Feasibility Study of a Turbo-Cracker

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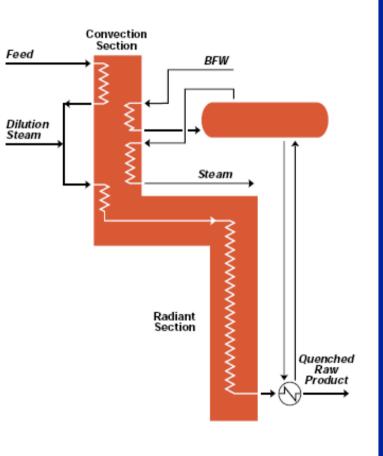
Thermal Cracking Furnace: Current Status

•Ethylene is the largest volume building block for many petrochemicals.

•Currently thermal cracking technology provided by ABB Lummus Global, Shaw Stone and Webster, Kellogg Brown & Root, Linde, and KTI.

 Thermal Cracking Furnace: tubular reactors where thermal cracking of hydrocarbon takes place.

Cracking reactions are endothermic.
 Lots of energy transferred through the tube wall.

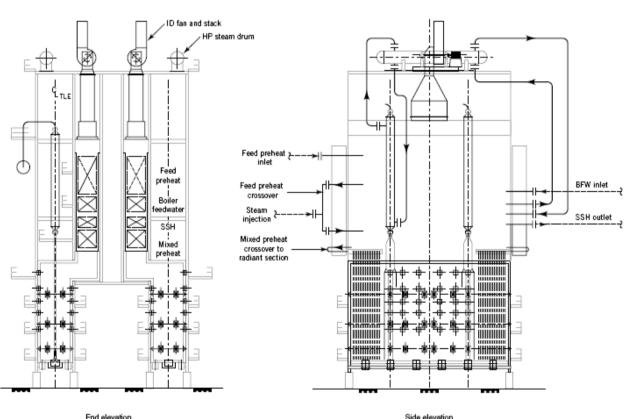


Thermal Cracking Furnace: Current Status

•Coke is formed during pyrolysis. Steam is added as a diluent to the feed.

 Product at the tubular reactor outlet must be cooled down quickly to avoid any further cracking.

•Currently transferline exchanger is used.



Thermal Cracking Furnace: Current Status

 Disadvantage/drawback in existing Thermal Cracking Furnace

- Since thermal cracking reaction is very sensitive to temperature, high heat flux through tube walls is required, high surface temperature is involved.
- At this high surface temperature, heavy coke deposits is formed during normal operation. De-coking every 40 to 100 days. Coking limits heat transfer, also reduces ethylene selectivity.
- Thermal Cracking Furnace is huge in volume (approximately 10,000 m³).

•Use Compressor and Turbine to replace the existing thermal cracking furnace, so-called Turbo-Cracker.

- •Heat is provided by compression in the Compressor of a Gas Turbine (GT).
- Product at the tubular reactor outlet is expanded with the Turbine of a Gas Turbine (GT).

The Idea of Turbo-Cracker

Advantages

- Higher temperature of the mixture of hydrocarbon and steam will increase the selectivity of ethylene.
- Residence time should be much shorter.
- Product at the tubular reactor outlet expands rapidly in the Turbine. This also helps to improve ethylene selectivity.
- Coking will be less severe since no heat is transferred across the tube wall.
- One GT (Compressor and Turbine only) can replace several existing thermal cracking furnaces due to its high processing capacity.

Reaction Scheme in Froment (1975)

