



Dissofilm: A Novel Approach for Delivery of Phenobarbital; Design and Characterization

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ABSTRACT

An attempt to develop and evaluate mouth-dissolving film of phenobarbital for quick effect in treatment of epilepsy occurring in pediatric population has been made in the present study. Suitable film formers and plasticizers are selected based on optimization studies. Effect of superdisintegrants in formulation of mouth dissolving films at different concentrations has been investigated. Films were prepared by solvent casting method. The prepared films were evaluated for physicochemical parameters, *in vitro* disintegration and dissolution time, *in vitro* release rate study, stability study, and *in vivo* animal safety study. The best formulation was found to be F3 showing 96.57% drug release in 14 min, following first-order kinetics. X-Ray diffraction studies show change in crystalline nature of drug in formulation. *In vivo* studies in hamster reports effective and safe use of formulation in animals.

Key words: Hydroxypropylmethyl cellulose, mouth-dissolving film, phenobarbital, sodium starch glycolate, X-ray diffraction

INTRODUCTION

Rapidly dissolving films (RDFs) have recently gained popularity in the form of breath fresheners. These films are placed in the mouth and dissolve quickly to release the flavor. Rapidly dissolvable films were developed on the technology of transdermal patch. RDFs are already being used in breath-freshening products from Warner Lambert and Wrigley's in the US and Europe. Zengen recently launched a chloraseptic relief strip in the US to

deliver benzocaine-a local anesthetic to treat sore throats. This delivery system is simply placed on a patient's tongue or any oral mucosal tissue. Instantly wet by saliva, the film rapidly hydrates and adheres onto the site of application. It then rapidly disintegrates and dissolves to release the medicament for mucosal absorption or, with modifications, allows oral gastrointestinal absorption with a quick-dissolving aspect.^[1]

Advantages of rapidly dissolving films

- Ease of administration to patients who cannot swallow, such as the elderly, stroke victims and bedridden patients; patients who should not swallow, such as renal failure patients; and who refuse to swallow, such as pediatrics, geriatric, and psychiatric patients.
- Patient's compliance for disabled bedridden patients and for traveling and busy people who do not have

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ready access to water.

- More rapid drug absorption from the pregastric area, i.e., mouth, pharynx, and oesophagus which may produce rapid onset of action.
- Pregastric absorption can result in improved bioavailability, reduced dose, and improved clinical performance by reducing side effects.

Epilepsy is a chronic condition characterized by the repeated attacks of epileptic seizures. Epileptic seizures can occur in nonepileptic patients subjected to a variety of stresses and stimuli. Epileptic seizures differ from other types of paroxysmal attacks by their clinical expression and by their mechanism. The epileptic seizures and paroxysmal attacks are expressed on the electroencephalogram by the epileptic discharge.^[2]

Phenobarbital, the longest-acting barbiturate, is used for its anticonvulsant and sedative-hypnotic properties in the management of all seizure disorders except absence. Absorbed in varying degrees following oral, rectal or parenteral administration. The salts are more rapidly absorbed than are the acids. The rate of absorption is increased if the sodium salt is ingested as a dilute solution or taken on an empty stomach. Phenobarbitone is the clinically effective anticonvulsant enhances the inhibitory effect of gamma aminobutyric acid (GABA) by facilitating the GABA-mediated opening of chloride channels.^[3]

An attempt was made in the present investigation to prepare fast dissolving drug delivery systems of phenobarbital by solvent-casting method for rapid therapy of epileptic seizures.

MATERIALS AND METHODS

Materials

Phenobarbital was a gift sample obtained from Anglo-French Drugs and Industries Ltd, Bangalore, India. Methocel – E 15, E 50 obtained from Dow Chemical Company, Mirland. Pectin and Pullulan were obtained from Lipoid GmbH, Germany. Hydroxypropyl cellulose, Gelatin, Xanthan Gum, Propylene Glycol and Aspartame were obtained from S.D. Fine-Chem Ltd., Mumbai, India. Sodium Starch Glycolate and Croscarmellose Sodium were obtained from DMV— Fonterra Excipients, Netherlands. INDION-414 was obtained from Ion Exchange India Ltd., Ankaleshwar, India. Other chemicals used were of analytical grade and procured from S.D. Fine Chemicals (Mumbai, India). Concentrations of Phenobarbital were measured with a UV–VIS spectrometer (UV-1700,

Shimadzu Corporation, Tokyo, Japan). Interaction between Phenobarbital and polymers was verified using FTIR and TLC techniques.

