



**Review Article** 

# JOURNAL OF APPLIED PHARMACEUTICAL RESEARCH | JOAPR www.japtronline.com ISSN: 2348 - 0335

# A BRIEF REVIEW ON RECENT ADVANCEMENT OF TABLET COATING TECHNOLOGY

Debgopal Ganguly<sup>1</sup>, Soumyadip Ghosh<sup>2</sup>\*, Pubali Chakraborty<sup>1</sup>, Sumit Mitra<sup>1</sup>, Shounak Chatterjee<sup>1</sup>, Sarmistha Panja<sup>1</sup>, Ananta Choudhury<sup>3</sup>

#### Article Information

Received: 23<sup>rd</sup> July 2021 Revised: 4<sup>th</sup> December 2021 Accepted: 10<sup>th</sup> January 2022 Published: 31<sup>st</sup> March 2022

#### Keywords

Conventional tablet coating techniques, advantages & disadvantages, modern techniques

#### ABSTRACT

A tablet can be defined as a solid unit dosage form. There are several reasons for coating of solid dosage form, the most important reason is to control the release profile & also to control the bioavailability parameters of the APIs (Active Pharmaceutical Ingredients). Tablets containing active pharmaceutical ingredients (API) can be coated with thin polymer-based film for various advantages. Generally, horizontal rotating pans are used for coating purposes & coating solution can be spread through the spraying systems over the surface of tablets. Bitter taste masking, odor masking, physical and chemical protection, and also environmental protection are all benefits of the tablet coating. Despite that, Tablet coating also plays an important role in controlling the action site. Sugar-coating, film coating, and enteric coating technologies is to eliminate the numerous disadvantages of solvent-based coating. Coating solution preferentially applied on the surface of solid dosage forms without the need for any solvent in these novel technologies. This review article provides information regarding the techniques of conventional tablet coating, the recent advancement of tablet coating procedures, and tablet coating components.

#### **INTRODUCTION**

Tablet is an example of unit dosage, which is being compressed after mixing of active constituents and another additive so that a proper shape may be given to the tablet. This is the medication in a compressed form. Solid measure formulations unit of measurement necessary measure forms in prescription drugs. Solid dosage form includes tablets, capsules, granules, sachets, powders, dry powder inhalers, and chewable. A unit dose of one or more medications

<sup>1</sup>School of Pharmacy, Seacom Skills University, Bolpur, Birbhum, West Bengal-731236 <sup>2</sup>\*Calcutta Institute of Pharmaceutical Technology and AHS, Uluberia, Howrah, West Bengal-711316 <sup>3</sup>Faculty of Pharmaceutical Science, Assam downtown University, Guwahati, Assam

# \*For Correspondence: ghoshs9764@gmail.com

## ©2022 The authors

This is an Open Access article distributed under the terms of the Creative Commons Attribution (CC BY NC), which permits unrestricted use, distribution, and reproduction in any medium, as long as the original authors and source are cited. No permission is required from the authors or the publishers. (https://creativecommons.org/licenses/by-nc/4.0/)

is contained in the solid measure kind. Binders, glidants, sweeteners, and other excipients are all examples of excipients [1].

## **Objective of tablet coating**

- 1. Increase shelf-life of drugs.
- 2. Loss of volatile ingredients can be reduced.
- 3. Enhancement of mechanical strength of solid dosage form.
- 4. Drug release rate can be modified through this process specifically delay release tablets or enteric-coated tablets and sustained-release medicines.

## **Drawbacks of tablet coating**

- 1. Tablet coating is expensive than normal formulation.
- 2. Sometimes, tablet coating leads to the degradation of active ingredients such as chipping, capping, mottling, and bridging.
- 3. Some drugs are much more sensitive to the coating that leading to serious adverse effects.

## **Coating process**

Coating pans that rotate are widely employed for this purpose. During the process of coating the tablets, liquid coating solutions are spread over the uncoated tablet beds inside the pan and after the transfer of air passes over the uncoated tablets, the liquid part of the coating solution is evaporated and leaving a solid coated layer over tumbling tablets. Various stages are mentioned below [2, 3]

- i. Batch identification and selection of the type of coating. (Film or Sugarcoating)
- ii. Dispensing (accurately dosing of all required raw materials)
- iii. Loading of tablets into the pan.
- iv. Warming of tablets.
- v. Spraying (application of coating materials and rolling of the tablet are carried out simultaneously)
- vi. Drying.
- vii. Cooling and unloading.

# **Types of Coating**

- 1) Sugar Coating.
- 2) Film Coating.
- 3) Enteric Coating.
- 4) Controlled Release Coating.
- 5) Specialized Coating.

