

e-ISSN: 2319-8753 | p-ISSN: 2347-6710

TEDIA

International Journal of Innovative Research in SCIENCE | ENGINEERING | TECHNOLOGY

INTERNATIONAL JOURNAL OF INNOVATIVE RESEARCH

IN SCIENCE | ENGINEERING | TECHNOLOGY

Volume 11, Issue 7, July 2022

INTERNATIONAL STANDARD SERIAL NUMBER INDIA

Impact Factor: 8.118

9940 572 462 🔊

6381 907 438





| e-ISSN: 2319-8753, p-ISSN: 2320-6710| <u>www.ijirset.com</u> | Impact Factor: 8.118|

Volume 11, Issue 7, July 2022

| DOI:10.15680/IJIRSET.2022.1107048 |

A SHORT ASSESSMENT ON "TABLETS" AND IT'S TYPES

Shubham Kumar, Dr. Kaushal Kr chandual, Dr.Gaurav Kumar Sharma, Ms. Sarita Sharma

B.Pharma Student(4th year), Department of Pharmacy, Mewar University Gangrar, Chittorgarh, India

Dean, Department of Pharmacy, Mewar University Gangrar, Chittorgarh, India

HOD, Department of Pharmacy, Mewar University Gangrar, Chittorgarh, India

Counselor, Department of Pharmacy, Mewar University Gangrar, Chittorgarh, India

ABSTRACT: Medicines are not only a science; it is also an art. It does no longer consist of compounding capsules and plasters; it deals with the very methods of life, which need to be understood earlier than theymay be guided. Pharmaceutical oral solid dosage varieties have been used broadly for a long timemainly due to their convenience of administration and their suitability for delivery fordelivery of drugs for systemic effects. The tablets can be made without delay from powders or from granules pellets, or from film-coated a couple of units. Tablets are now the most popular dosage form, accounting for some 70% of all moral pharmaceutical preparations produced. Tablets may be described as strong pharmaceutical dosage varieties containing drug substances with or without appropriate diluents and organized by using either compression or moulding methods. Hence,tablets can be widely labeled as compressed capsules and moulded tablets. Compressed tablets can be further categorised as directly compressible tablets, chewable capsules and pill triturates etc.

KEYWORDS: Binders, Coated Tablets, Compression, Granulation, Ingredients.

I. INTRODUCTION

Solid medicaments may also be administered orally as powders, pills, cachets, capsules or tablets. These dosage types incorporate a quantity of drug which is given as a single Unit and they are acknowledged jointly as solid unit dosage forms, even in the case of Sustained action preparations which, technically, incorporate the equal of several Normal doses of drug. The stringent components requirements of modern Medicaments, the many advantages of pill and pill medication, coupled With expanding health offerings and the dedication want for large- scale Economic manufacture, have led to a steady decline in the prescribing of powders And capsules. Tablets and capsules, on the different hand, currently account for nicely over Two third of the total range and price of drug treatments produced all over the world. Tablets are solid dosage form which is the conventional as nicely as have many advantages over other dosage forms. Tablets are the most popular dosage form; about 70% of the complete medicines are dispensed in the shape of tablet. Tablets had different shapes, sizes, as properly as weight depending on medicinal supplies and the meant mode of administration. In this paper the some blessings as wellas some risks of tablets, the fundamental ingredients that are often discovered itablets, techniques of tablet instruction and the quite a number sorts of the drugs are quickly reviewed.



| e-ISSN: 2319-8753, p-ISSN: 2320-6710| <u>www.ijirset.com</u> | Impact Factor: 8.118|

Volume 11, Issue 7, July 2022

DOI:10.15680/IJIRSET.2022.1107048



Definition

According to the Indian Pharmacopoeia Pharmaceutical capsules are solid, flat orbiconvex dishes, unit dosage form, prepared by using compressing a capsules or amixture of drugs, with or without diluents. Tablet is described as a compressed stable dosage shape containing medicaments with or except excipients. They range in form and range extensively measurement and weight, depending on amount of medicinal substances and the meant mode of administration.

Properties of Tablets

a) Should be dependent product having its very own identification while being free of defects such as chips, cracks, discoloration and contamination.

b) Should have power to withstand the rigors of shocks encountered in its production, packaging, delivery and dispensing.

c) Should have the bodily stability to maintain its bodily attributes over time.

d) Must be in a position to launch the medicament agent(s) in the physique in a predictable and reproducible manner.

e) Must have a suitable chemical stability over time so as no longer to allow alteration of the medicinal agent.

