

# Fabrication of Poly (Sebacic acid-co-ricinoleic-ester anhydride) with $\beta$ -cyclodextrin-loaded Doxorubicin Implants and *in vitro* Characterization

Nagaraja Sreeharsha<sup>1,9,\*</sup>, Jagadeesh G Hiremath<sup>2</sup>, Bandar E Al-Dhubiab<sup>1</sup>, Girish Meravanige<sup>3</sup>, Ranjith Kumar Karnati<sup>4</sup>, Mahesh Attimarad<sup>1</sup>, Anroop B Nair<sup>1</sup>, Katharigatta N Venugopal<sup>1,10</sup>, Mohamed A Morsy<sup>1,8</sup>, Shinu Pottathil<sup>5</sup>, Saleemmulla Khan<sup>6</sup>, Nagashekhara Molugulu<sup>7</sup>

<sup>1</sup>Department of Pharmaceutical Sciences, College of Clinical Pharmacy, King Faisal University, Al-Ahsa, SAUDI ARABIA.

<sup>2</sup>Department of Pharmaceutics, PA College of Pharmacy, Mangalore, Karnataka, INDIA.

<sup>3</sup>Department of Biomedical Sciences, College of Medicine, King Faisal University, Al-Ahsa, SAUDI ARABIA.

<sup>4</sup>Department of Chemistry, College of Science, King Faisal University, Al-Ahsa, SAUDI ARABIA.

<sup>5</sup>Department of Biomedical Sciences, College of Clinical Pharmacy, King Faisal University, Al-Ahsa, SAUDI ARABIA.

<sup>6</sup>Department of Pharmacognosy, PA College of Pharmacy, Mangalore, Karnataka, INDIA.

<sup>7</sup>School of Pharmacy, Monash University, Jalan Lagoon Selatan, Bandar Sunway, Selangor, MALAYSIA.

<sup>8</sup>Department of Pharmacology, Faculty of Medicine, Minia University, El-Minia 61511, EGYPT.

<sup>9</sup>Department of Pharmaceutics, Vidya Siri College of Pharmacy, Off Sarjapura Road, Bangalore, INDIA.

<sup>10</sup>Department of Biotechnology and Food Technology, Durban University of Technology, Durban, SOUTH AFRICA.

## ABSTRACT

**Background:** Doxorubicin is an excellent molecule for the formation of biodegradable implants using the poly (sebacic acid-co-ricinoleic-ester anhydride) 70:30 w/w (poly[SA-RA] 70:30 w/w PSRA 7/3) polymer. **Methods:** The cylindrical implants were successfully produced by means of hot melt extrusion. We used differential scanning calorimetric (DSC) and X-ray diffraction (XRD) methods to identify the melting state and crystal type of blank and drug-loaded implants. A study was conducted on PSRA 7/3 w/w blank and drug-loaded implants for *in vitro* hydrolytic degradation. The drug present in the remaining sample was estimated. **Results:** The hydrolytic degradation rate of the 10% w/w (F2)-loaded implant was relatively low when compared with the 20% w/w (F3) implant. **Discussion:** *In vitro* drug release studies illustrated that the drug-release rate was faster in association with increasing amounts of doxorubicin in the implant.

**Key words:** Poly(sebacic acid-co-ricinoleic-ester anhydride) 70:30 w/w, Doxorubicin,  $\beta$ -cyclodextrin, Cylinder, Biodegradable polymer, *in vitro*.

Submission Date: 29-02-2020;

Revision Date: 30-03-2020;

Accepted Date: 21-05-2020

DOI: 10.5530/ijper.54.3.135

**Correspondence:**

**Dr. Nagaraja SreeHarsha**

Department of  
Pharmaceutical Sciences,  
College of Clinical Pharmacy,  
King Faisal University,  
Al-Ahsa-31982, SAUDI  
ARABIA.

Phone: +96 653 548 5322

E-mail: sharsha@kfu.edu.sa

## INTRODUCTION

Biodegradable polymers are used in the production of implantable devices and drug carriers and as scaffolds in regenerative medicine. They are different from biodegradable polymer biomaterials since they degrade via hydrolysis or enzymatic processes and are eliminated or absorbed. They do not need to be replaced, nor do they require surgical intervention. Polyanhydrides have been utilized as carriers in the controlled delivery of drugs. Today, polyanhydride polymers are the “materials of advanced drug delivery systems.”<sup>1-3</sup>

In 1983, Langer recognized a polyanhydride class of hydrolytically unstable polymers that was suitable for controlled drug release.<sup>4,5</sup> Later, Domb *et al.* synthesized sebacic acid (SA), ricinoleic acid (RA) and poly ([carboxy phenoxy] propane-sebacic acid)(PCPP-SA).<sup>6</sup> The established pharmacological effects of polyanhydride polymer-formed drug carriers include their biocompatibility and safety, as well as their non-toxic, non-mutagenic and low-teratogenic properties.<sup>7</sup> Marketed products composed of hydroxy stearic acid (HPA)-based polyanhydride



