letter under which the sample was forwarded	SUDA/120/96 (Pt-III) dt. Nil
3.Name of Drug / Cosmetics / Raw material purporting to be contained in the sample.....	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 materials and drugs repacked).....	Kansas Laboratories Pvt. Ltd 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~~	Claim/~~ml~~
Trimethoprim I.P.	79.80mg	80.00mg
Sulphamethoxazole I.P.	391.37mg	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~~.

Analyst / Junior Analyst /

Signature of Officer In-Charge

18/1/2007

3

Computerised File Name: D&Creport2064 A/ 3573/00

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

Sample Receiving No:D/2147/00 Dt of Receipt: 15.12.00

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

Description :-A green scales.

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

Assay:--

	Found	Limit
Bromine Content	20.82%	18% to 21.30%
Mercury Content	24.95%	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~~.

Analyst / Junior Analyst / Signature of Officer In-Charge

3

Computerised File Name: D&Creport2064 A. 3588/07

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/2148/00 Dt of Receipt: 15.12.00

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

Description :-A green scales.

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

Assay:--

	Founded	Limit
Bromine Content	20.82%	18% to 21.30%
Mercury Content	24.95%	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~~.

Analyst / Junior Analyst / ..... Signature of Officer In-Charge

Computerised File Name: D&Creport2077 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 / Supplier's from whom sample received.....	Adviser (Health),State Urban Development Agency, Ilgus Bhawan, II-C Block, Sec-III, Bidhan Nagar, Cal-91,
2.Reference No. and date of Manufacturer / Supplier's letter under which the sample was forwarded	SUDA/120/96 (Pt-III) dt. Nil
3.Name of Drug / Cosmetics / Raw material purporting to be contained in the sample.....	Fesolic- L
4.Detail of Raw material/final product in bulk / final product (in finished pack) as obtained from the manufacturer for analysis.....	50 tabs
(a) Original manufacturer's name (in case of raw materials and drugs repacked).....	Pure Pharma Ltd, 41,42 &44, Industrial Estate, (Pologround), Indore-452015, Mfg Lics No.-30/64
(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:--

	Found/tab	Claim/tab	Limit
Dried Ferrous Sulphate I.P.	0.0573gm (85.61%)	0.067gm	80% to 90%
Folic Acid	0.45mg (90%)	0.5mg	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~~.

10/11/2009

Analyst / Junior Analyst / Signature of Officer In-Charge

Computerised File Name: D&Creport2077 A/ 3574/9

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 / Supplier's from whom sample received.....	Adviser (Health),State Urban Development Agency, Ilgus Bhawan, II-C Block, Sec-III, Bidhan Nagar, Cal-91,
2.Reference No. and date of Manufacturer / Supplier's letter under which the sample was forwarded	SUDA/120/96 (Pt-III) dt. Nil
3.Name of Drug / Cosmetics / Raw material purporting to be contained in the sample.....	Fesolic- L
4.Detail of Raw material/final product in bulk / final product (in finished pack) as obtained from the manufacturer for analysis.....	50 tabs
(a) Original manufacturer's name (in case of raw materials and drugs repacked).....	Pure Pharma Ltd, 41,42 &44, Industrial Estate, (Pologround), Indore-452015, Mfg Lics No.-30/64
(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:--

	Found/tab	Claim/tab	Limit
Dried Ferrous Sulphate I.P.	0.0573gm (85.61%)	0.067gm	80% to 90%
Folic Acid	0.45mg (90%)	0.5mg	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~~.

18/1/2001

Analyst / Junior Analyst / Signature of Officer In-Charge

Computerised File Name: D&Creport2076

A/ 3577/00

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

Sample Receiving No:D/2141/00 Dt of Receipt: 15.12.00

1.Name, address and license No. of Manufacturer / Supplier's from whom sample received.....	Adviser (Health),State Urban Development Agency, Ilgus Bhawan, II-C Block, Sec-III, Bidhan Nagar, Cal-91,
2.Reference No. and date of Manufacturer / Supplier's letter under which the sample was forwarded	SUDA/120/96 (Pt-III) dt. Nil
3.Name of Drug / Cosmetics / Raw material purporting to be contained in the sample.....	Fesolic- S
4.Detail of Raw material/final product in bulk / final product (in finished pack) as obtained from the manufacturer for analysis.....	50 tabs
(a) Original manufacturer's name (in case of raw materials and drugs repacked).....	Pure Pharma Ltd, 41,42 &44, Industrial Estate, (Pologround), Indore-452015, Mfg Lics No.-30/64
(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~~.

