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the free encyclopedia that anyone can edit.117,185 active editors 7,001,904 articles! Learn how you can take part in the encyclopedia's continued improvement. Members of the victorious Blondie crewThe Boat Race 2018 took place on
24March. Held annually, The Boat Race is a side-by-side rowing race between crews from the universities of Oxford and Cambridge along a 4.2-mile (6.8km) tidal stretch of the River Thames in south-west London, England. For the third time in the history of the event, the men's, the women's and both reserves' races were all held on the Tideway on
the same day. The women's race saw Cambridge lead from the start, eventually winning by a considerable margin to take the overall record to 4330 in their favour. In the women's reserve race, Cambridge's Blondie (crew pictured) defeated Oxford's
Isis by a margin of four lengths. The men's race was the final event of the day and completed a whitewash as Cambridge won, taking the overall record to 8380 in their favour. The races were watched by around 250,000 spectators live, and broadcast around the world. (Fullarticle...)Recently featured: Radar, Gun Laying, Mk.I and Mk.IIAndrea
NavageroNosy KombaArchiveBy emailMore featured articlesAboutKitty Marion... that Kitty Marion... that Kitty Marion (pictured) was force-fed over 200 times during a hunger strike?... that the North Korean destroyer Choe Hyon is the largest ship constructed for the Korean People's Navy?... that after the release of High and Low, director Akira Kurosawa received
telephone calls imitating his film that threatened to kidnap his daughter?... that May Bradford Shockley is why Silicon Valley is where it is?... that Joy Laking predicted in a school writing assignment that within ten years she would be making a living as an artist?... that
the Taiwanese restaurant chain Formosa Chang drew inspiration from McDonald's for its non-greasy atmosphere and corporate practices?... that "Steve's Lava Chicken" recently became the shortest song to enter the UK Top 40? ArchiveStart a new
articleNominate an articleNgg wa Thiong'o (pictured) dies at the age of 87. In sumo, nosato Daiki is promoted to yokozuna. In association football, Liverpool win the Premier League title. In motor racing, lex Palou wins the Indianapolis 500. In basketball, the EuroLeague concludes with Fenerbahe winning the
Final Four Playoff.Ongoing: Gaza warM23 campaignRussian invasion of UkrainetimelineSudanese civil wartimelineRecent deaths: Harrison Ruffin TylerPhil RobertsonMary K. GaillardPeter DavidAlan YentobGerry ConnollyNominate an articleMay 31: Dragon Boat Festival in China and Taiwan (2025); World No Tobacco DayBessarion455 Petronius
Maximus, the ruler of the Western Roman Empire, was stoned to death by a mob as he fled Rome ahead of the arrival of a Vandal force that sacked the city.1223 Mongol invasion of Kievan Rus': Mongol forces defeated a Kievan Rus' army at the Battle of the Kalka River in present-day Ukraine.1468 Cardinal Bessarion (pictured) announced his
donation of 746 Greek and Latin codices to the Republic of Venice, forming the Biblioteca Marciana. 1935 A magnitude-7.7 earthquake struck Balochistan in British India, now part of Pakistan, killing eight people and injuring
more than 150 others. Albertino Mussato (d.1329) Joseph Grimaldi (d.1837) Dina Boluarte (b.1962) Mbaye Diagne (d.1994) More anniversaries: May 30 May 31 June 1 Archive By email List of days of the year About Cucumis metuliferus, the African horned cucumber, is an annual vine in the cucumber and melon family, Cucurbitaceae. Its fruit has horn-like
spines, hence the name "horned melon". The ripe fruit has orange skin and lime-green, jelly-like flesh. It is native to Southern Africa, where it is a traditional food. Along with the gemsbok cucumber and the citron melon, it is one of the few sources of water during the dry season in the Kalahari Desert. This photograph, which was focus-stacked from
25 separate images, shows two C.metuliferus fruits, one whole and the other in cross-section. Photograph credit: Ivar LeidusRecently featured: Ignace TonenAustralian white ibisHell Gate BridgeArchiveMore featured picturesCommunity portal The central hub for editors, with resources, links, tasks, and announcements. Village pump Forum for
discussions about Wikipedia itself, including policies and technical issues. Site news Sources of news about wikipedia. Help desk Ask questions about using or editing Wikipedia. Reference desk Ask research questions about encyclopedic
topics. Content portals A unique way to navigate the encyclopedia. Wikipedia is written by volunteer editors and hosted by the Wikimedia Foundation, a non-profit organization that also hosts a range of other volunteer projects: CommonsFree media repository MediaWikiWiki software development Meta-WikiWikimedia project coordination
WikibooksFree textbooks and manuals WikidataFree knowledge base WikinewsFree-content news WikiquoteCollection of quotations WikisourceFree-content library WikispeciesDirectory of species WikiversityFree learning tools WikiversityFree learning tools
Wikipedias are available; some of the largest are listed below. 1,000,000+ articles Bahasa IndonesiaBahasa MelayuBn-lm-gCataletinaDanskEestiEsperantoEuskaraMagyarNorsk bokmlRomnSimple EnglishSloveninaSrpskiSrpskohrvatskiSuomiTrkeOzbekcha
50,000+ articles AsturianuAzrbaycancaBosanskiFryskGaeilgeGalegoHrvatskiKurdLatvieuLietuviNorsk nynorskShqipSlovenina Retrieved from "2This article is about the year 455. For other uses, see 455 (disambiguation). This article is about the year 455. For other uses, see 455 (disambiguation).
