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API 521 8th Edition Ballot Item

Thermal Cracking (Work Item 3)

Instructions to Voters/Commenters

- Please limit your comments to the red- or blue-lined portions of the ballot only. Changes represent changes from the prior ballot.
- If you are voting negative with multiple comments, please indicate which comment(s) is the reason for your negative vote, otherwise API's balloting system will categorize all of your comments as negative.

Thanks to Freeman Self and the work group for their efforts.

Melissa Marashi (Chevron) David Fenton (ExxonMobil)

API 521 Task Force Chairs

API-521 - 8th Edition - Expected 2025

This is the 7th Ed, 2020 as revised with suggested statements on cracking

| Г | 3 | 521 | 4.4.13.2. | New section on therm al cracking (modify previous draft | Draftmodifications/additions to several sections; | F. Self; Z. | Presentation |
|---|-----|----------|-----------|---|---|-------------|---------------|
| | | meeting- | 4.2 | section per comments) | See Fall 2020 meeting minutes | Kumana; V. | Fall 2020; |
| | | Spring | | | _ | Quiroga | update Spring |
| | | 2011 | | | | - | 2021 |
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API-521 Proposed Language for Ballot

Changes shown are from the previous ballot

New Definition:

Thermal Cracking:

Thermal cracking of hydrocarbons entails cracking of carbon-carbon bonds with hear but no catalyst or other reactants (e.g. steam). The free-radical reactions typically include cracking of long chain paraffins, splitting side chains from cycloparaffins, but generally no cracking of ring structures or combining reactions of lower molecular weight compounds to form higher molecular weight compounds.

4.4.13 Fire 4.4.13.2 Open Pool Fires 4.4.13.2.1 General

For the purposes of PRD sizing for equipment within the scope of this standard, the design fire scenario has been and continues to be an open pool fire. The recommended method was empirically derived to size PRDs for open pool fires involving hydrocarbons in a refinery environment that is typical for the facilities within the scope of this standard. The method is supported by full-scale test data (see A.2 and C.6 for details).

Both API 521 and API 2000 [^{111]} use open pool fires as the basis for sizing PRDs for the fire case. It is important to apply the appropriate standard when sizing for fire relief because there are differences in assumptions for the fire pool fire intensity, exposed area, and other factors specified in those standards. API 2000 is limited to aboveground liquid-petroleum or petroleum-products storage tanks and aboveground and underground refrigerated storage tanks designed for operation at gauge pressures from vacuum through 103.4 kPa (15 psi).

An open pool fire can affect multiple vessels simultaneously. See 5.3.2 for a discussion.

If the open pool fire involves other types of fuels (e.g. alcohols) that have significantly different radiative fluxes than fuels similar to gasoline, diesel, liquefied petroleum gas (LPG), etc., the method in Annex A can be used with adjusted variables (see Annex A for details).

Relieving temperatures are often above the design temperature of the equipment being protected. If the elevated temperature is likely to cause vessel rupture, additional protective measures should be considered (see 4.4.13.2.6 — and 4.4.13.2.7).

High-boiling materials may exhibit thermal cracking during fire contingencies. Thermal cracking may produce liquids and low-molecular weight gases. The relief temperature may be lowered by this phenomenon. The required relief rate may also be lowered by the thermal cracking process as a portion of the fire heat input is

<u>consumed by the thermal cracking process.</u> This may result in less heat available for vaporization and bulk <u>fluid/vapor temperature increases</u>. For fire contingencies for vessels containing high-boiling materials, the material may experience mild thermal cracking to produce low<u>er</u> molecular weight gases and liquids. This lowers the relief temperature and <u>T</u>the fire heat input utilized to crack material will not be used for vaporization or sensible heat, thus<u>lowering the relief temperature and typically</u> reducing the fire relief rate.