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drug carrier systems are available for the treatment of brain cancer and osteomyelitis and include agents such as Gliadel® and Septacin®.<sup>8-10</sup> Recently, polyanhydride polymers gained much interest in the preparation and development of controlled drug-delivery systems. These drug-delivery systems include implants<sup>11,12</sup> pastes (injectables),<sup>13</sup> microspheres<sup>14</sup> and nanomedicine<sup>15,5</sup> and target-specific drug-delivery systems.<sup>16</sup> It has been well described that without the need for any pharmaceutical excipients, fatty acid polymers of polyanhydride polymers undergo surface erosion due to the effects associated with unstable anhydride bonds and the presence of an unstable backbone, such as SA and RA. However, erosion is a rate-limiting step to ensuring drug release from the polymeric matrix.<sup>17,13</sup> The drug released from any carrier system is not dependent on its physicochemical properties and drug release can be controlled by alterations in SA or RA in different proportions, or it can be affected by modifications to the microstructure. The polyanhydrides are so important in developing drug-delivery systems. Specifically, it is predicted that polyanhydride-based drug carriers undergo surface erosion given the significant water lability of anhydride bonds inhibited water penetration thus results in bulking and hydrophobicity.<sup>18</sup> A reduction in the material thickness throughout the surface disintegration process of cylindrical shape approximately zero-order hydrolytic degradation kinetics illustrate the heterogeneous surface erosion.<sup>18</sup> One characteristic property of the high hydrolytic reactivity of anhydride linkages is its adaptability to manage degradation rates. This type of control can be manipulated by varying the type and proportions of the monomer used. The ability to synthesize particular kinds of polymers can result in the predictable release of a drug from a formulation in over the course of 1 week to several weeks. By changing the hydrophobic-hydrophilic balance of the polymer, basic changes in the polymer backbone occur and hydrolytic degradation can be controlled. A delay in degradation amounts of aliphatic and aromatic polyanhydride polymers were few days to months. Thus, polyanhydride provides a convenient method through which to produce a drug discharge occur by a sequence of the erosion of the matrix, diffusion through the polymer matrix and/or diffusion through pores formed in the matrix by the dissolution of the water-soluble additives.<sup>2,19,20</sup> PSRA7/3 w/w is a polyanhydride class of polymers used in controlled drug-carrier systems. Doxorubicin is an ideal molecule for the preparation of biodegradable implants using the PSRA7/3 w/w polymer. Doxorubicin is one of the most potent, broad-spectrum anticancer

agents to be used as a therapeutic option for solid tumors. Drugs have a short half-life due to this narrow therapeutic index and they also demonstrate significant increases in high-dose distributions to healthy normal tissues. Drug distribution leads to the onset of drug-related side effects, such as cardiotoxicity, myelosuppression and mucositis.<sup>21,22</sup> Hence, localized drug delivery systems are needed for doxorubicin, which has many applications and offers promising results as a potential solution to the significant drawbacks associated with some other currently available drug-delivery systems. Polyanhydride-based, localized drug-release systems (such as insoluble drugs in aqueous media) have low bioavailability and a short half-life, meaning that the drug can be delivered directly to the desired site. These delivery techniques are useful for drugs with a limited therapeutic index. Local drug delivery is suitable for medications that exert dosage-reliant action; this prevents the need to use other drugs that have adverse effects when delivered systemically. Local delivery further reduces dose-related problems that are typically encountered during animal studies and clinical studies with humans. With respect to anticancer drugs, intratumoral delivery is unrestricted by insufficient blood supply, which is induced by radiation therapy or surgery. Polyanhydrides also demonstrate reduced liability before delivering the active agent due to its surface or heterogeneous hydrolytic degradation.<sup>23</sup> Hence, the present study explores the preparation of DOXHCl-loaded biodegradable implants using a PSRA7/3 w/w polymer and also features *in vitro* characterization studies.

## MATERIALS AND METHODS

Doxorubicin (DOXHCl) was obtained as a free product from RPG Life Sciences Ltd. (Mumbai, India). Poly (sebacic acid-co-ricinoleic-ester anhydride) 70:30 w/w (poly[SA-RA]70:30 w/w, designated as PSAR 7/3 w/w) (monomer number [Mn]=21,000, molecular weight [Mw] of 60,000 and a melting temperature [mT°C]=68.8, data H[J]/gr°C), was procured from the School of Pharmacy, Faculty of Medicine, the Hebrew University of Jerusalem, Hadassah Ein Kerem Medical Center Campus, Jerusalem 91120, Israel. It was used without further purification. Complete information on the polymers used is available.<sup>13</sup> Beta  $\beta$ -cyclodextrin (mW 1,134.98) and all analytical grade solvents were procured from Sigma Aldrich Co. (St. Louis, MO, USA).