Optimization of the components for formulation of placebo fast dissolving films

The placebo films were prepared by solvent-casting method. Hydroxypropylmethyl cellulose (HPMC) is known for its good film forming properties and has excellent acceptability. Hence, HPMC namely Methocel E15 was evaluated to optimize the effective concentration keeping the other components of the formulations to a constant mean value. Similarly an identical approach was used to optimize the other components such as secondary film modifier (pectin), film modifier (Xanthan gum) and plasticizer (propylene glycol) using the previously optimized concentration of respective components. The general method for preparation of placebo film is as follows. Previously primary film former, secondary film formers and film modifiers were soaked in water for 4 h and then uniformly dispersed to obtain dispersion. Aspartame and plasticizers were uniformly dissolved in 5 ml of 95% (v/v) ethanol. Alcoholic solution and the polymeric dispersion were mixed to obtain a homogenous dispersion. Ten milliliters of the dispersion was cast onto each petri dish which was lubricated with olive oil. The dispersion was dried in a hot air oven at 40–45°C. The films were carefully removed from petri dishes and cut into strips of dimensions 3 × 3 cm and stored in an air tight glass bottle. The films were evaluated for imperfections and cuts, mechanical strength, thickness, and dissolution time.^[4]

Preparation of the phenobarbital containing fast dissolving films

Phenobarbital (PBT) containing fast dissolving films were fabricated as per the method described for the fabrication of blank fast dissolving films. PBT was incorporated in 5 ml of 95% (w/v) ethanolic solution and the rest of the procedure was same as that for the fabrication of blank fast dissolving films. Superdisintegrants (sodium starch glycolate or croscarmellose sodium) was uniformly dispersed in aqueous dispersion. The compositions of PBT containing films are shown in Table 1.

Evaluation of formulated mouth dissolving films

Film thickness

Thickness of films was measured using calibrated dial caliper.^[4]

Weight variation

The study was carried out on ten films obtained from

Table 1: Compositions of phenobarbital mouth-dissolving films

Ingredients	F1	F2	F3	F4	F5	F6	F7
Phenobarbital	117	117	117	117	117	117	117
METHOCEL- E 15	150	150	150	150	150	150	150
Pectin	50	50	50	50	50	50	50
HPC (low viscosity grade)	25	25	25	25	25	25	25
Menthol	40	40	40	40	40	40	40
Aspartame	3	3	3	3	3	3	3
Glycerol	125	125	125	125	125	125	125
Xanthan gum	10	10	10	10	10	10	10
Sodium starch glycolate	-	50	100	150	-	-	-
Croscarmellose sodium	-	-	-	-	50	100	150
Propylene glycol	100	100	100	100	100	100	100
Water	5	5	5	5	5	5	5
Ethanol	7	7	7	7	7	7	7

All the weights are taken in mg, except water and ethanol taken in ml

each formulation batch. The mean weight of film as well as the deviation from the mean was calculated and recorded.^[5]

Hydration study (water uptake/swelling study)

The film sample was weighed and placed on a preweighed stainless steel wire mesh. The wire mesh was then submerged in a petridish containing 20 ml distilled water. Increase in weight of the film was determined at regular time intervals until a constant weight was obtained.^[6,7]

The hydration ratio of the film was calculated using following formula:

$$\% \text{ Hydration ratio} = \frac{W_t - W_o}{W_o} \times 100,$$

where W_t = weight of film at time t and W_o = weight of film at zero time.

Moisture loss (moisture vapor transmission)

The percent moisture loss was determined by placing prepared film in desiccators containing anhydrous calcium chloride. After three days, the film was taken and reweighed. The percent moisture loss was calculated using following formula:^[8,9]

$$\text{Moisture loss (\%)} = \frac{W_o - W_t}{W_o} \times 100,$$

where W_o = initial weight and W_t = final weight.

Percent elongation at break

The prepared film was pulled by means of a pulley system.^[10] Weights were gradually added to the pan to increase the pulling force till the film was broken. The elongation was determined by noting the distance traveled by pointer before break of film on the graph paper. The percent elongation at break was calculated by using formula:

Percent elongation at break (mm^{-2})

$$= \frac{L_1}{L_o} \times \frac{100}{\text{cross-sectional area}},$$

where L_1 = increase in the length, L_o = initial length.

