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Preparation of anthranilic acid from phthalimide pdf

Preparation of phthalimide from phthalic acid. Synthesis of anthranilic acid from phthalimide. Preparation of anthranilic acid from phthalimide pdf. Gabriel phthalimide reaction is used for preparing. Synthesis of anthranilic acid from phthalimide mechanism. Preparation of anthranilic acid from phthalimide mechanism.

When everything is mixed and the exotic reaction is no longer observed, it can continue with a heating of the mixture. This can be a good OTC form of obtaining the right phthalimida of the fñic E;ido, whether purchased or obtained by hydrolisis of different phthalates. Finally, you will get a successful reaction with a decent performance. If you have the smell of ammonaco, it is likely that some incorrect reactions are being produced. Therefore, it probably led some by -products and an excess coloration of my final solution. So briefly what I did was 14 g of phthalimida + 10 g naoh + 80 ml of water + cooling ... to continue working on the site, start session if you are not registered, register the registration at the same time. Publishing a brief report on my lack of reorganization of the phtalimidation to the antrany and executing qualitative animals in the Pthalimida http://www.sciencemadness.org/talk/viewthread.php?Tid=19662 & ... For some reason I could smell a definite smell of De El Ammonaco leaving the solution, probably due to some inorganic ammonium salts like NH4Cl. After a few hours, a conversion of phthalimida to sodium phthalamate is completed. The US patent. 2653971 has some good instructions on reaction conditions. 21, 1445 (1956). If you cannot hydrolyze the phthalimida correctly: you will obtain a really poor anthranhcic performance, you will get a clear solution without precipitated, which is exactly what happened to me. At first I thought it was a good idea to filter the solution with a green Antranilate precipitate, avoiding the gray background layer that is relatively difficult to disturb. [Edited on 18-4-2015 by byko3y] [edited on 18-4-2015 by byko3y] solution color changed to murky green, so I would say that the first copper salt precipitate content was of copper, copper and copper acetate and copper Antranilate. Phtalimidas, such as succinimidas, can reversibly attract the hal of hal. Note that Vogel's procedure uses Hipobromito Recien prepared. You can precise it with a salt (ii) salt (CUCL2 or CUSO4). At least not nothing. The mo fuse point was like 224 ° C, but after melting the crystallization point it was like 232 ° C. It is possible to use the heat of chlorination for subsequent reorganization, but I am not so good in this reaction. + Acyic acid -> Collect precipitate. Next, it should not add more than a stochamomide amount of a cold hypochlorite solution that tries to keep the mixture below 40 ° C, otherwise, the Antranian time will begin to form, reacting to more with hypochlorite, forming chlorinated by -products. Vogel also mentions hydrolisis, but there are no instructions in this regard in the experimental part, they simply mix everything and let the reaction warm up in s. Actually, when adding a CuSO4 solution, many packages began to leave and I have two layers of precipitate (used a saturated solution CUSO4): heavy gray gray lay S light and is very difficult to precipitate, so that filter. In addition, the patent recommends ensuring that hypochlorite solution never heats up 15 ° C during its preparation and storage, but I have no way to meet the requirement because my hypochlorite solutions were stored at 20-25 ° C. I think it is important to pay attention to the smell of the reaction mixture above 40 ° C. This last one is a really strong oxidant that will oxidize and chlorinate the benzene ring, lowering the yields. The crystals obtained from Antrany have perhaps color and seem to have at least few impurities. Chem. The real compound experienced by reorganization is a base hydrolysis product: sodium phthalamate or 2 -Carbamoilbenzoate of sodium. After Filter the silido, I have a family solution of family sodium. familiar. What overheated and discard . Calefaccin ... Basically, you can follow the org. If you screwed something and your performance is really poor or not added enough academic, your solution will be clear. The solution becomes brown (third flask in the image). There is a question about the compound is and if any prison is formed at all. There is a good qualitative test for the antrany, a solution of Cu (II) (for example, CuSO4) in the aqual dilulte. Keep in mind that the color of the mixture did not change. You should not add hypochlorite that the stochamomose quantity, because when the hypochlorite without reacting will become a more stable mixture of chloride and chlorate. A usual benzoic? , but the Antranian (O-aminobenzoico) gives a green precipitate and the M-aminobenzoico E;ido only gives a blue color (not tested by M). 