

pared ocular insert film had good appearance with smooth surface. Ocusert films prepared were semi-transparent. Surface texture was smooth and uniform. Evaluation of ocular inserts for weight and thickness variation were carried out and analyzed by ANOVA.

Formulation Code	Rate controlling membrane				Ocusert reservoir			
	Polymer		Plasticizer	Solvent	Drug	Polymer	Plasticizer	Solvent
	HPMC % W/V	EC % W/V	DBP % W/W of polymer	Ethanol ml	Clotrimazole (mg)	Na CMC % W/V	PEG-400 % W/W	Water+Ethanol (1:1) ml
F1	2	-	30	15	50	2	30	15
F2	3	-	30	15	50	2	30	15
F3	4	-	30	15	50	2	30	15
F4	-	2	30	15	50	2	30	15
F5	-	3	30	15	50	2	30	15
F6	-	4	30	15	50	2	30	15
F7	1	1	30	15	50	2	30	15
F8	1.5	1.5	30	15	50	2	30	15
F9	2	2	30	15	50	2	30	15

Table 3: Composition of prepared ocuserts.

Formulation Code	Weight variation (mg) \pm SD*, (P value)	Thickness (mm) \pm SD*, (P value)	Folding endurance \pm SD#	Surface pH \pm SD#	Average drug content in each formulation (% of theoretical amount/ocusert = 0.9308) \pm SD#	% Moisture content \pm SD*, (P value)	% Moisture uptake \pm SD*, (P value)
F1	28.205 \pm 1.669 p<0.0001	0.0262 \pm 0.016 p<0.0001	78.33 \pm 6.02	7.16 \pm 0.404	92.42 \pm 3.309	3.19 \pm 0.056 p<0.0061	2.98 \pm 0.111 p<0.0246
F2	37.1 \pm 2.018 p<0.0001	0.326 \pm 0.02 p<0.0001	72 \pm 14.0	7.23 \pm 0.351	89.01 \pm 1.028	2.55 \pm 0.02 p<0.0007	3.17 \pm 0.036 p<0.0025
F3	47.165 \pm 2.499 p<0.0001	0.364 \pm 0.026 p<0.0001	64.33 \pm 4.16	7.6 \pm 0.5	90.72 \pm 2.138	3.19 \pm 0.053 p<0.0055	4.34 \pm 0.036 p<0.0025
F4	24.45 \pm 2.722 p<0.0001	0.224 \pm 0.021 p<0.0001	51 \pm 2.0	7.2 \pm 0.3	87.31 \pm 0.946	3.39 \pm 0.044 p<0.0037	2.42 \pm 0.177 p<0.0616
F5	34.38 \pm 2.177 p<0.0001	0.229 \pm 0.017 p<0.0001	45.66 \pm 2.51	7.46 \pm 0.416	83.9 \pm 0.882	4.57 \pm 0.046 p<0.0041	4.21 \pm 0.036 p<0.0025
F6	47.165 \pm 2.499 p<0.0001	0.354 \pm 0.027 p<0.0001	44.66 \pm 4.16	7.5 \pm 0.519	87.31 \pm 0.823	3.15 \pm 0.036 p<0.0025	3.55 \pm 0.053 p<0.0055
F7	24.78 \pm 2.268 p<0.0001	0.224 \pm 0.016 p<0.0001	95 \pm 3.60	7.066 \pm 0.208	89.01 \pm 2.67	3.40 \pm 0.026 p<0.0013	3.19 \pm 0.07 p<0.0097
F8	33.18 \pm 0.891 p<0.0564	0.308 \pm 0.014 p<0.0001	74 \pm 13.07	7.2 \pm 0.360	87.31 \pm 2.094	3.69 \pm 0.026 p<0.0013	3.08 \pm 0.044 p<0.0037
F9	44.395 \pm 2.096 p<0.0001	0.38 \pm 0.02 p<0.0001	78.66 \pm 3.51	7.166 \pm 0.35	89.01 \pm 1.612	4.60 \pm 0.026 p<0.0013	4.30 \pm 0.046 p<0.0041

Table 4: Physicochemical evaluation of Clotrimazole ocuserts.

*Average of twenty readings, #Average of three readings, * Average of five readings.



Figure 6: Formulated ocuserts F1 to F9.

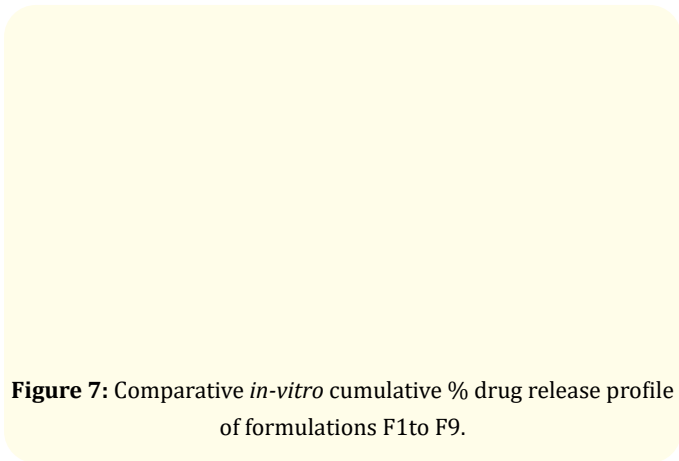


Figure 7: Comparative *in-vitro* cumulative % drug release profile of formulations F1to F9.

Conclusion

From current study we can conclude that by using different polymer in rate controlling membrane of an ocusert release rate of drug from ocusert can be controlled or altered. In present study polymers used in rate controlling membrane of ocusert showed following sequence of release rate (as shown by Figures) HPMC > 1:1 Combination of HPMC and EC > EC.

Furthermore, change in concentration of a polymer in rate controlling membrane alters the release rate of drug from ocusert and it was found that increasing the concentration of a polymer in rate controlling membrane retards the release rate as shown by (Figures) 2% polymer > 3% polymer > 4% polymer.

For achieving an effective required release of drug whether instantaneous or prolonged release, further study on different polymers in different concentration is suggested.

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