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(54) **METHOD OF SHUTTING DOWN A REACTOR**

VERFAHREN ZUM ABSCHALTEN EINES REAKTORS

PROCÉDÉ D'ARRÊT D'UN RÉACTEUR

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EP 2 969 172 B1

Description

CROSS-REFERENCE TO RELATED APPLICATIONS

[0001] This application claims priority to U.S. Provisional Patent Application Serial No. 61/781,531 filed March 14, 2013.

FIELD OF THE INVENTION

[0002] This application relates to a method of shutting down a dehydrogenation reactor having a catalyst bed containing a chromium-containing catalyst. More particularly, this application relates to a method of shutting down a dehydrogenation reactor having a catalyst bed containing a chromium-containing catalyst, wherein after the shut-down, at least some, and preferably a majority, of the chromium is in a reduced oxidation state.

BACKGROUND

[0003] Unsaturated hydrocarbons, e.g. olefins, may be derived by catalytic dehydrogenation of an alkane to create a product more useful and more valuable than the reactant. Olefins are feedstocks for many processes that yield increasingly valuable products through polymerization. Catalytic dehydrogenation of alkanes is a selective way to produce alkenes and was commercialized in the 1930's.

[0004] One such catalytic dehydrogenation process is the CATOFIN[®] process which produces alkenes, e.g. isobutylene, propylene and amylenes, from alkanes, e.g. isobutane, propane and isopentanes respectively. The CATOFIN[®] process is the most widespread catalytic dehydrogenation process employed in the production of propene, the precursor material to polypropylene. Another dehydrogenation process is the CATADIENE[®] process which results in the catalytic dehydrogenation of C₄ and C₅ hydrocarbons to produce diolefins, i.e. dienes, of the same carbon number. The CATOFIN[®] and CATADIENE[®] processes utilize a supported chromium catalyst in a series of adiabatic fixed bed reactors in cyclic operation with quickly alternating dehydrogenation and regeneration periods. Other catalytic dehydrogenation processes utilizing supported chromium catalysts are known in the art. Such processes typically employ frequent high temperature regeneration cycles.

[0005] Catalysts that are used for the dehydrogenation of light hydrocarbons typically comprise chromium oxide supported on the surface of an aluminum oxide carrier, i.e. chromium-alumina catalysts. Processes utilizing supported chromium catalysts such as chromia-alumina catalysts (Cr₂O₃/Al₂O₃) are well known and have been described in technical literature as well as in numerous patents. US 4 071 439 discloses a hydrosulfurization shut-down method. Dehydrogenation catalysts become spent after prolonged use through several alternating dehydrogenation and regeneration cycles. Spent catalyst must

be removed from the reactor and replaced with fresh catalyst. It is preferred that at least some and preferably a majority of the chromium in the spent catalyst be at a reduced oxidation state when the spent catalyst is removed from the reactor. The applicant has now surprisingly found a method for shutting down a dehydrogenation reactor which method results in at least some and preferably a majority of the chromium in the spent catalyst being at a reduced oxidation state.

SUMMARY OF THE INVENTION

[0006] A method for shutting down a dehydrogenation reactor having a catalyst bed with a chromium-containing catalyst operating at a first elevated temperature T₁ comprises cooling the catalyst bed with a first cooling gas to a second elevated temperature T₂ lower than T₁, removing the first cooling gas from the catalyst bed, treating the catalyst bed with a reducing gas to bring the chromium in the catalyst to a reduced oxidation state, cooling the catalyst bed with a second cooling gas from the second elevated temperature T₂ to a third elevated temperature T₃ lower than T₂, removing the reducing gas from the catalyst bed, cooling the catalyst bed with a third cooling gas from the third elevated temperature T₃ to a fourth elevated temperature T₄ lower than T₃, and cooling the catalyst bed to ambient temperature T_{amb}, whereby the dehydrogenation reactor is shut down and the spent catalyst can be removed. Advantageously, upon completion of the method of the present application, at least some, and preferably a majority, of the chromium in the chromium-containing catalyst will be at a reduced oxidation state.

[0007] The reducing gas is preferably introduced after the catalyst is brought from the first elevated temperature T₁ to the second elevated temperature T₂, to bring the catalyst to the desired reduced oxidation state. In one embodiment, the reducing gas and the second cooling gas are the same gas. In another embodiment, the reducing gas and the second cooling gas are different gases.

[0008] The reducing gas is preferably purged from the catalyst bed after the catalyst has reached the desired oxidation state. If the second cooling gas is different from the reducing gas, then the second cooling gas can serve as the reducing gas purge medium. Otherwise, the third cooling gas can serve as the reducing gas purge medium.