II. INGREDIENTS

In addition to energetic ingredients, tablet consists of a range of inert substances acknowledged as additives or excipients. Different excipients are: S.no. Ingredients

Examples

1.Diluents Calcium Phosphate; Carboxymethylcellulose Calcium; Cellulose; Dextrin; Lactose; Microcrystalline Cellulose; PR gelatinized Starch; Sorbitol; Starch

2.Binders Acacia; Alginic Acid; Carboxymethylcellulose; Cellulose; Dextrin; Gelatin; Liquid Glucose; Magnesium Aluminum Silicate; Maltodextrin; Methylcellulose; Povidone; Sodium Alginate; Starch; Zein.

3.Lubricants Calcium Stearate; Glyceryl Palmitostearate; Magnesium Oxide; Poloxamer; Polyvinyl Alcohol; Sodium Benzoate; Sodium Lauryl Sulfate; Sodium Stearyl Sulfate; Stearic Acid; Talc; Zinc Stearate

4. Glidant Magnesium Trisilicate; Cellulose; Starch; Talc; Tribasic Calcium Phosphate.

5.Anti –adherentsCorn Starch; Metallic Stearate; Talc

6.Disintegrants Alginic Acid; Carboxymethylcellulose; Cellulose; Colloidal

Silicon Dioxide; Croscarmellose Sodium; Crospovidone; Potassium Polacrilin; Povidone

7.ColoringagentsFD&C or D&C Dyes or Lake Pigments



| e-ISSN: 2319-8753, p-ISSN: 2320-6710| <u>www.ijirset.com</u> | Impact Factor: 8.118|

Volume 11, Issue 7, July 2022

| DOI:10.15680/IJIRSET.2022.1107048 |

8.FlavoringagentsEthyl Maltol; Ethyl Vanillin; Menthol; Vanillin

9. AbsorbentsKaolin; Magnesium Aluminum Silicate; Tricalcium Phosphate

9) Absorbents: The inclusion of absorbents in a tablet formulation is necessary if the product includes a substance with a excessive affinity to water. Hygroscopic materials, if present, render the combination moist and hard to handle for the duration of manufacture.

III. PREPARATION OF TABLETS

Tablets are prepared through three methods

- 1) Wet granulation method
- 2) Dry granulation method
- 3) Direct compression

Wet Granulation Method

It is the most frequent and extensively usedmethod. This technique entails a range osteps like weighing of ingredients, mixing, granulation, and screening ofdamp pass, drying, lubrication and compression of tablets. The majoractive ingredient, diluent, disintegrantare blended together, and then it is allowed to ignore through the sieve(sifting). Solutions of the binding agentare delivered to the initial combination withstirring. The amount of binding ageadded be sufficient, in order to avoid over wetting of the tablet.

If the powder is no longer wettedproperly, the granules will be too gentleand can be broken down for the duration oflubrication, which is tough in the course of compression of tablet. Tray drying ismost common approach of drying thetablet granules, Tray drying was themost broadly used method of dryingtablet granulations in the past, which might be changed by using fluid –bed dryersas a novel approach. After drying thegranules, they are allowed to pass by fluid- bed dryer as a new approach. After drying the granules they are allowed to pass thru the screen; typically 60-100 mesh nylon cloth is used. After dry granulation, lubricant is delivered as excellent powder, which is required for suitablefilling of the die cavity.

Dry Granulation Method

This method is used for pill preparation, in case pill substances are noticeably sensitive to moisture, or unable to with stand multiplied temperatures throughout drying, slugging may be used to form the granules. Dry granulation or double compression, generally eliminates various steps, which involves slugging of the powder mass. The activeingredient, diluent and lubricant are blended together, to structure the slug. Thus, the compressed slug is surpassed through the mesh or through the mill, and the remaining lubricant is delivered to the granulation, blended suitable and compressed to shape the capsules.

Direct Compression

Direct compression entails direct compressing the powdered cloth into tablets. Direct compression is adopted, if drug constitutes major portion of pill whole weight .Tablets containing 25% or less of drug supplies can be formulated, with a appropriate diluent which acts as a service or automobile for the drug. Tablets organized with the aid of above method are subjected to compression machine which may additionally be single station or more than one stations .