Unsourced material may be challenged and removed. Find sources: 455 news newspapers books scholar JSTOR (April 2019) (Learn how and when to remove this message) Calendar year Years 452453454455 456457458 vte 455 by
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(CDLV) was a common year starting on Saturday of the Julian calendar. At the time, it was known as the Year of the Consulship of Valentinianus and Anthemius (or, less frequently, year 1208 Ab urbe condita). The denomination 455 for this year has been used since the early medieval period, when the Anno Domini calendar era became the prevalent
method in Europe for naming years. March 16 Emperor Valentinian III, age 35, is assassinated by two Hunnic retainers of the late Flavius Aetius, while training with the bow on the Campus Martius (Rome), ending the Theodosian dynasty. His primicerius sacri cubiculi, Heraclius, is also murdered. March 17 Petronius Maximus, former domesticus
("elite bodyguard") of Aetius, becomes (with support of the Roman Senate) emperor of the Western Roman Empire. He secures the throne by bribing officials of the imperial palace. Maximus consolidates his power by a forced marriage with Licinia Eudoxia, widow of Valentinian III. Maximus appoints Avitus, most trusted general, to the rank of
magister militum and sends him on an embassy to Toulouse, to gain the support of the Visigoths. He elevates his son Palladius to Caesar and has him marry Eudocia, eldest daughter of Valentinian III.May 31 Maximus is stoned to death by an angry mob while fleeing Rome. A widespread panic occurs when many citizens hear the news that the Vandals
are plundering the Italian mainland. June 2 Sack of Rome: King Genseric leads the Vandals into Rome, after he has promised Pope Leo I not to burn and plunder the city. Genseric sacks the city for a period of two weeks. Eudoxia and Placidia, are taken hostage. The loot is sent to the harbour of Ostia and loaded into ships,
from whence the Vandals depart and return to Carthage. July 9 Avitus is proclaimed Roman emperor at Toulouse, and later recognised by the Gallic army. He restores the imperial authority in Noricum (modern Austria) and leaves a Gothic force under Remistus, Visigoth
general (magister militum), at Ravenna. The Ostrogoths conquer Pannonia and Dalmatia. Battle of Aylesford (Kent). Hengist and his son Oisc become king of Kent. Horsa and Catigern, brother of Vortimer, are killed. The Britons
withdraw to London (according to the Anglo-Saxon Chronicle). Skandagupta succeeds Kumaragupta I as ruler of the Gupta Empire (India). During his reign he crushes the Hun invasion; however, the expense of the wars drains the empire's resources and contributes to its decline. Gaero becomes king of the Korean kingdom of Baekje. [1] Earliest
recorded date at Chichen Itza on the Yucatn Peninsula (Mexico) (approximate date). Barter economy replaces organized trade as Romans and other citizens desert their towns for the countryside, where they will be less vulnerable to barbarian raids (approximate date). The city of Vindobona (Vienna) is struck by an epidemic that spreads through the
Roman provinces. The disease is probably streptococcus or a form of scarlet fever with streptococcus pneumoniae (approximate date). Rusticus, archbishop of Lyon (approximate date).
sacri cubiculi )May 31 Petronius Maximus, emperor of the Western Roman EmpireBiyu of Baekje, king of Baekje, k
Petronius Maximus (approximate date) Prosper of Aquitaine, disciple and Christian writer (approximate date) a b "List of Rulers of Korea". www.metmuseum.org. Retrieved April 20, 2019.Retrieved from " 30ne hundred years, from 301 to
400Millennia1stmillenniumCenturies3rdcentury4thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5thcentury5t
Hemisphere at the end of the 4th century CE. The 4th century was the time period from 301 CE (represented by the Roman numerals CCCI) to 400 CE (CD) in accordance with the Julian calendar. In the West, the early part of the century was shaped by Constantine the Great, who became the first Roman emperor to adopt Christianity. Gaining sole
reign of the empire, he is also noted for re-establishing a single imperial capital, choosing the site of ancient Byzantium in 330 (over the current capitals, which had effectively been changed by Diocletian's reforms to Milan in the West, and Nicomedeia in the East) to build the city soon called Nova Roma (New Rome); it was later renamed
Constantinople in his honor. The last emperor to control both the eastern and western halves of the empire was Theodosius I. As the century progressed after his death, it became increasingly apparent that the empire had changed in many ways since the time of Augustus. The two-emperor system originally established by Diocletian in the previous
century fell into regular practice, and the east continued to grow in importance as a centre of trade and imperial power, while Rome itself diminished greatly in importance due to its location far from potential trouble spots, like Central Europe and the East. Late in the century Christianity became the official state religion, and the empire's old pagan
culture began to disappear.[citation needed] General prosperity was felt throughout this period, but recurring invasions by Germanic tribes plagued the empire from 376[1][2] CE onward. These early invasions by Germanic tribes plagued the empire from 376[1][2] CE onward. These early invasions by Germanic tribes plagued the empire from 376[1][2] CE onward. These early invasions marked the beginning of the end for the Western Roman Empire. In China, the Jin dynasty, which had united the nation prior in 280, began
rapidly facing trouble by the start of the century due to political infighting, which led to the insurrections of the northern barbarian tribes (starting the Sixteen Kingdoms period), which quickly overwhelmed the empire, forcing the Jin court to retreat and entrench itself in the south past the Yangtze river, starting what is known as the Eastern Jin
dynasty around 317. Towards the end of the century, Emperor of the Former Qin, Fu Jin, united the north under his banner, and planned to conquer the Jin dynasty in the south, so as to finally reunite the land, but was decisively defeated at the Battle of Fei River in 383, causing massive unrest and civil war in his empire, thereby leading to the fall of
the Former Qin, and the continued existence of the Eastern Jin dynasty. According to archaeologists, sufficient archaeologists, sufficient archaeologists, sufficient archaeologists of the Roman Empire refer to the "Long
Fourth Century" to the period spanning the fourth century proper but starting earlier with the accession of the Emperor Diocletian in 284 and ending later with the death of Honorius in 423 or of Theodosius II in 450.[3]See also: Christianity in the 4th centuryGregory the Illuminator mosaic, converted Armenia from Zoroastrianism to
ChristianityContemporary bronze head of Constantine I (r. 306337 AD)Early 4th century Former audience hall now known as the Basilica, Trier, Germany, is built. Early 4th century The Gupta Empire is established. 301: Armenia first to adopt Christianity as state religion. 304439: The Sixteen Kingdoms in China begins. 306337: Constantine the Great
ends persecution of Christians in the Roman Empire (see also Constantinian shift) and constantinian shift s
to the Philippines is built.325328: The Kingdom of Aksum adopts Christianity.325: Constantine the Great calls the First Council of Nicaea to pacify Christianity in the grip of the Arian controversy.335380: Samudragupta expands the Gupta Empire.337: Constantine the Great is baptized a Christian on his death bed.350: About this time the Kingdom of
Aksum conquers the Kingdom of Kush.350400: At some time during this period, the Huns began to attack the Sassanid Empire.[2]350: The Kutai Martadipura kingdom in eastern Borneo produced the earliest known stone inscriptions in Indonesia known as the Mulavarman inscription written in the Sanskrit language using Pallava scripture.[5]Mid-4th
