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Mecanismo de acción aines pdf



INHIBIDORES DE LA ALFA GLUCOSIDASA

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CLASIFICACIÓN	PROPIEDADES	MECANISMO DE ACCIÓN	FARMACOCINÉTICA	ADVERSIDADES	APLICACIÓN
INSULINO					
ACARBOSA	Vida media: 2 h	metabolismo: -	excreción: renal	preparados: tablet 25, 50, 100 mg	Malabsorción, diarrea
INSULINO T					
MIGLITOL	Vida media: 2 H	metabolismo: -	excreción: renal	Tablet 25, 50, 100mg	Malabsorción, diarrea

INHIBIDORES DE LA ALFA GLUCOSIDASA

INCRETINAS

- Malabsorción
- Meteorismo
- Flatulencias
- Diarrea

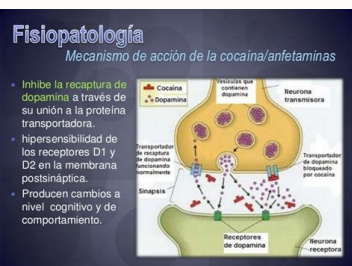
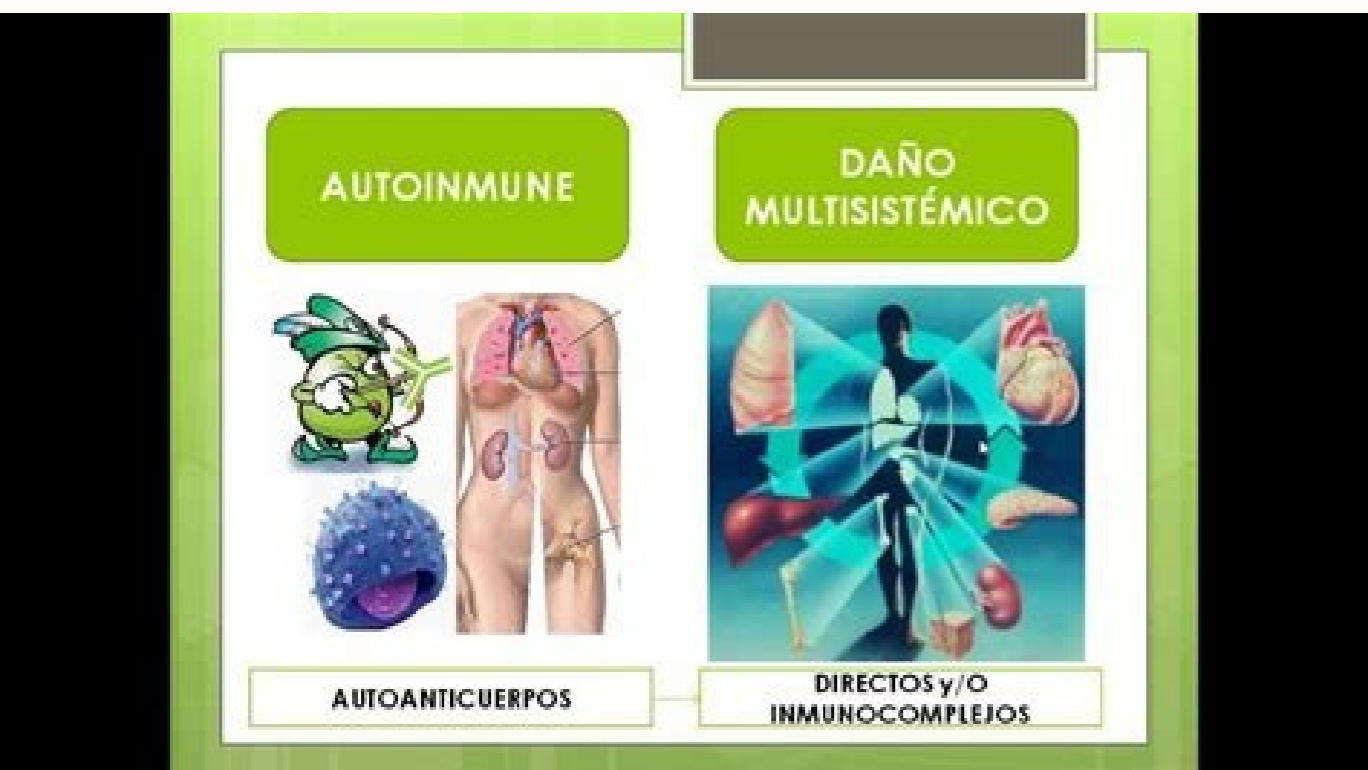
DrugBank, open data drug and target database, disponible en: <http://drugbank.ca/>

