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THE DEVELOPMENT OF PROPRANOLOL HYDROCHLORIDE MICROPOROUS OSMOTIC PUMP

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พิมพ์ตับฉบับบทคัดย่อวิทยานิพนธ์ภายในกรอบสีเขียวนี้เพียงแผ่นเดียว

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บัจจัยต่างๆที่มีผลต่อการปลดปล่อยตัวยาออกจากระบบนำส่งยาโปรปราโนลอล ไฮโดรคลอไรด์ออสโม-ติกปั๊มเช่น ชนิดของสารพลาสติกไซเซอร์, บริมาณของสารพลาสติกไซเซอร์, ชนิดของสารก่อแรงดันออสโมติก, ขนาด ของช่องนำส่งยา เป็นตัน จะถูกประเมินผลโดยการศึกษาลักษณะทางกายภาพ และด้วยการศึกษา ลักษณะในการ ปลดปล่อยตัวยาออกจากระบบนำส่งยา ซึ่งพบว่าการใช้สารโพลีเอทที่ลีน ไกลคอล 400 เป็นสาร พลาสติกไซเซอร์จะ ส่งผลให้ตัวยาถูกปลดปล่อยออกมามากกว่าการใช้สารไดบิวทิว พาทาเลท ภาพถ่ายจากกล้อง จุลทรรศน์อิเลคตรอน แสดงให้เห็นว่าผนังของพิล์มที่มีสารโพลีเอทที่ลีน ไกลคอล 400 เป็นองค์ประกอบจะมี ลักษณะที่เป็นรูพรุนภายหลังจาก การศึกษาลักษณะการปลดปล่อยตัวยา แต่ผนังของพิล์มที่มีสารไดบิวทิว พาทาเลทเบ็นองค์ประกอบจะมีลักษณะที่เรียบ เมื่อเพิ่มบริมาณของสารโพลีเอทที่ลีน ไกลคอล 400 พบว่าตัวยาจะถูกปลดปล่อยออกมามากขึ้น ผลของภาพถ่ายจาก กล้องจุลทรรศน์อิเลคตรอนแสดงให้เห็นว่า รูที่ผนังของพิล์มจะมีลักษณะที่ไหญ่มากขึ้น ผลเช่นนี้ยังคงเกิดขึ้นเมื่อมี การใช้สารโพลีเอทที่ลีน ไกลคอล 4,000 แต่เมื่อทำการเพิ่มบริมาณของสารไดบิวทิว พาทาเลทกลับพบว่าตัวยาที่ถูก ปลดปล่อยออกมากลับไม่แตกต่างกัน โดยผลจากภาพถ่ายซี้ว่า ผนังของพิล์มยังคงมีความเรียบไม่แตกต่างกันเมื่อ ทำการเติมสารก่อแรงดันออสโมติก ชนิดต่างๆในยาเม็ดผลที่ได้พบว่า หากสารที่เติมสามารถให้แรงดันออสโมติกที่สูง แล้วตัวยาที่ถูกปลดปล่อย ออกมาก็จะมีปริมาณมากตามไปด้วย ในแง่ของชนาดช่องนำส่งยานั้นพบว่า ขนาดของ ช่องนำส่งยา จะมีผลต่อการปลดปล่อยยาออกจากระบบนำส่งยาเมื่อปริมาณของสารโพลีเอทที่ลีน ไกลคอล 400 ใน ชั้นพิล์มมีมากขึ้น

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พิมพ์ตับฉบับบทกัดย่อวิทยานิพบธ์ภายในกรอบสีเขียวนี้เพียวแผ่นเดียว

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HYDROCHLORIDE MICROPOROUS OSMOTIC PUMP. THESIS ADVISOR:

ASSOC. PROF. GARNPIMOL C. RITHIDEJ, Ph.D., 160 pp. ISBN 974-638-299-3

There are several factors affecting the release of drug from the propranolol hydrochloride osmotic pump devices; such as the plasticizer type, the level of plasticizer in coated film, the type of osmotic agents within osmotic devices, the passageway size, etc. The effect of many factors was evaluated by using the scanning electron microscope and the release characteristic. The results displayed that Polyethylene glycol 400 supported the release of drug from osmotic devices whereas Dibutyl phthalate suppressed the release of drug. microporous membrane and sponge-like structure were created after the osmotic devices were coated with cellulose acetate plasticized with PEG 400 and were exposed in water. For the osmotic devices coated with cellulose acetate plasticized with DBP, the non-porous membrane was observed even when the osmotic devices was in contact attracted with water. The release of the drug from osmotic devices increased as the level of PEG 400 in coated film increased. The photomicrographs demonstrated that the size of the porosity of the film was increased with increasing level of PEG 400. In the case of PEG 4000, the results were also similar to PEG 400. The release of drug from osmotic devices was not altered when the levels of DBP in coated film increased. The photomicrographs of DBP plasticized film coated osmotic devices was the same although the level of DBP in coated film increased. However, the amount of drug release from osmotic devices increased with increasing osmotic pressure within the devices and decreased with increasing osmotic pressure of the dissolution medium. In consideration of the passageway size, the drug release rate from osmotic devices was indifferent although the size was increased from 400 to 1500 μm .

ภาควิชา	เภสัชอุตสาหกรรม	ลายมือชื่อนิสิต framchai Eamsiningpakhun
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LIST OF ABBREVIATIONS

°C degree celsius (centigrade)

cm centimeter (s)

DBP dibutyl phthalate

e.g. exampli gratia, for example

et al. Et alii, and others

hr hour (s)

i.e. id est, that is

TI thrrapeutic index

mcg microgram (s)

min. minute (s)

mg milligram (s)

ml milliliter (s)

mm millimeter (s)

N normality

nm nanometre (s)

pH the negative logarithm of the hydrogen ion concentration

qs. make to volume

rpm revolution per minute

SEM scanning electron microscope

USP The United States Pharmacopoeia

UV ultraviolet