| (1) $C_3H_8 \rightarrow C_2H_4 + CH_4$ | A=1.6E09 | E=44 kcal/mole |
|--|-----------|----------------|
| (2) $C_3H_8 \rightarrow C_3H_6 + H_2$ | A=2.0E09 | E=44 kcal/mole |
| (3) $2C_3H_8 \rightarrow C_2H_6 + C_4H_{10}$ | A=2.2E09 | E=54 kcal/mole |
| (4) $2C_3H_8 \rightarrow C_3H_6 + C_2H_6 + CH_4$ | A=1.1E09 | E=48 kcal/mole |
| (5) $C_2H_6 \rightarrow C_2H_4 + H_2$ | A=0.34E13 | E=60 kcal/mole |
| (6) $2C_2H_6 \rightarrow C_2H_4 + 2CH_4$ | A=3.9E12 | E=67 kcal/mole |
| (7) $2C_2H_6 \rightarrow C_3H_8 + CH_4$ | A=0.5E11 | E=50 kcal/mole |
| $(8) \ 2C_3H_6 \rightarrow 3C_2H_4$ | A=1.3E10 | E=50 kcal/mole |
| $(9) C_3H_6 + H_2 \rightarrow C_2H_4 + CH_4$ | A=1.0E15 | E=60 kcal/mole |
| $(10)C_{3}H_{6} \rightarrow C_{2}H_{2} + CH_{4}$ | A=1.4E10 | E=50 kcal/mole |
| (11) $C_2H_4 + H_2 \rightarrow C_2H_6$ | A=0.68E13 | E=52 kcal/mole |

Van Damme, P.S., Narayanan, S. and Froment, G.F. (1975), Thermal Cracking of Propane and Propane-Propylene Mixtures: Pilot Plant versus Industrial Data, *AIChE J*, Vol. 21, No. 6 (pp1065-1073) *Friday Nov. 21 2007*

Reaction Scheme in Froment (1975)

| (12) $C_2H_4 \rightarrow C_2H_2 + H_2$ | A=6.0E13 | E=76kcal/mole |
|---|---------------------------------------|---------------|
| (13) $3C_2H_4 \rightarrow 2C_3H_6$ | A=1.3E11 | E=45kcal/mole |
| (14) $2C_2H_2 + H_2 \rightarrow C_4H_6$ | A=6.0E13 | E=45kcal/mole |
| (15) $C_2H_2 + 2H_2O \rightarrow 2CO$ | + 3H ₂ A=3.5E ² | E=62kcal/mole |
| (16) $C_2H_2 + C_3H_6 \rightarrow C_5H_8$ | A=9.0E ² | E=64kcal/mole |

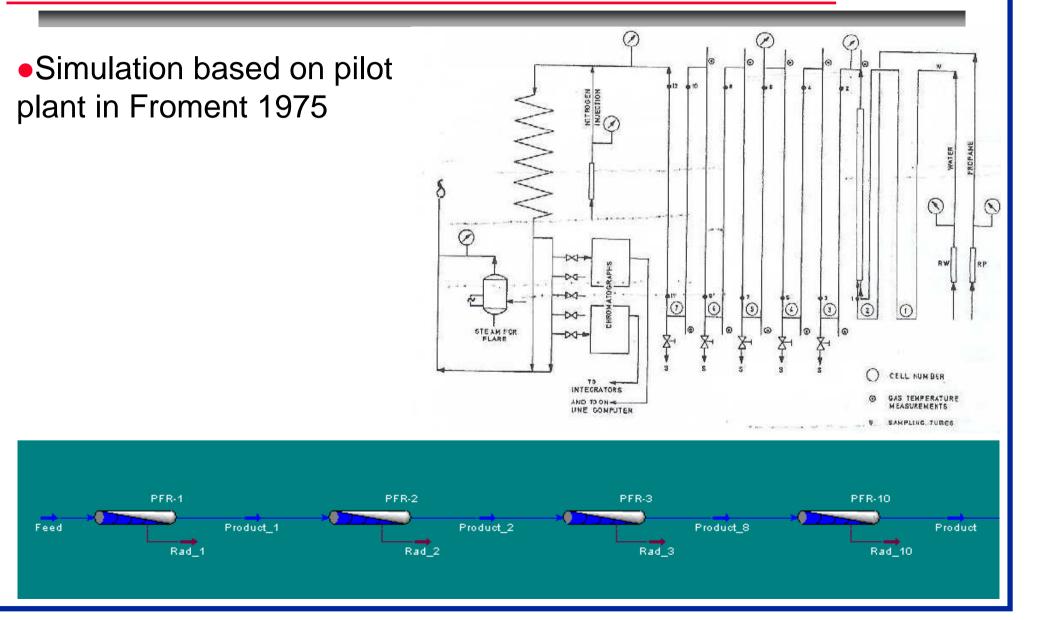
Of these 16 reactions,

(5) and (11) are forward and reverse reactions;

(8) and (13) are forward and reverse reactions;