DETAILED DESCRIPTION

[0009] The present application generally relates to a method of shutting down a dehydrogenation reactor used for manufacturing light olefins and di-olefins from light alkanes, the reactor having a catalyst bed containing a chromium-containing catalyst, such that at the conclusion of the shut-down method of the present application, at least some, and preferably a majority, of the chromium in the catalyst bed is in a reduced oxidation state.

[0010] One exemplary embodiment of such a method of shutting down a dehydrogenation reactor housing a catalyst bed containing a chromium-containing catalyst, with the catalyst bed operating at a first elevated temperature under standard dehydrogenation conditions, comprises:

- (a) cooling the catalyst bed with a first cooling gas from the first elevated temperature to a second elevated temperature lower than the first elevated temperature;
- (b) removing the first cooling gas from the catalyst bed while maintaining the catalyst bed at about said second elevated temperature;
- (c) treating the catalyst bed with a reducing gas for a period of time sufficient to cause chemical reduction of at least some of the chromium in the catalyst to a reduced oxidation state;
- (d) cooling the catalyst bed with a second cooling gas from the second elevated temperature to a third elevated temperature lower than the second elevated temperature;
- (e) removing the reducing gas from the catalyst bed;
- (f) cooling the catalyst bed with a third cooling gas from the third elevated temperature to a fourth elevated temperature lower than the third elevated temperature; and
- (g) cooling the catalyst bed with a fourth cooling gas from the fourth elevated temperature to ambient temperature, whereupon the reactor is shut down, and wherein at least some of the chromium in the catalyst is at the reduced oxidation state.

[0011] In one embodiment, upon completion of this shut-down method, more than 50% of the chromium is in a reduced oxidation state. In a preferred embodiment, upon completion of the shut-down method, at least 75% of the chromium is in a reduced oxidation state. In a more preferred embodiment, upon completion of the shut-down method, at least 90% of the chromium is in a reduced oxidation state. In a most preferred embodiment, upon completion of the shut-down method, at least 99% of the chromium is in a reduced oxidation state.

[0012] A dehydrogenation reactor system generally includes a dehydrogenation reactor vessel having a reactor inlet for receiving hydrocarbon feedstock, air, and reducing gas, and two reactor outlets for discharging reactor effluents, one for discharging hydrocarbons and reducing gas and the other for discharging air. The dehydrogenation reactor vessel defines a dehydrogenation reaction zone and can contain dehydrogenation catalyst pellets that are usually mixed with inert alpha-alumina and packed together to form the dehydrogenation catalyst bed.

[0013] In operation, a dehydrogenation feed is introduced into the dehydrogenation reactor that is operated under dehydrogenation reaction conditions with the dehydrogenation catalyst at a first elevated temperature.

During operation the dehydrogenation reactor is operated in the cyclic mode wherein the dehydrogenation part of the cycle is followed by steam purge for a period of time sufficient to purge hydrocarbon from the catalyst surface. The catalyst is then regenerated by air followed by evacuation of the catalyst bed and then reduction of the catalyst by means of a reducing gas. All of the foregoing steps are conducted at a first elevated temperature. Once reduction is complete, a new dehydrogenation cycle can begin.

[0014] As is known in the art, the dehydrogenation reactor system may comprise several parallel reactors. Depending on the size and production rate of the system, typically three to ten reactors operate in cyclic mode, whereby simultaneously some reactors are operated in the intermediate steps of the cycle such as reduction, evacuation or steam purge, other reactors are operated in the dehydrogenation step of the cycle, while still other reactors are being regenerated.

[0015] In the operation of a hydrocarbon dehydrogenation reactor, the catalyst, such as Al-Cr, may last up to several years. As the catalytic performance decreases there is a need to unload the spent catalyst and replace it with new catalytic material. The shut-down of the reactor system typically requires a procedure for cooling down the spent catalyst prior to its removal.

[0016] In accordance with the present application, a method for shutting down a dehydrogenation reactor having a catalyst bed with a chromium-containing catalyst is illustrated, wherein in Stage 1 the reactor is operating at a first elevated temperature T_1 , the method comprising the steps (a) cooling the catalyst bed with a first cooling gas to Stage 2 wherein the catalyst bed is at a second elevated temperature T_2 , (b) removing the first cooling gas from the catalyst bed while maintaining the catalyst bed at about said second elevated temperature, (c) treating the catalyst bed with a reducing gas for a period of time sufficient to cause chemical reduction of at least some of the chromium in the catalyst to a reduced oxidation state, (d) cooling the catalyst bed with a second cooling gas from the second elevated temperature T_2 to Stage 3 wherein the catalyst bed is at a third elevated temperature T_3 , (e) removing the reducing gas from the catalyst bed, (f) cooling the catalyst bed with a third cooling gas from the third elevated temperature T_3 to Stage 4 wherein the catalyst bed is at a fourth elevated temperature T_4 , and (g) cooling the catalyst bed to the Final Stage wherein the catalyst bed is at ambient temperature T_{amb} , whereby the dehydrogenation reactor is shut down and the spent catalyst can be removed. Advantageously, at the conclusion of this shut-down method, at least some, and preferably a majority, of the chromium in the catalyst bed will be at a reduced oxidation state.

