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Thermodynamics and reaction mechanism of urea decomposition †

Steffen Tischer,^{*a} Marion Börnhorst,^b Jonas Amsler,^b Günter Schoch,^b and Olaf Deutschmann^{a,b}

The selective catalytic reduction technique for automotive applications depends on ammonia production from a urea-water solution by thermolysis and hydrolysis. In this process, undesired liquid and solid by-products are formed in the exhaust pipe. The formation and decomposition of these by-products have been studied by thermogravimetric analysis and differential scanning calorimetry. A new reaction scheme is proposed that emphasizes the role of thermodynamic equilibrium of the reactants in liquid and solid phases. Thermodynamic data for triuret have been refined. The observed phenomenon of liquefaction and re-solidification of biuret in the temperature range 193–230 °C is explained by formation of a eutectic mixture with urea.

1 Introduction

Air pollution by nitrogen oxides from Diesel engines is a main problem concerning environment and society. Therefore, governments follow the need to regulate emissions by law (e.g. 715/2007/EG, "Euro 5 and Euro 6").¹ The favored method to reduce nitrogen oxides is selective catalytic reduction (SCR) using ammonia as reducing agent. Due to security issues, passenger cars cannot be equipped with an ammonia tank. Consequently, ammonia is provided in form of a urea-water solution, which decomposes through thermolysis and hydrolysis when dispersively dosed into the hot exhaust pipe. During the desired decomposition of urea at temperatures above 130 °C undesired intermediates and by-products in liquid and solid form occur and stick to the wall of the exhaust pipe due to the inevitable spray-wall interaction.^{2–4} These condensed deposits in the exhaust pipe bear the risk of increased pressure drop and insufficient conversion to ammonia. Thus there is a need to understand the decomposition kinetics of urea and its by-products, that has been extensively studied by many authors.^{5–11} Common experimental methods include Thermogravimetric Analysis (TG), Differential Scanning Calorimetry (DSC), High Performance Liquid Chromatography (HPLC) and Fourier transform-infrared spectroscopy (FT-IR).

A first detailed description of urea decomposition behavior was proposed by Schaber et al.⁸ based on TG, HPLC, FT-IR and ammonium ISE (ion-selective electrode) measurements. Concluding

from experimental results and literature data, 23 possible reactions including urea and its by-products biuret, cyanuric acid, ammelide, ammeline and melamine are presented. Further, cyanate and cyanurate salts and cyanamide are proposed as possible intermediates of high temperature urea decomposition. Triuret production and decomposition is not accounted for in this study. The authors classify urea decomposition into four temperature regions. The first temperature regime from room temperature to 190 °C comprises urea melting and vaporization starting from 133 °C. With increasing temperature, urea decomposes to ammonia and isocyanic acid, the latter leading to biuret, cyanuric acid and ammelide formation. The second temperature region of 190–250 °C is dedicated to biuret decomposition accompanied by several side reactions forming cyanuric acid and ammelide. At 225 °C the melt is observed to be converted into a sticky, solid matrix, which is assumed to originate from ionic formations of different by-products without evidence. Besides small amounts of ammelide, ammeline and melamine, cyanuric acid is the main component observed at 250 °C. The third temperature range from 250 to 360 °C represents the sublimation and decomposition of cyanuric acid. Ammelide, ammeline and melamine are proposed to gradually decompose at temperatures above 360 °C marking the fourth temperature region. The authors state elimination of ammelide at 600 °C and ammeline at 700 °C. High temperature residues are not investigated.⁸

Eichelbaum et al.¹⁰ propose a reaction network for urea decomposition consisting of nine major reactions based on simultaneous TG/DTA (Differential Thermal Analysis) coupled with GC/MS (Gas Chromatography/Mass Spectrometry) gas analyses. Decomposition reactions of ammelide, ammeline and melamine are defined and total decomposition is observed for temperatures above 625 °C. However, the proposed reaction scheme lacks in

^a Institute of Catalysis Research and Technology (IKFT), Karlsruhe Institute of Technology (KIT), Karlsruhe, Germany. Fax: +49 721 608 44805; Tel: +49 721 608 42114; E-mail: steffen.tischer@kit.edu

^b Institute for Chemical Technology and Polymer Chemistry (ITCP), Karlsruhe Institute of Technology (KIT), Karlsruhe, Germany.

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description of relevant parallel and equilibrium reactions of urea by-products. Acceleration of urea pyrolysis by different metal exchanged zeolites is demonstrated.¹⁰ A more detailed reaction scheme derived by flow reactor experiments using FTIR spectroscopy for gaseous and HPLC for solid reaction product analysis covers 15 decomposition reactions.¹¹ For the first time, triuret production and decomposition is included and several reactions are proposed to be equilibrium reactions.

Based on the proposed reaction schemes, a first kinetic model for evaporation and decomposition of urea water solution was developed by Ebrahimian et al.¹² The model describes urea decomposition to ammonia and isocyanic acid and the equilibrium reaction forming biuret. Reactions from biuret to cyanuric acid and from cyanuric acid to ammelide and isocyanic acid are included. Ammelide is assumed to decompose to gaseous by-products.

Current kinetic models are based on the reaction network proposed by Bernhard et al.¹¹ and validated against TG and HPLC experimental data. The model includes formation and decomposition reactions of the most relevant by-products and reproduces the characteristic decomposition stages of urea adequately.¹³ However, important physical and chemical processes are not accounted for. A *biuret matrix* species is defined to cover the effect of solidification at 220 °C. Ammelide decomposition is modeled as sublimation while further high molecular by-products are not included in the model. Further, recent investigations have shown that the current kinetic model lacks in reproduction of mass transfer effects at the sample surface and in the prediction of reactions including gas phase species.

Thermogravimetric measurements have shown a strong influence of the experimental boundary conditions on urea decomposition kinetics. Besides the sample heating rate^{5,6,8}, decomposition behavior is highly sensitive to the geometric arrangement of the samples and respective crucibles.^{6,10} Increased surface area of the sample is assumed to accelerate mass transport of gaseous products.^{10,11} Further, the presence of water is stated to decrease by-product formation due to isocyanic acid hydrolysis.^{10,11}

In a very recent publication by Wang et al.¹⁴ the gaseous and condensed products of the decomposition of urea have been analyzed in detail for the temperature range from 132.5 to 190 °C by means of online mass spectrometry and liquid chromatography mass spectrometry, respectively. They derived a reaction scheme where biuret and cyanuric acid are formed directly by self-combination reactions of urea.

So far, available kinetic schemes for chemical decomposition are not suitable for accurate description of reactions in and between all phases. This is our motivation to systematically investigate the decomposition of urea and its subsequent products biuret, triuret, cyanuric acid, and ammelide by TG and DSC experiments. Our numerical model is based on a consistent thermodynamic description of all phases. Using these data, a new kinetic scheme of urea decomposition is proposed.