century Dish, from Mildenhall, England, is made. It is now kept at the British Museum, London. Mid-4th century Wang Xizhi makes a portion of a letter from the Feng Ju album. Six Dynasties period. It is now kept at National Palace Museum, Taipei, Taiwan, Republic of China. 365: An earthquake with a magnitude of at least eight strikes the Eastern
Mediterranean. The following tsunami causes widespread destruction in Crete, Greece, Libya, Egypt, Cyprus, and Sicily.376: Visigoths appear on the Danube and are allowed entry into the Roman Empire in their flight from the Huns.378: Battle of Adrianople: Roman army is defeated by the Visigoth cavalry. Emperor Valens is killed.378395:
Theodosius I, Roman emperor, bans pagan worship, Christianity is made the official religion of the Empire. 378: Siyaj K'ak' conquers Waka on (January 8), Tikal (January 16) and Uaxactun. Wall painting of the Council of Constantinople reaffirms the Christian doctrine of
the Trinity by adding to the creed of Nicaea.383: Battle of Fei River in China.395: The Battle of Canhe Slope occurs.395: Roman emperor Theodosius I dies, causing the Roman Empire to split permanently. Late 4th century: Cubiculum of Leonis, Catacomb of Commodilla, near Rome, is made. Late 4th century: Atrium added in the Old St. Peter's
Basilica, Rome. For a more comprehensive list, see Timeline of historic inventions 4th century. The Stirrup was invented in China, no later than 322.[6][1]Kama Sutra, dated between c.400 BC to c. 300 AD.[7][8]Iron pillar of Delhi, India is the world's first Iron Pillar. [citation needed] Trigonometric functions: The trigonometric functions sine and versine
originated in Indian astronomy.[9]Codex Sinaiticus and the Codex Vaticanus Graecus 1209, are the earliest Christian bibles.[10][11]Book of Steps, Syriac religious discourses.[citation needed] a b "The invention and influences of stirrup". Archived from the original on December 3, 2008. a b Roberts, J: "History of the World". Penguin, 1994. The
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2013.Retrieved from 4The following pages link to 4th century External tools(link counttransclusion countsorted list) See help page for transcluding these entries howing 50 items. View (previous 50 | next 50) (20 | 50 | 100 | 250 | 500)List of decades, centuries, and millennia (links | edit)Religion in pre-Islamic Arabia (links | edit)Rosetta Stone (links | edit)Ro
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edit)510s (links | edit)View (previous 50 | next 50) (20 | 50 | 100 | 250 | 500)Retrieved from "WhatLinksHere/4th_century"Ever stumbled when pronouncing the name of a medication? Or watched a pharmaceutical ad on TV and wondered, Who comes up with these names?Theyre often a mouthful and may seem totally random. So, you might be
surprised to learn that just as theres a science and process behind developing medications, theres also a science and process behind naming them. A medications name plays a key role in patient receiving the wrong medication. This type of
medication error can be life-threatening.Dr. Vyas walks us through the science behind medication names.Whats in a drug name?The U.S. Food and Drug Administration (FDA) approved 55 new drugs in 2023. But before they hit the market, pharmaceutical companies had to take several steps to find names for each of them that met strict U.S. and
international requirements. Its a selection process that sometimes takes up to four years and a lot of money. According to some estimates, drug companies can pay outside consulting firms up to half a million dollars for their help naming new medications. And heres something else you probably didnt know: By the time a brand-name medication
becomes available, its actually on its third name. Thats because each medication name goes through a unique vetting process that includes a: Chemical drug nameGeneric drug nameGeneric drug name reflects its chemical makeup. For instance, N
(4-hydroxyphenyl) acetamide is the chemical name for: Acetaminophen (generic drug name) Tylenol (brand drug name) Unless you work in drug development or medicine, youll probably never hear the scientific name. Still, companies must follow rules set by the International Union of Pure and Applied Chemistry when creating a chemical drug
name. The science behind generic drug namesDrugs get their generic names when theyre ready for testing in a clinical trial, and the process for naming generic drugs has been in place since the early 1960s. Drug companies file three to six generic names suggestions with the United States Adopted Name (USAN) Council. Members of the American
Medical Association, American Pharmacists Association and U.S. Pharmacopeia make up this council. A generic drug always goes by the same name worldwide. Thats why the World Health Organizations International Nonproprietary Names (INN) expert group also has a say. A universal generic name ensures that healthcare providers anywhere in the
world will know the drug, even if the brand name is different, Dr. Vyas explains. It lowers the risk of a patient getting the wrong medication if theyre in a different country. Choosing a generic drug suffixComing up with a name for a generic drug is a bit easier than naming a brand drug. Generic drug names follow a formula, Dr. Vyas shares. The drug
names suffix, or stem, must reflect how the drug works on the body. For instance, generic drugs that end in:-oxetine (fluoxetine and paroxetine) are selective serotonin reuptake inhibitors (SSRIs) that treat mood disorders.-prazole (omeprazole) are proton pump inhibitors for acid reflux. Choosing a generic drugs prefix Companies can
get more creative with the one- to two-syllable prefix. But there are some things the prefix cant do, like: Have the letters H, J, K, W or Y because some world languages dont use those lettersLook or sound similar to other generic drugs that share the same suffix Reflect or promote the name of the drug company Use medical terms, like onco for a cancer
 (oncology) drugApproving a generic drug nameBefore granting approval, USAN and INN consider whether a generic drug name: Is easy to pronounce and rememberHas a suffix that accurately reflects what the drug doesTranslates well into non-English languagesHow brand drugs get their namesPharmaceutical companies invest a lot of resources
into developing a new drug. Theyre like the companys baby and just like expectant parents, they scrap hundreds of potential names, weeding out ones that arent quite right. Why? Because the brand name matters. Healthcare providers and patients will associate the name with that trademarked drug for life. Some drug names like Prozac and Viagra
 have become so commonplace that they now appear in dictionaries, notes Dr. Vyas. Theyre now seen as synonymous with a particular health condition. Focus groups and other research methods help companies home in on their top choice from as many as 200 drug names under consideration. It might not always seem like it, but companies try to
choose names that are easy to pronounce in different languages. They also make sure that a translated name doesnt mean something offensive. Often, companies try to come up with a brand name that somehow reflects what the drug does. For example: Viagra, which treats erectile dysfunction, is a combination of vigor (a word that reflects potency
and stamina) plus Niagara (a reference to a steady stream or Niagara Falls). Lyrica, which treats nerve pain, evokes soothing music lyrics (like the soothing of nerves). Approving a brand drug nameDrug companies can submit only one brand trug nameDrug companies can submit only one brand trug nameDrug companies can submit two names to the European Medicines Agency (EMA).