Analyst / Junior Analyst /

18/1/2001

Signature of Officer In-Charge

Computerised File Name: D&Creport207o

A. 3577/01

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/2141/00 Dt of Receipt: 15.12.00

1.Name, address and license No. of Manufacturer / Supplier's from whom sample received.....	Adviser (Health),State Urban Development Agency, Ilgus Bhawan, II-C Block, Sec-III, Bidhan Nagar, Cal-91,
2.Reference No. and date of Manufacturer / Supplier's letter under which the sample was forwarded	SUDA/120/96 (Pt-III) dt. Nil
3.Name of Drug / Cosmetics / Raw material purporting to be contained in the sample.....	Fesolic- S
4.Detail of Raw material/final product in bulk / final product (in finished pack) as obtained from the manufacturer for analysis.....	50 tabs
(a) Original manufacturer's name (in case of raw materials and drugs repacked).....	Pure Pharma Ltd, 41,42 &44, Industrial Estate, (Pologround), Indore-452015. Mfg Lics No.-30/64
(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~~.

Analyst / Junior Analyst /

Signature of Officer In-Charge

3575/07 3

Computerised File Name: D&Creport2078 A/ 3575/07

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/2146/00 Dt of Receipt: 15.12.00

1.Name, address and license No. of Manufacturer / Supplier's from whom sample received.....	Adviser (Health),State Urban Development Agency, Ilgus Bhawan, II-C Block, Sec-III, Bidhan Nagar, Cal-91,
2.Reference No. and date of Manufacturer / Supplier's letter under which the sample was forwarded	SUDA-120/96 (Pt-II)/633 dt. 24.12.00
3.Name of Drug / Cosmetics / Raw material purporting to be contained in the sample.....	Antiseptic Lotion
4.Detail of Raw material/final product in bulk / final product (in finished pack) as obtained from the manufacturer for analysis.....	2X200ml
(a) Original manufacturer's name (in case of raw materials and drugs repacked).....	Hindusthan Drugs 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:--

	Found	Claim
Benzal Konium Chloride	2.10%	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.

18/1/2007

Analyst / Junior Analyst / Signature of Officer In-Charge

Computerised File Name: D&Creport2078 A/ 3573/07

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/2146/00 Dt of Receipt: 15.12.00

1.Name, address and license No. of Manufacturer / Supplier's from whom sample received.....	Adviser (Health),State Urban Development Agency, Ilgus Bhawan, II-C Block, Sec-III, Bidhan Nagar, Cal-91,
2.Reference No. and date of Manufacturer / Supplier's letter under which the sample was forwarded	SUDA-120/96 (Pt-II)/633 dt. 24.12.00
3.Name of Drug / Cosmetics / Raw material purporting to be contained in the sample.....	Antiseptic Lotion
4.Detail of Raw material/final product in bulk / final product (in finished pack) as obtained from the manufacturer for analysis.....	2X200ml
(a) Original manufacturer's name (in case of raw materials and drugs repacked).....	Hindusthan Drugs 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:--

	Found	Claim
Benzal Konium Chloride	2.10%	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.

18/1/2007

Analyst / Junior Analyst / Signature of Officer In-Charge

Benzyl Benzoate Application IP.

B.no - ~~10071~~
11

Nitrofurazone Cream.