[0017] In one exemplary embodiment, in one or more of the cooling steps the rate of cooling is controlled such that the rate of change of the temperature of the catalyst in the bed is no more than 50°C per hour, preferably no more than 20°C per hour, and more preferably no more

than 10°C per hour. In a preferred embodiment, in each of the cooling steps the rate of change of the temperature of the catalyst in the bed is no more than 50°C per hour, preferably no more than 20°C per hour, and more preferably no more than 10°C per hour.

[0018] In one exemplary embodiment, when the catalyst bed is at Stage 1 and prior to the first cooling of the catalyst bed to the second elevated temperature T_2 , the introduction of the dehydrogenation feed into dehydrogenation reactor at the first elevated temperature T_1 is terminated and the catalyst at the first elevated temperature T_1 is subjected to a standard dehydrogenation-regeneration operating procedure such as described above. Upon completion of the dehydrogenation-regeneration process, the dehydrogenation catalyst is at a first elevated temperature T_1 that approximates the reactor temperature conditions existing immediately prior to the dehydrogenation feed termination and last regeneration, and a substantial proportion of the chromium in the catalyst is at a reduced oxidation state.

[0019] The first elevated temperature T_1 of the dehydrogenation catalyst bed can range from 500°C to 800°C, preferably from 525°C to 750°C, and most preferably from 550°C to 700°C.

[0020] In accordance with the present application, the dehydrogenation catalyst at the first elevated temperature T_1 is cooled with a first cooling gas to Stage 2 characterized by a second elevated temperature T_2 . The second elevated temperature T_2 is less than the first elevated temperature T_1 of the dehydrogenation catalyst bed as described above, and can be in the range of from 300°C to just below the first elevated temperature, preferably from 325°C to 500°C, and most preferably from 350°C to 450°C.

[0021] Cooling of the catalyst bed from the first elevated temperature T_1 to the second elevated temperature T_2 can be accomplished by introducing a first cooling gas at a desired temperature through the inlet. In one embodiment, the first cooling gas is air which passes through air heaters before entering the reaction zone through the inlet. The pressure of the first cooling gas in the dehydrogenation reactor can range from below atmospheric pressure to 40 psia or greater. The temperature of the air heaters can be gradually reduced at a predetermined rate, such that the rate of change of the temperature of the catalyst in the bed is no more than 50°C per hour, preferably no more than 20°C per hour, and more preferably no more than 10°C per hour. The slow rate of change results in a gradual cooling down of the dehydrogenation catalyst bed inside of the reactor while avoiding thermo-shocks of the dehydrogenation reactor and air pre-heater. This cooling continues until the catalyst bed achieves the desired second elevated temperature as described above, as gauged by the temperature of the cooling gas at the reactor outlet. In some embodiments of the present application, and depending on the scale of the reactor, this first cool down step can take from several hours to several days. Once the desired second

elevated temperature T_2 is reached, the flow of the first cooling gas is terminated.

[0022] After the flow of the first cooling gas has been terminated and Stage 2 has been reached, a substantial portion of the residual first cooling gas is removed from the reactor. This removal can be achieved by different methods. In one embodiment, the first cooling gas is removed by evacuation, with an evacuation pressure of 0.5 atmospheres or less being preferred. In another embodiment, the first cooling gas is removed from the reactor by purging with a non-air purging medium for a period of time, which can last from about ten seconds up to a few minutes, sufficient to eliminate the first cooling gas. Preferably, the removal of the first cooling gas is sufficient to remove traces of the first cooling gas from the catalyst pores and from any voids in the dehydrogenation reactor. The temperature of the purging medium is maintained in the range of the second elevated temperature T_2 to maintain the catalyst bed at the second elevated temperature during the purging. The choice of purging media can depend on factors such as the availability of such media at each reactor site and the compatibility of such media with the reactor and catalyst. Steam and nitrogen are each acceptable purging media typically used in dehydrogenation facilities, and therefore likely to be readily available. Upon completion of the purge, the introduction of the purging medium is terminated.