IV. TYPES OF TABLETS

1. The a range of kinds of drugs described as pr comply with :-

a.Compressed Tablets

Compressed drugs signify a considerable share of tablets that are clinically used to provide systemic administration of therapeutic agents both in a of the uncoated nation (i.e., in their simplest form) or in a covered state. These pills are designed to provide rapid disintegration in the gastric fluid following ingestion hence, allowing fast release of



| e-ISSN: 2319-8753, p-ISSN: 2320-6710| <u>www.ijirset.com</u> | Impact Factor: 8.118|

Volume 11, Issue 7, July 2022

| DOI:10.15680/IJIRSET.2022.1107048 |

the drug and, ultimately, systemic absorption of the dosage form. Compressed pills are formed with the aid of compression of powdered, crystalline, or granular materials into the required geometry via the application of high pressures, using metal punches and die. In addition to the Active Pharmaceutical Ingredient(s) (APIs), compressed capsules normally include a range of pharmaceutical excipients e.g., bulking agents, disintegrants, binders, lubricants, controlled-release polymers and different miscellaneous adjuncts such as colourants and flavourants which serve distinctive and specialised reason for the duration of pill manufacture, storage, and use. Examples of compressed drugs consist of tablets for oral, buccal, sublingual, or vaginal administration.

b. Sugar Coated Tablet

These are compressed pills that have been lined with focused sugar solution to improve patient's compliance, extend aesthetic appeal, masks objectionable tastes or odours, enlarge stability and/or regulate the launch of therapeutic agent(s). Sugarcoating was as soon as pretty frequent however misplaced industrial appeal due to the time and understanding required in the coating process, the increase in dimension and weight of lined tablets, excessive value of procedure validation and shipping.

c.Film Coated Tablets

Film-coated tablets are conventional pills covered with a skinny layer of polymer (e.g., hydroxypropyl methylcellulose, hydroxypropyl cellulose) or a combination of polymers (e.g., Eudragit E100) capable of forming a skin-like film. The movie is usually colored and additionally affects the same common characteristics as sugar coating with the delivered gain of being greater durable, much less bulky, and much less time-consuming to apply. By its composition, the coating is designed to spoil and expose the core tablet at the desired location in the gastrointestinal tract.

Advances in material science and polymer chemistry have made these coatings the first desire for system scientists. Examples of Film-coated tablets consist of Curefenac a hundred – Diclofenac potassium USP 100mg (Unicure Pharmaceutical Ltd), Valsartan 320mg Film-coated Tablets (Actavis UK Ltd), etc.

d. Effervecsent Tablets

Effervescent drugs are uncoated capsules that commonly contain natural acids (such as tartaric or citric acid) and sodium bicarbonate in addition to the medicinal substance or API. They react swiftly in the presence of water via releasing carbon dioxide which acts as a disintegrator to produce both a drug suspension or an aqueous solution. These tablets are prepared by way of compressing granular effervescent salts (organic acid and bicarbonate) with the medicinal substances. A ordinary instance of this tablet kind is Ca C1000 Sandoz effervescent tablet (Novartis).

e. Enteric Coated Tablets

Enteric-coated pills are compressed drugs that have delayed-release properties. They are covered with polymeric resources (such as cellulose acetate phthalate/cellulose acetate butyrate; hydroxypropylmethylcellulose succinate; and methacrylic acid copolymers) that withstand answer in gastric fluid but fall apart and permit drug dissolution and absorption in the intestine. Enteric coatings are specifically employed when the drug substance is inactivated or destroyed by using gastric acid (e.g., erythromycin) or is especially hectic to the gastric mucosa (e.g., non-steroidal anti-inflammatory drugs) or when pass by of the stomach substantially enhances drug absorption. Example of enteric-coated tablets consists of Lofnac one hundred – Diclofenac sodium delayed-release pill USP 100mg (bliss GVS Pharma Ltd), Ecotrin pills and caplets (GlaxoSmithKline Beecham).

f. Chewable Tablets

Chewable tablets are huge sized capsules which are hard to swallow and thus, are chewed inside the buccal cavity prior to swallowing. They are specially beneficial for administration of giant capsules to children and adults who have subject swallowing conventional capsules or antacid formulations in which the dimension of the tablet is usually large and the neutralisation efficacy of the pill is associated to particle size within the stomach.



| e-ISSN: 2319-8753, p-ISSN: 2320-6710| <u>www.ijirset.com</u> | Impact Factor: 8.118|