While drug companies hope the two organizations agree on the same name, they dont have to. In India, for instance, the statin drug Lipitor goes by the name Atocor. The FDA and EMA require that a brand drug works faster
than othersUse a generic suffix or too closely resemble the drugmakerIf the FDA or EMA rejects a drug name, pharmaceutical companies must go back to the drawing board. Selecting another name lt might seem like
pharmaceutical companies are trying to create tongue-twisters, but in fact, the opposite is true. Companies want a drug name that youll tell your friends about the new medication thats worked for you. IBRANCE. Xeljanz
Sildenafil. Viagra. Most Americans have heard of at least one of those drug names. But where do those monikers come from? Is it scientific? Metaphorical? Both? The drug-naming process is one thats long and involved, whether a company is devising a generic name or a brand name (there are different processes for each). In fact, according to Michae
Quinlan, who is senior manager, trademark development, within the Customer Analytics & Insights group with Pfizer, naming a drug can be a long and laborious process that begins before the drug, itself, has been approved by the FDA. In most industries, you create a name for the product and as long as the trademark is considered available you can
start using that name on your product, says Quinlan. But the drug name has to get reviewed and be considered safe before itll be approved to be used on that potential product. In some cases, Quinlan says it can take four years to go through the name selection and approval process. Still, he says, the exercise can be fun. Youre getting to name the
baby, he says. And that name could be around for generations. In this two-part series, well look at how drugs get both their generic and brand names. The early days t starts with a compound. Like anyone or anything, drugs need labels in order to distinguish them from one another. Marie-Claire Peakman, PhD., executive director of the Primary
Pharmacology Group in Worldwide Research & Development with Pfizer explains that in the early days, chemists register a newly synthesized compound in a database, labeling it with PFwhich stands for Pfizerfollowed by 10 numbers (for example, PF-04965842-01). When the chemists first make up compounds, they have to register them in the
database as soon as theyre identified so that we can identify them and keep track of their performance in our studies, says Peakman. If a compound shows enough promise to make it through early experiments and head towards clinical trials, two naming processes begin to devise a generic name and a brand name for the future drug. How generic
drugs get their generic names The first step in coming up with a name for a drug is selecting its generic, or non-proprietary name. The generic names came about because of the world growing smaller, says
Quinlan. People were traveling abroad more frequently, and it became clear that in other countries, their drugs might be known by another name than back home and would not be able to be identified. Today, two different organizations must approve the names of generic drugs the United States AdoptedNames(USAN) Council and the World Health
Organization (WHO) INN Programmeso that regardless of where someone is located, patients and health care professionals will be able to safely communicate about medications. Generic names are, in part, based on a formula. The suffix, or, as Quinlan calls it, the family name, important piece of information to health care professionals will be able to safely communicate about medications.
about how the substance works in the body. Take Viagra, for example. Its generic name is sildenafil. The suffix, afil, explains the way it works, says Quinlan, referring to its role as a PDE 5 inhibitor, meaning it helps control blood flow. Afil is also the suffix of other erectile dysfunction generics, such as tadalafil(Cialis), vardenafil (Levitra and Staxyn
and avanafil (Stendra). The prefix gets a little more creative. We look for syllables that obviously are different from other existing generic name, says Quinlan. Celecoxib. Quinapril. Ziprasidone. When devising those
names, there are a number of rules that apply. Quinlan shared some of those rules: It must avoid certain letters. The generic drug name is created using the Roman alphabet, and the goal is to create a name that can be communicated
globally. Because the letters Y, H, K, J, and W arent used in certain languages that use the Roman alphabet, they arent used in the creation of the prefix of the name. It can't be considered marketing. Using the companys name within the drugs name must be avoided. Also, its important to stay away from superlatives or laudatory terms (best, new,
fastest, strongest) that could be considered promotional. It avoids medical terminology. You dont want to imply that a drug is intended only for one particular function, because in time, if it is also helpful for another purpose, the name could be reductive. Say you were developing a treatment for oncology indications and you launched a product for
those indications, but over time in further research you discovered it worked on inflammation and immunology indications. If you had something like Onc- in the beginning of your generic name that would be very limiting, says Quinlan. When the team has three to six names they like, it submits them to USAN Council, which is made up of
representatives from the American Medical Association (AMA), United States Pharmacopeia (USP) and the American Pharmacists Association (APhA). Sometimes, its a success and one of the names is accepted. Other times, USAN declines to accept the names, and counter proposes a name of its choosing, which Pfizer can then choose to screen and
accept. When a name is accepted, USAN then submits it on behalf of Pfizer to the WHO, where a committee reviews it and decides whether to accept a name, its published on a proposed International Nonproprietary Names (INN) list, and, over the course of four months, the publication and the course of four months and decides whether to accept a name, its published on a proposed International Nonproprietary Names (INN) list, and, over the course of four months, the publication accepts a name, its published on a proposed International Nonproprietary Names (INN) list, and, over the course of four months, the publication accepts a name, its published on a proposed International Nonproprietary Names (INN) list, and, over the course of four months, the publication accepts a name, its published on a proposed International Nonproprietary Names (INN) list, and, over the course of four months, the publication accepts a name, its published on a proposed International Nonproprietary Names (INN) list, and over the course of four months, the publication accepts a name of the course of four months accepts a name of the course of four months accepts a name of the course of four months accepts a name of the course of four months accepts a name of the course of four months accepts a name of the course of 
can come forward and object to the name. If no one objects, it publishes to a recommended INN list and Pfizer can start referring to the drugs are named? Check out part two on how branded drugs are named. For millions of patients around the world, the development
of a new treatment or vaccine can be life-changing. For millions of patients around the world, the development of a new treatment or vaccine can be life-changing. Pfizer is proud of its commitment to advancing hemophilia care for more than 40 years, dating back to the introduction of recombinant therapies. Pfizer is proud of its commitment to
advancing hemophilia care for more than 40 years, dating back to the introduction of recombinant therapies. About Pfizer's initiative to make clinical trials more participant-friendly through the decentralized "Clinical trials more participant-friendly through the decentral trials more participant-friendly through the decentral trials more participant-friendly through the decentral trial
units to reduce the need for in-person visits. About Pfizer's initiative to make clinical trials more participant-friendly through the decentralized "Clinical Trial Anywhere" model; a suite of solutions like remote sample collection, alternative study conduct locations, home health, and mobile units to reduce the need for in-person visits. Artificial
intelligence (AI) has quietly become a part of our daily lives through personalized recommendations, virtual assistants, or smart devices, we barely notice it anymore. Artificial intelligence (AI) has quietly become a part of our daily lives through personalized recommendations, virtual assistants, or smart devices, we barely notice it anymore. Despite through personalized recommendations, virtual assistants, or smart devices, we barely notice it anymore.
