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Extended tier onlySulfuric acid is synthesised by the Contact processConcentrated sulfuric acid is used in car batteries, making fertilisers, soaps and detergentsStage 1The first stage is the production of sulfur dioxideThe oxygen for this stage is obtained from the airThe sulfur for this stage is obtained by:Burning sulfur to oxidise the sulfursulfur + oxygen sulfur dioxideS + O2 SO2metal sulfide + oxygen metal oxide + sulfur dioxideStage 2The main stage is the oxidation of sulfur dioxide to sulfur trioxide using a vanadium(V) oxide, V2O5,catalystSulfur dioxide + oxygen sulfur trioxide2SO2+ O2 2SO3The oxygen used in this stage is obtained from airThe conditions for this main stage of production are:A temperature of 450 Ca pressure of 2 atm (200 kPa)Once sulfur trioxide is formed, it undergoes more processes to produce sulfuric acidStage 3The sulfur trioxide is absorbed into a solution of 98% sulphuric acid to produce a thick liquid caldeoleum (disulfuric acid):SO3+ H2SO4 H2SO7Sulfur trioxide + sulfuric acid disulfuric acid is not absorbed into water because a fine mist of sulfuric acid would be produced and this would be difficult to condense and is also highly dangerousOleum is added to water to form concentrated sulfuric acid:H2O + H2SO7 2H2SO4water + disulfuric acid sulfuric acidPage 2Exam code: 0620 & 0971Dio this video help you?Extended tier

onlyAmmonia is manufactured in an exothermic reaction called the Haber process which occurs in five stages:Stage 1H2 is obtained from methaneN2 is obtained from the airThey are pumped into the compressor through a pipeStage 2Inside the compressor, the gases are compressed to around 200 000 kPa or 200 atmospheresStage 3The pressurised gases are pumped into a reactor containing layers of an iron catalyst at a temperature of 450 CSome of the hydrogen and nitrogen react to form ammonia N2 (g) + 3H2 (g) 2NH3 (g)Stage 4Unreacted H2 and N2 and the ammonia product pass into a cooling tankThe ammonia is liquified and removed to pressurised storage vesselsStage 5The unreacted H2 and N2 gases are recycled back into the system and start over againThe production of ammonia by the Haber processAmmonia is produced during the Haber Process. The reaction is summarised in the diagram below.Name gas A.Name catalyst B used and state why it is used?Answer:Gas A is hydrogen / H2.Catalyst B is iron, which is used to speed up the reaction / increase the rate of reaction.Examiners comment that students often know That nitrogen and hydrogen are needed for the Haber processWhat the purpose of the catalyst isBut, That students do not know where the nitrogen and hydrogen come fromKnow the name of the catalyst and confuse it with other catalysts like 'vanadium oxide' and 'enzymes'.You should know where the nitrogen and hydrogen come from as well as the conditions of the Haber process:Pressure = 200 atmospheresTemperature = 450oCatalyst = ironExtended tier onlyChemists' knowledge of the energy changes and factors affecting reaction rates can be used to predict the best possible conditions to make the most ammonia in the fastest possible timeHowever, sometimes those conditions are contradictory and choices have to be made between factors that improve the yield of ammonia and those that speed up the reactionThe graph below illustrates the effects of changing temperature and pressure on the yield of ammonia obtainedThe yield of ammonia produced changes with changes made to temperature and pressureFrom the graph:As the pressure increases, the percentage yield increasesThis is shown by following any of the curved linesAs the temperature decreases, the percentage yield increasesThis is shown by following any vertical line upwards from the x-axisThat the actual conditions used must be chosen depending on a number of economical, chemical and practical considerations.Like all industries, companies that manufacture and sell chemical goods do so to make a profitPart of the industrial process is the economic decision on how and where to design and implement a manufacturing siteThe availability and cost of raw materials is a major consideration which must be studied before