[0023] Following the first purge, the catalyst bed is treated with a reducing gas for a period of time sufficient to cause reduction of at least some of the chromium in the catalyst bed from a higher oxidation state to a lower oxidation state. At the beginning of this treatment, the catalyst bed is at the second elevated temperature T_2 , which is a temperature high enough to allow the reaction of the reducing gas with the chromium in the catalyst to take place. The choice of the reducing gas will depend on the availability of the gas at the reactor site and compatibility of the reducing gas with the reactor and the catalyst. Suitable reducing gases include, without limitation, dehydrogenation hydrocarbons, such as isobutane, propane, n-butane, propylene, natural gas, hydrogen mixed with C_1 - C_4 hydrocarbon, carbon monoxide; and mixtures of any of the foregoing. The treatment of the catalyst bed with the reducing gas continues until the desired level of reduction of the chromium in the catalyst bed has been achieved.

[0024] In one exemplary embodiment, the reducing gas has a heat capacity that allows it to also serve as a second cooling gas such that during the reducing gas treatment, the temperature in the catalyst bed can be lowered from the second elevated temperature T_2 of Stage 2 to a third elevated temperature T_3 of Stage 3. If the reducing gas does not have the heat capacity to cool the catalyst bed, then a separate second cooling gas is used to bring the catalyst bed from the second elevated temperature T_2 of Stage 2 to a third elevated temperature T_3 of Stage 3. In one embodiment, the third elevated temperature T_3 is in the range of 155°C - 320°C. In a

preferred embodiment, the third elevated temperature T_3 is in the range of 230°C - 320°C. In a more preferred embodiment, the third elevated temperature T_3 is in the range of 280°C - 320°C.

[0025] Once the reduction of the chromium has reached a desired level of completeness and the third elevated temperature T_3 of the catalyst bed is reached as measured by the outflow of the reactor gas, then introduction of the reducing gas is terminated. If the reducing gas does not serve as the second cooling gas, introduction of the reducing gas will be terminated when the reduction of the chromium has reached a desired level of completeness, and then a separate second cooling gas will be introduced until the third elevated temperature T_3 of the catalyst bed is reached as measured by the outflow of the reactor gas.

[0026] If the reducing gas does not serve as the second cooling gas, then the second cooling gas may be selected from N_2 , CO_2 , or propylene or isobutylene, each of which may be advantageously stored on site in the liquid state and converted into the gas state upon introduction into the reactor.

[0027] Next, the reducing gas is purged from the reactor. If the second cooling gas is different from the reducing gas, then the second cooling gas may serve as the purge medium. If the reducing gas and the second cooling gas are the same gas, then a different purge medium may be used to remove the reducing gas. The purging medium is preferably chemically inert and can again be steam, nitrogen, or other suitable non-reactive media, depending on the availability of such media at each reactor site and the compatibility of the media with the catalyst. The purging medium for the reducing gas purging step may also advantageously serve as a third cooling gas to cool the catalyst from the third elevated temperature T_3 of Stage 3 to a fourth elevated temperature T_4 at Stage 4 that is lower than the third elevated temperature, is in the range of 50°C - 150°C. The time required for this purge and cooling will depend on the magnitude of the difference between the third and fourth elevated temperatures. When the desired fourth elevated temperature T_4 is reached, as indicated by the temperature of the effluent gas, the flow of the purge medium/third cooling gas is terminated. The fourth elevated temperature T_4 is low enough so that any oxygen that is thereafter introduced into the reactor will not react with the catalyst to re-oxidize the reduced chromium in the catalyst back to a higher oxidation state.

[0028] After the reducing gas has been removed and the catalyst bed has been cooled to the fourth elevated temperature T_4 , the catalyst bed is cooled to the Final Stage of ambient temperature T_{amb} , or any temperature that permits the safe removal of the catalyst from the reactor. This cooling can be accomplished by a fourth cooling medium, which can be air or any other suitable cooling medium.

[0029] The dehydrogenation catalyst of the dehydrogenation reactor system operated in accordance with the

method of the present application can be any known alumina - chromium catalyst that can be used in the dehydrogenation of hydrocarbons. Such dehydrogenation catalysts include those catalysts that comprise chromium and alumina oxides. The chromium oxide of the dehydrogenation catalyst may be in any form and obtained from any source or by any method that provides a suitable chromium oxide material for use in the chromium aluminum catalyst. The aluminum oxide of the dehydrogenation catalyst may be in any form and obtained from any source or by any method that provides a suitable aluminum oxide material for use in the chromium aluminum catalyst including aluminum oxide that can be represented by gamma, theta, eta, kappa and alpha-alumina phase.