significant advancements in hemophilia treatments over the past several decades, balancing management of the disorder while maintaining regular day-to-day activities remains a challenge. Despite significant advancements in hemophilia treatments over the past several decades, balancing management of the disorder while maintaining regular day-to-day activities remains a challenge.
to-day activities remains a challenge. Medication used to treat a viral infection used for treat a viral infection. "Antiviral drugs for HIVAntiviral drugs are a class of medication used for treating viral infections. [1]
Most antivirals target specific viruses, while a broad-spectrum antiviral drugs are a class of antimicrobials, a larger group which also includes antibiotic (also termed antibacterial), antifungal and antiparasitic drugs, [3] or antiviral drugs based on monoclonal antibiodies. [4] Most antivirals are
considered relatively harmless to the host, and therefore can be used to treat infections. They should be distinguished from virucides are produced by some plants such as eucalyptus and Australian tea trees. [5] Most of the antiviral virucides are produced by some plants such as eucalyptus and Australian tea trees.
drugs now available are designed to help deal with HIV, herpes viruses, and influenza A and B viruses. [6] Viruses use the host organism's cells. Moreover, the major difficulty in
developing vaccines and antiviral drugs is due to viral variation.[7]The emergence of antivirals is the product of a greatly expanded knowledge of the genetic and molecular function of viruses, major advances in the techniques for finding new drugs, and the
pressure placed on the medical profession to deal with the human immunodeficiency virus (HIV), the cause of acquired immunodeficiency syndrome (AIDS).[8]The first experimental antivirals were developed in the 1960s, mostly to deal with herpes viruses, and were found using traditional trial-and-error drug discovery methods.[9] Researchers grew
cultures of cells and infected them with the target virus. They then introduced into the cultures chemicals which they thought might inhibit viral activity and observed whether the level of virus in the cultures chemicals which they thought might inhibit viral activity and observed whether the level of virus in the cultures chemicals which they thought might inhibit viral activity and observed whether the level of virus in the cultures chemicals which they thought might inhibit viral activity and observed whether the level of virus in the cultures chemicals which they thought might inhibit viral activity and observed whether the level of virus in the cultures chemicals which they thought might inhibit viral activity and observed whether the level of virus in the cultures chemicals which they thought might inhibit viral activity and observed whether the level of virus in the cultures chemicals which they thought might inhibit viral activity and observed whether the level of virus in the cultures chemicals which they are the constant of the cultures of the
procedure, and in the absence of a good knowledge of how the target virus worked, it was not efficient in discovering effective antivirals which had few side effects. Only in the 1980s, when the full genetic sequences of viruses began to be unraveled, did researchers begin to learn how viruses worked in detail, and exactly what chemicals were needed
to thwart their reproductive cycle.[12]This section needs additional citations for verification. Please help improve this article by adding citations to reliable sources in this section needs additional citations for verification. Please help improve this article by adding citations to reliable sources in this section.
design is to identify viral proteins, or parts of proteins, or parts of proteins, that can be disabled.[11][13] These "targets" should generally be as unlike any proteins or parts of proteins in humans as possible, to reduce the likelihood of side effects and toxicity.[8] The targets should also be common across many strains of a virus, or even among different species of virus in
the same family, so a single drug will have broad effectiveness. For example, a researcher might target a critical enzyme synthesized by the virus, but not by the patient, that is common across strains, and see what can be done to interfere with its operation. Once targets are identified, candidate drugs can be selected, either from drugs already known
to have appropriate effects or by actually designing the candidate at the molecular level with a computer-aided design program. The target protein into bacteria or other kinds of cells. The cells are then cultured for mass
production of the protein, which can then be exposed to various treatment candidates and evaluated with "rapid screening" technologies. Viruses consist of a genome and sometimes called an 'envelope'). Viruses cannot reproduce
on their own and instead propagate by subjugating a host cell to produce copies of themselves, thus producing the next generation. Researchers working on such "rational drug design" strategies for developing antivirals have tried to attack viruses at every stage of their life cycles. Some species of mushrooms have been found to contain multiple
antiviral chemicals with similar synergistic effects. [14] Compounds isolated from fruiting bodies and filtrates of various mushrooms have broad-spectrum antiviral is a long way away. [15] Viral life cycles vary in their precise details depending on the type of
virus, but they all share a general pattern: Attachment to a host cell. Release of viral genes and possibly enzymes into the host cell. Replication of viral components into complete viral particles. Release of viral particles to infect new host cells. One antiviral strategy is to interfere with the ability of a
virus to infiltrate a target cell. The virus must go through a sequence of steps to do this, beginning with the virus "uncoating" inside the cell and releasing its contents. Viruses that have a lipid envelope must also fuse their envelope with the target cell, or with a
vesicle that transports them into the cell before they can uncoat. This stage of viral replication can be inhibited in two ways: Using agents which mimic the virus-associated protein (VAP) and bind to the cellular receptor, and anti-receptor antibodies. Using agents which