Tensile strength

Film strip of dimension $5 \times 2 \text{ cm}^2$ and free from air bubbles or physical imperfections was held between two clamps positioned at a distance of 3 cm apart. A cardboard was attached on the surface of the clamp via a double sided tape to prevent the film from being cut by the grooves of the clamp. During measurement, the strips were pulled at the bottom clamp by adding weights in pan till the film breaks. The force was measured when the films broke. Results from film samples, which broke at and not between the clamps were not included in calculations:^[11]

Tensile strength (kg / mm^2)

$$= \frac{\text{Breaking force (F)}}{\text{Cross-sectional area of sample (A)}}$$

Folding endurance

This parameter was determined by repeatedly folding one film at the same place till it broke. The number of times the film could be folded at the same place without breaking/cracking give the value of folding endurance.^[8,9]

Drug content and content uniformity

The drug content and content uniformity test was performed to ensure uniform and distribution of drug. 3 cm^2 films was cut, weighed and dissolved in 100 ml methanol. Three milliliters of supernatant aqueous solution was withdrawn and transferred in 100 ml volumetric flask. Final dilution up to mark was done using simulated saliva and analyzed spectroscopically at 254 nm wavelength. The concentration of phenobarbital (in $\mu\text{g}/\text{ml}$) was calculated by using standard calibration curve of phenobarbital. Content uniformity of phenobarbital films were done by selecting randomly five 3 cm^2 films from different batches and performing study same as drug content.^[4]

In vitro disintegration and dissolution time

The disintegration time is the time when a film starts to break or disintegrate. The dissolution time is the time when the film completely dissolves. The *in vitro* disintegration and dissolution time of fast-dissolving films was determined visually in a glass dish of 25 ml distilled water with swirling every 10 s. Disintegration and dissolution time was measured.^[4]

In vitro dissolution study (drug release rate study)

The *in vitro* dissolution test was carried out in USP basket type

apparatus.^[12,13] Samples of PBT-loaded films were equivalently containing 15 mg (3 cm²) was cut and placed basket. The dissolution medium consisted of 500 ml freshly deionized simulated saliva (pH 6.8), maintained at 37 ± 1°C and stirred at 100 r/min. Samples of 1 ml were withdrawn at predetermined time intervals and replaced with fresh medium. The samples were filtered through Whatman filter paper and PBT concentrations were assayed spectrophotometrically at 254 nm after final appropriate dilution.

The dissolution profile of conventional marketed tablet LUMINAL[®] containing 15 g equivalent phenobarbital was also determined using same experimental setup.

Surface morphology study using scanning electron microscopy

The surface morphology of formulated PBT film was assessed using a scanning electron microscope. Samples were mounted on round brass stubs (12 mm diameter) using double-backed adhesive tape and then sputter coated for 8 min under argon atmosphere with gold before examination under the scanning electron microscope. Pictures were then taken at an excitation voltage of 15 kV.^[9]

Stability studies

Stability tests are the series of tests designed to obtain information on the stability of the pharmaceutical product in order to define its shelf life and utilization period under specified packaging and storage conditions.

From the seven batches of PBT films, optimized formulation was tested for stability studies. Films of optimized formula were stored at two different storage conditions namely 30°C ± 0.5/60 ± 1% RH and 40°C ± 0.5/75 ± 1% RH. Each film was wrapped in a butter paper followed by aluminium foil and placed in an aluminium pouch, which was heat-sealed at the end. The films were evaluated for weight, PBT content and *in vitro* drug release after storage for 10, 20, and 30 days. The values for *in vitro* PBT release from the films were calculated and were compared for change in the dissolution profile.^[4]

In-vivo animal safety study (irritation study)

The *in vivo* animal safety study is carried out to find safety of films in oral cavity. Hamster cheek pouch was selected as model for study.^[14]

Either sex of Hamster of 100–150 g was selected for study. They were housed individually in an environmentally controlled room with 12 h light/dark cycle and had free access to food and water. After 7 days acclimatization period, they were randomly selected for different experimental groups.

All the experimental procedures were carried out in accordance with committee for the purpose of control and supervision of experiments on animal (CPCSEA) guidelines. The study was reviewed and approved by the Institutional Animal Ethics Committee (JNMC/IAEC/2/28/2008). J.N. Medical College, KLE University, Belgaum, India.

Experimental study design in hamsters

Group: I Normal or placebo film treated group.

Group: II PBT film treated group.