2 Experiments

Kinetic and thermodynamic data on the decomposition of urea and its by-products are derived from thermogravimetric (TG) analysis and differential scanning calorimetry (DSC). For thermo-

Table 1 Experiments used for this study

Type	Crucible	Substance	Initial weight (mg)	Ramp (K/min)
TG	plate	urea	6.23	2
TG	cylinder	urea	6.18	2
TG	cylinder	urea	60.3	2
TG	cylinder	urea	10.8	10
TG	cylinder	32.5 wt-% UWS	27.5	10
TG	plate	biuret	5.24	2
TG	cylinder	biuret	5.18	2
TG	cylinder	biuret	9.8	10
TG	cylinder	biuret	95.8	2 (to 195 °C)
TG	cylinder	biuret	94.6	2 (to 200 °C)
TG	cylinder	biuret	94.4	2 (to 210 °C)
TG	cylinder	triuret	6.98	2
TG	cylinder	triuret	10.31	10
TG	plate	cyanuric acid	5.87	2
TG	cylinder	cyanuric acid	5.37	2
TG	cylinder	cyanuric acid	10.1	10
TG	plate	ammelide	5.58	2
TG	cylinder	ammelide	8.72	2
TG	cylinder	ammelide	9.34	10
DSC	cylinder	urea	14.2	2
DSC	cylinder	biuret	12.8	2
DSC	cylinder	triuret	5.9	2
DSC	cylinder	cyanuric acid	16.4	2

gravimetric analysis, a Netzsch STA 409 C was equipped with the thermal controller TASC 414/2. The following standard procedure is conducted for each deposit sample and for reference measurements. Representative samples are ground and placed in a corundum crucible with an initial sample mass of 5–100 mg. The samples are heated from 40 to 700 °C at a constant heating rate of 2 or 10 K/min. TG analysis is performed in synthetic air (20.5% O₂ in N₂) using a purge gas flow rate of 100 mL/min. Different geometries of corundum crucibles are used to hold the samples during measurement: a cylinder-type crucible with an inner diameter of 6 mm and a height of 12 mm and a plate-type crucible of 15 mm inner diameter and a height of 5 mm. Cylinder geometry, sample mass and heating rate are systematically varied for different samples in order to derive a large database for kinetic modeling. Pure urea (Merck, ≥ 99.5%), biuret (Sigma Aldrich, ≥ 98%), triuret (Sigma Aldrich, ≥ 95%), cyanuric acid (Sigma Aldrich, ≥ 98%), ammelide (Dr. Ehrenstorfer GmbH, 99%), ammeline (Sigma Aldrich, 97.9%) and a 32.5 wt-% urea water solution (UWS) are used for the measurements. TG analysis gives information about the mass loss of a sample by evaporation and reactions under specified conditions and therefore gives information about the decomposition behavior.

DSC is a thermo-analytical measurement technique to determine the difference in heat required to increase the sample temperature compared to a reference sample. A Mettler DSC 30 is used to measure the thermal properties of urea and its by-products. The calorimetry is operated under air flow applying a heating rate of 2 K/min from 25 to 600 °C. Initial sample weight amounts to 10–15 mg. Aluminium crucibles are used as sample holder and reference. All other experimental conditions were kept the same as in TG experiments. This allows correlation of measured thermal and kinetic properties and yields valuable data for model development.

A list of all experiments used in this study is given in Table 1.



3 Numerical model

The TG and DSC experiments are simulated by a zero-dimensional batch-type reactor model. It was implemented as DETCHEM^{MPTR} (MPTR = multi-phase tank reactor) into the DETCHEMTM program package.¹⁵

The reactor model consists of a set of species S_i grouped into sets of phases P_j . Each species belongs to exactly one phase, i. e. a phase transition of a chemical substance is handled by two different species. Each species is associated with thermodynamic data in form of the NASA polynomials.¹⁶ The molar heat capacity, molar enthalpy and molar entropy are computed as function of temperature based on 7 coefficients $a_{1i} \dots a_{7i}$.

$$\frac{c_{p,i}}{R} = a_{1i} + a_{2i}T + a_{3i}T^2 + a_{4i}T^3 + a_{5i}T^4 \quad (1)$$

$$\frac{H_{m,i}}{R} = a_{1i}T + \frac{a_{2i}}{2}T^2 + \frac{a_{3i}}{3}T^3 + \frac{a_{4i}}{4}T^4 + \frac{a_{5i}}{5}T^5 + a_{6i} \quad (2)$$

$$\frac{S_{m,i}}{R} = a_{1i} \ln(T) + a_{2i}T + \frac{a_{3i}}{2}T^2 + \frac{a_{4i}}{3}T^3 + \frac{a_{5i}}{4}T^4 + a_{7i} \quad (3)$$

The molar volume of a species is either defined by ideal gas-law for gaseous species $V_{m,i} = RT/p$ or by assuming a constant density for condensed species $V_{m,i} = \rho_i/M_i$. The phases are separated. Each phase occupies a volume

$$V_j = \sum_{S_i \in P_j} V_{m,i} \quad (4)$$

The concentrations are expressed locally with respect to the corresponding phase, i. e. $c_i = n_i/V_j$. The reaction rates are mostly given in terms of an extended Arrhenius expression (see also appendix A in supplemental information). For reaction R_k we can write a molar rate

$$r_k = A_k T^{\beta_k} \exp\left(-\frac{E_{a,k}}{RT}\right) \prod_{S_i \in R_k} c_i^{\nu_{ik}} \quad (5)$$

Homogeneous reactions are reactions with reactants from the same phase. However, the products of homogeneous reactions may be released to a different phase. Let \dot{n}_{ik} be the rate of production of species S_i by reaction R_k . Thus,

$$\dot{n}_{ik} = V_j \nu_{ik} r_k \quad \text{reactants} \subset P_j. \quad (6)$$

Heterogeneous reactions are assumed to occur at the interface of two phases. The reactants can come from both phases, but can also come from only one of them. In our case, the contact area between the phases shall be the cross-sectional area A of the crucible, i. e. the phases are considered to be stacked on top of each other in the cylindrical reactor. For an Arrhenius type of reaction, the production rate is likewise

$$\dot{n}_{ik} = A \nu_{ik} r_k. \quad (7)$$

All phases are considered to be ideal mixtures. Thus, the chemical activity a_i of a species is p_i/p^\ominus for the gas-phase and the molar fraction for all other phases. The activities can also be ex-

pressed in terms of concentrations as $a_i = c_i/c_i^\ominus$ with a reference concentration $c_i^\ominus = p^\ominus/RT$ for gas-phase species and $c_i^\ominus = \rho_i/M_i$ for condensed species. Then the rate of a reverse reaction can be linked to the equilibrium constant

$$K_{p,k} = \exp\left(-\frac{\Delta R_k G^\ominus}{RT}\right), \quad (8)$$

$$K_{c,k} = K_{p,k} \cdot \prod_{S_i \in R_k} c_i^{\nu_{ik}}, \quad (9)$$

$$r_k^{\text{reverse}} = \frac{r_k}{K_{c,k}}. \quad (10)$$

A special case of a heterogeneous process is the phase transition between liquid and gas. For the condensation, we may assume that molecules hitting the phase boundary stick with an accumulation factor α_c . Thus, kinetic gas theory yields

$$r_k^{\text{condensation}} = \alpha_c \sqrt{\frac{RT}{2\pi M_i}} c_i^{\text{gas}}. \quad (11)$$

Applying the definition of the reverse rate (Eq. 10), we get the Herz-Knudsen equation for evaporation¹⁷

$$r_k^{\text{evaporation}} = \alpha_c \sqrt{\frac{RT}{2\pi M_i}} \frac{c_i^{\text{liq}}}{h} \quad (12)$$

with the Henry constant

$$h = \frac{\rho_i^{\text{liq}}}{\rho_i^{\text{vap}}} \frac{RT}{M_i}. \quad (13)$$

The batch-type reactor model consists of conservation equations for species and enthalpy:

$$\frac{dn_i}{dt} = \sum_{R_k} \dot{n}_{ik} \quad (14)$$

$$\frac{dH}{dt} = A k_W (T^{\text{extern}} - T) \quad (15)$$

where temperature and total enthalpy are linked by

$$H = \sum_{S_i} n_i \cdot H_i(T) \quad (16)$$

k_W is a heat transfer coefficient. Since both, the TG and DSC experiments are driven by an external temperature profile, the value of the heat transfer coefficients is not very sensitive. It just has to be finite, otherwise the solution would jump in case of phase transitions. For the simulation reported here a value of $k_W = 200 \text{ W m}^{-2} \text{ K}^{-1}$ has been chosen.