### Preparation of DOXHCl Implants

DOX·HCl-loaded cylindrical polyanhydride-based implants were prepared using a method published previously.<sup>24</sup> In short, we employed a laboratory-based

method to produce the implants. For the first step, the drug and  $\beta$ -cyclodextrin were mixed; a pasty polymer was uniformly blended and brought into a cylindrical mold (7.0 mm in diameter and 8.0 mm in length). It was set on and attached to a cylinder (rod) of 6.9 mm in diameter (portable). The full component was heated to 50°C using a heating mantle. The polymer was plasticized and homogeneously mixed using a steel rod (length 10 mm  $\times$  breath 2 mm). After homogenous mixing, the full unit was placed in a freezer (-4°C). After 5 min, the unit was removed and a plunger was introduced into the mold; a cylindrical shape was created and manually compressed using a flexible plunger, which was pushed 1.0–7.0 mm into the mold. The cylinder was returned to the freezer for 20 min. After 20 min, the whole unit was left to return to room temperature. At this point, the plunger and rod were drawn out, which extruded the solidified implant from the mold.

### Physicochemical analysis of the implants

The area (cm<sup>2</sup>), thickness (h,cm), volume (cm<sup>3</sup>) and surface area (cm<sup>2</sup>) of the implants were measured using a digital vernier caliper. From the obtained values, the above dimensions were measured using the mathematical equations<sup>17</sup> shown in Table 1. We used differential scanning calorimetry (DSC) and X-ray diffraction (XRD) methods to detect the melting state and crystal structure of the blank and drug-loaded implants. DSC employing the Mettler Toledo DSC822e (Columbus, OH, USA) was used to investigate the thermal properties of the samples at a rate of 10°C per min, with temperatures ranging from 10°C to 230°C. It used a 4 in empty aluminum skillet was used as a reference. XRD examinations of the implants were carried out using a Philips PW1820 (Amsterdam, the Netherlands) with Cu K $\alpha$  and K $\alpha$  radiation to identify the crystal patterns. The lyophilized samples were sputtered coated with gold and the morphology of the implants was visualized by scanning electron microscope (SEM; JSM-848, JEOL, Tokyo, Japan). The generated surface morphology of the coated cross-section samples was examined under magnification.

### DOX·HCl content analysis in the implants

Implants loaded with 10 mg of DOX·HCl were dissolved in 5 mL of chloroform (CHL) combined with 10 mL of MilliQ water until it turned an opaque white color to ensure precipitation of the polymer. The samples were centrifuged at 4°C (Remi, Bengaluru, India) at 12,000 rpm for 12 min. The collected supernatant was clear and subsequently evaporated. After evaporating the CHL, 5 mL of phosphate buffer saline (PBS; pH 7.4) was added

and vortex stirring was performed for 5 min. A 0.22  $\mu$ m filter was used (Millipore India). The DOX·HCl content was analyzed with an ultraviolet visible (UV-VIS)-spectrophotometer (UV-1800; Shimadzu, Kyoto, Japan) at 250 nm. This analysis procedure was standardized and validated. The correlation coefficient ( $r^2$ ) is 0.9998 and the equation was  $y = 0.012x + 0.004$  with a slope of 0.012 and an intercept of 0.004. This method to assess DOX·HCl has been developed and was found to be highly precise and valid. This method was successfully adopted in alternative evaluations.

### Hydrolytic degradation of implants

An investigation was carried out on PSARA 7/3 w/w blank and drug-loaded implants to examine *in vitro* hydrolytic degradation and to investigate the amount of drug present in the remaining sample. The implants were placed in 20 mL of PBS (pH 7.4) in 50 mL screw-capped vials. These vials were maintained in a horizontal shaker and vibrated at a velocity of 50 rpm to activate the hydrolytic degradation for 20 days. At precise intervals, the blank and drug-loaded implants were collected from the media. These samples were lyophilized the implant weight was subsequently determined. For each sample, the vehicle in the flask was replaced with new buffer to promote the sink condition. The hydrolysis of the implant was measured by means of weight reduction and at each stage, the DOX·HCl content assembled in the implant (residue) was analyzed using the UV-VIS spectrophotometric method described above.

### *In vitro* PSRA7/3 w/w release studies of DOX·HCl-loaded implants

*In vitro* PSRA7/3 w/w release studies of the hydrolytic, degradable drug carrying the implants were carried out using PBS (pH 7.4) as a dissolution medium. The implants were placed in 20 mL of PBS (pH 7.4) in 50 mL screw-capped vials. These vials were kept in a horizontal shaker and the samples were vibrated at a velocity of 50 rpm to promote drug release for 40 days. At each sampling time, the dissolution media were changed in the flask with fresh buffer to continue the sink condition. The DOX·HCl content was determined using the UV-VIS spectrophotometric method described above. The cumulative amount released versus time was plotted. Finally, we determined the amount of drug present based on kinetic models.