- a. Compressed Coating,
- b. Electrostatic Coating.
- c. Dip Coating.
- d. Vacuum Coating.

# Sugar Coating:

The sugar-coating process includes five major steps:

a. Waterproofing/Sealing: creates a wetness barrier while conjointly hardening the pill surface.

b. Sub-coating permits the pill size to quickly increase and also the pill edges to spherical off.

c. Grossing/Smoothing: The surface of the tablets is smoothing out and pills are adjusted to the required sizes.

d. Coloring determines the tablet's final color and scale.

e. Sprucing provides a shiny look [5].

# Film Coating:

Major requirements for film coating ingredients are:

i) Selection of the coating solution depends upon the solubility profile of the active ingredients.

ii) Film coating also depends upon some other solubilities such as free pH-dependent solubility, slow water solubility, and free water solubility.

iii) Able to supply sublime products.

iv) Highly stable at different environmental conditions such as air, heat, moisture, and light.

- v) No such impact on color, style, or odor.
- vi) Highly compatible with alternative coating.

vii) Non-toxic, non-irritant, and no such interaction with active pharmaceutical ingredients.

viii) Cracking can be overcome through this process.

ix) Bridging and filling problems can be overcome through this process.

x) Printing process can be done easily with this coating process [6, 7].

# Enteric Coating:

Ideal properties for enteric coating materials are:

i) Resistance to internal organ fluids.

ii) Susceptible/permeable to enteral fluid.

iii) Compatibility with most coating resolution parts and therefore the medication substrate.

- iv) Formation of continuous film.
- v) Nontoxic, low-cost and easy application.
- vi) Ability to be promptly written.

Different polymers involved in the enteric coating process are as follows:

1) Acrylate polymers.

- 2) Polyvinyl acetate phthalate.
- 3) Cellulose acetate phthalate (CAP).
- 4) Hydroxypropyl methylcellulose phthalate [8], [9].

# TABLE 1: PROPERTIES OF SUGAR COATING

ТҮРЕ	PROPERTIES	SUGARCOATING	
<b>T 11</b> 4	Appearance	With a serious level of cleanliness, it's adjusted.	
Tablet	The weight has expanded Due to the covering material	30-50%	
	Logo or 'break lines'	It is not possible.	
Process	Administrator preparation is required	Considerable	
	Flexibility to Good Manufacturing Practices (GMP)	Difficulties could emerge.	
	Phases of the strategy	A multi-stage system	
	Coatings with utilitarian properties	This is generally unrealistic, except for enteric covering.	

# TABLE 02: MATERIALS USED IN FILM COATING

S. NO	MATERIALS	TYPES	USES	EXAMPLES
1	Film Former	Enteric Non-enteric	To control the release of the drug	Hydroxy Propyl Methylcellulose (HPMC), Methyl Hydroxy Ethyl Cellulose (MHEC).
2	Solvents		To dissolve or disperse the polymers	Iso Propyl alcohol (IPA) and Methyl Chloride.
3.	Plasticizer	Internal plasticizer	It pertains to the chemical modification of the basic polymer that alters the physical properties of the polymer	Glycerol, Propylene glycol, PEG 200- 6000 grades.
		External Plasticizer	It fused with the essential polymeric film previous, changes the adaptability, Rigidity, or bond property of the subsequent film.	Diethyl phthalate (DEP), Dibutyl phthalate (DBP), and Tributyl citrate (TBC)
4	Colorant	Inorganic Materials	For Light Shade: Concentration of less than 0.01% may be used	Iron oxides
		Natural coloring Materials	For Dark Shade: A concentration of more than 2.0% may be required.	Anthocyanins, Caramel.
5	Opaquant – extenders		For more pastel tones and expanding film inclusion detailing must be given	Titanium dioxide, Silicate (Talc & aluminum silicates), Carbonates (Magnesium Carbonates)

ITEMS	CHARACTERISTICS	IMPORTANCE
	Appearance	Retain contour of original core. Not as shiny as the sugar coat type.
Tablet	Weight increase because of coating materials	2-3%
	Logo of 'Break lines'	Possible
Process	Operator training required	The process tends to automation and operator training.
	Adaptability to GMP	High
	Process Stages	Usually, Single-stage.
	Functional Coating	Easy adaptable for controlled release.