Volume 11, Issue 7, July 2022

| DOI:10.15680/IJIRSET.2022.1107048 |

Chewable pills are not conventionally used if the drug has problems regarding style acceptability. Examples of chewable pills include Danacid – compound magnesium trisilicate pill B.P. (Dana Pharmaceuticals Limited), Gestid – tasty chewable antacid (Ranbaxy) etc.

g. Buccal and Sublingual Tablets

Buccal and sublingualdrugs are small, flat,oval tablets that are supposed to be dissolved in the buccal pouch(buccal tablets) or beneath the tongue (sublingual tablets) for absorption via the oral mucosa to produce a systemic effect. These tablets are employed to acquire both fast absorption into the systemic circulation e.g. glyceryl trinitrate sublingual capsules or, alternatively, to enable oral absorption of pills that are destroyed with the aid of the gastric juice.

2. Molded Tablets

a. Hypodermic Tablets:These are one type of sterile preparations. In these tablets are dissolved in the WFI or sterile water to inject before the authentic injection in the hypodermic cavity. They are meant to be added in WFI of sterile water to structure a clear solution which is to be injected parentally. They are widely used via rural doctor due to its portability. It can be used for medicaments whose stability in water is very poor. Their use in this manner should be discouraged, because the resulting solutions are not sterile.

b. Dispensing or Soluble Tablets: Theyare to be added to water or differentsolvents to make a solution containinga constant awareness of API. Shouldcontain no insolublematerials(including Glidants, binders etc.), due to the fact thatthey will be made into clear solution. A cloth incorporated in allottingtablets includes slight silver proteinate, bichloride of mercury and quaternaryammonium compounds. These drugsare exceedingly poisonous if taken orally with the aid ofmistake. These capsules furnish aconvenient quantity of mighty drug.

c. Tablet Triturate:Tablet triturates aresmall, normally cylindrical, Molded orcompressed tablets. The drugs employed in such products have been usually pretty amazing and were combined with lactose and perchancea bonder, such as powder acacia. Tabletriturates are typically gentle and friable. Manyof the drugs employed in these tables were highly potent and drug migration should occur as the alcohol evaporated. Only minimal stress is utilized throughout theirmanufacturing, due to the fact that they should be without problems and totally soluble in water.

3.Structure Wise

a) Divisible Tablet: It is every so oftennecessary to administer one-half oone-fourth of a tablet and beneath such circumstances pills are commonly scored once in the center or twice withlines perpendicular to one another. Vshaped double layer pills with scoring in the middle havebeen designed.

b) Aperture Tablets: Designed with aview to reap fidelity in the surface area in the course of disintegration & dissolution.

c) Concave-convex Tablets: Thesetablets have been designed with a viewto hold floor location of the structure relatively constant during the dissolution process. Area is misplaced on the convex surfaces and gained at the concavities.

d) **Core Tablets:** These pills have acentral core over which every other layer of fabric is compressed and aregenerally made with the aid of two successivecompressions. Separate incompatibleing redients.

ACTION WISE

1) **Modified Release Tablet:** Release themedicament slowly for lengthy timeduration after administration of asingle tablet. Used to target the websitespecific releases. Comparison of bloodconcentration vs. time any adjuvantthat can alter water uptake rate, swelling, and gelling characteristicscan alter the release fee of API. Thedrug launch can be



| e-ISSN: 2319-8753, p-ISSN: 2320-6710| <u>www.ijirset.com</u> | Impact Factor: 8.118|

Volume 11, Issue 7, July 2022

| DOI:10.15680/IJIRSET.2022.1107048 |

modified by usingproviding appropriate micro environmentpH in the pill .Inclusion of alkalinepolymers results in perfect drugrelease of acidic drugs.

V. CONCLUSION

As a stable dosage form, capsules are famousamong sufferers and practitioners alike asthey furnish a potential of selfadministration. The components of a pillcontains, in addition to the API, a range of substances to assure suited delivery of the API to the patient. With advancement intechnology and amplify in focustowards modification in fashionable tablet toachieve higher acceptability as well asbioavailability, newer and greater environment friendlytablet dosage varieties are being developed. The important motives behind formulation of different types of tablets are to create adelivery device that is incredibly easand cheaper to manufacture. Provide dosage structure that is handy frompatient's perspective and utilize anapproach that is unlikely to addcomplexity for the duration of regulatory approval process. To apprehend each dosage form, tablets here are categorized by their route administration and by the type of drugdelivery machine they symbolize inside thatroute.