Van Damme, P.S., Narayanan, S. and Froment, G.F. (1975), Thermal Cracking of Propane and Propane-Propylene Mixtures: Pilot Plant versus Industrial Data, *AIChE J*, Vol. 21, No. 6 (pp1065-1073)

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Simulation based on pilot plant in Froment (1975)

Feed conditions

| | Steam dilution rate | 0.4 kg steam / kg C3H8 | (i.e. steam mass fraction 0.2857) |
|---------|---------------------|------------------------|-----------------------------------|
| | Pressure | 3 bar | |
| | Temperature | 600 °C | |
| | Mass Flowrate | 0.7655 kg/s | |
| Produc | t conditions | | |
| | Pressure | 2 bar | |
| | Temperature | 838 °C | |
| | Mass Flowrate | 0.7655 kg/s | |
| Plug Fl | ow Reactor details | | |
| | Length | 95m | |
| | Diameter | 0.108m | |
| | Wall thickness | 0.008m | |
| | | | |

Modelling of Propane Pyrolysis in HYSYS

Conclusions from simulating the pilot plant in Froment (1975):

- •At the same operating conditions (P, T and flowrate) for the PFR as described in Froment (1975)
- The product composition is close to those data published in Froment (1975)

•This shows the HYSYS model can be used as the basis of various case studies.

Case Studies

•To investigate the relationship between tubular reactor outlet temperature and the residence time

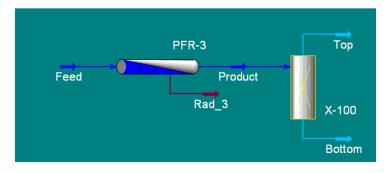
- •The criterion is to achieve the same propane conversion.
- •To investigate the pressure impact

The reaction kinetics in Froment (1975) were used. These were obtained from 650 °C to 900 °C and under pressure about 3 bars.
Actually wider pressure range and temperature range are used in the Cases studied.

Case Studies

Base Case Design

- •Use the Reaction Scheme in Froment (1975)
- Feed same as before
- •PFR details (D=0.108m, L=95m and one long tube only)
- Operating conditions slightly changed
 - Pressure decreases linearly from 3 bar to 2 bar
 - •Temperature increases linearly from 600 C to 880 C (Since only outlet temperature at 880 C, can the propane conversion and ethylene selectivity be similar as before).
 - •Residence time: about 970 ms (calculated with average density)



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Case Studies

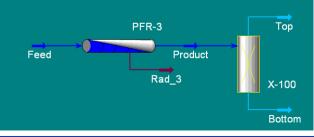
Summary of results from cases studied

- Same mass flowrate for mixture of propane and steam
- Temperature still around 3 bar
- •Reactor length varied from 95 m to 25 m, 6m and 2.5 m
- •Residence time decrease about 0.9s to 0.28s, 0.06s and 0.027s

•Outlet temperature increased 880C to 965 C, 1084 C and 1170 C to achieve the same propane conversion.

•For 2.5m length reactor, when pressure increased from 3 bar to 6 bar, 9 bar, outlet temperature required from 1170 C decrease to 1099 C, 1061 C and the product yield for ethylene and propylene increased.

This means increasing pressure helps to speed up reaction.



Simulation of Turbo-Cracker

A standard Compressor is chosen with

- Compressor Pressure Ratio 23.3:1
- 17 stages
- Mass Flow rate 84.5 kg/s (for air)
- Diameter approximately 2 m and 2.5 m in length
- A standard Turbine is chosen with
 - •6 stage power turbine
 - Diameter approximately 1.5 m and 1.0 m in length
- The same feed is used as in Froment (1975)
- The results from the simulation indicates that
 - •Slightly more steam is required to achieve the same propane conversion.
 - •One Compressor and Turbine can process 4-5 times of the flowrate of existing thermal cracking furnaces.

Conclusions

 Preliminary simulation indicates that Turbo-Cracker concept worth exploring in more detail.

•The main advantage is that one Turbo-Cracker can replace 4-5 conventional Thermal Cracking Furnaces and is much smaller.

Main Challenges

•The kinetics of thermal cracking of propane and naphtha under higher pressure are not clear.

- Information on existing commercial thermal cracking furnace not enough.
- •Compressor requires more power than Turbine can generate.