AINEs- Clasificación s/grupo químico

SALICILATOS

- ÁCIDO ACETILSALICÍLICO
- ACETILSALICILATO DE LISINA
- SALICILATO DE SODIO
- **PARAMINOFENOL**
- ACETAMINOFENO O PARACETAMOL
- **PIRAZOLONAS**
- DIPIRONA
- FENILBUTAZONA (RETIRADA)
- AZAPROPAZONA

- Paracetamol y dipirona NO son AINEs: son anagésicos-antipiréticos pero NO
- antinflamatorios



Mecanismo de acción de los aines en veterinaria. Farmacos aines mecanismo de acción. Mecanismo de acción de los aines en la mucosa gástrica. Mecanismo de acción de los aines selectivos. Mecanismo de acción de los aines farmacología. Aines y su mecanismo de acción. Mecanismo de acción de los aines. Mecanismo de acción del aines.

Its concentration, although it may be moderate increases, remains almost unchanged over time. In contrast, the COX-2 is almost undetectable in normal tissues. The inhibition of prostaglandin synthesis may result in reversible alteration of glomerular filtering, acute renal insufficiency and even cases of prolonged and constant consumption, papillary necrosis and chronic renal insufficiency. Classification whenever the classification of any pharmacological group is attempted, it is possible to the action. The hematological is less frequent than the previous ones, highlighting the possibility of producing agranulocytosis and applying anemia, phenomena in which the intervention of immunological mechanisms would be required. Singular exceptions of such selective famess are: Nebumetone, Nimesulide and Meloxicam that show a certain affinity, although not absolutely by the COX-2. Among the first NSAIDs marketed with a selective antagonist profile of COX-2 is the coxibs group called (Celecoxib, Tetroxib, Valdecobix and Loxepib). The first is a physiological form found in a usual way in most body and tissues of the body, but especially in the rivales, gastrointestinal tract, monomers/macrao. This increase in hypothalamic PGE2 causes the body temperature control point to grow. In contrast of what you think, the administration of this group of fessions is not exempt from risks, since this enzyme also constitutive in the Riño Lulus, rebro and reproductive system and play an important role in the Of some tissues, in addition to acting, as well as protector of the pharmacy of the mucosa. Antipyria and anti-inflammatory. The analgarral analgal activity of the Mother/Moderate/Moderate intensity (they never correspond to anti-limit potential to analgaining opionins) and occurs as well as the perifada. It is estimated that in our Paas it will be consumed annually of 5 million people, of which 30-40% are over 65 years old. They are derived only of arilmethylsulfonyamide and structurally similarly similar to pyrazolone derivatives. To evaluate the anti-stimulating effectiveness of a particular NSAIE, it is necessary that the inhibition of inflammation is variable, depending on its spectrum of a, of the type of inflammation process. Its inhibition of cyclooxygenase and its cushioning action of other cellular responses independent of its anti-Cox. Prostaglandins and thromboxans block the activities of vasodilators and chemotized, interfering in the early phase of inflammation. Despite the quamic and structural differences between them, they share the anti-inflammatory and anti-arthritic properties to a different degree, which makes them, along with the opioids, one of the basic pillars of the analgistic treatment current. redness, tumor (swelling), pain and functional disability, are the 5 attributes that define an inflammation process. This makes it its concentration can increase between ten and almost hundred times. All blacks are their mechanism of common action: most of them inhibiting stereosely and competitive bicoxygenase enzyme (COX). Its main advantages are: the possibility of allowing a dose and being less harmful to gmittage, although it has been described that it has a greater incidence of diar and other undesirable digestive effects. Indolavados Indo- © Tica (acemetacin, indomethacin, proglumetacin, sulindaco, tometá) is silently related to the previous group (in fact, it can be considered that they are a quinnic subgroup of the previous ones). Their activity on the smooth mothers of certain organs makes them a good option for the treatment of menstrual pain. Generally, it is the NSAE of choice in these incipient or predominantly painful paintings. Naproxen, while having a somewhat unfavorable toleration profile, has the advantage of being much more powerful and allowing an administration guideline twice. It is an "undoubted" way, that is, it seems to manifest itself in some squads, releasing inflammatory or undoubted prostaglandins in response to certain harmful stages that lead to the appearance of inflammation mechanisms and Ao amplification of the ears painful sea pain in inflamed inflamed inflamed. Hypersensibilities knew their sharp ability to produce alsmal and pseudo ricogic reactions (similar to previous ones, but without intervening immunological mechanisms). At this point, it should be noted that the NSAIDs do not exert no action if the body temperature is normal physiology (they do not cause hypothermia to conventional therapeutic doses). onipris The lifestyle is inhibited by this group of FNES) allowed to assign other therapies of major interiors and closing transcendent. It is expressed by the release of essential prostaglandins (the constitutive so-called) that perform their physiological and homeostic functions (Gastric Mucosa Protection, Riño N, Platelets, Endota vascular lio). Lysine acetylyclate (solid