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4.4.13.2.4 Fire-relief Loads 4.4.13.2.4.1 General

It is typically assumed that the vessel is isolated during a fire in order to simplify the analysis, although a more detailed analysis can be warranted in certain cases. Crediting for flow paths that remain open during an overpressure event is generally an acceptable practice. However, it should be recognized that operators and/or emergency responders may attempt to isolate certain lines and vessels during a fire condition in order to limit the fire spread and to safely shutdown the unit. There can also be actuated valves that fail in the closed condition when exposed to a fire. It can be difficult to establish with a degree of certainty whether a particular line will indeed remain open under all fire conditions. Further, consideration should be given to the potential that the fire-relief flow in the flow path will overpressure other equipment. Hence, it can be necessary to add the fire-relief load elsewhere. Ultimately, the user shall decide whether a scenario is credible or not.

4.4.13.2.4.2 provides the heat absorption equations for vessels containing liquids, and 4.4.13.2.4.3 provides the equations to determine open-pool-fire-relief requirements for vessels containing only gases/vapors.

Either the vapor thermal-expansion-relief load or the boiling-liquid vaporization-relief load, but not both, should be used. It is a practice that has been used for many years. There are no known experimental studies where separate contributions of vapor thermal expansion versus boiling-liquid vaporization have been determined. When sizing the PRD for fire exposure, the contribution of vaporizing liquid compared with vapor expansion is generally governing unless, for example, the wetted surface has external insulation in accordance with 4.4.13.2.7 and the unwetted surfaces are not insulated. If thermal cracking occurs with high-boiling materials, credit aAccounting for thermal cracking may result in provide-lower calculated relief temperatures, changed vapor compositions, and lower required relief rates and fluid temperatures.

See Annex I for a discussion of thermal cracking of refinery resid from atmospheric and vacuum columns.

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4.4.13.2.5.2 Vapor

For pressure and temperature conditions below the critical point, the rate of vapor formation (a measure of the rate of vapor relief required) is equal to the total rate of heat absorption divided by the latent heat of vaporization. The vapor to be relieved is the vapor that is in equilibrium with the liquid under conditions that exist when the PRD is relieving at its accumulated pressure.

The latent heat and relative molecular mass values used in calculating the rate of vaporization, should pertain to the conditions that can generate the maximum vapor rate.

If credit for thermal cracking of high-boiling point materials is taken, the amount of fire heat input <u>may be used</u> is utilized to crack the material and produce low molecular weight gases and liquids, <u>reducing heat available</u> <u>forsupply sensible heating to raise the liquid's temperature and provide latent heat offor liquid vaporization.</u> <u>Consideration for thermal cracking of high-boiling point materials can be included in the analysis. This can result</u> in lower molecular weight gases and liquids, lower relief temperature, and a different latent heat of liquid vaporization. Additionally, a portion of the fire heat input goes into sensible heat to raise the liquids temperature.

The vapor and liquid composition can change as vapors are released from the system. As a result, temperature and latent-heat values can change, affecting the required size of the PRD. On occasion, a multicomponent liquid can be heated at a pressure and temperature that exceed the critical temperature or pressure for one or more of the individual components. For example, vapors that are physically or chemically bound in solution can be liberated from the liquid upon heating. This is not a standard latent-heating effect but is more properly termed degassing or dissolution. Vapor generation is determined by the rate of change in equilibrium caused by increasing temperature.

For these cases involving vapors physically or chemically bound in solution, and other multicomponent mixtures that have a wide boiling range, such as or thermal cracking of high-boiling materials during an external fire scenario, it might may be necessary to develop a time-dependent model where the total heat input to the vessel not only causes vaporization but also raises the temperature of the remaining liquid, keeping it at its boiling range.

The recommended practice of finding a relief vapor flow rate from the heat input to the vessel and from the latent heat of liquid contained in the vessel becomes invalid near the critical point of the fluid, where the latent heat approaches zero and the sensible heat dominates. If no accurate latent heat value is available for these hydrocarbons near the critical point, a minimum value of 115 kJ/kg (50 Btu/lb) is sometimes acceptable as an approximation. If pressure-relieving conditions are above the critical point, the rate of vapor discharge typically depends on the rate at which the fluid expands as a result of the heat input because a phase change does not occur.