B.no - NF 1234

Ref no. PPDC/Lab/1012 dt. 21-12-2000

3

Computerised File Name: D&Creport1747

A/ 3215

(2)

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 / Supplier's from whom sample received.....	Adviser (Health),State Urban Development Agency, Ilgus Bhawan, II-C Block, Sec-III, Bidhan Nagar, Cal-91,
2.Reference No. and date of Manufacturer / Supplier's letter under which the sample was forwarded	SUDA-15/98 (Pt-II)/633 dt. 24.11.00
3.Name of Drug / Cosmetics / Raw material purporting to be contained in the sample.....	Benzyl Benzoate Application I.P.
4.Detail of Raw material/final product in bulk / final product (in finished pack) as obtained from the manufacturer for analysis.....	2X100ml
(a) Original manufacturer's name (in case of raw materials and drugs repacked).....	Hindusthan Drugs, 139, Shyam Nagar Road, Cal-700 056 DL No.-1069M
(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:**

Method : I.P.

Description :-White viscous emulsion in glass bottle

Identification Positive

Assay:--

	Found	Claim	Limit
Benzyl Benzoate I.P.	25.89%	25.0%	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~~.

Analyst / Junior Analyst /

Signature of Officer In-Charge

Computerised File Name: D&Creport1747

3

A. 9457/M

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 / Supplier's from whom sample received.....	Adviser (Health),State Urban Development Agency, Ilgus Bhawan, II-C Block, Sec-III, Bidhan Nagar, Cal-91,
2.Reference No. and date of Manufacturer / Supplier's letter under which the sample was forwarded	SUDA-15/98 (Pt-II)/633 dt. 24.11.00
3.Name of Drug / Cosmetics / Raw material purporting to be contained in the sample.....	Benzyl Benzoate Application I.P.
4.Detail of Raw material/final product in bulk / final product (in finished pack) as obtained from the manufacturer for analysis.....	2X100ml
(a) Original manufacturer's name (in case of raw materials and drugs repacked).....	Hindusthan Drugs, 139, Shyam Nagar Road, Cal-700 056 DL No.-1069M
(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:

Method : I.P.

Description :-White viscous emulsion in glass bottle

Identification Positive

Assay :-

	Found	Claim	Limit
Benzyl Benzoate I.P.	25.89%	25.0%	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~~.

Analyst / Junior Analyst / Satisfactory

Signature of Officer In-Charge 19/12/00

Computerised File Name: D&Creport1749

A/ 3246/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/2141/00 Dt of Receipt: 15.12.00

1.Name, address and license No. of Manufacturer / Supplier's from whom sample received.....	Adviser (Health),State Urban Development Agency, Ilgus Bhawan, II-C Block, Sec-III, Bidhan Nagar, Cal-91,
2.Reference No. and date of Manufacturer / Supplier's letter under which the sample was forwarded	SUDA-15/98 (Pt-II)/633 dt. 24.11.00
3.Name of Drug / Cosmetics / Raw material purporting to be contained in the sample.....	Nitrofurazone cream U.S.P.
4.Detail of Raw material/final product in bulk / final product (in finished pack) as obtained from the manufacturer for analysis.....	10X15gm
(a) Original manufacturer's name (in case of raw materials and drugs repacked).....	Pilco Pharma Pvt. Ltd 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:–

	Found	Claim	Limit
Nitrofurazone	0.22%	0.2%	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 advice

Analyst / Junior Analyst /

19/12/00

Signature of Officer In-Charge

Computerised File Name: D&Creport1749

A/ 3216/00

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

Sample Receiving No:D/2141/00 Dt of Receipt: 15.12.00

1.Name, address and license No. of Manufacturer / Supplier's from whom sample received.....	Adviser (Health),State Urban Development Agency, Ilgus Bhawan, II-C Block, Sec-III, Bidhan Nagar, Cal-91,
2.Reference No. and date of Manufacturer / Supplier's letter under which the sample was forwarded	SUDA-15/98 (Pt-II)/633 dt. 24.11.00
3.Name of Drug / Cosmetics / Raw material purporting to be contained in the sample.....	Nitrofurazone cream U.S.P.
4.Detail of Raw material/final product in bulk / final product (in finished pack) as obtained from the manufacturer for analysis.....	10X15gm
(a) Original manufacturer's name (in case of raw materials and drugs repacked).....	Pilco Pharma Pvt. Ltd 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:--

	Found	Claim	Limit
Nitrofurazone	0.22%	0.2%	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~~.