The system of differential-algebraic equations is solved by the solver LIMEX.¹⁸ The heat signal measured by DSC equals the (negative) enthalpy change of the condensed phases in the crucible. Since gas-phase reactions shall not be considered in this study, this enthalpy change equals the change of total enthalpy minus the power to heat the gas.

$$P^{\text{DSC}} = -\frac{dH}{dt} + \left(\sum_{S_i^{\text{gas}}} n_i c_{p,i} \right) \frac{dT}{dt} \quad (17)$$



4 Thermodynamic data

Previous models^{10,13} did not pay much attention to the thermodynamics of the system. TG experiments can be simulated in batch-type reactor models by kinetic models without solving a conservation equation for heat. With the availability of DSC data, thermodynamic aspects of the mechanism can be studied in detail. This was the motivation for the authors to gather information about the thermodynamic properties of all involved species.

For fundamental gaseous species, namely water H₂O(g) and ammonia NH₃(g), as well as for liquid water H₂O(l) the standard values from the thermodynamic database of DETCHEM¹⁵ have been used. For other species, values from literature were taken as given in Table 2. NASA coefficients for these species were fitted for a temperature range from 300 K to 800 K (see supplemental material).

Condensation of isocyanic acid

In the reaction mechanism by Brack et al.¹³, isocyanic acid plays a major role. Even though isocyanic acid evaporates at 23.5 °C, reactions occurring at temperatures exceeding 133 °C were formulated in the condensed phase. For a thermodynamically consistent formulation it is therefore necessary to look at the gas-liquid-equilibrium of isocyanic acid. Both are in equilibrium if

$$\mu_{\text{HNCO(l)}} = \mu_{\text{HNCO(g)}} \quad (18)$$

$$\mu_{\text{HNCO(l)}}^0 = \mu_{\text{HNCO(g)}}^0 + RT \ln \left(\frac{p_{\text{HNCO}}^{\text{vap}}}{p^\ominus} \right) \quad (19)$$

where $\mu_i^0 = H_{m,i} - TS_{m,i}$ is the chemical potential of the undiluted species. From the temperature dependency of the chemical potential of HNCO(l), we can derive the coefficients of the thermodynamic polynomials by linear regression. The vapor pressure follows an Antoine equation based on experimental data by Linhard et al.²⁸

$$\log_{10} \left(\frac{p_{\text{HNCO}}^{\text{vap}}}{\text{bar}} \right) = 4.69 - \frac{1252.195}{T/\text{K} - 29.167} \quad (20)$$

Aqueous phase

Urea, ammonia and isocyanic acid were also considered to be soluble in liquid water. In our model, these species form an ideally mixed aqueous phase (aq). Using literature data about the solubility, thermodynamic data were derived. At point of saturation, the chemical potentials of a species must be the same in both phases, i. e. for urea:

$$\mu_{\text{urea(aq)}} = \mu_{\text{urea(s)}} \quad (21)$$

$$\mu_{\text{urea(aq)}}^0 + RT \ln(X_{\text{urea(aq)}}) = \mu_{\text{urea(s)}}^0 \quad (22)$$

$$\mu_{\text{urea(aq)}}^0 = \mu_{\text{urea(s)}}^0 - RT \ln(X_{\text{urea(aq)}}) \quad (23)$$

Here, it is assumed that the mixture is ideal without interaction of the molecules (fugacity factor is unity). Experimental data^{29–32} were fitted in a temperature range of 0–70 °C yielding the satu-

ration mass fraction:

$$Y_{\text{urea(aq)}}(T) = 5.1687 \cdot 10^{-3} \cdot \frac{T}{\text{K}} - 1.0108 \quad (24)$$

There are no experimental data for higher temperatures. The simulation, however, requires a continuous polynomial throughout the entire interval (300–800 K). The extrapolation obeys two constraints: a non-negative heat capacity for the new species and an upper limit for the solubility to assure a mole fraction of urea(aq) below unity.

Likewise, the chemical potentials of NH₃(aq) and HNCO(aq) were determined. In these cases the reference chemical potentials are given by their respective gas-phase species.

$$\mu_{i(\text{aq})}^0 = \mu_{i(\text{g})}^0 - RT \ln(X_{i(\text{aq})}) \quad (25)$$

The temperature dependency of solubility of ammonia was fitted to literature data^{33–35} for temperature interval from 0 to 100 °C.

$$Y_{\text{NH}_3}(T) = 1.055 \cdot 10^{-7} \cdot \left(\frac{T}{\text{K}} \right)^3 - 6.81 \cdot 10^{-5} \cdot \left(\frac{T}{\text{K}} \right)^2 + 6.549 \cdot 10^{-3} \cdot \frac{T}{\text{K}} + 1.61 \quad (26)$$

For isocyanic acid the temperature dependency of Henry's coefficient $K_{\text{H}} = c_{\text{HNCO(aq)}}/p_{\text{HNCO(g)}}$ was approximated for 25 to 100 °C by van't Hoff's extrapolation³⁶

$$K_{\text{H}}(T) = K_{\text{H}}^\ominus \cdot \exp \left(\frac{\Delta_{\text{sol}}H^\ominus}{R} \left(\frac{1}{T} - \frac{1}{T^\ominus} \right) \right) \quad (27)$$

with a reference Henry coefficient³⁷ $K_{\text{H}}^\ominus = 26 \text{ mol L}^{-1} \text{ atm}^{-1}$ and enthalpy of solution $\Delta_{\text{sol}}H^\ominus = -34 \text{ kJ mol}^{-1}$.

Melting of urea and biuret

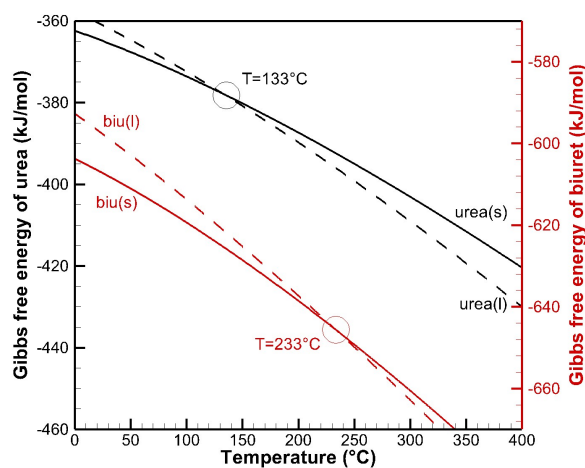


Fig. 1 Gibbs free energy of urea and biuret

With the literature data in Table 2 we can look at the phase change from solid to liquid. For the pure substances the chemical potential equals the Gibbs free energy. The phase transition



Table 2 Thermodynamic properties of secondary products of urea decomposition in different phases (g=gas, l=liquid, s=solid)