[0030] Typical alumina-chromium catalysts comprise from 10 to 40% of chromium oxide calculated as Cr_2O_3 , and 55-90% of alumina oxide calculated as Al_2O_3 . Moreover, typical Al-Cr catalyst comprises from 0.3-5% of alkali or alkali-earth metals selected from the group consisting of Na, Li, K, Cs, Mg, Sr, Ca, Ba and combinations thereof. The alumina-chromium dehydrogenation catalyst may further comprise one or more promoters that are usually in form of oxides. These promoter metals may be selected from the group consisting Sc, La, Mo, W, Zr, Sn, Mn, Cu, and mixtures thereof.

[0031] Temperatures at various stages of the method of the present application are expressed herein as temperatures of the catalyst bed. Those skilled in the art will recognize, however, that in commercial dehydrogenation processes carried out in adiabatic reactors, the temperature across the dehydrogenation catalyst bed can vary between the top part of the bed and the bottom part of the bed by as much as 150°C, but more typically from 10 to 120°C. As used herein, the temperature of the catalyst is indicated by the temperature of the effluent gas, whether it is a cooling medium, reducing gas, or other effluent.

EXAMPLE

[0032] In a test scale reactor, 30 g of CATOFIN STD catalyst was heated to 600°C in air. The catalyst was cooled with air flow at atmospheric pressure to a temperature of 350°C. The air was evacuated to 0.5 atm. The catalyst was treated with propane gas for 30 minutes, which was then purged from the reactor by low pressure steam to cool the catalyst to 150°C. The reactor was purged with N_2 for 5 minutes, then cooled in air to ambient temperature. Four catalyst samples were identically treated. At the end of the treatment, it was determined that more than 99% of the chromium in the catalyst was in a reduced oxidation state.

[0033] Reasonable variations, modifications and adaptations can be made within the scope of the described disclosure and appended claims without departing from the scope of the invention.

Claims

1. A method of shutting down a dehydrogenation reactor, the dehydrogenation reactor housing a catalyst bed containing a chromium-containing catalyst, the catalyst bed operating at a first elevated temperature under standard dehydrogenation conditions, the method comprising:

(a) cooling the catalyst bed with a first cooling gas from the first elevated temperature to a second elevated temperature lower than the first elevated temperature;

(b) removing the first cooling gas from the catalyst bed while maintaining the catalyst bed at about said second elevated temperature;

(c) treating the catalyst bed with a reducing gas for a period of time sufficient to cause chemical reduction of at least some of the chromium in the catalyst to a reduced oxidation state;

(d) cooling the catalyst bed with a second cooling gas from the second elevated temperature to a third elevated temperature lower than the second elevated temperature;

(e) removing the reducing gas from the catalyst bed;

(f) cooling the catalyst bed with a third cooling gas from the third elevated temperature to a fourth elevated temperature lower than the third elevated temperature; and

(g) cooling the catalyst bed with a fourth cooling gas from the fourth elevated temperature to ambient temperature, whereupon the reactor is shut down, and wherein at least some of the chromium in the catalyst is at the reduced oxidation state.

2. The method of claim 1, wherein the rate of cooling of the catalyst bed from the first elevated temperature to the second elevated temperature is no greater than 50°C per hour.

3. The method of claim 1, wherein the first cooling gas is heated air.

4. The method of claim 1 wherein the first elevated temperature is in the range of 500°C to 800°C.

5. The method of claim 1 wherein the removal of the first cooling gas is accomplished by evacuation of the reactor to a pressure of 0.5 atm. or less.

6. The method of claim 1 wherein a gas selected from steam, nitrogen, and mixtures thereof, is used to remove said first cooling gas by purging.

7. The method of claim 1, wherein the second elevated temperature is in the range of greater than 300°C

and less than the first elevated temperature.

8. The method of claim 1, wherein the reducing gas comprises a gas selected from dehydrogenation hydrocarbons, propylene, natural gas, hydrogen mixed with C₁-C₄ hydrocarbons, carbon monoxide, and mixtures of any of the foregoing.

9. The method of claim 1, wherein the reducing gas also serves as the second cooling gas to reduce the temperature of the catalyst bed from the second elevated temperature to the third elevated temperature.

10. The method of claim 9 wherein the third cooling gas serves both to cool the catalyst bed to the fourth elevated temperature and to purge the reducing gas from the catalyst bed.

11. The method of claim 1 wherein the second cooling gas is introduced after the catalyst bed is at the desired oxidation state, and serves to purge the reducing gas from the catalyst bed.