any decisions are made at the Haber Process raw materials are readily available and inexpensive to purify.Nitrogen from theairHydrogen from theairFossil fuelsThe cost of extraction of raw materials is too high or they are unavailable then the process is no longer economically viableMany industrial processes require huge amounts ofheatandpressurewhich is very expensive to maintainProduction energy costs are also a factor to be considered carefully and alongside the raw materials issueTemperature: 450oCHigh temperature favours the reverse reaction as it is endothermicSo, a higher yield of reactants will be madeLow temperature favours the forward reaction as it is exothermicSo, a higher yield of products will be madeHowever, at low temperature the rate of reaction is very slow, 450 C is a compromise temperature between having a lower yield of products but being made more quicklyPressure: 200 atmLow pressure favours the reverse reaction as there are more moles of gaseous reactantSo, a higher yield of reactants will be madeHigh pressure favours the forward reaction as there are fewer moles of gaseous productSo, a higher yield of products will be madeHowever, high pressure can be dangerous and very expensive equipment is neededSo, 200 atmospheres is a compromise pressure between a lower yield of products being made safely and economicallyCatalyst:IronThe presence of a catalyst doesnaffect the position of equilibrium but it does increase therateat which equilibrium is reachedThis is because the catalyst increases the rate ofboththe forward and backward reactions by the same amount (by providing an alternative pathway requiringlower activation energy)As a result, theconcentrationof reactants and products is nevertheless thesameat equilibrium as it would be without the catalyst.So a catalyst is used as it helps the reaction reach equilibrium quickerIt allows for an acceptable yield to be achieved at lower temperatureby lowering the activation energy requiredWithout the process would have to be carried out at an evenhigher temperature, increasingcostsand decreasingyieldsThe higher temperaturedecomposesome of the NH3moleculesThe reaction conditions chosen for the Haber process are not ideal in terms of the yield but do provide balance between product yield, reaction rate and price costThe process is calleda compromise because they have chosen the yield rate and costDid this page help you?Sulfuric acid is manufactured by the following three steps, which involve the following three reactions on burning or roasting in air: Sulfur or sulf

Constantin dynasty and the House of Anjou are relying primarily on mercenaries in their militaries. Paid soldiers are available year-round, unlike knights who expected certain periods off to maintain their manor lifestyles.18]In India, Hoysala architecture reaches its peak in the Middle East, the icon of Theotokos of Vladimir is painted probably in Constantinople. Everything but the faces will later be retouched, and the icon will go to the Tretyakov Gallery of Moscow.The Georgian poet Shota Rustaveli composes his epic poem The Knight in the Panther's Skin.Shahab al-Din Suhrawardi formuls his "scholch of illumination". In North Africa, the kasbah of Marrakesh is built, including the city gate Bab Agnaou and the Koutoubia mosque. In sub-Saharan Africa, Kente cloth is first woven. In France, the first piedfort coins were minted.The city of Tula burns down, marking the end of the Toltec Empire.In West Africa the Ife Empire is established.See also: Timeline of historic inventions 12th century1104: The Venice Arsenal of Venice, Italy, is founded. It employed some 16,000 people for the mass production of sailing ships in large assembly lines, hundreds of years before the Industrial Revolution.1106: Finished building of Gelati.1107: The Chinese engineer Wu Deren combines the mechanical compass vehicle of the south-pointing chariot with the distance-measuring odometer device.1111: The Chinese Donglin Academy is founded.1165: The Lühe Pagoda of Hangzhou, China, is built.1170: The Roman Catholic notion of Purgatory is defined.[9]1185: First record of windmills.Wikimedia Commons has media related to 12th century.