mimic the cellular receptor and bind to the VAP. This includes anti-VAP antibodies, receptor anti-idiotypic antibodies, extraneous receptor and synthetic receptor and synthetic receptor and synthetic receptor anti-idiotypic antibodies, extraneous receptor and synthetic receptor anti-idiotypic antibodies, extraneous receptor and synthetic receptor anti-idiotypic antibodies is partly trial and error, it can be a relatively slow process until
an adequate molecule is produced. Main article: Entry inhibitor very early stage of viral infection is viral entry, when the virus attaches to and enters the host cell. A number of "entry-inhibiting" or "entry-blocking" drugs are being developed to fight HIV. HIV most heavily targets a specific type of lymphocyte known as "helper T cells", and identifies
these target cells through T-cell surface receptors designated "CD4" and "CCR5". Attempts to interfere with the binding of HIV to the CCR5 receptor in hopes that it will be more effective. HIV infects a cell
through fusion with the cell membrane, which requires two different cellular molecular participants, CD4 and a chemokine receptor (differing depending on the cell type). Approaches to blocking this virus/cell fusion have shown some promise in preventing entry of the virus into a cell. At least one of these entry inhibitorsa biomimetic peptide called
Enfuvirtide, or the brand name Fuzeonhas received FDA approval and has been in use for some time. Potentially, one of the benefits from the use of an effective entry-inhibiting agent is that it potentially may not only prevent the spread of the virus within an infected individual but also the spread from an infected to an uninfected
individual. One possible advantage of the therapeutic approach of blocking viral entry (as opposed to the currently dominant approach of viral enzyme inhibition) is that it may prove more difficult for the virus to develop resistance to this therapy than for the virus to mutate or evolve its enzymatic protocols. Inhibitors of uncoating have also been
investigated.[16][17]Amantadine and rimantadine have been introduced to combat influenza. These agents act on penetration and uncoating process. This pocket is similar in most strains of rhinoviruses
monkeys, researchers reported in Nature Communications in 2016.[20]Rhinoviruses are the most common could; other viruses such as respiratory syncytial virus, parainfluenza virus and adenoviruses can cause them too.[21] Rhinoviruses are the most common cold; other viruses such as respiratory syncytial virus, parainfluenza virus and adenoviruses can cause them too.[21] Rhinoviruses are the most common cold; other viruses such as respiratory syncytial virus, parainfluenza virus and adenoviruses can cause them too.[21] Rhinoviruses are the most common cold; other viruses such as respiratory syncytial virus, parainfluenza virus and adenoviruses can cause them too.[21] Rhinoviruses are the most common cold; other viruses such as respiratory syncytial virus, parainfluenza virus and adenoviruses can cause them too.[21] Rhinoviruses are the most common cold; other viruses are the most cold; other viruses are the most cold; other viruses are the most cold; other
drift to the same degree that influenza viruses do. A mixture of 50 inactivated rhinovirus types should be able to stimulate neutralizing antibodies against all of them to some degree. [22] A second approach is to target the processes that synthesize virus components after a virus invades a cell. One way of doing this is to develop nucleotide or nucleoside
analogues that look like the building blocks of RNA or DNA, but deactivate the enzymes that synthesize the RNA or DNA, but deactivate the enzymes that synthesize the RNA or DNA, but deactivate the enzymes that synthesize the RNA or DNA, but deactivate the enzymes that synthesize the RNA or DNA, but deactivate the enzymes that synthesize the RNA or DNA, but deactivate the enzymes that synthesize the RNA or DNA, but deactivate the enzymes that synthesize the RNA or DNA, but deactivate the enzymes that synthesize the RNA or DNA, but deactivate the enzymes that synthesize the RNA or DNA, but deactivate the enzymes that synthesize the RNA or DNA once the analogue is incorporated.
nucleoside analogue, and is effective against herpesvirus infections. The first antiviral drug to be approved for treating HIV, zidovudine (AZT), is also a nucleoside analogue. An improved knowledge of the action of reverse transcriptase has led to better nucleoside analogues to treat HIV infections. One of these drugs, lamivudine, has been approved to
treat hepatitis B, which uses reverse transcriptase as part of its replication process. Researchers have gone further and developed inhibitors that do not look like nucleosides, but can still block reverse transcriptase. Another target being considered for HIV antivirals include RNase Hwhich is a component of reverse transcriptase that splits the
synthesized DNA from the original viral RNA. Another target is integrase, which integrate the synthesized DNA into the host cell, it then generates messenger RNA (mRNA) molecules that direct these original viral RNA. Another target is integrase, which integrates messenger RNA (mRNA) molecules that direct these original viral RNA. Another target is integrase, which integrates messenger RNA (mRNA) molecules that direct these original viral RNA.
synthesis of viral proteins. Production of mRNA is initiated by proteins known as transcription factors to viral DNA. Genomics has not only helped find targets for many antivirals, it has provided the basis for an entirely new type of drug, based on "antisense"
molecules. These are segments of DNA or RNA that are designed as complementary molecule to critical sections of these antisense segments to these target sections blocks the operation of those genomes, and the binding of these antisense drug named fomivirsen has been introduced, used to treat opportunistic eye
infections in AIDS patients caused by cytomegalovirus, and other antisense antivirals are in development. An antisense structural type that has proven especially valuable in research is morpholino antisense antivirals are in development. An antisense structural type that has proven especially valuable in research is morpholino antisense antivirals are in development. An antisense structural type that has proven especially valuable in research is morpholino antisense.