## RESULTS AND DISCUSSION

### Implant Characterization

Figure 1 provides photographs of the drug-loaded implants prepared using the melting method. The

implants that were produced were of cylindrical shape with a smooth surface. Table 1 that the weight of the implants was as follows:  $120.46 \pm 1.98$  mg (F1),  $142.86 \pm 1.24$  mg (F2) and  $160.84 \pm 2.46$  mg (F3). The implant thickness was as follows:  $2.06 \pm 0.01$  mm,  $2.24 \pm 0.02$  mm and  $3.06 \pm 0.01$  mm. The drug content uniformity was found to be 98%–99%. Figure 2 depicts the DSC studies that indicated a sharp endothermic peak of the drug measured at  $209.83^\circ\text{C}$ ; the PSRA7/3 w/w endothermic peak was generated at  $62.83^\circ\text{C}$  and the  $\beta$ -cyclodextrin short, wide melting peak occurred at  $58.86^\circ\text{C}$ . Figure 3 shows that based on the XRD analysis, the  $2\theta$  scattered angle peaks of the crystalline drug occurred at  $17.27$  and  $18.46$  and that their conforming peak intensities were 1,346 and 1,408, respectively. XRD analysis of the blank implant showed that the  $2\theta$  scattered angle peaks occurred at  $21.73$  and  $21.91$ , with peak intensities at 248 and 196, respectively, while the  $\beta$ -cyclodextrin  $2\theta$  scattered angle peak appeared at  $5.38$  with an intensity peak at 4,568.

### Hydrolytic polymer weight loss and drug accumulation

Figure 4 shows the investigated hydrolytic degradation and drug accumulation in the remaining drug sample carried by the PSAR 7/3 w/w implants. The blank implants exhibited a faster hydrolytic degradation rate of basic mass loss (4.78%) and at the end of 20 days,  $96.63 \pm 1.9\%$  degraded. The hydrolytic degradation rate of the 10% w/w (F2)-loaded implant was relatively lower when linked to the 20% w/w (F3) implant. After 2 days, degradation mass loss took place ( $2.64 \pm 0.9\%$  and  $6.64 \pm 1.2\%$ ). During the course of this degradation, we found that the quantity of the drug was  $99.45 \pm 0.8$  and  $98.46 \pm 0.4$ . At the end of a 20-day study, the resulting mass loss of the drug implant (F2) produced a quantity of  $76.96 \pm 1.2$ ; however, at this stage,

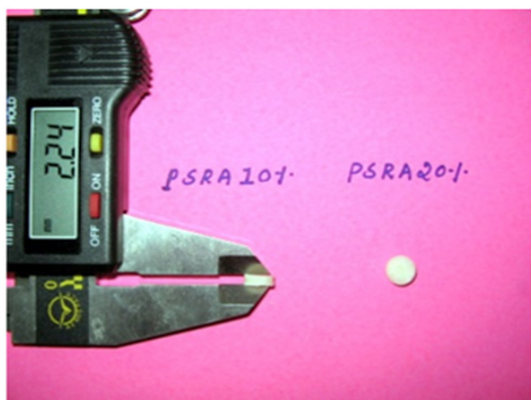


Figure 1: Illustration of the DOX HCL HCL-loaded 10% w/w and 20%w/w PSRA7/3 implants.

the acquired drug was analyzed to be  $36.94 \pm 0.6\%$ . It was determined that the F3 implant degradation mass loss after 20 days was  $90.59 \pm 1.6\%$ . A similar result was presented in separate studies that illustrated how the erosion grades of the P (FAD-SA) formulations were measured as weight loss that was prompted by the physicochemical equities of the blended compound. The hydrophilic compound accelerated erosion, while the hydrophobic compound minimized the destruction of P (FAD-SA) materials.<sup>17</sup>

### Dissolution studies

Figure 4 presents the results of the dissolution studies of the DOX·HCl-loaded PSRA7/3 w/w implants. Drug release was carried out in PBS (pH 7.4) as a dissolution medium at  $37^\circ\text{C}$ . The 10% w/w loaded implant's mean cumulative drug-release rate in the first 2 days was  $0.84 \pm 0.5\%$ . Similarly, the 20% w/w drug-loaded implant showed a drug-release rate of  $4.04 \pm 2.4\%$ . As the amount of DOX·HCl increased in the implant, the drug-release rate increased correspondingly. From day 4, the release curve appeared to show an upturn at a comparatively faster pace until day 20, with an average,