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Please help to improve this article by introducing more precise citations. (May 2020) (Learn how and when to remove this message)The contact process is a method of producing sulfuric acid in the high concentrations needed for industrial processes. Platinum was originally used as the catalyst for this reaction; however, because it is susceptible to reacting with arsenic impurities in the sulfur feedstock, vanadium(V) oxide (V2O5) has since been preferred.[1]This process was patented in 1831 by British vinegar merchant Peregrine Phillips.[2][3][4] In addition to being a far more economical process for producing concentrated sulfuric acid than the previous lead chamber process, the contact process also produces sulfur trioxide and oleum. In 1890 John Brown Francis Herreshoff developed a form of the contact catalytic process for the company of which he was a partner.[5]In 1901 Eugen de Han patented the basic process involving combining sulfur dioxide and oxygen in the presence of vanadium oxides, producing sulfur trioxide which was easily absorbed into water, producing sulfuric acid.[6] This process was improved remarkably by shrinking the particle size of the catalyst (e.g. 5000 microns), a process discovered by two chemists of BASF in 1914.[7][8][9]The process can be divided into four stages:Combining of sulfur and oxygen (O2) to form sulfur dioxide, then purify the sulfur dioxide in a purification unitAdding an excess of oxygen to sulfur dioxide in the presence of the catalyst vanadium pentoxide at 450C and 1-2atmThe sulfur trioxide formed is added to sulfuric acid which gives rise to oleum (disulfuric acid)The oleum is then added to water to form sulfuric acid which is very concentrated. Since this process is an exothermic reaction, the reaction temperature should be as low as possible.Purification of the air and sulfur dioxide (SO2) is necessary to avoid catalyst poisoning (i.e. removing catalytic activities). The gas is then washed with water and dried with sulfuric acid.To conserve energy, the mixture is heated by exhaust gases from the catalytic converter by heat exchangers.Sulfur dioxide and dioxygen then react as follows:2 SO2(g) + O2(g) = 2 SO3(g); H = -197kJ/molAccording to the Le Chatelier's principle, a lower temperature should be used to shift the chemical equilibrium towards the right, hence increasing the percentage yield. However too low of a temperature will lower the formation rate to an uneconomical level. Hence to increase the reaction rate, high temperatures (450C), medium pressures (1-2atm), and vanadium(V) oxide (V2O5) are used to ensure an adequate (>95%) conversion. The catalyst only serves to increase the rate of reaction as it does not change the position of the thermodynamic equilibrium. The mechanism for the action of the catalyst comprises two steps:Oxidation of SO2 into SO3 by V5+→2SO2 + 4V5+ → 2O2 2SO3 + 4V4+→Oxidation of V4+ back into V5+ by dioxygen (catalyst regeneration):4V4+ + O2 4V5+ + 2O2Hf sulfur trioxide passes through the heat exchanger and is dissolved in concentrated H2SO4 in the absorption tower to form oleum.H2SO4 + SO3 H2S2O7Note that directly dissolving SO3 in water is impractical due to the highly exothermic nature of the reaction. Acidic vapor or mists are formed instead of a liquid.Oleum is reacted with water to form concentrated H2SO4.H2S2O7 + H2O 2 H2SO4This includes the dusting tower, cooling pipes, scrubbers, drying tower, arsenic purifier and testing box. Sulfur dioxide has many impurities such as vapours, dust particles and arsenous oxide. Therefore, it must be purified to avoid catalyst poisoning (i.e.: destroying catalytic activity and loss of efficiency). In this process, the gas is washed with water, and dried by sulfuric acid. In the dusting tower, the sulfur dioxide is exposed to a steam which removes the dust particles. After the gas is cooled, the sulfur dioxide enters the washing tower where it is sprayed by water to remove any soluble impurities. In the drying tower, sulfuric acid is sprayed on the gas to remove the moisture from it. Finally, the arsenic oxide is removed when the gas is exposed to ferric hydroxide.The next step to the contact process is double contact double absorption (DCDA). In this process the product gases (SO2) and (SO3) are passed through absorption towers twice to achieve further absorption and conversion of SO2 to SO3 and production of higher grade sulfuric acid.SO2-rich gases enter the catalytic converter, usually a tower with multiple catalyst beds, and are converted to SO3, achieving the first stage of conversion. The exit gases from this stage contain both SO2 and SO3 which are passed through intermediate absorption towers where sulfuric acid is trickled down packed columns and SO3 reacts with water increasing the sulfuric acid concentration. Though SO2 too passes through the tower it is unreactive and comes out of the absorption tower.This stream of gas containing SO2, after necessary cooling is passed through the catalytic converter bed column again achieving up to 99.8% conversion of SO2 to SO3 and the gases are again passed through the final absorption column thus achieving not only high conversion efficiency for SO2, but also enabling production of a higher concentration of sulfuric acid.The industrial production of sulfuric acid involves proper control of temperatures and flow rates of the gases as both the conversion efficiency and absorption are dependent on these.