### TABLE 03: CHARACTERISTIC OF FILM COATING

## MODERN TABLET COATING TECHNIQUES

### 1) Aqueous Film coating technology:

The glossing over the measure is exceptionally tedious and it is relying upon the abilities to cover administrator, this method has been supplanted by film covering innovation. This procedure was begun with the utilization of natural solvents like methylene chloride however presently has been supplanted with fluid film administrative covering because of ecological and contemplations. Additionally, the expense of any natural dissolvable is undeniably more than the expense of cleansed water. Day by day watery-based framework or aqueous film coating technology has been gaining too much interest to the researchers. After getting some disadvantages of the aqueous coating technology, film covering innovation has now progressed to a high level. "The successful introduction of a large type of liquid primarily based film coating merchandise (below the name INSTACOAT) has resulted in straightforward conversion from organic solvent-based coatings to liquid film coating for many companies; several of them still use the standard coating instrumentation"[10],[11].

#### Development of film coating formulation: -

Improvement of the coating adhesion properties to the core materials required the optimization of the film coating formulation, reducing intagliation bridging, raising coating hardness, or improving any other attribute that the formulator considers insufficient. The development scientist must take into account three major factors that can affect film quality: elasticity, tensile strength, and film–tablet surface interaction [12].

## 2) Electrostatic Dry Coating

For the first time, an electrostatic dry powder coating process was developed in a pan coater gadget system for solid dosage forms. This process helps to provide the tablet with a smooth surface bed, excellent coating uniformity, and release at a particular solvent grade. The electricity coating method played a very important role in several manufacturing industries such as paint technology, food technology, metal coating process, and pharmaceutical industries to the coating of solid dosage forms. The principle involved in the electrostatic dry coating process, directly spreading of particle and polymer mixture on the surface of tablet bed without the addition of solvent and heating will be applied until its form a film over the surface of tablets [13], [14]. Mainly two types of charging units can be found out based on charging mechanism as, a) Corona charging mechanism, b) Tribo charging mechanism.

### Mechanism of Tribo Charging: -

The principle involved in the Tribo charging is friction charging conjugated with dielectric properties and there are no such free ions or electrical fields present in between the spray gun and grounded substance. Electrical forces present in the Tribo charging gun are the repulsive force that exists in between the particles. Charged particles are able to enter into the space adjacent where attraction forces exist between ground substrate that leads to deposition of the particles on the substrate. Mechanical forces lead to the uniformity of the charged particles onto the earthen substrate [15].

At last, Forces are equal as repulsion force equals the attraction force, which leads to particles cannot adhering to the substrate and thickness of coating materials do not enhance. Electrostatic coating of pharmaceutical tablet core and electrically nonconducting substance is very much difficult than other coating processes. To protect the core of the tablet, powder transformed into the film without damaging the core materials of the tablet, especially organic materials [16].

## 3) Super-Cell Coating Technology

Supercell coating technology is an amazing coating technology of the cutting-edge tablet that can withstand extremely hygroscopic materials and friable coating ingredients can be deposited into it. Sometimes this technology gives nonhomogenous output due to imperfections and inconsistency. The process involved in the supercell coating technology is that edges of tablets are ground off and corners of the tablet are not coated with the same thickness applied in tablet faces otherwise tablets can be stacked in the rotating pans and airflow cannot pass throughout the pan. That's why the modified release of coating is limited due to the deposition of the coating materials [17]. Supercell coating technology was invented by Niro Pharma Systems which helps with various problems employing a small modular design.

## Features of Super-Cell coating technology

 Coating of multiple layers over the surface bed of solid dosage forms.

- 2) Modular designs are easily flexible.
- 3) Coating can be continuous and easily adaptable.
- Production capacity is more & can be 6 cells coats up to 120mg
- 5) Can be used in the R & D department withstand of the minimum batch size range 30mg
- 6) Much more accurate than other technology.
- 7) Having a low humidity process that is suitable for low moisture-sensitive materials.
- 8) Enabling technology.
- 9) Friable tablets.

# **Defects in tablet coating**

Tablet coating plays an important role in the stability of the core materials from environmental conditions. Several problems can find out during the manufacturing process and the majority of the visual faults caused by insufficient technical skills, improper machine settings, and might be moisture contained in granules before compression. Several defects can be found during the manufacturing process as follows [18], [19]. [20].