Species	Symbol(Phase)	$\Delta_f H^\ominus$ (kJ/mol)	S^\ominus (J/mol/K)	c_p (J/mol/K)	Reference
Water	H ₂ O(l)	-285.828	69.939	4 th order polynomial (75.351 at 298 K)	15,16
	H ₂ O(g)	-241.825	188.828	4 th order polynomial (33.588 at 298 K)	15,16
Ammonia	NH ₃ (g)	-45.567	192.474	4 th order polynomial (34.597 at 298 K)	15,16
Urea	urea(s)	-333.599	105.9	93	19,20
	urea(l)	-319.7	140.15	93	21
	urea(g)	-235.55	282.94	$26.84 + 0.2T - 1 \cdot 10^{-4}T^2$	19
Biuret	biu(s)	-563.70	146.1	131.3	21,22
	biu(l)	-537.06	203.27	93	21,22
	biu(g)	-437.30	354.31	$72.198 + 0.237T - 8.847 \cdot 10^{-5}T^2 - 1.455 \cdot 10^6T^{-2}$	19
Triuret	triu(s)	-746.7	146.1	131.3	23
Cyanuric acid	cya(s)	-703.5	142.20	130.0	24,25
	cya(g)	-564.1	339.37	$15.74 + 0.42T - 2 \cdot 10^{-4}T^2$	24
Isocyanic acid	HNCO(g)	-101.7	238.229	$15.34 + 0.126T - 9.523 \cdot 10^{-5}T^2 + 6.49 \cdot 10^4T^{-2}$ ($T \leq 400K$) $45.24 + 0.0307T - 7.619 \cdot 10^{-6}T^2 - 8.96 \cdot 10^5T^{-2}$ ($T \geq 400K$)	19
Ammelide	amm(d)(s)	-492.9	149.10	$94.958 + 0.262T - 8.444 \cdot 10^{-5}T^2 - 2.639 \cdot 10^6T^{-2}$	26
	amm(d)(g)	-303.9	429.87	$15.74 + 0.42T - 2 \cdot 10^{-4}T^2$	27
Ammeline	ammn(s)	-300.0	149.10	$94.958 + 0.262T - 8.444 \cdot 10^{-5}T^2 - 2.639 \cdot 10^6T^{-2}$	26

occurs when

$$\mu_{i(s)} = \mu_{i(l)} \quad (28)$$

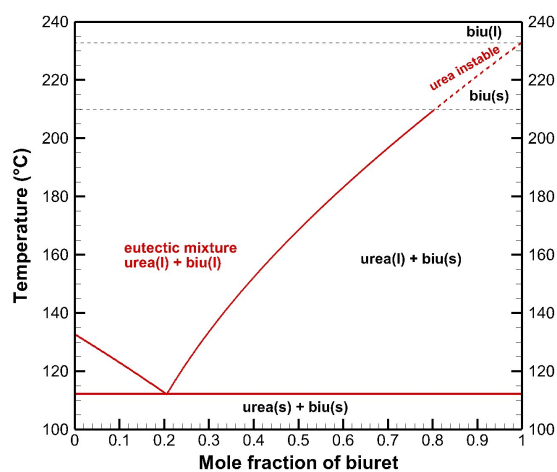
Figure 1 plots the Gibbs free energy of urea and biuret in solid and liquid phases. As expected for urea, the two graphs intersect at $T = 133$ °C. However, in case of biuret the predicted melting temperature is 233 °C. This seems to be contradictory, because the common literature value^{8,38} is between 190 and 193 °C. However, it was also observed by Schaber⁸ and Brack¹³ that biuret starts decomposing at 193 °C when it turned into a liquid. The decomposition slowed down around 210 °C and the melt formed a foam-like structure. Between 220 and 230 °C the consistence was described as a matrix-like aggregate before a second decomposition step sets in. Analysis of the substance revealed that it still mainly consisted of biuret. Thus Brack introduced two phases of biuret, namely biu(melt) and biu(matrix) to model this behavior.

The authors want to give a thermodynamic explanation for this observation. It can be explained by an eutectic mixture of urea and biuret. For varying mole fraction, we can plot the temperatures where

$$\mu_{i(s)}^0 = \mu_{i(l)}^0 + RT \ln(X_{i(l)}) \quad (29)$$

Figure 2 shows the phase diagram of the binary urea/biuret system. It agrees well with the results by Voskov et al.³⁹ This means that biuret is a liquid at 193 °C if the liquid phase consists of 67% biuret and 33% urea. As decomposition or evaporation of urea becomes faster at temperatures above 210 °C, biuret will become solid again. As we will see later, there is another thermodynamic argument for the apparent melting point of biuret.

This interpretation yields a qualitative understanding of experimental DSC data for urea and biuret. Figure 3 shows a comparison of the DCS signals. Together with TG data, we can identify several processes. The sharp peak at 133 °C indicates the melting of urea. The decomposition of urea starts more or less with the phase change. Biuret as a reaction product will remain in the liquid phase. At 193 °C the decomposition process also starts for biuret. The reaction slows down at around 210 °C, when a foam-like structure is formed. Presumably, there is no liquid urea present anymore. Endothermic reactions may lead to follow-up

**Fig. 2** Phase diagram of an eutectic mixture of urea and biuret

products like triuret, cyanuric acid or ammelide. Around 230 °C, the biuret becomes liquid again and a second decomposition step is observable in TG measurements. The third decomposition step between 330 and 400 °C is associated with the sublimation of cyanuric acid. The remaining solid substances decompose at temperatures above 400 °C, for which we include a reaction path through ammelide, but this shall not be the focus of this paper.

5 Reaction mechanism

Cyanuric acid

In Brack's model¹³ cyanuric acid decomposes directly to isocyanic acid ($\text{cya}(s) \rightarrow 3\text{HNCO}(g)$) by a zero-th order reaction. In order to investigate this hypothesis, TG and DSC experiments were run with pure cyanuric acid as initial sample. The TG experiment (Fig. 4) was conducted using two crucibles with different diameters. If the reaction were a zero-th order reaction in the homogeneous phase, the size of the crucible would not matter. However, the reaction is faster in a flat crucible with larger surface area than in a tall crucible. This suggests that the reaction



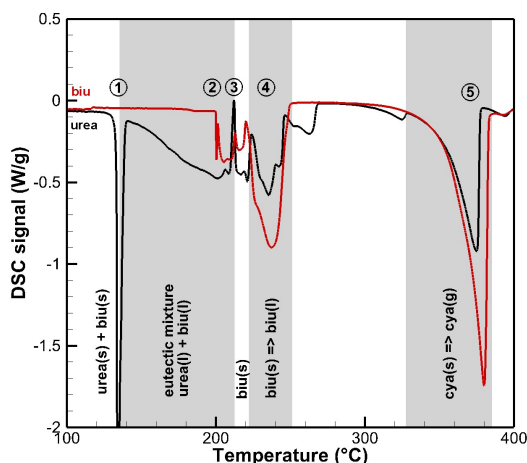


Fig. 3 Qualitative comparison of DSC signals for urea and biuret decomposition (1 - melting of urea, 2 - melting of biuret in a eutectic mixture, 3 - urea disappears, 4 - melting of biuret, 5 - sublimation of cyanuric acid)

takes place at the surface of the sample. Indeed, both TG curves can be modeled by a first-order surface reaction with an activation energy of 141.3 kJ/mol. However, this immediately raises the question about the resulting products, because the direct decomposition to isocyanic acid is an endothermic process consuming 352 kJ/mol. A comparison between experimental and expected DSC signal (Fig. 5) shows that for a direct decomposition the DSC signal should be larger by a factor of 2.5. The activation energy obviously agrees with the heat of sublimation of cyanuric acid. Therefore, it must be concluded that cyanuric acid evaporates directly into the gas-phase.