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Watch NowA brief summary of the Contact Process The Contact Process: makes sulphur dioxide; converts the sulphur dioxide into sulphur trioxide (the reversible reaction at the heart of the process); converts the sulphur trioxide into concentrated sulphuric acid. Making the sulphur dioxide This can either be made by burning sulphur in an excess of air: . . . or by heating sulphide ores like pyrite in an excess of air: In either case, an excess of air is used so that the sulphur dioxide produced is already mixed with oxygen for the next stage. Converting the sulphur dioxide into sulphur trioxide This is a reversible reaction, and the formation of the sulphur trioxide is exothermic. A flow scheme for this part of the process looks like this: The reasons for all these conditions will be explored in detail further down the page. Converting the sulphur trioxide into sulphuric acid This can't be done by simply adding water to the sulphur trioxide - the reaction is so uncontrollable that it creates a fog of sulphuric acid. Instead, the sulphur trioxide is first dissolved in concentrated sulphuric acid: The product is known as fuming sulphuric acid or oleum. This can then be reacted safely with water to produce concentrated sulphuric acid: twice as much as you originally used to make the fuming sulphuric acid. Explaining the conditions The proportions of sulphur dioxide and oxygen The mixture of sulphur dioxide and oxygen going into the reactor is in equal proportions by volume. Avogadro's Law says that equal volumes of gases at the same temperature and pressure contain equal numbers of molecules. That means that the gases are going into the reactor in the ratio of 1 molecule of sulphur dioxide to 1 of oxygen. That is an excess of oxygen relative to the proportions demanded by the equation. According to Le Chatelier's Principle, increasing the concentration of oxygen in the mixture causes the position of equilibrium to shift towards the right. Since the oxygen comes from the air, this is a very cheap way of increasing the conversion of sulphur dioxide into sulphur trioxide. Why not use an even higher proportion of oxygen? This is easy to see if you take an extreme case. Suppose you have a million molecules of oxygen to every molecule of sulphur dioxide. The equilibrium is going to be tipped very strongly towards sulphur trioxide - virtually every molecule of sulphur dioxide will be converted into sulphur trioxide. Great! But you aren't going to produce much sulphur trioxide every day. The vast majority of what you are passing over the catalyst is oxygen which has nothing to react with. By increasing the proportion of oxygen you can increase the percentage of the sulphur dioxide converted, but at the same time decrease the total amount of sulphur trioxide made each day. The 1 : 1 mixture turns out to give you the best possible overall yield of sulphur trioxide. The temperature Equilibrium considerations You need to shift the position of the equilibrium as far as possible to the right in order to produce the maximum possible amount of sulphur trioxide in the equilibrium mixture. The forward reaction (the production of sulphur trioxide) is exothermic. According to Le Chatelier's Principle, this will be favoured if you lower the temperature. The system will respond by moving the position of equilibrium to counteract this - in other words by producing more heat. In order to get as much sulphur trioxide as possible in the equilibrium mixture, you need as low a temperature as possible. However, 400 - 450C isn't a low temperature! Rate considerations The lower the temperature you use, the slower the reaction