The placebo or PBT film was placed twice a day for 4–5 consecutive days. The prepared PBT film sample was placed in cheek pouch of hamster for 10 min and then rinsed using distilled water. Then pouch was observed for any adverse effects (irritation), redness immediately and after 24 h interval.

RESULTS AND DISCUSSION

Thickness

The thicknesses of formulated films were found to be in range of 13.76 ± 0.539 to 14.3334 ± 0.0506 mm. The mean values are tabulated in Table 2. The values are almost uniform in all formulations. Obtained results has shown that increase in film thickness decreases tensile strength while increases % elongation. Further increases in thickness of film increases crystallinity of film and decrease dissolution rate.

Weight variation test

The percentage weight variation for all the formulation is tabulated in Table 2. All the films passed weight variation test as the % weight variation was within the pharmacopoeial limits of ±7.5%. It was found to be in range of 110.2 ± 0.20 to 127.2 ± 0.30 mg. The weight of all the films was found to be uniform.

Hydration study (water uptake/swelling study)

The hydration ratio defines the ability of films to absorb water from environment or from dissolution media. It is also termed as water uptake study. Hydration ratio gives an idea of films dissolution time and disintegration time. Higher values of hydration ratio indicate that films dissolve faster and easily. Hydration ratio values of all formulation are shown in Table 2. Increase in concentration of superdisintegrant increases hydration ratio or water uptake and decrease disintegration and dissolution time. F3 and F6 formulations showed higher hydration ratio while F1 formulation shows low hydration ratio.

Table 2: Thickness, mean weight (mg), drug content (mg), %hydration ratio, %moisture loss of mouth dissolving films of phenobarbital

Formulation	Thickness	Mean weight (mg) ^a	Drug content (mg) ^b	% Hydration ratio ^a	% Moisture loss ^a
F1	13.9166 ± 0.045	110.2 ± 0.20	13.65 ± 0.05	0.808 ± 0.004	2.995 ± 0.002
F2	14.3054 ± 0.0986	117.3 ± 0.25	16.29 ± 0.076	0.727 ± 0.003	1.024 ± 0.004
F3	14.22 ± 0.0239	121.8 ± 0.95	18.71 ± 0.26	0.616 ± 0.002	1.426 ± 0.005
F4	13.76 ± 0.539	119.3 ± 0.30	17.13 ± 0.13	0.518 ± 0.005	1.973 ± 0.003
F5	13.9722 ± 0.050	127.2 ± 0.30	14.50 ± 0.076	0.701 ± 0.001	2.203 ± 0.003
F6	14.3334 ± 0.050	123.6 ± 0.20	19.75 ± 0.17	0.505 ± 0.001	2.346 ± 0.001
F7	14.1388 ± 0.0326	119.1 ± 0.20	13.92 ± 0.047	0.462 ± 0.007	2.016 ± 0.002

^aSignificance value $P > 0.0001$, ^bSignificance value $P > 0.001$, $n =$ average of triplicate was determined

Moisture loss (moisture vapor transmission)

MVT is defined as the quantity of moisture transmitted through unit area of film in unit time. The moisture loss study gives an idea about films stability nature and ability of films to withstand its physicochemical properties under normal conditions. It also gives idea about hydrophilicity of film formulations. The obtained results are tabulated in Table 2. The obtained values are almost uniform and ranges from $1.024 \pm 0.004\%$ to $2.995 \pm 0.002\%$. F1, F5, and F6 formulation showed high % moisture loss while F2 and F3 formulations showed low % moisture loss.

Tensile strength

The tensile testing gives an indication of the strength and elasticity of the film, reflected by the parameters, tensile strength (TS), elastic modulus (EM) and elongation at break (E/B). A soft and weak polymer is characterized by a low TS, EM, and E/B; a hard and brittle polymer is defined by a moderate TS, high EM and low E/B; a soft and tough polymer is characterized by a moderate TS, low EM and high E/B; whereas a hard and tough polymer is characterized by a high TS, EM and E/B. Tensile strength of all prepared formulation is shown in Table 3. Results revealed that formulations showed better tensile strength and less to moderate % elongation.

Tensile strength was found in range of 0.462 ± 0.007 to 0.808 ± 0.004 kg/mm². There is no significant change in the tensile strength of all formulations. Addition of superdisintegrants (SD) affects tensile strength and also % elongation. An optimum concentration of superdisintegrant showed improvement of tensile strength, but at higher concentration there was no further increase in tensile strength. Higher concentration of superdisintegrant increases thickness and crystallinity of films which causes decrease in tensile strength as well as percent elongation.