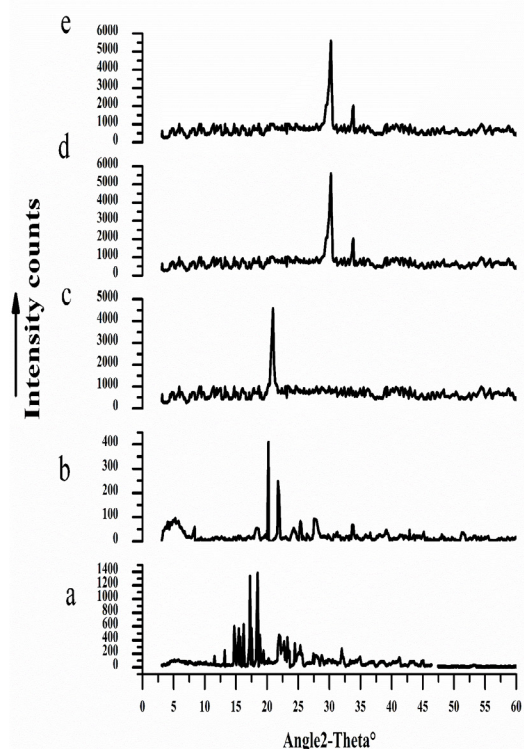
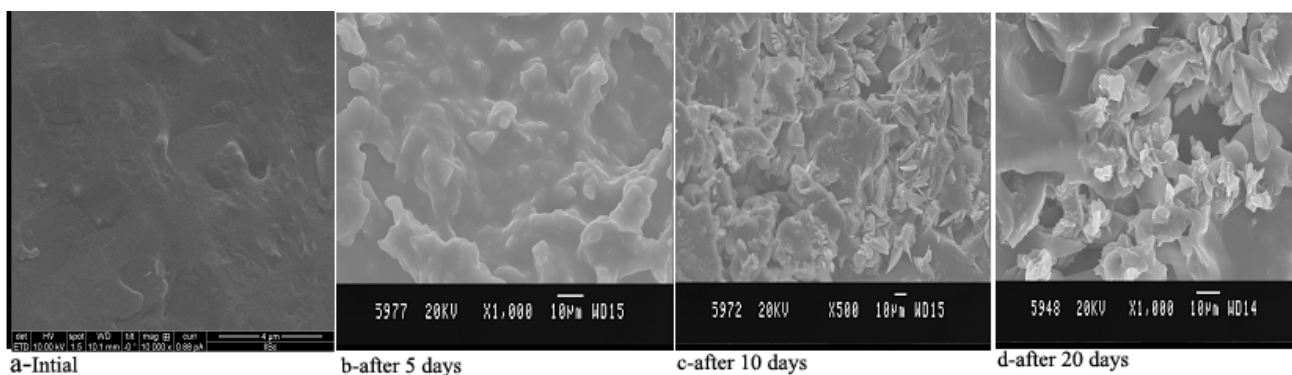
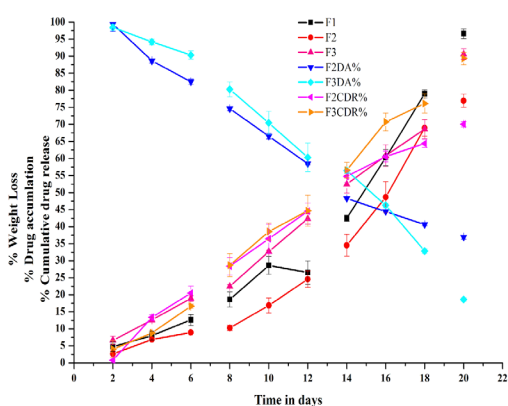


Figure 2: The X-ray diffraction patterns of DOX HCL (a), PSRA7/3 (b),  $\beta$ -cyclodextrin (c) and drug-loaded implants F2 (d) and F (e). The equipment was operated using a step-scan technique of  $0.02$  ( $2\theta$ ) increments to measure the crystalline character of the patterns. With respect to the formulations, the implants were powdered well and the crystalline quality of the samples was tested.



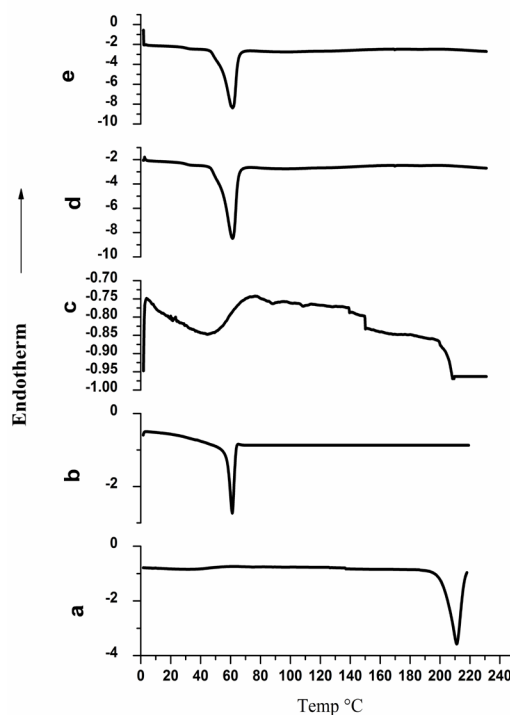
**Figure 3:** SEM photographs of the *in vitro* hydrolytic cleavage samples of the 20% w/w drug-loaded sample at the initial time point (a) before being exposed to the degradation subject, after day 5 (b), after 10 day (c) and at day 20 (d). The patterns showed the progression of hollow water cavities or large, uneven thicknesses of pores, which might be influenced by the presence of  $\beta$ -cyclodextrin. This might solubilize the drug during the cleavage of PSRA 7/3 w/w surface erosion; the hydrolytic degradation was investigated. The SEM images were obtained at suitable magnification.