12. The method of claim 1 wherein the third cooling gas is chemically inert.

13. The method of claim 1, wherein the third elevated temperature is in the range of 155°C to less than 320°C.

14. The method of claim 1 wherein the fourth cooling gas is air.

15. The method of claim 14 wherein the fourth elevated temperature of the reactor is sufficiently low such that, when the air is introduced into the reactor, the chromium in the catalyst is not oxidized and the fourth elevated temperature is in the range of 50°C - 150°C.

Patentansprüche

1. Ein Verfahren zum Abschalten eines Dehydrierungsreaktors, wobei der Dehydrierungsreaktor ein Katalysatorbett aufnimmt, das einen chromhaltigen Katalysator enthält, wobei das Katalysatorbett bei einer ersten erhöhten Temperatur unter Standard-Dehydrierungsbedingungen arbeitet, wobei das Verfahren umfasst:

(a) Abkühlen des Katalysatorbetts mit einem ersten Kühlgas von der ersten erhöhten Temperatur auf eine zweite erhöhte Temperatur, die niedriger ist als die erste erhöhte Temperatur,

(b) Entfernen des ersten Kühlgases aus dem Katalysatorbett, während das Katalysatorbett

- bei etwa dieser zweiten erhöhten Temperatur gehalten wird,
- (c) Behandeln des Katalysatorbetts mit einem reduzierenden Gas für einen ausreichenden Zeitraum, um eine chemische Reduktion von wenigstens einem Teil des Chroms im Katalysator zu einem reduzierten Oxidationszustand zu bewirken,
- (d) Abkühlen des Katalysatorbetts mit einem zweiten Kühlgas von der zweiten erhöhten Temperatur auf eine dritte erhöhte Temperatur, die niedriger ist als die zweite erhöhte Temperatur,
- (e) Entfernen des reduzierenden Gases aus dem Katalysatorbett,
- (f) Abkühlen des Katalysatorbetts mit einem dritten Kühlgas von der dritten erhöhten Temperatur auf eine vierte erhöhte Temperatur, die niedriger ist als die dritte erhöhte Temperatur, und
- (g) Abkühlen des Katalysatorbetts mit einem vierten Kühlgas von der vierten erhöhten Temperatur auf Umgebungstemperatur, wonach der Reaktor abgeschaltet ist, und wobei wenigstens ein Teil des Chroms im Katalysator sich im reduzierten Oxidationszustand befindet.
2. Das Verfahren nach Anspruch 1, wobei die Abkühlgeschwindigkeit des Katalysatorbetts von der ersten erhöhten Temperatur auf die zweite erhöhte Temperatur nicht mehr als 50°C pro Stunde beträgt.
 3. Das Verfahren nach Anspruch 1, wobei das erste Kühlgas erhitzte Luft ist.
 4. Das Verfahren nach Anspruch 1, wobei die erste erhöhte Temperatur im Bereich von 500°C bis 800°C liegt.
 5. Das Verfahren nach Anspruch 1, wobei das Entfernen des ersten Kühlgases durch Evakuieren des Reaktors auf einen Druck von 0,5 atm oder weniger erfolgt.
 6. Das Verfahren nach Anspruch 1, wobei ein Gas, ausgewählt aus Dampf, Stickstoff und Mischungen davon, zum Entfernen des ersten Kühlgases durch Spülen verwendet wird.
 7. Das Verfahren nach Anspruch 1, wobei die zweite erhöhte Temperatur im Bereich von mehr als 300°C und weniger als die erste erhöhte Temperatur liegt.
 8. Das Verfahren nach Anspruch 1, wobei das reduzierende Gas ein Gas umfasst, ausgewählt aus Dehydrierungs-Kohlenwasserstoffen, Propen, Erdgas, Wasserstoff im Gemisch mit C₁-C₄-Kohlenwasserstoffen, Kohlenmonoxid und Mischung aus beliebigen der obigen Gase.
 9. Das Verfahren nach Anspruch 1, wobei das reduzierende Gas auch als das zweite Kühlgas zur Verringerung der Temperatur des Katalysatorbetts von der zweiten erhöhten Temperatur auf die dritte erhöhte Temperatur dient.
 10. Das Verfahren nach Anspruch 9, wobei das dritte Kühlgas sowohl zum Abkühlen des Katalysatorbetts auf die vierte erhöhte Temperatur als auch zum Entfernen des reduzierenden Gases aus dem Katalysatorbett durch Spülen dient.
 11. Das Verfahren nach Anspruch 1, wobei das zweite Kühlgas erst eingebracht wird, wenn das Katalysatorbett sich im erwünschten Oxidationszustand befindet, und zum Entfernen des reduzierenden Gases aus dem Katalysatorbett durch Spülen dient.
 12. Das Verfahren nach Anspruch 1, wobei das dritte Kühlgas chemisch inert ist.
 13. Das Verfahren nach Anspruch 1, wobei die dritte erhöhte Temperatur im Bereich von 155°C bis weniger als 320°C liegt.
 14. Das Verfahren nach Anspruch 1, wobei das vierte Kühlgas Luft ist.
 15. Das Verfahren nach Anspruch 14, wobei die vierte erhöhte Temperatur des Reaktors ausreichend niedrig ist, so dass, wenn die Luft in den Reaktor eingebracht wird, das Chrom im Katalysator nicht oxidiert wird, und die vierte erhöhte Temperatur im Bereich von 50°C - 150°C liegt.