REFERENCES

1) Leon Lachman, Herbert A. Lieberman, Joseph L. Kanig: The theory and Practice of Industrial Pharmacy, Varghese ebook house, 3rdedition, 1990, 293-373.

2) Herbert A. Liberman, Martin M.Rieger and Gilbert S. Banker, pharmaceutical dosage forms: Tablets; volume-I. 49-52

3) Al-Achi A (2019) Tablets: A BriefOverview. Journal of Pharm Practice and Pharmaceutical Science 2019.(1).

4) Nagashree K. Solid dosage forms: Tablets. Research and Reviews: Journal of Pharmaceutical Analysis 2015.

5) Kaur Harbir. International ResearchJournal of Pharmacy. 2012.3(7)

6) G. Hymavathi, J. Adilakshmi, K.Dwarathi, M. Kavya, G. Pravallika.Review Article on In method Problemsand Evaluation Tests of TabletManufacturing. International Journalof Research in Pharmaceutical andNano Sciences. 2012.3(7).

7) D. M. Jariwala, H. P. Patel, C. T. Desai, S. A. Shah and D. R. Shah. A Review on Multiple Compressed Tablets. Journal of pharmaceutical sciences and boiscientific research. 2016, 6(3). 371-379.

8) Purushottam R. Patil, Vaibhav D. Bobade, Pankaj L. Sawant and Rajendra P. Marathe. Emerging Trends In Compression Coated Tablet Dosage Forms: A Review. International journal of pharmaceutical sciences and research. 2016, 7(3). 930-938.

9) Bhavjit Kaur, Shivani Sharma, GeetikaSharma, Rupinder Saini, SukhdevSingh, Meenu Nagpal, Upendra K Jain, Mandeep Sharma. A Review of Floating Drug Delivery System. Asianjournal of biomedical and Pharmaceutical Sciences. 2013. 3(24) ,1-6.

10) Anita , Anil Singh and Ankit Dabral. A Review On Colon Targeted DrugDelivery System. International journal of pharmaceutical sciences and research. 2019; Vol. 10(1): 47-56.

11) Renu, Jyoti Dahiya, Pawan Jalwal, Balvinder Singh. The PharmaInnovation Journal 2015; 4(5): 100-105.

12) Nilesh Lahanu Gawade, RaosahebSopanrao Shendge. A Review onChewable Tablet. Journal of EmergingTechnologies and Innovative Research(JETIR). March 2020, Volume 7, Issue.

13) Cottrell, J.; Koenig, K.; Perfekt, R.; Hofmann, R.; For the Loperamide–Simethicone Acute Diarrhoea Study Team (2015). "Comparison of Two Forms of Loperamide–Simeticone and a Probiotic Yeast (Saccharomyces boulardii) in the Treatment of Acute Diarrhoea in Adults: A Randomised Non-Inferiority Clinical Trial". Drugs in R&D. 15 (4): 363–373.

14) "Compressed Sugar Coated Caplet Dosage Form (Code C69002)". NCI Thesaurus. Retrieved 2017-12-06.

15) "Lubricant Excipients | American Pharmaceutical Review". www.americanpharmaceuticalreview.com. Retrieved 2021-09-22.

16). "Extrusion Speheronisation". PharmaCMC.com. Archived from the unique on 1 October 2016. Retrieved 26 September 2016.

17). "Articles :: ГК "Аптека-95"". Archived from the original on 2014-01-27.



| e-ISSN: 2319-8753, p-ISSN: 2320-6710| <u>www.ijirset.com</u> | Impact Factor: 8.118|

Volume 11, Issue 7, July 2022

| DOI:10.15680/IJIRSET.2022.1107048 |

18). Gendre C., Genty M., César da Silva J., Tfayli A., Boiret M., Lecoq O., Baron M., Chaminade P., Péan J-M., Comprehensive find out about of dynamic curing impact on tablet coating structure, Eur. J. Pharm. Biopharm., 81 (2012), 657-665

19). Coating machines Archived 2012-03-28 at the Wayback Machine, apteka95.com.





Impact Factor: 8.118





INTERNATIONAL JOURNAL OF INNOVATIVE RESEARCH

IN SCIENCE | ENGINEERING | TECHNOLOGY





www.ijirset.com