**Figure 4:** Hydrolytic degradation of the polymer; percent weight loss for F1 (■), F2 (●) and F3 (▲), as well as the percent of drug accumulated during weight loss for F2 (▼) and F3 (◆). Also presented is the cumulative percent of drug release versus time for F2 (◀) and F3 (▶). The implants were placed in 20 mL of phosphate buffered saline (PBS; pH 7.4) in 50 mL screw-capped bottles. These vials were kept in a horizontal shaker and continuously vibrated at an acceleration of 50 rpm to activate the hydrolytic degradation, surface erosion, or bulk erosion over 20 days.

sustained drug release of  $70.06\% \pm 1.9\%$  (F2 10% w/w) and  $89.36\% \pm 1.8\%$ .

Table 1 shows the macroscopic properties of the blank and drug-loaded implants. Drug-loaded PSRA7/3 with yellow  $\beta$ -cyclodextrin cylinders, created using a melting procedure, featured a set amount of polymers that varied in the amount of drug that was loaded; they produced 10% w/w and 20% w/w. The change in the weight and thickness of the implants was recognized when correlated to the initial amount of drug present; polymers were taken to prepare the formulations. This attribute because of the chain reorientation of the polymeric molecules during heating/melt process, direct freezing and insertion of a plunger to make a



**Figure 5:** The X-ray diffraction patterns of DOX HCL (a), PSRA7/3 (b),  $\beta$ -cyclodextrin (c) and drug-loaded implants F2 (d) and F (e). The equipment was operated using a step-scan technique of 0.02 ( $2\theta$ ) increments to measure the crystalline character of the patterns. With respect to the formulations, the implants were powdered well and the crystalline quality of the samples was tested.

compact uniform shape accompanied by solidification of implants. Rapid cooling could influence the implants' solid or rigid structure. Various dimensions of the implant, such as its area, volume and surface area played a significant role in the hydrolytic degradation of the polymer, ultimately influencing the drug's ability to discharge from the cylinder. The differences in the thickness and weight values of the implants are

displayed in Table 1.  $\beta$ -cyclodextrin was incorporated in the manufacturing process to enhance the drug's solubility within a polymeric surface and to generate pores that could easily influence water penetration into the implants, while activating the polymer to induce the surface or bulk hydrolytic erosion of the polymer. The DSC analysis (Figure 5) shows that the drug peak disappeared in the process of implants F2 and F3 exhibiting the drug existence in the presence of dissolution, semi crystalline, or drug adsorbed at the amorphous polymer surface. These results were again confirmed with XRD studies (Figure 3), which indicated that the crystalline scattered angle peaks disappeared in the drug-loaded implants. Hydrolytic degradation is the transformation of polymer chain cleavage. They hydrolyze into dicarboxylic acids and hydroxy acid monomers when placed in an aqueous vehicle. Polyanhydride belongs to a class of water-insoluble hydrophobic polymers. It has a hydrophobic backbone with a hydrolytically labile anhydride and/or ester links. Erosion is the method of transfer that can lead to the loss of polymer mass or monomers.<sup>25,26</sup> The blank implants showed a faster hydrolytic degradation rate when compared with the drug-loaded implants. The erosion rates of the cylinders were determined to be physicochemical properties of the blended compound, which prompted the observed weight loss. Figure 4 presents the results of the dissolution studies of the DOX·HCl-loaded PSRA7/3 w/w implants. Drug release was carried out in PBS (pH 7.4) as the dissolution

medium at 37°C. As Since the percentage of the drug is higher in the polymer, the amount of drug released is faster due to the influence of  $\beta$ -cyclodextrin polymer-induced drug solubility<sup>27</sup> and the creation of more water channels. Further, high water uptake produced faster hydrolytic polymer degradation which, in turn, controlled the drug released from cylinder.<sup>28</sup> This means that DOX·HCl is easily soluble in water and normal saline. Degradation transforms the surface (homogenous) and bulk heterogeneous eroding polymers. The time at which the degradation of the full polymer quantity is for high water penetration than the amount of polymer cleavage. It is a continual disposing control cleavage of oligomers into monomers throughout the polymer matrix. Surface cleavage (breakdown) is the hydrolysis of the polymer to the outer surface and the interior matrix remains an under process of cleavage of oligomers followed by monomers. It selects the particular surface erosion process the polymer cleavage should be quicker than water uptake or diffusibility or dissolution. The hydrophobic polyanhydride polymers show how surface disintegration occurs only when the highest water penetration should be at the appearing surface than at the interior of the bulk.<sup>29</sup> We observed the SEM results of the *in vitro* release types of the drug-loaded 20% w/w typical F3 formulation at periodic time intervals (a). Figure 4 presents the surface morphology of the intact implant before the *in vitro* hydrolytic degradation occurred and was characterized as a stable, continuous, smooth and uniform surface (a). The surface