Revendications

1. Procédé d'arrêt d'un réacteur de déshydrogénation, le réacteur de déshydrogénation présentant un lit catalytique qui contient un catalyseur au chrome, le lit catalytique fonctionnant à une première température élevée dans des conditions de déshydrogénation classiques, le procédé comprenant :
 - (a) le refroidissement du lit catalytique avec un premier gaz de refroidissement de la première température élevée à une deuxième température élevée plus basse que la première température élevée ;
 - (b) l'élimination du premier gaz de refroidissement du lit catalytique tout en maintenant le lit catalytique à ladite deuxième température élevée environ ;
 - (c) le traitement du lit catalytique par un gaz réducteur sur une période de temps suffisante pour causer la réduction chimique d'au moins une partie du chrome du catalyseur et son pas-

- sage à un état d'oxydation réduit ;
- (d) le refroidissement du lit catalytique avec un deuxième gaz de refroidissement de la deuxième température élevée à une troisième température élevée plus basse que la deuxième température élevée ;
- (e) l'élimination du gaz réducteur du lit catalytique ;
- (f) le refroidissement du lit catalytique avec un troisième gaz de refroidissement de la troisième température élevée à une quatrième température élevée plus basse que la troisième température élevée ; et
- (g) le refroidissement du lit catalytique avec un quatrième gaz de refroidissement de la quatrième température élevée à la température ambiante, sur quoi le réacteur est arrêté et où au moins une partie du chrome du catalyseur est à l'état d'oxydation réduit.
2. Procédé de la revendication 1, où la vitesse de refroidissement du lit catalytique de la première température élevée à la deuxième température élevée ne dépasse pas 50 °C par heure.
 3. Procédé de la revendication 1, où le premier gaz de refroidissement est un air chauffé.
 4. Procédé de la revendication 1, où la première température élevée est dans la plage qui va de 500 °C à 800 °C.
 5. Procédé de la revendication 1, où l'élimination du premier gaz de refroidissement est effectuée par évacuation du réacteur à une pression de 0,5 atm ou moins.
 6. Procédé de la revendication 1 où un gaz sélectionné parmi la vapeur, l'azote et des mélanges de ceux-ci est utilisé pour éliminer ledit premier gaz de refroidissement par une purge.
 7. Procédé de la revendication 1, où la deuxième température élevée est dans la plage qui dépasse 300 °C et est plus basse que la première température élevée.
 8. Procédé de la revendication 1, où le gaz réducteur comprend un gaz sélectionné parmi des hydrocarbures obtenus par déshydrogénation, le propylène, le gaz naturel, l'hydrogène mélangé à des hydrocarbures en C₁-C₄, le monoxyde de carbone et des mélanges de quelconques de ceux-ci.
 9. Procédé de la revendication 1, où le gaz réducteur sert également de deuxième gaz de refroidissement pour abaisser la température du lit catalytique de la deuxième température élevée à la troisième tempé-
- rature élevée.
10. Procédé de la revendication 9, où le troisième gaz de refroidissement sert à la fois au refroidissement du lit catalytique à la quatrième température élevée et à la purge du gaz réducteur hors du lit catalytique.
 11. Procédé de la revendication 1, où le deuxième gaz de refroidissement est introduit après que le lit de catalyseur est à l'état d'oxydation désiré et sert à la purge du gaz réducteur hors du lit catalytique.
 12. Procédé de la revendication 1, où le troisième gaz de refroidissement est chimiquement inerte.
 13. Procédé de la revendication 1, où la troisième température élevée est dans la plage qui va de 155 °C à moins de 320 °C.
 14. Procédé de la revendication 1, où le quatrième gaz de refroidissement est l'air.
 15. Procédé de la revendication 14, où la quatrième température élevée du réacteur est suffisamment basse pour que le chrome du catalyseur ne soit pas oxydé à l'introduction de l'air dans le réacteur et où la quatrième température élevée est dans la plage qui va de 50 °C à 150 °C.