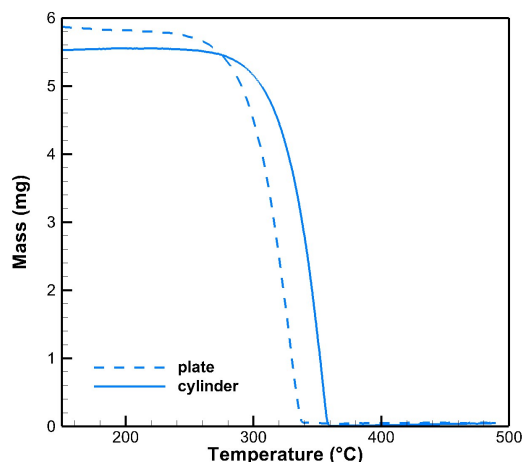
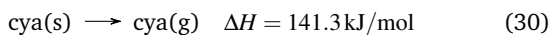


Fig. 4 Experimental TG data for decomposition of cyanuric acid in cylinder (diameter 6 mm) and plate-type (15 mm) crucibles

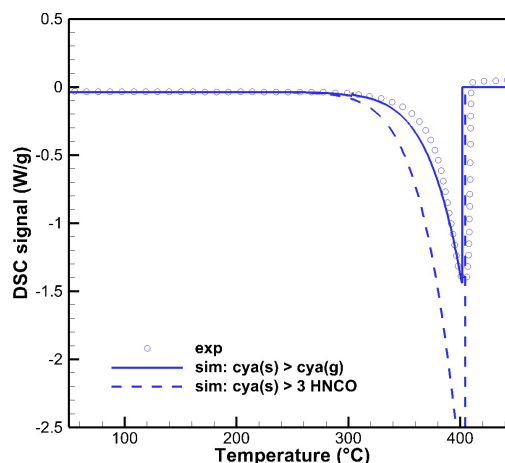


Fig. 5 Comparison of DSC signal of decomposition of cyanuric acid (model 1: sublimation, model 2: direct decomposition to isocyanic acid)

Triuret

In the previous mechanism by Brack¹³, triuret was a side product of biuret reacting with isocyanic acid. It could have been omitted without significantly changing the predicted amounts of cyanuric acid or ammelide. Looking at the chemical structures of the latter two, there is good argument that triuret may decompose to cyanuric acid by separation of ammonia and to ammelide by separation of water. Thus, instead of going directly from biuret to cyanuric acid and ammelide by complex multi-molecule reactions, we may assume that triuret is an intermediate species of these processes.

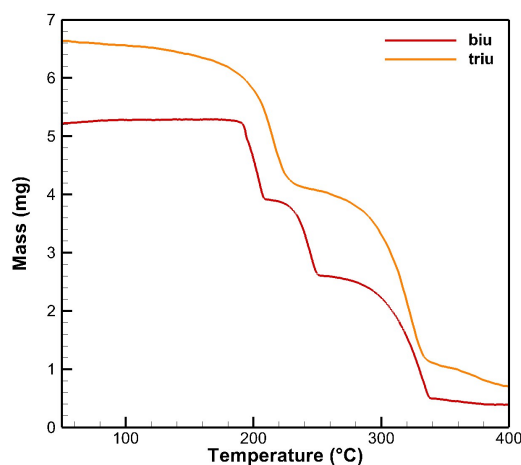


Fig. 6 Experimental TG data for decomposition of biuret and triuret

Figures 6 and 7 show a comparison of experimental TG and DSC data for biuret and triuret. It can be noted that there are parallels in decomposition steps for both substances. A major decomposition step is observed for temperatures above 192 °C. The mass loss is mainly due to the release of isocyanic acid and am-



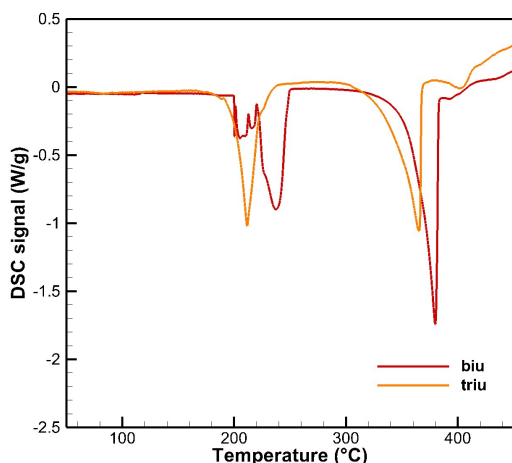


Fig. 7 Experimental DSC data for decomposition of biuret and triuret

monia. For biuret, as stated before, a second decomposition step appears at around 235 °C. This is not present for triuret. The possible reactions to cyanuric acid and ammelide would both yield a mass loss of about 12% and almost no heat signal of 8 and 22 kJ/mol, respectively. The decomposition step beyond 260 °C can mainly be associated with the sublimation of cyanuric acid. A shift in temperature can be observed with respect to the decomposition steps of both experiments. This is attributed to the difference of the initial sample mass.

Triuret is a highly unstable substance. The TG data (Fig. 6) already show a mass loss for temperatures below 192 °C. Since there is no significant DSC signal (besides the heat capacity of triuret) for these low temperatures, the mass loss might have been due to kinetically limited reactions to isocyanic acid or ammelide, which are almost isoenthalpic. However, all fits of kinetic parameters over-predicted the produced amounts of these two products for higher temperatures. Therefore it was concluded that the underlying process is a thermodynamically controlled equilibrium. Using the program DETCHEM^{EQUIL}¹⁵, equilibrium calculations for an ensemble consisting of urea(s), urea(l), biu(s), biu(l), triu(s), HNCO(l), HNCO(g) and N₂ were carried out. According to the literature data (Table 2), triuret would have remained stable up to 192 °C. However, literature data for triuret are very insufficient. Thus, we tried to reverse-engineer thermodynamic data for triuret, by fitting the values of Gibbs free energy of triuret in such a way that the calculated equilibrium composition shows the same mass loss (due to release of HNCO(g)) as the TG experiment (Fig. 6). The calculation becomes sensitive to the dilution of HNCO(g) by inert gas (e. g. N₂). Figure 8 shows the results of the fitting for varying degrees of dilution. For the calculation of thermodynamic coefficients for triu(s), we have chosen a dilution of 90%, i. e. the initial mixture consisted of 1 mol triuret and 9 mol N₂. As result we get standard enthalpy of formation $\Delta_f H^\ominus = -708.9$ kJ/mol and standard entropy $S^\ominus = 359.1$ J/mol/K. The molar heat capacity was kept constant at $c_p = 131.3$ J/mol/K.

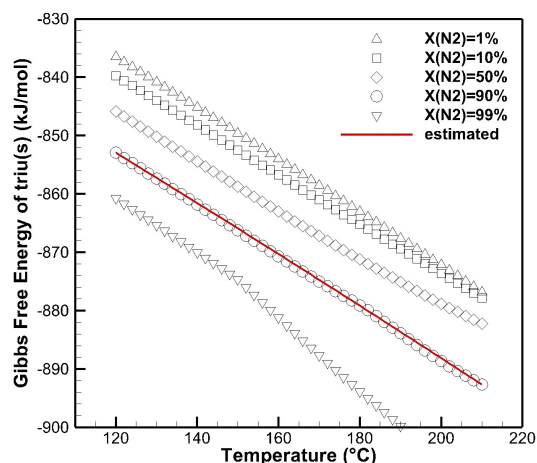


Fig. 8 Fitting of thermodynamic data of triuret such that equilibrium calculation yield the same mass loss as TG experiments

Biuret

As seen for triuret, we cannot look at the decomposition of one substance independently. One always has to keep in mind the thermodynamical ensemble consisting of isocyanic acid, urea, biuret and triuret. In order to better understand the processes of biuret decomposition for temperatures between 193 and 210 °C, we conducted TG experiments where biuret was heated up to 195, 200 and 210 °C, respectively. Initially a constant temperature ramp of 2 K/min was applied. After reaching the target temperature, it was kept constant. Figure 9 shows the result for the case 195 °C.