**Table 1: Formulations and evaluation of poly (SA-RA) 70:30 w/w DOX·HCL implants.**

Batch	Polymer (mg)	DOX·HCL (mg)	$\beta$ -cyclodextrin (mg)	Drug loading %	Implant weight (mg)	Drug content %
F1	180	-	20	-	120.46±1.98	-
F3	180	20	20	10	142.86±1.24	98.96±0.4
F4	180	40	20	20	160.84±2.46	99.84±0.8
Batches	Appearance	Color	Area (cm <sup>2</sup> )	Thickness (h) (cm)	Volume (cm <sup>3</sup> )	Surface area cm <sup>2</sup>
F1	Cylinder	Yellow	0.346±0.4	2.06±0.01	6.27±0.02	7.01±0.05
F2	Cylinder	Yellow	0.468±0.2	2.24±0.02	6.48±0.04	7.06±0.02
F3	Cylinder	Yellow	0.586±0.1	3.06±0.01	9.51±0.8	7.22±0.03
Batches	Zero order $Q_t = Q_0 + K_0 t$	First order $\ln Q_t = \ln Q_0 + K_1 t$	Higuchi $Q_t = K_H \text{sq.}tR^2$	Korsmeyer-Peppas $Q_t/Q_{\infty} = K_k t^n R^2$	Korsmeyer-Peppas $Q_t/Q_{\infty} = K_k t^n R^2$ n=values	P (P-value)
F2	0.934	0.845	0.863	0.9444	(n=0.69)	<0.001
F3	0.912	0.824	0.856	0.9064	(n=0.64)	<0.003

The following mathematical equations were used to determine the physical parameters of the implants: (cm<sup>2</sup>)A=2 $\pi$ rh+2 $\pi$ r<sup>2</sup>. Volume (cm<sup>3</sup>)= $\pi$  X r<sup>2</sup> X h, Surface area (cm<sup>2</sup>) = [( $\pi$  X r<sup>2</sup> X 2) + (2 X  $\pi$  X r X h)].

morphology of the sample after day 5 slightly influenced the bulk erosion (b). After day 10 and at the conclusion of day 20, the surface morphology showed greater bulk erosion and surface erosion. Highly porous, extend in some hollow carries with an effect of  $\beta$ -cyclodextrin with a drug in aqueous solution, the compounds readily dissociate and free drug molecules persist in equilibrium with the molecules confined within the  $\beta$ -cyclodextrin hollow carries may start to PSRA7/3 polymer surface eroding, with further loss of material from the exterior surface and samples diminished further in their size with controlled drug deliver for a prolonged stage of polymer degradation was explained. The same approach used in separate studies indicated that drug delivery is independent of its physicochemical properties and is effectively commanded by erosion. The breakdown of the polymer matrices depends on transfers such as the rate of polymer cleavage, the acceleration of water uptake (at which point the polymer swells), the resulting pore structure and the momentum of oligomer and monomer dispersion from the matrices. Such erosion maintains a continuous surface field and leads to zero-order drug release.<sup>26</sup> Thus, the variation in matrix degradation observed between particular combinations of polymer and drug amounts could be in response to the polyanhydride degradation that occurs by surface dissolution and other physicochemical factors that involve transformation and shape degradation. For criterion, the monomers and their structure determine the significant associates.<sup>30,31</sup>

It is actually important to identify the visible and atomic degradation properties of polyanhydrides at a molecular level. The geometry of cylindrical, rod and rectangular dimensions influences the cleavage of polymers (erosion). As the devices increase in thickness, the polymer is slower to cleave to smaller surface areas and accommodates finite water penetration. This is involved in the delayed, controlled drug release that occurs by surface erosion.<sup>31</sup> There is a chance that this process varies based on hydrophilic high drug loading, the drug's solubility, the molecular weight of the drug and the polymer's solubility, which can intensify the co-adjuvant polymer or the excipients incorporated in the matrices or devices. The drug-release kinetics were established based on *in vitro* release data by plotting the percent cumulative drug release in association with time; this indicated the gradual and continuous drug release-patterns that were observed and which were independent of concentration. The Korsmeyer Peppas design showed that drug release occurs through diffusion and erosion process. A value between 0.64–0.69 indicates irregular and relaxation kinetics processes.

## CONCLUSION

Doxorubicin-loaded biodegradable polymeric implants using PSRA 7/3 and other  $\beta$ -cyclodextrin polymers were blended to enhance drug diffusion. The melting method successfully produced implants. Changes in the dimensions of the implants were observed when correlated to the initial amount of drug and polymers used for formulation development. The XRD and DSC studies of cylinders F2 and F3 showed the presence of the drug, the dissolution behavior, the semicrystalline structure and the drug adsorbed at the amorphous polymer surface. The *in vitro* drug-release studies explained that as the amount of DOX-HCl increased in the implant, the drug-release rate increased correspondingly. However,  $\beta$ -cyclodextrin promoted the release of the drug from the cylindrical device since the percentage of the drug was higher in the polymer; as such, the amount of drug released is higher because the influence of  $\beta$ -cyclodextrin-induced drug solubility was revealed. The *in vitro* drug release subjected samples morphology using SEM studies proved that surface morphology of the intact implant before succeeding *in vitro* hydrolytic degradation characterized as a stable continuous smooth uniform surface. The surface morphology of the sample after day 5 influenced bulk erosion. After day 10 and at the end of day 20, the surface morphology showed greater bulk and surface erosion. The structure was porous and presented some water cavities<sup>2</sup>-cyclodextrin was carried out. The drug-release kinetics investigations demonstrated that this occurred independent of concentration and the Korsmeyer Peppas design provided a good prepared for the drug release implants with greater regression coefficients in corresponding to alternative designs. These prepared implants may function as local drug-delivery systems using polyanhydride polymers as a safe substitute to transport anticancer agents to tumors. Investigations with *in vitro* cell lines and animal subjects will be carried out in future studies.