REFERENCES CITED IN THE DESCRIPTION

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Patent documents cited in the description

- US 61781531 A [0001]
- US 4071439 A [0005]

Szabadalmi igénypontok

1. Eljárás egy dehidrogénező reaktor leállítására, a dehidrogénező reaktor magába foglal egy krómtartalmú katalizátort tartalmazó katalizátorágyat, a katalizátorágy egy első magas hőmérsékleten, sztenderd dehidrogénezési körülmények között működik, az eljárás az alábbiakat tartalmazza:

(a) a katalizátorágy hűtése egy első hűtőgázzal az első magas hőmérsékletről egy második magas hőmérsékletre, amely alacsonyabb, mint az első magas hőmérséklet;

(b) az első hűtőgáz eltávolítása a katalizátorágyból, ez alatt a katalizátorágyban körülbelül a második magas hőmérsékleten való tartása;

(c) a katalizátorágy kezelése egy redukálógázzal olyan időtartamig, amely elegendő a katalizátorban lévő króm legalább egy részének redukált oxidációs állapotba történő kémiai redukációjához;

(d) a katalizátorágy hűtése egy második hűtőgázzal a második magas hőmérsékletről egy harmadik magas hőmérsékletre, amely alacsonyabb, mint a második magas hőmérséklet;

(e) a redukálógáz eltávolítása a katalizátorágyból;

(f) a katalizátorágy hűtése egy harmadik hűtőgázzal a harmadik magas hőmérsékletről egy negyedik magas hőmérsékletre, amely alacsonyabb, mint a harmadik magas hőmérséklet; és

(g) a katalizátorágy hűtése egy negyedik hűtőgázzal a negyedik magas hőmérsékletről környezeti hőmérsékletre, ahol a reaktor leáll, és ahol a katalizátorban lévő króm legalább egy része redukált oxidációs állapotban van.

2. Az 1. igénypont szerinti eljárás, ahol a katalizátorágy hűtésének sebessége az első magas hőmérsékletről a második magas hőmérsékletre legfeljebb 50°C óránként.

3. Az 1. igénypont szerinti eljárás, ahol az első hűtőgáz melegített levegő.

4. Az 1. igénypont szerinti eljárás, ahol az első magas hőmérséklet az 500-800°C tartományban van.

5. Az 1. igénypont szerinti eljárás, ahol az első hűtőgáz eltávolítását a reaktor 0,5 atm vagy annál alacsonyabb nyomásra történő kiszivattyúzásával érjük el.

6. Az 1. igénypont szerinti eljárás, ahol egy gőz, nitrogén és ezek keverékei köréből választott gázt alkalmazunk az első hűtőgáz eltávolítására kibajtással.

7. Az 1. igénypont szerinti eljárás, ahol a második magas hőmérséklet a 300°C-nál magasabb és az első magas hőmérsékletnél alacsonyabb tartományban van.

8. Az 1. igénypont szerinti eljárás, ahol a redukálógáz tartalmaz egy dehidrogénező szénhidrogének, propilén, természetes gáz, 1-4 szénatomos szénhidrogénekkel kevert hidrogén, szén-monoxid és ezek bármelyikének keverékei köréből választott gázt.
9. Az 1. igénypont szerinti eljárás, ahol a redukálógáz második hűtőgázként is szolgál a katalizátorágy hőmérsékletének csökkentésére a második magas hőmérsékletről a harmadik magas hőmérsékletre.
10. A 9. igénypont szerinti eljárás, ahol a harmadik hűtőgáz mind a katalizátorágy hűtésére szolgál a negyedik magas hőmérsékletre, mind a redukálógáz kihajtására a katalizátorágyból.
11. Az 1. igénypont szerinti eljárás, ahol a második hűtőgázt akkor vezetjük be, miután a katalizátorágy elérte a kívánt oxidációs állapotot, és arra szolgál, hogy a redukálógázt kihajtsa a katalizátorágyból.
12. Az 1. igénypont szerinti eljárás, ahol a harmadik hűtőgáz kémiaiilag inert.
13. Az 1. igénypont szerinti eljárás, ahol a harmadik magas hőmérséklet a 155°C és a kevesebb mint 320°C közötti tartományban van.
14. Az 1. igénypont szerinti eljárás, ahol a negyedik hűtőgáz levegő.
15. A 14. igénypont szerinti eljárás, ahol a reaktor negyedik magas hőmérséklete elegendően alacsony ahhoz, hogy amikor levegőt vezetünk be a reaktorba, a katalizátorban lévő króm nem oxidálódik és a negyedik magas hőmérséklet az $50\text{-}150^{\circ}\text{C}$ tartományban van.