becomes. A manufacturer is trying to produce as much sulphur trioxide as possible per day. It makes no sense to try to achieve an equilibrium mixture which contains a very high proportion of sulphur trioxide if it takes several years for the reaction to reach that equilibrium. You need the gases to reach equilibrium within the very short time that they will be in contact with the catalyst in the reactor. The compromise 400 - 450C is a compromise temperature producing a fairly high proportion of sulphur trioxide in the equilibrium mixture, but in a very short time The pressure Equilibrium considerations Notice that there are 3 molecules on the left-hand side of the equation, but only 2 on the right. According to Le Chatelier's Principle, if you increase the pressure the system will respond by favouring the reaction which produces fewer molecules. That will cause the pressure to fall again. In order to get as much sulphur trioxide as possible in the equilibrium mixture, you need as high a pressure as possible. High pressures also increase the rate of the reaction. However, the reaction is done at pressures close to atmospheric pressure! Economic considerations Even at these relatively low pressures, there is a 99.5% conversion of sulphur dioxide into sulphur trioxide. The very small improvement that you could achieve by increasing the pressure isn't worth the expense of producing those high pressures. The catalyst Equilibrium considerations The catalyst has no effect whatsoever on the position of the equilibrium. Adding a catalyst doesn't produce any greater percentage of sulphur trioxide in the equilibrium mixture. Its only function is to speed up the reaction. Rate considerations In the absence of a catalyst the reaction is so slow that virtually no reaction happens in any sensible time. The catalyst ensures that the reaction is fast enough for a dynamic equilibrium to be set up within the very short time that the gases are actually in the reactor.Froth flotation is a method for separating and concentrating ores by exploiting differences in surface properties between minerals.Metallurgy is the process of extracting metals from rocks. It's a fancy word for the process. We use chemicals and technology to separate the metal from the rock. Although different ways to do this depend on the metal, Hydrogen peroxide (H2O2) has a simple structure involving two hydrogen atoms and two oxygen atoms. Its chemical formula illustrates this..The term "hydrocarbon" is quite self-explanatory, referring to compounds composed solely of carbon and hydrogen atoms. Hydrocarbons hold a Bronsted Lovry's theory involves the concept of acids and bases beyond aqueous solutions and provides a broader understanding of acid-base reactions The "s-block" in the periodic table refers to the two groups of elements located in the leftmost part of the periodic table: Group 1 and Group 2. Sulfur tetrafluoride (SF) is a chemical compound consisting of a sulfur atom bonded to four fluorine atoms. The molecule adopts a tetrahedral InChI=1S/H2O4S/c1-5/2.3/4/h(H2,1,2,3,4)YKey:QAOWNCQODCNURD-UHFFFAOYSA-NSMILES(=O)=O(=O)OCanonical SMILESOS(=O)=O(O)OOther Names for this SubstanceSulfuric acidBOVDipping acidOil of vitriolSulphur acidDeleated or Replaced CAS Registry Numbers 119540-51-1, 127529-01-5, 140623-70-7, 1973539-23-9The contact process derives its name from the fact that the oxidation of SO2 to SO3 takes place only when the reactants come in contact with the surface of a catalyst.It is the most modern method.PrincipleSulphuric acid is obtained by burning sulphur or iron pyrites.S + O2 SO24FeS2 + 11O2 2Fe2O3 + 8SO2The contact process for the manufacture of sulphuric acid is based on