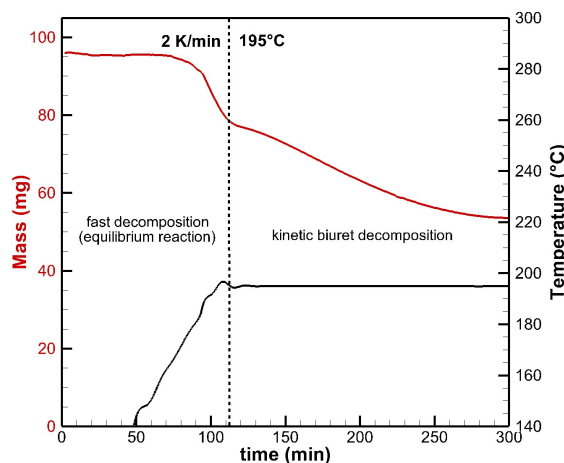


Fig. 9 Experimental TG data for biuret that was first heated to 195 °C with a constant heating ramp and then temperature was kept constant

For this sample, decomposition already starts below 193 °C. Most notable feature of this graph is that its slope is larger for the temperature ramp than for the case of constant temperature. This indicates that the decomposition is not only kinetically con-



trolled. Otherwise the reaction rate (and thus the slope of the graph) should be lower for lower temperatures. Therefore again, the mass loss during the temperature ramp is dominated by thermodynamic equilibrium processes. A fact that is not well taken into account in experiment design is that this equilibrium depends on the dilution of $\text{HNCO}(\text{g})$ in the gas phase. Thus, the decomposition of biuret may start for temperatures as low as $170\text{ }^\circ\text{C}$ for high purging rates with inert gas. On the other hand, if we calculate the equilibrium composition based on a lower dilution for an initial mixture of 1 mol biuret and 1 mol nitrogen using $\text{DETCHEM}^{\text{EQUIL}}^{15}$, we get a surprisingly nice result (Figure 10). Here, we see a phase change at $192.5\text{ }^\circ\text{C}$ when an eutectic mixture of liquid biuret and urea plus some triuret and gaseous isocyanic acid becomes thermodynamically favored. This explains the apparent melting point of biuret in literature.

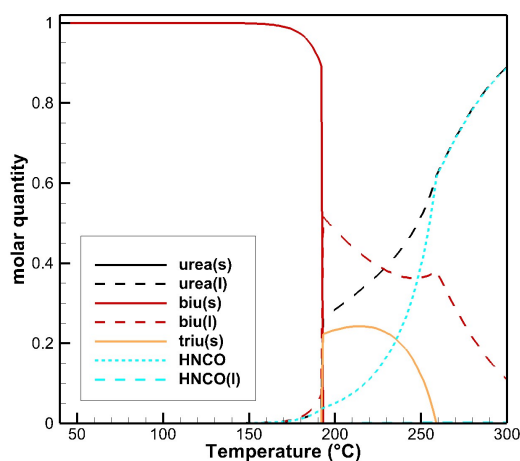
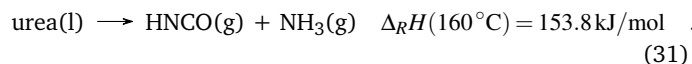


Fig. 10 Equilibrium composition of an initial mixture of 1 mol biuret and 1 mol N_2

Urea

Finally, we need to look at the initiation of urea decomposition, which starts with melting at $132\text{ }^\circ\text{C}$. Figure 11 shows measured TG and DSC data. Between 140 and $180\text{ }^\circ\text{C}$ the apparent reaction rate increases nearly linearly. According to the Brack mechanism¹³ this is mainly due to the reaction $\text{urea}(\text{l}) \rightarrow \text{HNCO}(\text{l}) + \text{NH}_3(\text{g})$. At this temperature, the vapor pressure of isocyanic acid is already more than 26 bar according to Equation 20. Unless $\text{HNCO}(\text{l})$ reacts very fast to form biuret or triuret, it would evaporate, leaving no caloric contribution of $\text{HNCO}(\text{l})$ in a DSC experiment. A direct decomposition would lead to a net reaction



According to Figure 11, we have a mass loss of 2.25 mg ($3.75 \cdot 10^{-5}$ mol urea) for the temperature interval 140 – $180\text{ }^\circ\text{C}$. The DSC signal increases nearly linearly from 0.13 to 0.375 W/g within 20 min. This accounts for a total heat consumption of

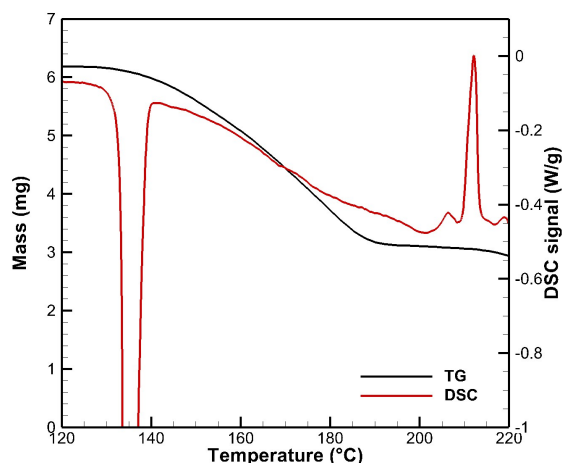
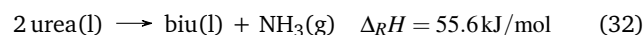


Fig. 11 Comparison of TG and DSC data for the initiation of urea decomposition

1.87 J . However, the direct decomposition would have required 5.77 J . All numerical simulations containing a direct decomposition reaction showed that TG and DSC signal could not be satisfied at the same time. They differed by a factor of 3. Thus, we had to discard the direct decomposition reaction.

Several ideas have been tried to initiate urea decomposition. Since the rate is sensitive to the size of the crucible, it was concluded that the process occurs at the gas-liquid interface. The net reaction that best fitted both, the DSC and the TG data in the interval 140 – $180\text{ }^\circ\text{C}$ was



as was also recently suggested by Wang et al.¹⁴

Kinetic parameters

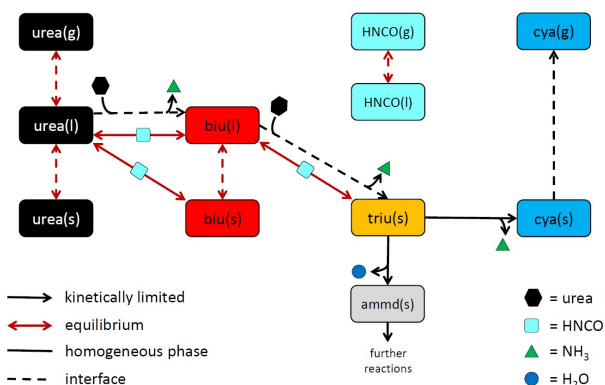


Fig. 12 Proposed reaction scheme of urea decomposition

As a result of the aforementioned investigations, the authors came up with the following reaction scheme (see Figure 12): The urea decomposition is mainly driven by a thermodynamic equilibrium in the system consisting of urea, biuret, triuret and isocyanic



acid. Ammonia is produced by a kinetically limited process with the net reaction $2 \text{urea(l)} \rightarrow \text{biu(l)} + \text{NH}_3(\text{g})$. In analogy, a reaction to form triuret has been added: $\text{biu(l)} + \text{urea(l)} \rightarrow \text{triu(s)} + \text{NH}_3(\text{g})$. Triuret then reacts further to form the solid deposits cyanuric acid and ammelide. Cyanuric acid sublimates for temperatures above 300 °C. This process is described here by a non-reversible surface process. However, it would make sense to describe it as an equilibrium process if reliable thermodynamic data for solid cyanuric acid would be available. For sake of completeness, 6 global reaction steps for the decomposition of ammelide were added (see supplemental material).