Analyst / Junior Analyst /

19/12/00

Signature of Officer In-Charge

Tab. Mebomidazole - 200 mg

K. - 1472

Tab. Bromhexine 8mg.

G 1414

Tab. Paracetamol -

K - 1459

K - 1460

K - 1468

Ref on PPSC/Lab/1056 dt 4-1-2001

Received on 05/01/2001

3

Computerised File Name: D&Crepo1t1806

A/ 3326/01

5

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/2362/00 Dt of Receipt: 26.12.00

1.Name, address and license No. of Manufacturer / Supplier's from whom sample received.....	Adviser (Health),State Urban Development Agency, Ilgus Bhawan, II-C Block, Sec-III, Bidhan Nagar, Cal-91,
2.Reference No. and date of Manufacturer / Supplier's letter under which the sample was forwarded	SUDA/120/96 (Pt-III) dt. Nil
3.Name of Drug / Cosmetics / Raw material purporting to be contained in the sample.....	Metronidazole tab I.P. 200mg
4.Detail of Raw material/final product in bulk / final product (in finished pack) as obtained from the manufacturer for analysis.....	50X2
(a) Original manufacturer's name (in case of raw materials and drugs repacked).....	Kansas Laboratories Pvt. Ltd 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:--

	Found/tab	Claim/tab	Limit
Metronidazole I.P.	206.5mg	200.00mg	190mg to 210mg

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

Computerised File Name: D&Creport1865

A/ 3326/01

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/2362/00 Dt of Receipt: 26.12.00

1.Name, address and license No. of Manufacturer / Supplier's from whom sample received.....	Adviser (Health),State Urban Development Agency, Ilgus Bhawan, II-C Block, Sec-III, Bidhan Nagar, Cal-91,
2.Reference No. and date of Manufacturer / Supplier's letter under which the sample was forwarded	SUDA/120/96 (Pt-III) dt. Nil
3.Name of Drug / Cosmetics / Raw material purporting to be contained in the sample.....	Metronidazole tab I.P. 200mg
4.Detail of Raw material/final product in bulk / final product (in finished pack) as obtained from the manufacturer for analysis.....	50X2
(a) Original manufacturer's name (in case of raw materials and drugs repacked).....	Kansas Laboratories Pvt. Ltd 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:--

	Found/tab	Claim/tab	Limit
Metronidazole I.P.	206.5mg	200.00mg	190mg to 210mg

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

3

Computerised File Name: D&Creport1870

A/ 3327/01

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/2156/00 Dt of Receipt: 15.12.00

1.Name, address and license No. of Manufacturer / Supplier's from whom sample received.....	Adviser (Health),State Urban Development Agency, Ilgus Bhawan, 11-C Block, Sec-III, Bidhan Nagar, Cal-91,
2.Reference No. and date of Manufacturer / Supplier's letter under which the sample was forwarded	SUDA/120/96 (Pt-II)/631 dt. 24.11.00
3.Name of Drug / Cosmetics / Raw material purporting to be contained in the sample.....	Bromhexine Tablet B.P. 8mg
4.Detail of Raw material/final product in bulk / final product (in finished pack) as obtained from the manufacturer for analysis.....	10 tabs X 5 strips
(a) Original manufacturer's name (in case of raw materials and drugs repacked).....	Kansas Laboratories Pvt. Ltd 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.	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~~.

Analyst / Junior Analyst /

4/1/2001

Signature of Officer In-Charge

Computerised File Name: D&Creport1870

A/ 3327/1

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/2156/00 Dt of Receipt: 15.12.00

1.Name, address and license No. of Manufacturer / Supplier's from whom sample received.....	Adviser (Health),State Urban Development Agency, Ilgus Bhawan, II-C Block, Sec-III, Bidhan Nagar, Cal-91,
2.Reference No. and date of Manufacturer / Supplier's letter under which the sample was forwarded	SUDA/120/96 (Pt-II)/631 dt. 24.11.00
3.Name of Drug / Cosmetics / Raw material purporting to be contained in the sample.....	Bromhexine Tablet B.P. 8mg
4.Detail of Raw material/final product in bulk / final product (in finished pack) as obtained from the manufacturer for analysis.....	10 tabs X 5 strips
(a) Original manufacturer's name (in case of raw materials and drugs repacked).....	Kansas Laboratories Pvt. Ltd 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.	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~~..