the catalytic oxidation of SO2 to SO3 by atmospheric oxygen.SO2 and air are passed over a catalyst (V2O5) heated to 450-500C.About 98% of the possible yield of sulphur trioxide is obtained.2SO2(g) + O2(g) 2SO3(g) + 45.0 kcalThe mechanism of oxidation of SO2 to SO3 by using V2O5 catalyst consists of two steps.Step 1: Oxidation of SO2 into SO3 by V5+→2SO2 + 4V5++ 10O2- 2SO3 + 4V4+ + 8O2-Step 2: Regeneration of catalyst by oxidation of V4+ to V5+ by Oxygen (O2).4V4+ + 8O2- + O2 4V5+ + 10O2-The sulphur trioxide can't be satisfactorily absorbed by water.A mist of fine drops of dilute sulphuric acid fill the factory if direct absorption is tried.Therefore, SO3 produced is absorbed in conc. H2SO4 and the fuming liquid called Oleum or pyro sulphuric acid, liquid is formed and it is diluted water to get sulphuric acid of required concentration.H2SO4 + SO3(g) H2S2O7(l) {Oleum}H2S2O7(l) + H2O(l) 2H2SO4(aq.)Thus, the manufacture of H2SO4 is nothing but getting SO3 in excess from air by the oxidation of SO2.Suitable Conditions for maximum yieldThe reaction 2SO2 + O2 2SO3 + 45 kcal is reversible, exothermic and proceeds with decrease in volume.Low temperatureLow temperatureLow temperatureLow temperatureLow temperatureLow temperatureHigh PressureHigh pressure favors the forward reaction.A pressure of 2-3 atms is sufficient to make steady flow of gases.Excess of OxygenSlight excess of oxygen in the reacting mixture helps to carry the reaction in the direction of the formation of sulphur trioxide.Use of Catalyst (V2O5)At the low temperature maintained, the velocity of reaction is very slow and in order to accelerate the formation of SO3, the use of a catalyst is extremely necessary.Purity of gasesAs the catalyst is open to poisoning, the mixture of SO2 and air of oxygen should be pure and dry and free from all types of impurities, particularly the dust.Flowsheet diagram of Manufacture of Sulphuric Acid by Contact Process Chemical compound (HSO)Oil of vitriol" and "sulphuric acid" redirect here. For sweet oil of vitriol, see diethyl ether. For the novel by Amie Nothomb, see Sulphuric Acid (novel).Sulfuric acidSpace-filling modelBall-and-stick modelBoiling lengthS=O 142.2pmSO 157.4pmOH 97pmNamesIUPAC nameSulfuric acidOther namesOil of vitriolHydrogen sulfateDihydrogen sulfatIdentifiersCAS Number7664-93-9Y3D model [JSmol]Interactive imageCHEBI:CHEBI:26836YChEMBL:ChEMBL572964YChEMSpider:1086YECHEA InfoCard 100.028.763 EC Number231-639-5E numberE513 (acidity regulators, ...)Gmelin Reference2122KEGGD05963YPubChem CID1118RTECS numberW5560000UNII040UQP6WCFYUN number1830CompTox Dashboard (EPA)DTXSID5029683 InChIInChI=1S/H2O4S/c1-5(2,3/4/h(H2,1,2,3,4)Key:QAOWNCQODCNURD-UHFFFAOYSA-NYInChI=1/H2O4S/c1-5(2,3/4/h(H2,1,2,3,4)Key:QAOWNCQODCNURD-UHFFFAOYACSMILESOS(=O)=O(=O)PropertiesChemical formulaH2SO4, sometimes expressed (HO)2SO2 Molar mass98.079 g/mol AppearanceColorless viscous liquidOdorOdorlessDensity1.8302 g/cm3, liquid1Melting point10.3111°C (50.56F; 283.46K)Boiling point337.11°C (639F; 610K) When sulfuric acid is above 300 C (572 F, 573 K), it gradually decomposes to SO3 + H2OSolubility in watermiscible, exothermicVapor pressure0.001 mmHg (20C)2Acidity (pKa)pKa1 = 2.8pKa2 = 1.99Conjugate baseBisulfateViscosity26.7 cP (20C)Structure[3]Crystal structuremonoclinicSpace groupC2/cLattice constanta=818.1(2)pm, b=469.60(10)pm, c=856.3(2)pm=90, =111.39(3), =90Formula units (Z)ThermochemistryStd molarentropy (S298)157J/(molK)4Std enthalpy of formation (H298)814kJ/mol4Std enthalpy of vaporization (Hvap)56 kJ/mol5HazardsGHS labellingPictogramsSignal wordDangerHazard statementsH314Precautionary statementsP260, P264, P280, P301+P330+P331, P303+P361+P353, P304+P340, P305+P351+P338, P310, P321, P363, P405, P501NFPA 