The kinetic parameters were adjusted manually until the main features of DSC and TG experiments were satisfied. A following automated parameter optimization did not significantly improve the results. Therefore, we use the manually fitted parameters for this publication (Table 3) in order to avoid pretending higher precision of the kinetic parameters. The biggest uncertainty is still the thermodynamic data of biuret and triuret.

6 Simulation results

The quality of the proposed reaction mechanism can be evaluated by comparison of the experimental data with the simulation results. Here, we will give results for some of the experiments listed in Table 1. The other cases will be shown in supplemental material.

One of the consequences of the equilibrium processes involving isocyanic acid is that the model becomes sensitive to the gas-phase concentration of HNCO(g) . This has not been taken into account during reactor model development when a zero-dimensional approach was chosen to describe TG experiments. The concentration gradient in the gas phase should be modeled as well. To guarantee a sufficient dilution of the evaporating HNCO(g) , the gas-phase volume was assumed to be a cylinder with the diameter of the crucible and a height of 2 m.

Cyanuric acid

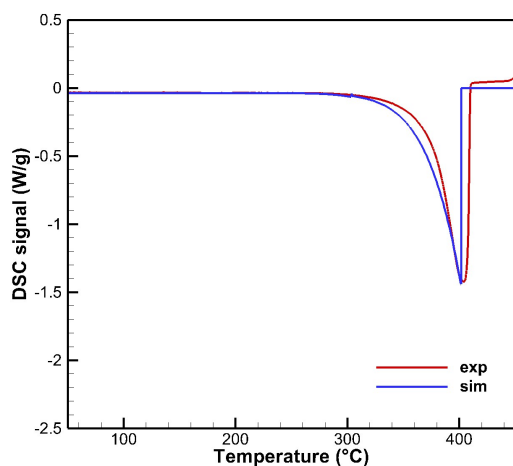


Fig. 13 DSC of cyanuric acid (cylinder crucible)

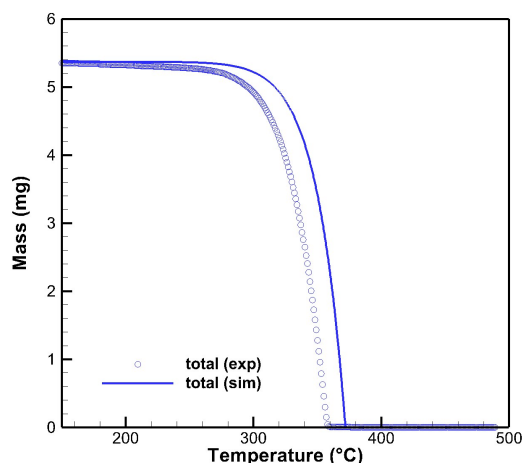


Fig. 14 TG of cyanuric acid (cylinder crucible, 2 K/min) with simulated composition of the mixture

The experiments with cyanuric acid are well modeled by the sublimation reaction. The DSC signal (Fig. 13) and TG data (Fig. 14) show good agreement. For the plate-type crucible, the sublimation is shifted correctly by ca. 40 K to lower temperatures. In reality, the dependency on the diameter of the crucible is less pronounced than in the simulation, because the surface area of the solid sample does not decrease linearly with the size of the crucible. However, if the model is applied to thin layers of deposits of cyanuric acid, it should be valid that the rate of sublimation is proportional to the size of the deposit.

Triuret

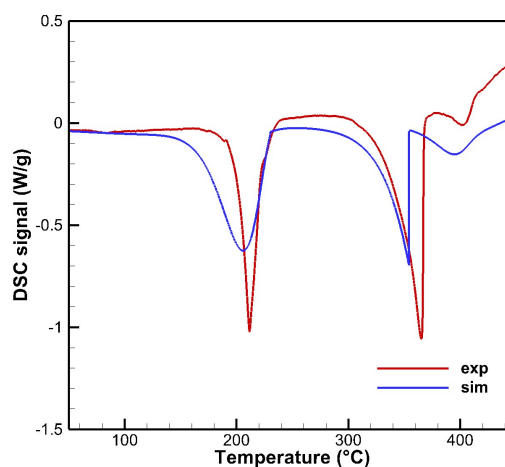


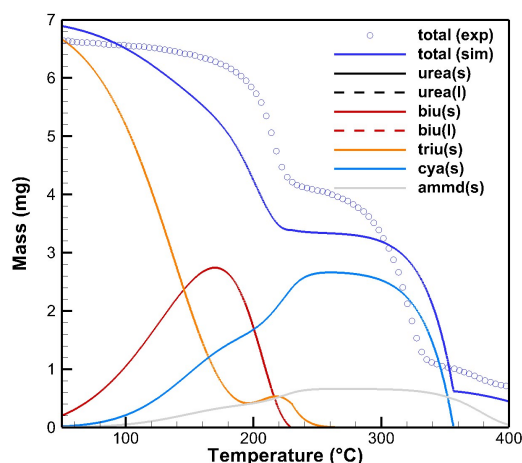
Fig. 15 DSC of triuret (cylinder crucible)

A comparison of experimental data and simulation results for triuret decomposition is shown in Figures 15 and 16. In both diagrams, two main decomposition steps can be identified: a step involving biuret for temperature between 190 and 230 °C and



Table 3 Kinetic parameters of the proposed reaction scheme of urea decomposition

Reaction	A_k (in SI units)	β_k	$E_{a,k}$ (in kJ/mol)	Notes
Homogeneous phase:				
urea(l) + HNCO(l) \rightleftharpoons biu(s)	$1 \cdot 10^{-4}$	0	0	These three reactions feature HNCO(l) even though the net product of the reverse reaction is HNCO(g).
urea(l) + HNCO(l) \rightleftharpoons biu(l)	$1 \cdot 10^{-4}$	0	0	
biu(l) + HNCO(l) \rightleftharpoons triu(s)	$1 \cdot 10^{-4}$	0	0	
triu(s) \rightarrow cya(s) + NH ₃ (g)	$1.2 \cdot 10^2$	0	45	
triu(s) \rightarrow ammd(s) + H ₂ O(g)	$3 \cdot 10^1$	0	45	
Surface:				
2 urea(l) \rightarrow biu(l) + NH ₃ (g)	$3.5 \cdot 10^0$	0	99	
biu(l) + urea(l) \rightarrow triu(s) + NH ₃ (g)	$2 \cdot 10^2$	0	116.5	
cya(s) \rightarrow cya(g)	$3 \cdot 10^4$	0	141.3	
Phase change (surface):				
H ₂ O(g) \rightleftharpoons H ₂ O(l)	$8.6 \cdot 10^{-2}$	0.5	0	The condensation-evaporation equilibria are based on Herz-Knudsen model (Eq. 11).
urea(g) \rightleftharpoons urea(l)	$4.7 \cdot 10^{-2}$	0.5	0	
HNCO(g) \rightleftharpoons HNCO(l)	$5.5 \cdot 10^{-2}$	0.5	0	Adjusted to DSC signal of urea melting.
urea(l) \rightleftharpoons urea(s)	$1 \cdot 10^{-6}$	0	0	
biu(l) \rightleftharpoons biu(s)	$1 \cdot 10^{-6}$	0	0	