Analyst / Junior Analyst /

4/1/2001

Signature of Officer In-Charge

3

A/3324/01

Computerised File Name: D&Creport1869

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 Dt of Receipt: 15.12.00

1.Name, address and license No. of Manufacturer / Supplier's from whom sample received.....	Adviser (Health),State Urban Development Agency, Ilgus Bhawan, II-C Block, Sec-III, Bidhan Nagar, Cal-91,
2.Reference No. and date of Manufacturer / Supplier's letter under which the sample was forwarded	SUDA/120/96 (Pt-III)/633 dt. 24.12.00
3.Name of Drug / Cosmetics / Raw material purporting to be contained in the sample.....	Paracetamol tablets I.P.
4.Detail of Raw material/final product in bulk / final product (in finished pack) as obtained from the manufacturer for analysis.....	10 tabs X 5 strips
(a) Original manufacturer's name (in case of raw materials and drugs repacked).....	Kansas Laboratories Pvt. Ltd 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 circular scored tablet in blister pack.

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

Assay:--

	Found/tab	Claim/tab	Limit
Paracetamol I.P.	479.0mg	500.0mg	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 / .....

4/1/2001

Signature of Officer In-Charge

Computerised File Name: D&Creport1869

A/3324/01

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 Dt of Receipt: 15.12.00

1.Name, address and license No. of Manufacturer / Supplier's from whom sample received.....	Adviser (Health),State Urban Development Agency, Ilgus Bhawan, II-C Block, Sec-III, Bidhan Nagar, Cal-91,
2.Reference No. and date of Manufacturer / Supplier's letter under which the sample was forwarded	SUDA/120/96 (Pt-III)/633 dt. 24.12.00
3.Name of Drug / Cosmetics / Raw material purporting to be contained in the sample.....	Paracetamol tablets I.P.
4.Detail of Raw material/final product in bulk / final product (in finished pack) as obtained from the manufacturer for analysis.....	10 tabs X 5 strips
(a) Original manufacturer's name (in case of raw materials and drugs repacked).....	Kansas Laboratories Pvt. Ltd 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 circular scored tablet in blister pack.

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

Assay:--

	Found/tab	Claim/tab	Limit
Paracetamol I.P.	479.0mg	500.0mg	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 / Parsenjeet

Signature of Officer In-Charge

3

A/ 3323/01

Computerised File Name: D&Creport1868

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/2159/00 Dt of Receipt: 15.12.00

1.Name, address and license No. of Manufacturer / Supplier's from whom sample received.....	Adviser (Health),State Urban Development Agency, Ilgus Bhawan, II-C Block, Sec-III, Bidhan Nagar, Cal-91,
2.Reference No. and date of Manufacturer / Supplier's letter under which the sample was forwarded	SUDA/120/96 (Pt-III)/633 dt. 24.12.00
3.Name of Drug / Cosmetics / Raw material purporting to be contained in the sample.....	Paracetamol tablets I.P.
4.Detail of Raw material/final product in bulk / final product (in finished pack) as obtained from the manufacturer for analysis.....	10 tabs X 5 strips
(a) Original manufacturer's name (in case of raw materials and drugs repacked).....	Kansas Laboratories Pvt. Ltd 8/1, Lal Bazar Street, Cal-700 001, DL No. – DL 1446M
(b) Batch Number	K-1460
(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 circular scored tablet in blister pack.