Reference [72] gives an example of a time-dependent model used to calculate relief requirements for a vessel that is exposed to fire and that contains fluids near or above the critical range.

Annex I Thermal Cracking of Refinery Crude Resid in Atmospheric and Vacuum Columns During Fire

This annex describes thermal cracking for fire on vessels containing heavy components (e.g. crude resid in refinery crude atmospheric columns, crude vacuum columns and other equipment). During a fire, the material may experience mild thermal cracking to produce low molecular weight gases and produce liquid and lighter hydrocarbon fluids (e.g. gasoline, distillate and gas oil) in vapor and liquid form.

Two towers typically found in refineries are the crude atmospheric column and crude vacuum column. The atmospheric column is the first major separation of crude oil and operates at the bottom of the column at 1 barg [15 psig] and approximately 700°F [370°C]. The <u>atmospheric tower bottoms</u> product is historically called <u>typically boils at 650°F [340°C] resid since that is the typical temperature</u>. The bottoms from the atmospheric column are fed to the vacuum column, which operates at approximately 0.4 psi-vacuum [0.027 bar-vacuum] and between 750-770°F [400-410°C]. The pressure depends on whether steam is used to lower the partial pressure. The vacuum <u>tower bottoms product</u> is historically called <u>typically boils at 1050°F [565°C] resid</u>, which is the typical temperature.

During a fire, the column pressure will increase to the relief pressure. For both columns, a typical relief pressure with 21% of external fire accumulation is between 60-90 psig [4.1-6.2 barg]. Thermal cracking is a function of time and temperature. If the resid is held at approximately 700°F [370°C] for 60

minutes, then 1% of the resid may crack. At higher temperatures, 1% of the resid will crack If the resid is held at approximately 750°F [400°C] for only 10 minutes.

The resid is expected to crack into liquid gas oil, distillate, gasoline and light gases.-<u>After cracking</u> begins, some of the fire heat input will be used for the thermal cracking and formation of lower <u>molecular weight products</u> A significant portion of fire heat input goes to cracking. The remaining fire heat input goes to sensible heat to increase the liquid temperature and vaporization of liquids to form vapor relief. Thus the relief rate is reduced. Since the cracked liquid products have lower boiling points, the relief temperature will be reduced <u>compared to direct boiling of uncracked components</u> (Reference [use reference "Quiroga" below])(Ref XX).

In contrast, if no thermal cracking is assumed to occur at 60 psig, the atmospheric crude resid relief temperature may reach approximately 900°F [480°C]. Similarly, the vacuum resid may reach approximately 1,300°F [700°C]. These temperatures <u>-may</u> require high-temperature metallurgy and piping stress analysis.

The following factors should be considered when determining the relief rate for fire during a cracking reaction:

- Initial Cconditions of temperature and time when the fire occurs and duration.
- Types of products and yields of products formed by cracking, including gases and liquids
- Calculation of heat of cracking reaction
- Distribution of the fire heat input into heat of cracking, sensible heat and heat of vaporization
- Determination of a<u>A</u>mount of cracked liquid vaporized
- The a<u>A</u>mount of relief due to gases cracked from the resid and the amount of relief vapor due to vaporization of cracked liquids

For estimation purposes, <u>approximate parameters</u> may be obtained from reference [use reference <u>"Quiroga" below]</u>, the following may be sometimes acceptable as approximate parameters. These are based a refinery atmospheric column with atmospheric resid at the given conditions and should be adjusted for different situations. The parameter estimations are from reference [use reference <u>"Quiroga" below]</u>.

- Relief pressure and temperature is established at 60 psig and 750°F
- Total relief rate [lb/hour] is the Fire Heat Input [btu/hour] divided by 350 [btu/lb of relief]
- Vapor relief molecular weight is 60 [lb/lb-mole]

Bibliography

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