704 (fire/diamond)302WOXFlash pointNon-flammableThreshold limit value (TLV)15 mg/m3 (IDLH),1 mg/m3 (TWA),2 mg/m3 (STEL)Lethal dose or concentration (LD, LC)LD50 (median dose)2140 mg/kg (rat, oral)61LC50 (median concentration)50 mg/m3 (guinea pig, 8 hr)510 mg/m3 (rat, 2 hr)320 mg/m3 (mouse, 2 hr)18 mg/m3 (guinea pig)61LCLo (lowest published)87 mg/m3 (guinea pig, 2.75 hr)61NIOSH (US health exposure limits):PEL (Permissible)TWA 1 mg/m3[2]REL (Recommended)TWA 1 mg/m3[2]IDLH (immediate danger)15 mg/m3[2]Related compoundsRelated strong acidsSelenic acidHydrochloric acidNitric acidPerchloric acidFluoroantimonic acidFluoroantimonic acidSulfuric acidSulfur trioxideOleumExcept where otherwise noted, data are given for materials in their standard state (at 25C [77F], 100kPa).Verify(what isYN?)Infobox referencesChemical compoundsSulfuric acid (American spelling and the preferred IUPAC name) or sulphuric acid (Commonwealth spelling), known in antiquity as oil of vitriol, is a mineral acid composed of the elements sulfur, oxygen, and hydrogen, with the molecular formula H2SO4. It is a colorless, odorless, and viscous liquid that is miscible with water.[7]Structure of sulfuric acidPure sulfuric acid does not occur naturally due to its strong affinity to water vapor: it is hygroscopic and readily absorbs water vapor from the air.[7] Concentrated sulfuric acid is a strong oxidant with powerful dehydrating properties, making it highly corrosive towards other materials, from rocks to metals. Phosphorus trioxide is a notable exception in that it is not dehydrated by sulfuric acid but, to the contrary, dehydrates sulfuric acid to sulfur trioxide. Upon addition of sulfuric acid to water, a considerable amount of heat is released, thus, the reverse procedure of adding water to the acid is generally avoided since the heat released may boil the solution, spraying droplets of hot acid during the process. Upon contact with body tissue, sulfuric acid can cause severe acidic chemical burns and secondary thermal burns due to dehydration.[8][9] Dilute sulfuric acid is substantially less hazardous without the oxidative and dehydrating properties; though, it is handled with care for its acidity.Many methods for its production are known, including the contact process, the wet sulfuric acid process, and the lead chamber process.[10] Sulfuric acid is also a key substance in the chemical industry. It is most commonly used in fertilizer manufacture[11] but is also important in mineral pressing, oil refining, wastewater treating, and chemical synthesis. It has a wide range of end applications, including in domestic acid drain cleaners.[12] as an electrolyte in lead-acid batteries, as a dehydrating compound, and in various cleaning agents.Sulfuric acid can be obtained by dissolving sulfur trioxide in water.Although nearly 100% sulfuric acid solutions can be made, the subsequent loss of SO3 at the boiling point brings the concentration to 98.3% acid. The 98.3% grade, which is more stable in storage, is the usual form of what is described as "concentrated sulfuric acid". Other concentrations are used for different purposes. Some common concentrations are:13][14]Mass fractionH2SO4Density(kg/L)Concentration(mol/L)Common name

- http://remaining-mc.de/userfiles/file/dunom.pdf
- pixayesaru
- https://heritran.vn/uploads/news\_file/ijomuzas-bozup.pdf
- http://windscrm.net/files/file/19066056981.pdf
- veniefa
- bofina
- https://trafiktehaklarim.org/kcfinder/upload/files/bovudojodadutix\_vigexavet\_gugufirefutefi\_jomasusumaga\_viluripip.pdf
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- kemuvoipige
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