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

Assay:--

	Found/tab	Claim/tab	Limit
Paracetamol I.P.	509.79mg	500.0mg	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 / Signature of Officer In-Charge

Computerised File Name: D&Creport1868

A/ 7323/99

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/2159/00 — Dt of Receipt: 15.12.00

1.Name, address and license No. of Manufacturer / Supplier's from whom sample received.....	Adviser (Health),State Urban Development Agency, Ilgus Bhawan, II-C Block, Sec-III, Bidhan Nagar, Cal-91,
2.Reference No. and date of Manufacturer / Supplier's letter under which the sample was forwarded	SUDA/120/96 (Pt-III)/633 dt. 24.12.00
3.Name of Drug / Cosmetics / Raw material purporting to be contained in the sample.....	Paracetamol tablets I.P.
4.Detail of Raw material/final product in bulk / final product (in finished pack) as obtained from the manufacturer for analysis.....	10 tabs X 5 strips
(a) Original manufacturer's name (in case of raw materials and drugs repacked).....	Kansas Laboratories Pvt. Ltd 8/1, Lal Bazar Street, Cal-700 001, DL No. – DL 1446M
(b) Batch Number	K-1460
(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 circular scored tablet in blister pack.

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

Assay:--

	Found/tab	Claim/tab	Limit
Paracetamol I.P.	509.79mg	500.0mg	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 / .....

Signature of Officer In-Charge

3

Computerised File Name: D&Creport1867

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 Dt of Receipt: 15.12.00

1.Name, address and license No. of Manufacturer / Supplier's from whom sample received.....	Adviser (Health),State Urban Development Agency, Ilgus Bhawan, II-C Block, Sec-III, Bidhan Nagar, Cal-91,
2.Reference No. and date of Manufacturer / Supplier's letter under which the sample was forwarded	SUDA-120/96 (Pt-III)/633 dt. 24.11.00
3.Name of Drug / Cosmetics / Raw material purporting to be contained in the sample.....	Paracetamol tablets I.P.
4.Detail of Raw material/final product in bulk / final product (in finished pack) as obtained from the manufacturer for analysis.....	10 tabs X 5 strips
(a) Original manufacturer's name (in case of raw materials and drugs repacked).....	Kansas Laboratories Pvt. Ltd 8/1, Lal Bazar Street, Cal-1, DL No. –DL 1446M
(b) Batch Number	1459
(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.

Description :-White circular scored tablet in blister pack.

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

Assay:-

	Found/tab	Claim/tab	Limit
Paracetamol I.P.	481.8mg	500.00mg	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-..

Does not comply with respect to Cresol Content.

Analyst / Junior Analyst / Pasupur.

Signature of Officer-In-Charge

Tab. Furazolidine -
K-1470

Tab - Chlorpheniramine
maleate.
J- 1458

Rg No PPDE/Lab 1045 dt. 2-1-2001 Rec. on 5th Jan. 2001

Computerised File Name: D&Creport1826

A/ 3286/07

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

1.Name, address and license No. of Manufacturer / Supplier's from whom sample received.....	Adviser (Health),State Urban Development Agency, Ilgus Bhawan, II-C Block, Sec-III, Bidhan Nagar, Cal-91,
2.Reference No. and date of Manufacturer / Supplier's letter under which the sample was forwarded	SUDA-120/96 (Pt-III)/512 dt. 13.12.00
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 product (in finished pack) as obtained from the manufacturer for analysis.....	50 tablets
(a) Original manufacturer's name (in case of raw materials and drugs repacked).....	M/s. Kansas Labs (P) Ltd 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:--

	Found/tab	Claim/tab	Limit
Furazolidone I.P.	94.83mg	100.00mg	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.

Analyst / Junior Analyst /

Signature of Officer-in-Charge

2/1/2007

Computerised File Name: D&Creport1826

A/ 3286/07

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

1.Name, address and license No. of Manufacturer / Supplier's from whom sample received.....	Adviser (Health),State Urban Development Agency, Ilgus Bhawan, II-C Block, Sec-III, Bidhan Nagar, Cal-91,
2.Reference No. and date of Manufacturer / Supplier's letter under which the sample was forwarded	SUDA-120/96 (Pt-III)/512 dt. 13.12.00
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 product (in finished pack) as obtained from the manufacturer for analysis.....	50 tablets
(a) Original manufacturer's name (in case of raw materials and drugs repacked).....	M/s. Kansas Labs (P) Ltd 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:--

	Found/tab	Claim/tab	Limit
Furazolidone I.P.	94.83mg	100.00mg	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.