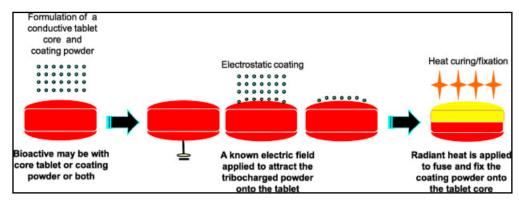


Figure 01 -Diagrammatic representation of Electrostatic Dry Coating System

Table 04 - Disadvantages of the coating of the tablet

Defects	Definition	Reason	Remedies
Blistering	It is a local detachment of film from substrate forming blister.	Entrapment of gases in the film due to overheating either during spraying or at the end of the coating run	Milder drying conditions are warranted in this
Chipping	It is a defect where the film becomes chipped and dented, usually at the edges of the tablet.	Decrease in fluidizing air or speed of rotation of the drum in pan coating	Be careful not to over-dry the tablets in the preheating stage. That can make the tablets brittle and promote capping.
Picking	It is a defect where isolated areas of the film are pulled away from the surface when the tablet sticks together and then parts.	produce an overly wet tablet bed	A reduction in the liquid application rate or increase in the drying air temperature and air temperature and air volume usually solves this

Defects	Definition	Reason	Remedies
-			problem. Excessing tacking may be
			an indication of a poor formulation.
			Assuming you don't wish to change
			the tablet shape, you can solve this
			problem by balancing the pan speed
			and spray rate. Try reducing the spray
Twinning	This is the term for two tablets	A common problem with capsule- shaped tablets	rate and increasing pan speed. In
Iwinning	that stick together.		some case, it is necessary to modify
			the design of tooling by very slightly
			changing the radius. The change is
			almost impossible to see, but it
			prevents the twinning problem.
	It is a defect whereby pits occur	The temperature of the tablet core is	
Pitting	on the surface of a tablet core	greater than the melting point of the	Control the temperature of the tablet
Titting	without any visible disruption of	materials used in the tablet	core during the formulation.
	the film coating.	formulation.	
		The coating solution penetration the	
	It is the defect of film coating	surface of the tablet, often at the	
Cratering	whereby volcanic-like craters	crown where the surface is more	
crutoring	appear exposing the tablet	porous, causing localized	
	surface.	disintegration of the core and	
		disruption of the coating.	
	It is a defect where coating becomes dull immediately or	It is due to collection on the surface	
		of low molecular weight ingredients	
Blooming	after prolonged storage at a high	included in the coating formulation.	
	temperature	In most circumstances, the ingredient	
		will be a plasticizer.	
	It is a defect best described as whitish specks or haziness in the film	It is through to be due to precipitated	
D1 1		polymer exacerbated by the use of	
Blushing		high coating temperature at or above	
		the thermal gelation temperature of	
		the polymers	A reformation with different
Coler	A defect that involves variation	Alternation of the frequency and duration of appearance of tablats in	
Color variation	in color of the film.	duration of appearance of tablets in the spray zone or the size/ shape of	plasticizers and additives is the best
variation		the spray zone or the size/ shape of the spray zone	way to solve the film instabilities caused by ingredients.
	It is a defect in which the film	the spray zone	caused by ingredients.
Cracking or splitting	either crack across the crown of		The tensile strength of the film can be
	the tablet (Cracking) or splits	Internal stress in the film exceeds the	increased by using higher molecular
	around the edges of the tablet	tensile strength of the film.	weight polymers or polymer blends.
	(splitting)		
	It is a defect that renders the	The inability of foam, formed by air	Judicious monitoring of fluid
Infilling	integrations indistinctness	spraying of polymer solution, to	application rate and thorough mixing
		spraying or porymer bolution, to	"rrneadon fale and thorough mixing

Defects	Definition	Reason	Remedies
		break. The foam droplets on the	of the tablets in the pan can prevent
		surface of the tablet break down	filling
		readily due to attrition but the	
		ingredients from a protected area	
		allow the foam to accumulate and	
		"set". Once the foam has accumulated	
		to a level approaching the outer	
		contour of the tablet surface, normal	
		attrition can occur allowing the	
		structure to be covered with	
		continuous film.	
	It is a surface defect resulting in		
Orange peel/	the film being rough and	Inadequate spreading of coating	Thinning the solution with additional
roughness	nonglossy. Appearance is similar	solution before drying.	solvent may correct this problem.
	to that of an orange.		
	Mottling is an uneven	It is mainly due to different coloration on excipient or the degradation	Coating solution prepares properly insufficient quantity.
Mottling	distribution of the color on the		
Mottling	surface of the tablet, with dark		
	and light patches on it.	product of tablet is colored.	

#### **CONCLUSION**

Coating of the pharmaceutical dosage form is a wonderful advantage and remarkable development in recent decades to the enhancement of the quality of the solid dosage form.

Several development techniques have come to market to elegance the look, reduction of the error, stability of the tablet and easy way to control and operate. Each technique has its own set of benefits and drawbacks. This technology has undergone significant development and advancement in terms of energy consumption, film processing, and drying quality. In the future, there is still a lot of opportunities for improvement in coating technique. More research into better coating solvents, drying techniques, and spaying methods is required.