Percent elongation

Percent elongation is mainly based on tensile strength

of films. The nature of polymers affects tensile strength and % elongation. Soft and brittle polymer increases tensile strength and decreases % elongation while hard and tough polymer increases tensile strength as well as % elongation. % Elongation of all formulation is tabulated in Table 3. The values of % elongation are in range of $4.108 \pm 0.001\%$ to $6.143 \pm 0.001\%$. Increase in concentration of superdisintegrant decreases % elongation which indicates that elasticity and softness of film decreases which helps in fast dissolution of films.

Folding endurance

Folding endurance measures the ability of patch to withstand rupture, higher the folding endurance lower will be chances of film to rupture easily and vice versa. The folding endurance of all formulations is tabulated in Table 3. Increase in concentration of superdisintegrants increases folding endurance of films but after specific concentration increase in concentration of superdisintegrants decreases folding endurance. This was due to increase in crystallinity of film and also film thickness. More the thickness of lower will be folding endurance. F3 and F6 formulation showed high folding endurance 55.67 ± 1.15 and 46.33 ± 0.57 , respectively.

Drug content and content uniformity study

The drug content and content uniformity test was performed to ensure uniform and accurate distribution of drug. The content uniformity was performed for all the seven formulations and results are tabulated in Table 2. Five trials from each formulation were analyzed spectrophotometrically. The mean value and standard deviation of all the formulations were calculated. The results indicated that in all the formulations the drug content was uniform. The cumulative percentage drug released by each film to the *in vitro* release studies was based on the mean content of the drug present in the respective film.

In vitro disintegration time dissolution time test

The disintegration time is the time when a film starts to break or disintegrate. The dissolution time is the time when

Table 3: Tensile strength, %elongation, folding endurance, disintegration time, dissolution time of mouth-dissolving films of phenobarbital

Formulation	Tensile strength ^a kg/mm ²	% Elongation ^a at break	Folding endurance ^a	Disintegration time ^a (s)	Dissolution time ^a (s)
F1	0.808 ± 0.004	6.143 ± 0.001	45 ± 1	31.33 ± 0.577	126 ± 0.58
F2	0.727 ± 0.003	5.144 ± 0.001	50 ± 2	27 ± 1.00	94.00 ± 1.0
F3	0.616 ± 0.002	5.132 ± 0.001	55.67 ± 1.1	23.33 ± 1.155	73.67 ± 1.528
F4	0.518 ± 0.005	4.114 ± 0.001	44.00 ± 2	23.33 ± 1.00	72.33 ± 1.528
F5	0.701 ± 0.001	5.141 ± 0.001	40.67 ± 0.57	28 ± 1.00	99.33 ± 1.528
F6	0.505 ± 0.001	4.118 ± 0.001	46.33 ± 0.57	26 ± 0.577	79.33 ± 1.528
F7	0.462 ± 0.007	4.108 ± 0.001	37.67 ± 0.57	24.33 ± 1.155	76.00 ± 2.0

^a: significance value $P > 0.0001$, n = Average of triplicate was determined

the film completely dissolves. Disintegration time and dissolution time of formulated films is tabulated in Table 3. Results revealed that addition of superdisintegrants decreases the disintegration time and dissolution time. Disintegration time and dissolution time of all formulation were in range of 23.33 ± 1.00 to 31.33 ± 0.577 s and 72.33 ± 1.528 to 126 ± 0.58 s, respectively.

In vitro dissolution study (drug release rate study)