Analyst / Junior Analyst /

Signature of Officer In-Charge

• Computerised File Name: D&Creport1844

A/ 3329/01

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/2157/00 Dt of Receipt: 15.12.00

1.Name, address and license No. of Manufacturer / Supplier's from whom sample received.....	Adviser (Health),State Urban Development Agency, Ilgus Bhawan, II-C Block, Sec-III, Bidhan Nagar, Cal-91,
2.Reference No. and date of Manufacturer / Supplier's letter under which the sample was forwarded	SUDA-15/98 (Pt-II)/720 dt. 14.12.00
3.Name of Drug / Cosmetics / Raw material purporting to be contained in the sample.....	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 materials and drugs repacked).....	Kansan Laboratories Pvt. Ltd 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

5.**Result of Analysis Recorded Below:**

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:--

	Found/tab	Claim/tab	Limit
C.P. Maleate I.P.	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~~..

Analyst / ~~Senior Analyst~~

Signature of Officer In-Charge

Computerised File Name: D&Creport1844

A/ 3 3329/01

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/2157/00 Dt of Receipt: 15.12.00

1.Name, address and license No. of Manufacturer / Supplier's from whom sample received.....	Adviser (Health),State Urban Development Agency, Ilgus Bhawan, II-C Block, Sec-III, Bidhan Nagar, Cal-91,
2.Reference No. and date of Manufacturer / Supplier's letter under which the sample was forwarded	SUDA-15/98 (Pt-II)/720 dt. 14.12.00
3.Name of Drug / Cosmetics / Raw material purporting to be contained in the sample.....	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 materials and drugs repacked).....	Kansan Laboratories Pvt. Ltd 8/1, Lal Bazar Street, Cal-700 001, DL 14o4M
(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' 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:--

	Found/tab	Claim/tab	Limit
C.P. Maleate I.P.	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~~.

Analyst

Signature of Other

R.C.H

SUDA

STATE URBAN DEVELOPMENT AGENCY

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

Ref No..... *Date*.....

SUDA-120/96(Pt.III)/.... . 2000

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,

Yours faithfully,
18/1/2000
Adviser (Health)
SUDA

LH/p-78

Tel No : 358-6403 / 6421 / 5767,Fax No. - 358-5800

SUDA

List of samples to be tested

Sl. No.	Name of Items	Mfg. By	Qty. of Sample	Batches Of items	Mfg. Date	Exp. Date
1.	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

Adviser (Health)
SUDA

LH/p-79

SUDA

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)/631** *Date* **24.11.2000**

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,

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

A list 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)

24/11/2000

Labtst/LH/p17

SUDA

List of samples to be tested

Sl. No.	Name of Items	Mfg. By	Qty. of Sample	Batches Of items	Mfg. Date	Exp. Date
1.	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
4.	Paracitamol IP 500 mg.	Kansas Lab	3 x 50 tab	K-1459 K-1460 K-1468	11/2000	10/2003
5.	Cotrimoxazole IP S/S	Kansas Lab	2 x 50 tab	~~K-1462~~ ~~K-1463~~ K-1463	11/2000	10/2003

Adviser (Health)
SUDA

LH/p-72

SUDA

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

Date 24.11.2000

From: Adviser(Health)
SUDA

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Adviser (health)
24/11/2000

Labsn/LH/p17

Tel/Fax No.: 359-3184

SUDA

List of samples to be tested

Sl. No.	Name of Items	Mfg. By	Qty. of Sample	Batches Of items	Mfg. Date	Exp. Date
1.	Compound Magnesium Trisilicate tab	Avron Lab	10 x 100 tab	AS-(4140 to 4144)	Nov.2000	Oct.2003
2.	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
4.	Oral Rehidration Salt	Pure Pharma Ltd	16 x 4 pes.	0021 to 0024	Nov.2000	Oct.2002
5.	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	11	Nov.2000	Oct.2002
10.	Absorbent Cotton 50 gm	Jaya & Co.	2(4 x 50) gm	43,46	Nov.2000	3 years