### FINANCIAL ASSISTANCE Nil

## CONFLICT OF INTEREST

The authors declare no conflict of interest

### AUTHOR CONTRIBUTION

Soumyadip Ghosh, Debgopal Ganguly, Pubali Chakraborty and Ananta Choudhury designed the work and made the required corrections and revisions in the manuscript. Debgopal Ganguly and Soumyadip Ghosh collected the content and performed literature survey and also contributed in designing the manuscript. All the authors designed the final manuscript.

#### **REFERENCES**

- Kamble N, Chaudhari SP, Oswal RJ, Kshirsagar SS, Antre RV.Innovations in tablet coating technology. A review. *International Journal of Applied Biology and Pharmaceutical Technology*, 2, 214-218(2011).
- [2] Lachman Leon et al. The Theory and Practice of Industrial Pharmacy. Second edition. Fourth Indian Reprint, Bombay: *Published by Varghese Publishing house* (1991)
- [3] Lippincott Williams and Wilkins.Remington's The Science and Practice of Pharmacy. Volume-I. 21<sup>st</sup>ed. Indian Edition (2005).
- [4] Piccoli G, Onofri F, Cirnaru MD, Kaiser CJ, Jagtap P, Kastenmüller A, Pischedda F, Marte A, Von Zweydorf F, Vogt A, Giesert F. Leucine-rich repeat kinase 2 binds to neuronal vesicles through protein interactions mediated by its C-terminal WD40 domain. *Molecular and cellular biology*. 34(12), 2147-61 (2014)
- [5] Qiao M, Zhang L, Ma Y, Zhu J, Chow K.A Novel Electrostatic Dry Powder Coating Process for Pharmaceutical Dosage Forms: Immediate Release

Coatings for Tablets. *European J Pharm Biopharm*,**3**, 304-310(2010).

- [6] Pawar A, Deepak VB, Vineeta VK, Vilasrao JK.Advances in Pharmaceutical Coatings. *International Journal of Chem Tech Research*, 2, 733-737(2010).
- [7] Mazumder M, Sims R, Biris A, Sriramaa PK, Sainia D, YurteriCU.Twenty-first century research needs in electrostatic processes applied to research industry and medicine. *Chem Eng Sci*,**61**, 2192-2211(2006).
- [8] Ramlakhan M, Chang Yu Wu, Satoru Watano, Rajesh N. Dave, Robert Pfeffer.Dry particle coating using magnetically assisted impaction coating: modification of surface properties and optimization of system and operating parameters. *Powder Technol*, **112**, 137-148(2011).
- [9] Singh P, Solanky TKS, Mudryy R, Pfeffer R, Dave R. Estimation of Coating Time in the Magnetically Assisted Impaction Coating Process. *Powder Technology*, **121**, 159-167 (2001).
- [10] Lachman L, Lieberman HA, Joseph LK. The Theory and Practice of Industrial Pharmacy. Varghese Publishing House; Mumbai; Third Edition: 297-321.
- [11] Lachman L, Liberman H, Kanig J. The Theory and Practice of Industrial Pharmacy; **Third Edition**, 293-345: 346-373.
- [12] Aulton M. Pharmaceutics: The Science of Dosage Form Design. International Student **3<sup>rd</sup>Edition**: 304-321: 347-668.
- [13] Vyas S, Khar R. Controlled Drug Delivery Concepts and Advances; **First Edition**: 219-256.

- [14] Ansel H, Allen L, Popovich N. Ansel's Pharmaceutical Dosage Forms and Drug Delivery Systems; Eighth Edition: 227-259.
- [15] Remington J. Remington: The Science and Practice of Pharmacy, 2, 1615-1641.
- [16] Porter SC. Novel drug delivery: Review of recent trends with oral solid dosage forms. *American Pharmaceutical Reviews*,4, 28-35 (2001).
- [17] Vinay V, Sivakumar T, TamizhmaniT.Colon targeting drug delivery system: A review on recent approaches. *International Journal of Pharmaceutical and Biomedical Science*.; 2: 11-19(2011).
- [18] Anil K. Philip, Betty Philip.Colon Targeted Drug Delivery Systems: A Review on Primary and Novel Approaches. Oman Medical Journal, 25, 70-78(2010).
- [19] Raju D, Padmavathy J, Sai Saraswathi V, Saravanan D, Aparna Lakshmi I.Formulation and development of enteric coated tablets of prednisolone as a colon targeted drug delivery. *IJPSR*, 2, 685-690(2011).
- [20] Zaid AN, QaddomiA.Development and stability evaluation of enteric coated diclofenac sodium tablets using sureteric. *Pak. J. Pharm. Sci*, 25, 59- 64(2011).