[24]dengue[25]HCV[26]coronaviruses[27]Yet another antiviral technique inspired by genomics is a set of drugs based on ribozymes are designed to cut RNA
and DNA at sites that will disable them. A ribozyme antiviral to deal with HIV.[29] An interesting variation of this idea is the use of genetically modified cells that can produce custom-tailored ribozymes. This is part of a broader effort to create genetically modified cells that can produce custom-tailored ribozyme antivirals are being developed to deal with HIV.[29] An interesting variation of this idea is the use of genetically modified cells that can produce custom-tailored ribozyme antivirals are being developed to deal with HIV.[29] and ribozyme antiviral to deal with hepatitis C has been suggested, [28] and ribozyme antiviral same being developed to deal with HIV.[29] and ribozyme antiviral same being developed to deal with hepatitis C has been suggested.
modified cells that can be injected into a host to attack pathogens by generating specialized proteins that block viral replication at various phases of the viral life cycle. Interference with post translational modifications or with targeting of viral proteins in the cell is also possible. [30] Some viruses include an enzyme known as a protease that cuts viral replication at various phases of the viral life cycle. Interference with post translational modifications or with targeting of viral proteins in the cell is also possible.
protein chains apart so they can be assembled into their final configuration. HIV includes a protease inhibitors became available in the 1990s and have proven effective, though they can have unusual side effects, for
example causing fat to build up in unusual places.[32] Improved protease inhibitors are now in development. Protease inhibitors have also been seen in nature. A protease inhibitor was isolated from the shiitake mushrooms (Lentinus edodes).[33] The presence of this may explain the Shiitake mushrooms in other antiviral activity in vitro.[34] Most viruses
produce long dsRNA helices during transcription. In contrast, uninfected mammalian cells generally produce dsRNA helices of fewer than 24 base pairs during transcription. DRACO (double-stranded RNA activated caspase oligomerizer) is a group of experimental antiviral drugs initially developed at the Massachusetts Institute of
Technology. In cell culture, DRACO was reported to have broad-spectrum efficacy against many infectious viruses, including dengue flavivirus, Amapari and Tacaribe arenavirus, Guama bunyavirus, H1N1 influenza and rhinovirus, and was additionally found effective against influenza in vivo in weanling mice. It was reported to induce rapid apoptosis
selectively in virus-infected mammalian cells, while leaving uninfected cells unharmed.[35] DRACO effects cell death via one of the last steps in the apoptosis signalling molecules simultaneously bind multiple procaspases. The procaspases transactivate via cleavage, activate additional
caspases in the cascade, and cleave a variety of cellular proteins, thereby killing the cell.[citation needed]Rifampicin acts at the assembly phase.[36]The final stage in the life cycle of a virus is the release of completed viruses from the host cell, and this step has also been targeted by antiviral drug developers. Two drugs named zanamivir (Relenza)
and oseltamivir (Tamiflu) that have been recently introduced to treat influenza prevent the release of flu viruses, and also seems to be constant across a wide range of flu strains. Main article: immunostimulantRather than attacking viruses directly, a second
category of tactics for fighting viruses involves encouraging the body's immune system to attack them. Some antivirals of this sort do not focus on a specific pathogen, instead stimulating the immune system to attack a range of pathogen, instead stimulating the immune system to attack a range of pathogen, instead stimulating the immune system to attack a range of pathogen, instead stimulating the immune system to attack a range of pathogen, instead stimulating the immune system to attack a range of pathogen.
form of human interferon named "interferon alpha" is well-established as part of the standard treatment for hepatitis B and C,[38] and other interferons are also being investigated as treatments for various diseases. A more specific approach is to synthesize antibodies, protein molecules that can bind to a pathogen and mark it for attack by other
elements of the immune system. Once researchers identify a particular target on the pathogen, they can synthesize quantities of identical "monoclonal" antibodies to link up that target. A monoclonal drug is now being sold to help fight respiratory syncytial virus in babies, [39] and antibodies purified from infected individuals are also used as a
treatment for hepatitis B.[40] Antiviral resistance can be defined by a decreased susceptibility to a drug caused by changes in viral genotypes. In cases of antiviral resistance, drugs have either diminished or no effectiveness against their target virus.[41] The issue inevitably remains a major obstacle to antiviral therapy as it has developed to almost all
specific and effective antimicrobials, including antiviral agents.[42]The Centers for Disease Control and Prevention (CDC) inclusively recommends anyone six months and older to get a yearly vaccination to protect them from influenza A viruses (H1N1) and (H3N2) and up to two influenza B viruses (depending on the vaccination).[41] Comprehensive
protection starts by ensuring vaccinations are current and complete. However, vaccines can be limited based on financial or locational reasons which can prevent the effectiveness of herd immunity, making effective
antivirals a necessity.[41]The three FDA-approved neuraminidase antiviral flu drugs available in the United States, recommended by the CDC, include: oseltamivir (Rapivab).[41] Influenza antiviral resistance often results from changes occurring in neuraminidase and hemagglutinin proteins on the viral
surface. Currently, neuraminidase inhibitors (NAIs) are the most frequently prescribed antivirals because they are effective against both influenza A and B. However, antiviral resistance is known to develop if mutations to the neuraminidase proteins prevent NAI binding.[43] This was seen in the H257Y mutation, which was responsible for oseltamivir
resistance to H1N1 strains in 2009.[41] The inability of NA inhibitors to bind to the virus allowed this strain of virus with additional antiviral drugs
including zanamivir. This finding was based on a performance evaluation of these drugs supposing the 2009 H1N1 'Swine Flu' neuraminidase (NA) were to acquire the oseltamivir-resistance (His274Tyr) mutation, which is currently widespread in seasonal H1N1 strains.[44]The genetic makeup of viruses is constantly changing, which can cause a virus
to become resistant to currently available treatments.[45] Viruses can become resistant through spontaneous or intermittent mechanisms throughout the course of an antiviral treatments. [41] Immunocompromised patients, more often than immunocompromised patients, more often than immunocompromised patients.
resistance during treatment. [41] Subsequent to exposure to someone else with the flu, those who received oseltamivir for "post-exposure prophylaxis" are also at higher risk of resistance. [46] The mechanisms for antiviral resistance development depend on the type of virus in question. RNA viruses such as hepatitis C and influenza A have high error
rates during genome replication because RNA polymerases lack proofreading activity. [47] RNA viruses also have small genome sizes that are typically less than 30 kb, which allow them to sustain a high frequency of mutations. [48] DNA viruses, such as HPV and herpesvirus, hijack host cell replication machinery, which gives them proofreading
capabilities during replication. DNA viruses are therefore less error prone, are generally less diverse, and are more slowly evolving than RNA viruses reproduce, which provides more opportunities for mutations to occur in successive replications. Billions of
viruses are produced every day during the course of an infection, with each replication giving another chance for mutations that encode for resistance to occur.[49] Multiple strains of one virus can be present in the body at one time, and some of these strains may contain mutations that cause antiviral resistance.[42] This effect, called the quasispecies
model, results in immense variation in any given sample of virus, and gives the opportunity for natural selection to favor viral strains with the highest fitness every time the virus is spread to a new host. [50] Recombination, the joining of two different viral variants, and reassortment, the swapping of virus among viruses in the same cell,
also play a role in resistance, especially in influenza. [48] Antiviral resistance has been reported in antivirals for herpes, HIV, hepatitis B and C, and influenza, but antiviral resistance is a possibility for all viruses. [42] Mechanisms of antiviral resistance vary between virus types. [citation needed] National and international surveillance is performed by the
CDC to determine effectiveness of the current FDA-approved antiviral flu drugs.[41] Public health officials use this information to make current recommendations about the use of flu antiviral medications. WHO further recommendations to make current recommendations to make current recommendations about the use of flu antiviral medications.