Adviser (Health)
SUDA

24/11/2000

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Sl.No.	Name of Item	Mfg. Name.	Batch No.	Mfg.Date	Expy.Date	Quantity
1.	Compound Magnesium Trisilicate Tab.	Avron Labaratory	AS4140- AS4141- AS4142- AS4143- AS4144-	11/2000 11/2000 " " "	10/2003 10/2003 " " "	2x100 Tabs. " " " "
	Mag. Trisilicate - 250 mg Dried Aluminium Hydroxide Gel-250 mg					
2.	Fesolic-L	Pure Pharma Ltd.	0017	10/2000	9/2002	1x50
3.	Fesolic-S	-do-	0009	10/2000	9/2002	1x50
4.	Oral Rehidration Salt	-do-	0021 0022 0023 0024	11/2000 -do- -do- -do-	10/2002 -do- -do- -do-	4 (4 PC) " " "
5.	Antiseptic Lotion	Hindustan Drugs	LE/06	Apr-99	MAR-2004	2x200
6.	Merbromin 20 gm.	C.D.Pharmeutical Works	377	-	-	2x20gm.
7.	Nitrofurantoin Skin Cream 15gm. tube	Pilco Pharma Pvt.Ltd.	NF-1234	11/2000	10/2002	10 Tubes
8.	Adhesive Plaster (10cmx5mm:	Precision Coatings LTD.	010779 010776 010780	OCT.2000	3 Years	1 Reel " "
9.	Benzyl Banzoate Application	Hindustan Drugs	11	NOV-2000	Oct. 2002	(2) 2x100ml.
10.	Absorbent Cotton 50gm.	Jaya & Co.	43 46	NOV-2000 NOV-2000	3 years 3 years.	4x50gm. 4x50gm.

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FOR KANSAS LABS. P. LTD.

Sl.No.	Name of the items.	Mfg. by :	Batch No.	Mfg.Date	Exp.Date	Quantity
1.	Mebendazole IP-100mg	Kansas Lab.	H 1427	08/2000	07/2003	1x50 Tabs.
2.	Bromhexine Hcl.BP.8mg	-do-	G 1414	07/2000	06/2003	1x50 Tabs.
3.	Chlorphniramine Maleate IP 4mg.	-do-	J 1458	10/2000	09/2003	1x50 Tabs.
4.	Paracetamol IP 500mg	-do-	K 1459	11/2000	10/2003	1x50 Tabs.
	-do-	-do-	K 1460	11/2000	10/2003	1x50 Tabs.
	-do-	-do-	K 1468	11/2000	10/2003	1x50 Tabs.
5.	Cotrimoxazole IP S/S Trimethoprim - 80 mg & Sulphamethoxazole 400	-do-	K 1462	11/2000	10/2003	1x50 Tabs. X Not supplied
	-do-	-do-	K 1463	11/2000	10/2003	1x50 Tabs.

Office of Adviser Health Stds
TAIF

23 NOV 2000

FOR KANSAS LABS. P. LTD.

SL.NO.	NAME OF ITEM	MFG.NAME	BATCH NO.	MFG.DATE	EXPY. DATE	QUANTITY
1.	FURAZOLIDONE TABLETS NFI III 100mg.	Kansas Laboratories Pvt.Ltd.	K 1470	11/2000	10/2003	2 x 50 Tabs.
2.	METRONIDAZOLE TABLETS I.P. 200mg.(Film Coated	-"-	K 1472	11/2000	10/2003	2 x 50 Tabs.
3.	ASPIRIN TABLETS I.P 300mg.	-"-	K 1477	11/2000	4/2002	3 x 50 Tabs.
4.	OXYPHENONIUM BROMIDE TABLETS I.P 5mg.	-"-	K 1476	11/2000	10/2003	10x10 Tabs.
5.	CHLORAMPHENICOL EYE OINTMENT I.P 1% W/W (EYE Applicaps)	Jyoti Capsules	JGC 21010	NOV.2000	APRIL 2002	2 x 50 Caps.

For KANSAS LABORATORIES PVT.

Tab. Fersolic -L

B.no. 0017

Tab. Fersolic -S

B no. 0009

Antiseptic LOZ -

LE-06

Mebendazole tab. 100 mg

H - 1427

Aspirin tab K-1477

Oxyphenonium Bromide - 500 mg.

K - 1476

Chloramphenicol Eye Applicap.

~~J-G~~ JBC 21010