## FORMULASI MATRIKS TRANSDERMAL PENTAGAMAVUNON-0 DENGAN KOMBINASI POLIMER PVP K30 DAN HIDROKSIPROPIL METILSELULOSA

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Abstract	Abstract: Transdermal delivery system is one of the delivery system for Pentagamavunon-0 (PGV-0) to $\tilde{A}f\hat{A}, \tilde{A}, \hat{A}$ avoid the high intensity of first pass metabolism of PGV-0 in peroral route. The purpose of this research $\tilde{A}f\hat{A}, \tilde{A}, \hat{A}$ was to optimize the formula of PGV-0 transdermal matrix with a combination of PVP K30 and HPMC $\tilde{A}f\hat{A}, \tilde{A}, \hat{A}$ polymers. The simplex lattice optimization approach of the transdermal matrix formulas was performed by $\tilde{A}f\hat{A}, \tilde{A}, \hat{A}$ using Design Expert 7.1.5 software. The visual appearance, weight, thickness, moisture content, moisture $\tilde{A}f\hat{A}, \tilde{A}, \hat{A}$ uptake, folding endurance, drug content, and dissolution efficiency of the release profil of PGV-0 from the $\tilde{A}f\hat{A}, \tilde{A}, \hat{A}$ matrix for 6 hours were evaluated as responses to determine optimum formula of matrix. The result $\tilde{A}f\hat{A}, \tilde{A}, \hat{A}$ showed that a combination of PVP K30 and HPMC polymers had a significant influence on the visual $\tilde{A}f\hat{A}, \tilde{A}, \hat{A}$ appearance, moisture content, and dissolution efficiency of PGV-0. Combination of 1.98% of PVP K30 $\tilde{A}f\hat{A}, \tilde{A}, \hat{A}$ and 4.52% of HPMC as the optimum formula could produce homogeneous and flexible matrix with $\tilde{A}f\hat{A}, \tilde{A}, \hat{A}$ moisture content of 3.21%. The dissolution efficiency was 9.11%, indicating that 101.93 $\tilde{A}f\hat{A}, \tilde{A}, \hat{A}$ of PGV-0 was $\tilde{A}f\hat{A}, \tilde{A}, \hat{A}$ released from the optimum formula during 6 hours. Keywords: Pentagamavunon-0, Transdermal matrix, PVP K30, HPMC
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