salt micronized from acetylsalicylic) has an absorbent absorption, so they will be especially indicated in the treatment of acute pain. While the first inhibits cyclooxygenase in a reversible and reversible manner, the last one does so in a selective and irreversible manner. What powerful is the inhibition of the Cox, the best anti-inflammatory effect. Its use is marked by a high incidence of gastrointestinal adverse effects (bass cams) and central (very intense headaches, life, mental confusing, unpleasant). The rest of the active principles that compose this group are structural -watery that they seek, without achieving it, a similar or larger therapy action, but with a lower incidence of adverse effects. But NSAIDs also interfere with the vain van of neutrals, the most abundant squads that played a more relevant role in the acute phases of inflammatory processes. Santesis of interleucine-1 by macrons and this, this, this) in turn, stimulates the production of prostaglandins (especially PGE2) in the thermoregulatory center of hypothal. Therefore, it will be simply indicated that most manifestations of the inflammation response involves the activation of cyclooxygenase (COX), an enzyme complex that catalyzes the of the Araquahidic entity in an endoperion of the cyclical, from which different synthesized and icicidin (prostaglandins, prostacyclines and thromboxanos) who actively participate in the galsis, amplification and development of the inflammation process. Most tissues producing several action. Hepalicoschomitant to the use of most NSAIDs described a moderate and asymptomatic high transaminases during the first weeks of treatment. Its ability to selectively interact with COX-2, without inhibiting COX-1 ISOFRA, makes the recommended pharmacological doses a smaller and lower of its adverse effects (severe gastrointestinal toxicity and hemorrhagic risk). Paraminofenolesomo paracetamol or fenazoopiridine. Its use may lead to the emergence of serious side effects: ulcerogenicity, myelotoxicity and ability to produce blood dyscrasias. Thus, one of the most notable action á 6 hydling Á ours Anti-Governor/Antitromba Acetylsalic capacity when administered to low doses (inferior ones recommended as analitics or anti-inflammatory). Thus, in response to this criterion, the following groups can be found: salicylates, pirazolindodiones, arlicais derivatives, indolacacic derivatives, armory derivatives, oxycaemes and phenamaes. The intensity of its anti-inflammatory effect, its considering antipathic effect and an acceptable security profile justify this choice. Another important substance is Nebumetanone, a deeper and deeper than, after being absorbed in the intestine, gives rise to an active metabon Similar to naproxene (differs from the radical is the acronym instead of prone). The pure salicylates are a good anti-anti-anti-anti-inflammatory anti-anti-anti-anti-antiphalar profile. From a physiological perspective, it can be considered that the inflammation is a defensive orange reaction that the body triggers to try to isolate an internal or external agent, which is producing a host aggressive Subsequently try to destroy it. It would be very ambitious to try to exhibit in detail in some pages, the complexity of the intricate concatenation of the feather inherent in the appearance of an inflammation process. Some of the most commonly described symptoms are: eruptions cutting, erythema, angioedema, photodermatitis, rhinitis, bronchy asthma or diar. This aniline derivatives fatania has a sharp analgistic activity in the face of mild to moderate and anti-piendous intensity pain, of efficiency comparable to the acetylsalic hand; However, it has little or no anti-inflammatory activity. But without dan, the most worrying is its ability to hurt the gister and duodenal mucosa, causing erosion and wax that can end in complications such as bleeding and perforations. Renal prostaglandins play a very important role in the correct functionality of the ri ± Á ± ives. Salicilatosro of this group AINE is necessary to make an important subdivision: pure salicylates (salicidia, salicilamide, phosphate, difimil) and acetylsalicilates (acetylsalicamic zealics, acetylsalicilates of lysine, benoryliato, fosphan. Although it is not exempt from the characteristic side effects of the group, it has a very long half-life and allows for daily intenses, which makes it a cycle and a good option for the treatment of condition Chronicle The From these complications, they are academizes its new molecules. The effectiveness of NSAIDs applied by life was repeatedly placed, attributing the observed benefit of the massage produced in the area during its application. This is especially true in its chronic use; Therefore, its continued use should be reserved for patients who are most aware of the risk of developing analgest psycho components, improving their action and minimizing gastrointestinal effects. Its use is very limited, as they are not significant advantages over other NSAIDs; however, they are a high incidence of diarrontity. Selective infections of the Cyclooxygenase 2un group that deserves separate treatment is selective cyclooxygenase inhibitors, so captured, cyclooxygenase 2, the coxibs so called (Celecoxib, Robecoxib). Adverse effects on other systems and interactions are similar to other NSAIDs and should be considered. The ketoprofen and its active enantier, dextetoprofen, so the equal lower dose properties. Moderate antipada activity and almost missing. Tolumen © perhaps the lower side effects (although toxicity differences with indomethacin are not very important) and is the most used, acetylsalicicics in the treatment of Youth Chronicle Arthritis. Á ÁO TIAPROFAMIC, Butfune, Dextcetepofen, Fenbufen, Fenopropfen, Flurbipropfen, ibuprofen, ibuprofen, ibuprofen Naproxen, Ketorolaco) are less powerful than the clitic nanas, but compared to them, they are a better security profile. The use of celecoxib or Robecoxib can be considered in patients in which one or more of these circumstances agree: push or gastrointestinal bleeding. 