Cumulative drug release and cumulative % drug retained were calculated on the basis of drug content of phenobarbital present in the respective film. The results obtained in the *in vitro* drug release for the formulations F1 to F7 and marketed tablet "LUMINAL[®]" is tabulated in Table 4. The plots are depicted in Figure 1 (a and b) for % cumulative drug release versus time. Formulation F1, F2, F3, shows drug release up to 87.836%, 89.282%, 96.574%, respectively, at end of 14 min. Rapid drug dissolution was observed in F4, F5, F6, and F7 which release 96.307%, 88.442%, 93.735%, and 95.114%, respectively, at end of 14 min. F3 formulation shows highest percent of drug release (96.574%) than other formulations. The observed rapid dissolution might be due to fast disintegration of films and rapid dissolution of drug. Increase in the concentration of superdisintegrants increases dissolution rate of drug which was observed in F3, F4, F6 formulation, but further increase in concentration of superdisintegrants will not increase dissolution rate much more which was observed in film F4, F7. Also increase in the concentration of superdisintegrants causes precipitation of films and decreases tensile strength and folding endurance. This is observed at concentration of $\geq 2\%$ of the film. The drug release was completely achieved in a shorter duration of time. *In vitro* release rate study of F3 formulation vs. conventional marketed tablet LUMINAL[®] containing 15 mg phenobarbital has shown that F3 release found to be faster and complete within 15 min. *In vitro* release of LUMINAL[®] was found to be 15% in 14 min and up to 46% at half an hour. The correlation coefficient values obtained are shown

Table 4: In vitro release profile of phenobarbital mouth dissolving film formulations

Time (min)	Cumulative % drug released							
	F1	F2	F3	F4	F5	F6	F7	LUMINAL [®]
1	16.781	18.935	23.117	22.672	17.648	20.592	21.668	0.264
2	28.420	29.225	37.169	35.966	27.640	27.744	33.679	0.794
3	40.257	39.181	47.094	46.891	37.553	33.631	45.624	1.591
4	47.997	45.665	55.892	55.974	45.648	41.376	54.993	2.393
5	56.048	54.404	66.182	65.513	53.921	49.814	64.147	3.463
6	66.290	67.891	76.921	70.759	64.645	60.643	73.747	4.407
7	72.022	74.791	84.697	84.721	72.886	67.230	78.642	5.488
8	77.785	78.333	90.806	89.120	79.434	77.632	88.187	6.707
9	80.355	83.329	92.349	91.233	83.475	83.006	91.175	7.932
10	82.937	84.563	93.636	92.404	84.595	87.496	92.196	9.165
11	84.054	86.062	94.401	93.307	85.719	90.444	93.088	10.932
12	85.578	87.045	95.168	94.621	86.847	91.451	93.850	12.312
13	86.839	88.162	95.805	95.39	87.577	92.461	94.614	13.700
14	87.836	89.282	96.574	96.307	88.442	93.735	95.114	14.831

in Table 5. The results were found to be linear for first-order release. It is concluded that release of drug from formulations F1–F7 followed first order.

Surface morphology study using scanning electron microscopy

SEM was performed on the films to assess changes in their surface morphology prior to and after dissolution testing and is depicted in Figure 2 (a–c). A smooth and compact surface with crystals of phenobarbital was noted at time 0 min for the optimized film F3. As dissolution time progressed to the first minute, film appeared porous. After 1 min the surface morphology of films showed significant changes in texture, to the extent that the film developed clearly visible pores with solubilization of phenobarbital crystals. SM of pure drug was also taken and it showed crystalline nature of phenobarbital.

Stability studies

F3 formulation was selected for stability studies on the basis of high cumulative % drug release and also results of *in vitro* disintegration time. From these results it was concluded

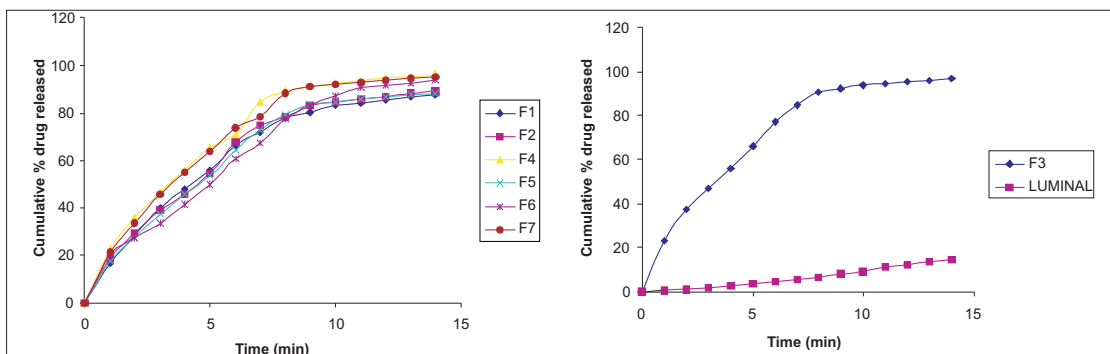


Figure 1: (a) *In vitro* release profile of phenobarbital mouth dissolving film formulations F1, F2, and F4–F7 (b) *In vitro* release profile of phenobarbital mouth-dissolving film formulation F3 and LUMINAL