## ACKNOWLEDGEMENT

The authors acknowledge the Deanship of Scientific Research at King Faisal University for providing financial support under the Nasher Track (grant number: 186248).

## CONFLICT OF INTEREST

The authors declare that there are no conflicts of interest.

This article does not contain any studies with human or animal subjects performed by any of the authors.

## ABBREVIATIONS

**(DOX·HCl):** Doxorubicin; **(FTIR) spectroscopy:** Fourier Transform Infrared; **(XRD):** X-ray diffraction; **DSC:** Differential scanning calorimetry; **SEM:** Scanning Electronic Microscopy; **SA:** Synthesized sebacic acid; **RA:** Ricinoleic acid; **PCPP-SA:** poly ([carboxy phenoxy] propane-sebacic acid); **HPA:** Hydroxy stearic acid.

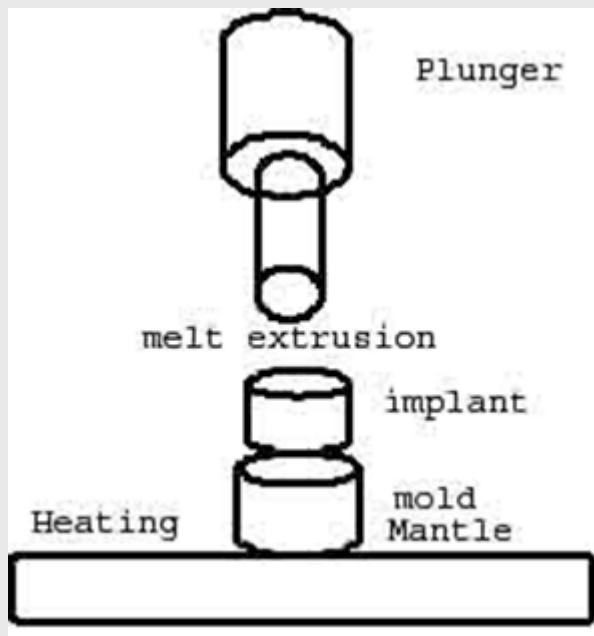
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## SUMMARY

Doxorubicin-loaded biodegradable polymeric implants using PSRA 7/3 and other  $\beta$ -cyclodextrin polymers were blended to enhance drug diffusion. The melting method successfully produced implants. Changes in the dimensions of the implants were observed when correlated to the initial amount of drug and polymers used for formulation development. *In vitro* drug release studies illustrated that the drug-release rate was faster in association with increasing amounts of doxorubicin in the implant.

### PICTORIAL ABSTRACT



### About Authors

**Dr. Sree Harsha** received his (ranked top 5) Master of Pharmacy Degree and subsequently earned a doctorate in Pharmaceutics from Rajiv Gandhi University of Health Sciences, Bangalore, India in 2006. He came to King Faisal University in 2007 as an assistant professor in the Department of Pharmaceutical Sciences, bringing with him several years' worth of teaching experience in fundamentals of pharmaceutics and drug delivery systems. He was actively participated in Accreditation Council of Pharmacy Education (ACPE) and Canadian Council for the Accreditation of Pharmacy Programs (CCAPP). His primary area of focus is pharmaceutical technology and novel/ targeted drug delivery systems. For this research, he received grants (30 number) from Deanship of Scientific research, King Faisal University. The author contributed so far to 80 peer-reviewed full papers on a variety of topics in lung targeting, topical drug delivery and mucoadhesive drug delivery systems, He has contributed in writing a book chapter titled "Targeted Drug Delivery System" and "Microspheres" in Textbook of Industrial Pharmacy, Publisher-Orient Longman Private Ltd. In addition, he is an Ad-hoc reviewer for scientific journals. He has attended many seminars and Workshop both national and international on Pharmaceutical Technology and Public health issues.

**Cite this article:** Sreeharsha N, Hiremath JG, Al-Dhubiab BE, Meravanige G, Karnati RK, Attimarad M, Nair AB, Venugopal KN, Morsy MA, Pottathil S, Khan S, Molugulu N. Fabrication of Poly (sebacic acid-co-ricinoleic-ester anhydride) with  $\beta$ -cyclodextrin-loaded doxorubicin Implants and in vitro characterization. Indian J of Pharmaceutical Education and Research. 2020;54(3):826-34.