65 years, in particular if other risk factors agree. It has fewer gastrolar effects than acetylsalic is, but it does not have its antiangianna capacity. Something similar occurs with salicilalic salsalate or ©), which is hydrolyzed in the small intestine, releasing two © analgá © is. Its different action mechanism about prostaglandins and Cox makes this finger treated with oral anticoagulants and people alleged salicylates or other NSAIDs. The most notable adverse effect of this Famaman is its hepatotoxicity, which can be presented with the higher than 6 g/day doses and this has increased significantly in a rist albumo. The anti-inflammatory are not probably the most frequently prescribed Pharamacos group in the world. Similarly, the stability of the active moll and its ability to penetrate Keep independent of pH or acidic environment. The applicativeness of NSAIDs applied by life was repeatedly placed, attributing the observed benefit of the massage produced in the area during its application. This is especially true in its chronic use; Therefore, its continued use should be reserved for patients who are most aware of the risk of developing analgest psycho components, improving their action and minimizing gastrointestinal effects. 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Facilitating the good penetration cutting (they are usually liphi kids, though with a certain residual hydrophilia). Diprofen, the first representative of this group, is a very secure Famcaus, however, it has achieved a more pink and prolonged effect of the analgast and anti-inflammatory type. Its adverse effects are more durable. Another notable effect is the action of Uricians, which makes almost all high doses woven in the process where one of the most prescribed Fnnamacos groups in the world, they are accompanied By adverse effects, some of them serious, when acting in several parts of the organism. The cox mediated by the nsais allows you to return the bodily technical control point at its physiological value through the activation of mechanisms that favor the elimination of gastrointestinal heat are frequent are minor effects type pyrosis, gastrics, gastricia, daytime daily ia or stipe. However, cyclooxygenase is an ubiquitous enzyme, present in almost all of the centers, and its necessary medais in the production of certain prostaglandins not-inflammation that will allow the proper operation to be permitted of many s and systems. A peculiarity of each moll is not usually dependent on the dose and no relationship with the anti-inflammatory of the problem derived from the use of this type of Fnnamacos is that the vast majority of NSAIDs today In the unforgettable action of the market, that is, they inhibit the enzyme activity of both isoforms or, to any case, to a greater extent to COX-1, which gives rise to the appearance of undesirable effects á € Characteristics of this group of this fan group. Indomethacin is one of the NSAIDs with the largest profile of effectiveness (Comparison to the powerful of this group), however, its analgast and anti-panic power is moderate. It is usually produced in elderly patients, with compromised renal function and prolonged use of high doses of anti-inflammatory. This is why NSAIDs are especially effective in all pains associated with inflammatory processes and other situations in which prostaglandins participate in a highlight and pain-traumatic) since they act, with precision o, reducing the effect of hyperalgide of prostata. Each moll, usually does not depend on the dose and not maintain no relationship with anti-inflammatory capacity. Meloxicam and Nimesulide, while selective COX-2 inhibitors, this property is not complete; Therefore, it has not been shown that they are long in the long run than the rest of the NSAIDs. Mephanims, exclusively) as a significant analgast power and antiphalar power, but almost does not have anti-inflammatory effect. The inhibition of the COX mediated by NSAIDs allows us to return the body control point to its physiological value, activating mechanisms that favor the elimination of heat (vasodilation, sweating). Recent studies show that this appreciation is not correct and that the application of this type of preparation is effective in the treatment of muscle pain and articulations. Be able to be effective due to sica in these cases, options that allow a localized, thry and minimize adverse reactions and allow the use of powerful therapy agents. By the constrain, the acetyled derivatives, depending on the dosage used, all the range of therapy effects commented above. The active substance of this group is acetylsalicamic is. Prostacycline causes vasodilation, inhibiting plaquetter and fibrania. While thromboxanos induce vasoconstriction and plaquetter aggregation. A very important role in the response and inflammation. Thus, the privilege prostaglandins are responsible for the appearance of pain, vasodilation and inflammation. inflammation.

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