1922, 2, 75. org. The most likely that it does not add enough acic and carbamate will not be completely hydrolyzed (perhaps even 20% of the performance), that is why more co2 came out in This step and reacted with the cupbearer. Therefore, you can add a ammonaco solution to anything you have, except salts and alkaline stems, and after boiling the water and reaching the temperature above the fusing points of all the fñic compounds possible (300 ° C) will obtain a reasonable pure phthalim. Therefore, there is a mixture of some copper salts in the precipitate. Reacted to the same sequence as the of patents, while the instructions of the rhodium's pgina seem insufficient. In addition, after filtering the antrany's precipitate, it will obtain a clear marine solution. clear. A small amount of anthranilico Ercido in   L + 150 ml of cooling Naocl + ... + HCl ... that is why it is important to assess hypochlorite (for example, iodine-tiosulfate). I think that the precipitate of the Rhodium procedure is the phtalimida without reacting, but we ensure that it is completely hydrolyzed. The Antranian entry can be purified by recrystallization of the water-metra mixture. The patent says that 60 ° C is sufficient, and I have a great investigation of the Hofmann reorganization that says that only a brief warming is needed to complete the reorganization, although there is no great damage when reaching 75-80   C for 1-2 minutes to ensure that the reorganization is complete. See "Recrystallization of Organic Compounds of the detergent systems" Sugihara, Newman, J. Synth. The truth is that you cannot reorganize hofmann in the phtalimide. Which is almost the same as the performance in the Vogel procedure through the hypobromite and rhodium procedure, which is an adoption of the Vogel procedure using hypochlorite, although by some reason I could not reproduce the results using hypochlorite. DOI: 10.15227/orSYN.002.0075 Sources: Some Internet articles and "Chemistry of Encyclopedia Concise" by Mary Eagleson (1994), p. Ammonium carbonate will also work, but not if the urea is a good idea, because the reaction can become violent and the urea cannot remove the heat from the mixture unlike the ammonium carbonate and the water solution of ammonaco. A good book called "Organic Chemistry" by Fieser, Fiester (1956) helped me to rule out what is really happening in the hofmann reorganization of the phthalimide. As, the reaction mixture becomes the pages as the patent (second flask in the image). It is important to ensure that the temperature of the solution is always That 20 ° C, otherwise, the phthalimida will be hydrolyzed in the fñic E;ido (the patent says that the pty hydrolisis begins at 35 ° C). The good news is: the phthalimide is is The most stable of all the fñenic cuimentos in non- alkaline high temperature conditions. DOI: 10.1021/fo01118A032 or you can simply use hot water as Vogel says. I am quite sure that I have up to 5-10% of the by-products, while a 99% compound will be white. That is why I probably want to dissolve it in a icy naoh solution, add phthalimida in portions and keep the reaction flask in a refrigerator. Then, after filtering, washing with water and drying the precipitate, I have 6 g of raw Antrannium, corresponding to 45% of yield more copper salt could give Apex 2 g of more than more Antrany acid. It is important to keep the  Then, after cooling, carefully adjusting the pH to 7-8 and almost did not get precipitated from something without reacting (less than 1 g), because at least the stochamomide amount of the hypochlorite. First dissolve the phthalimida in the solution of 80 ml of 10 g of naoh. Another possible reason for which I obtained bad performance in my first attempts is a dirty phthalim. 832-833 (you can read on Google Books) the set of orsyn mentioned above says that it can make the phthalimida form ammonium phthalate, but the performance is in doubt. But when you have a lot of free Antranian, they will begin to precipitate as it does in the fourth flask in the image. Then, the last step is a careful acidification with an aceticism accompanied by a large number of CO2 bubbles that leaves the solution. This last one will discover that the green layer has almost no Antranian. In my last attempt to use a regenerated flimide, I could not notice a slight smell of ammonaco in this step. Next, hydrolylo It will probably also at Acid. This provides a cloth yellow solution (first flask in the image). You must use an excess of phthalimida if you want a reasonably pure product, but I have no idea how recovery instruction. instructions.

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