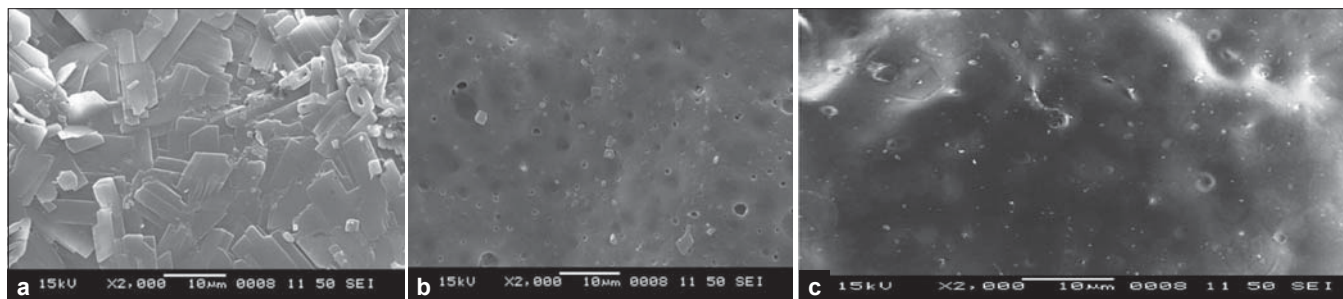


Figure 2: Scanning electron microscope image of (a) pure phenobarbital, (b) formulation F3 at t = 0 min, (c) formulation F3 at t = 1 min

Table 5: Model fitting of the release profile using five different models (R value)

Formulation code	Mathematical models (Kinetics)			
	Zero order	First order	Higuchi	Best fit kinetic
F1	0.882	0.978	0.912	First order
F2	0.885	0.961	0.927	First order
F3	0.844	0.974	0.891	First order
F4	0.855	0.979	0.901	First order
F5	0.890	0.955	0.927	First order
F6	0.939	0.979	0.952	First order
F7	0.862	0.971	0.887	First order

that, formulations F3 is stable and retained their original properties with minor differences. The *in vitro* release profile of F3 at 30°C/60% RH and 40°C/75% RH conditions after 30 days was 95.114% and 92.316%, respectively, has indicated that no or minor alteration after storage.

***In vivo* animal safety study (irritation study)**

The *in vivo* animal safety study is carried out to ensure safe use of films in oral cavity. Hamster cheek pouch was selected as model for study. Images of cheek pouch area were taken before and after placing of the films. The observation of images obtained of cheek pouch area of hamster before and after placing film shows no irritation or redness. Therefore, F3 formulation has shown good acceptability for oral use in treatment of epileptic attack or seizures. Obtained results have been shown in Figure 3 (a and b).

CONCLUSIONS

Mouth dissolving films of Phenobarbital could be formulated with available low viscosity film formers viz. HPMC E15, Pectin, HPC, Xanthan gum. Phenobarbital, a poorly water soluble drug could be successfully incorporated in mouth dissolving film with the help of propylene glycol as solubilizer and plasticizer. In the present study successful optimization was done to study the influence of film formers, film modifier, plasticizers and superdisintegrants (SDs) in mouth dissolving films. Use of plasticizers in combination could give better results to films in respect to physicochemical parameter, such as tensile strength, %elongation, folding endurance and flexibility. Incorporation of superdisintegrants in the mouth dissolving film could be useful for rapid disintegration and dissolution time. Amongst all the developed formulations, phenobarbital mouth-dissolving film formulated using sodium starch glycolate (SSC) as superdisintegrant in concentration range of 1% (w/v), showed good *in vitro* disintegration and dissolution time.

In comparison with available marketed formulation, it has found that formulated films showed improved dissolution.

The *in vitro* drug release, *in vitro* stability evaluation and physicochemical/mechanical results obtained in this work, confirm the potentials of the mouth dissolving film of phenobarbital as a promising candidate for quick relief from epileptic seizures and attacks with better patient



Figure 3a: Placebo film safety study on hamsters



Figure 3b: PBT F3 optimized film safety study on hamsters

compliance, with all class of patients. From the *in vivo* animal safety studies, it was concluded that the optimized formulation had no irritation or redness on cheek pouch area of hamster and had good acceptability for oral use in treatment of epileptic attack or seizures.

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