progression.[51] As novel treatments and detection techniques to antiviral resistance are enhanced so can the establishment of strategies to combat the inevitable emergence of antiviral resistance. [52] If a virus is not fully wiped out during a regimen of antiviral resistance.
there is a chance that a resistant strain may repopulate the host. [53] Viral treatment mechanisms must therefore account for the selection of resistant viruses is combination therapy, which uses multiple antivirals in one treatment regimen. This is thought to decrease the likelihood that
one mutation could cause antiviral resistance, as the antivirals in the cocktail target different stages of the viral life cycle. [54] This is frequently used in retroviruses like HIV, but a number of studies have demonstrated its effectiveness against influenza A, as well. [55] Viruses can also be screened for resistance to drugs before treatment is started.
This minimizes exposure to unnecessary antivirals and ensures that an effective medication is being used. This may improve patient outcomes and could help detect new resistance mutations during routine scanning for known mutants. [53] However, this has not been consistently implemented in treatment facilities at this time. The term Direct-acting
antivirals (DAA) has long been associated with the combination of antiviral drugs used to treat hepatitis C are taken orally, as tablets, for 8 to 12 weeks. [56] The treatment
depends on the type or types (genotypes) of hepatitis C virus that are causing the infection.[57] Both during and at the end of treatment, blood tests are used include:[58]Harvoni (sofosbuvir and ledipasvir)Epclusa (sofosbuvir and
velpatasvir)Vosevi (sofosbuvir, velpatasvir, and voxilaprevir)Zepatier (elbasvir and grazoprevir)Mavyret (glecaprevir and pibrentasvir)The United States Food and Drug Administration approved DAAs on the basis of a surrogate endpoint called sustained virological response (SVR).[59] SVR is achieved in a patient when hepatitis C virus RNA remains
undetectable 1224 weeks after treatment ends. [60][61] Whether through DAAs or older interferon-based regimens, SVR is associated with improved health outcomes and significantly decreased mortality. [62][63][64] For those who already have advanced liver disease (including hepatocellular carcinoma), however, the benefits of achieving SVR may
be less pronounced, though still substantial.[64]Despite its historical roots in hepatitis C research, the term "direct-acting antivirals" is becoming more broadly used to also include other anti-viral drugs with a direct viral target such as aciclovir (against herpes simplex virus), letermovir (against cytomegalovirus), or AZT (against human
immunodeficiency virus). In this context it serves to distinguish these drugs from those with an indirect mechanism of action such as immune modulators like interferon alfa. This difference is of particular relevance for potential drug resistance mutation development. [65] Guidelines regarding viral diagnoses and treatments change frequently and limit
quality care.[66] Even when physicians diagnose older patients with influenza, use of antiviral treatment can be low.[67] Provider knowledge of antiviral therapies can improve patient care, especially in geriatric medicine. Furthermore, in local health departments (LHDs) with access to antivirals, guidelines may be unclear, causing delays in
treatment.[68] With time-sensitive therapies, delays could lead to lack of treatment. Overall, national guidelines, regarding infection control and management, standardize care and improve healthcare worker and patient safety. Guidelines, regarding infection control and management, standardize care and improve healthcare worker and patient safety.
caused by the H1N1 virus, recommend, among other things, antiviral treatment regimens, clinical assessment algorithms for coordination of care, and antiviral treatment regimens, clinical assessment algorithms for coordination of care, and antiviral treatment regimens, clinical assessment algorithms for coordination of care, and antiviral treatment regimens, clinical assessment algorithms for coordination of care, and antiviral treatment regimens, clinical assessment algorithms for coordination of care, and antiviral treatment regimens, clinical assessment algorithms for coordination of care, and antiviral treatment regimens, clinical assessment algorithms for coordination of care, and antiviral treatment regimens, clinical assessment algorithms for coordination of care, and antiviral treatment regimens, clinical assessment algorithms for coordination of care, and antiviral treatment regimens, clinical assessment algorithms for coordination of care, and antiviral treatment regimens, clinical assessment algorithms for coordination of care, and antiviral treatment regimens, clinical assessment algorithms for coordination of care, and antiviral treatment regimens, clinical assessment algorithms for coordination of care, and antiviral treatment regimens, and antiviral treatment regimens regimens, and antiviral treatment regimens regimens.
Strategic National Stockpile Public Health Emergency Preparedness initiatives are managed by the CDC via the Office of Public Health Preparedness and Response. [71] Funds aim to support communities in preparing for public Health Preparedness and Response.
consists of bulk quantities of medicines and supplies for use during such emergencies. [72] Antiviral stockpiles prepare for shortages of antiviral medications in cases of public health departments was unclear, revealing gaps in antiviral planning. [68] For
example, local health departments that received antivirals from the SNS did not have transparent guidance on the use of the treatments. The gap made it difficult to create plans and policies for their use and future availabilities, causing delays in treatments. The gap made it difficult to create plans and policies for their use and future availabilities, causing delays in treatments.
development of CCR5 receptor antagonists (for HIV)Monoclonal antibodyList of antiviral drugsAntiprion drugs and AstemizoleDiscovery and development of NS5A inhibitorsCOVID-19 drug repurposing research Antiviral agents. 2012. PMID31643973. Rossignol JF (2014). "Nitazoxanide: a first-in-class broad-spectrum antiviral agent". Antiviral Res
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