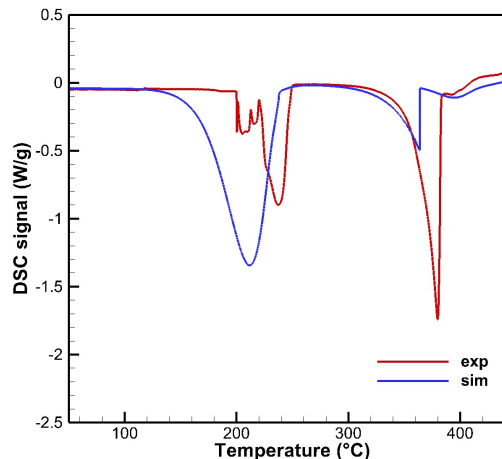
**Fig. 16** TG of triuret (cylinder crucible, 2 K/min) with simulated composition of the mixture

the sublimation of cyanuric acid. The simulation broadens the first step to lower temperatures. This is because the TG data up to 200 °C have been used to adjust thermodynamic data of triuret on the assumption of an equilibrium in the urea-biuret-triuret-HNCO ensemble. However, this does not account for the mass loss due to the kinetically limited reactions that release ammonia. Therefore, the total mass loss using the full mechanism is overestimated and shifted to lower temperatures. Here, a better agreement could only be achieved by improving the thermodynamic data of biuret and triuret.

The simulation of TG experiments (Fig. 16) also gives an insight into the composition of the mixture in the crucible. No significant amounts of liquid urea seem to be formed. Therefore, biuret also stays solid until complete decomposition at around 230 °C. Since the first decomposition step is slightly overestimated, the amount of cyanuric acid is then underestimated. Ammelide formation is matched well when assuming that production of cyanuric acid and ammelide are in the ratio 4:1. The TG results show even better agreement for a temperature ramp of 10 K/min

because thermodynamically controlled equilibrium processes are more dominant than kinetically controlled reactions.

Biuret

**Fig. 17** DSC of biuret (cylinder crucible)

Unfortunately, no good agreement could yet be achieved for biuret (Figures 17 and 18). The interplay of thermodynamics and reaction kinetics does not resolve the effects in Fig. 3. There should be three decomposition steps: the eutectic urea-biuret phase, the biuret-triuret phase and the sublimation of cyanuric acid. For the same reason as the triuret decomposition is overestimated at lower temperatures, the triuret production is now underestimated in the first reaction step. Thus, the triuret decomposition does not contribute much to the TG signal in the simulation, even though triuret is present at the expected temperature range 210–230 °C. Thermodynamic data should shift here the equilibrium of urea-biuret-triuret-HNCO towards triuret. Better agreement was achieved for the 10 K/min experiment when decomposition is shifted to higher temperatures and the two decomposition steps also merge in reality.



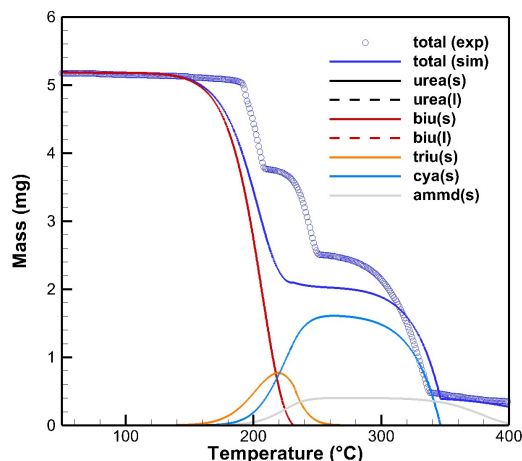


Fig. 18 TG of biuret (cylinder crucible, 2 K/min) with simulated composition of the mixture

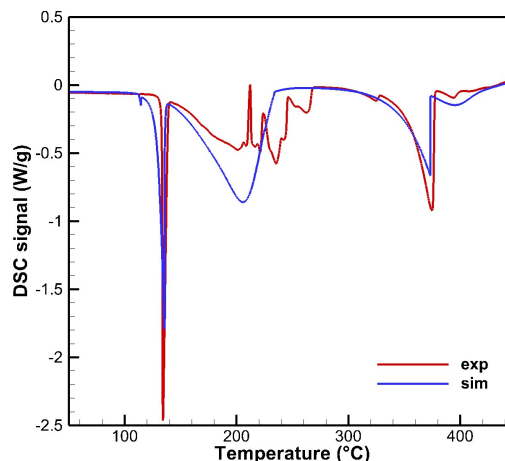


Fig. 20 DSC of urea (cylinder crucible)

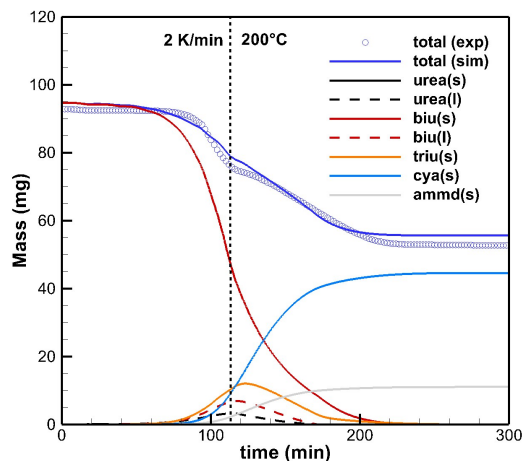


Fig. 19 TG of biuret maintaining a constant temperature of 200 °C

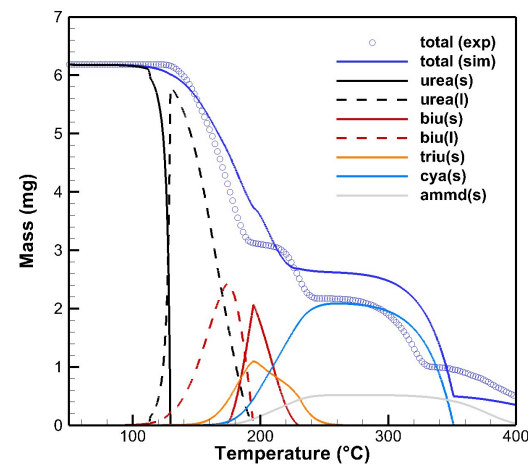


Fig. 21 TG of urea (cylinder crucible, 2 K/min) with simulated composition of the mixture

On the other hand, simulations of TG experiments with biuret where heating was stopped at 195, 200 and 210 °C show very good agreement (Fig. 19). It should be noted that these experiments used a 15 times higher initial mass of biuret. Thus, the contribution of volumetric equilibrium reactions is increased over the ammonia producing surface reactions.

Urea

For practical simulations of SCR systems, we are mainly interested in deposits resulting from urea decomposition. Here, the situation looks better than for biuret. Experiments show three decomposition steps: a step in liquid urea-biuret mixture, a biuret-triuret step and the sublimation of cyanuric acid. The initiation of the decomposition is well matched in DSC and TG (Figures 20 and 21). The biuret-triuret step occurs in simulation again shifted to lower temperatures, but there is a phase change at around 200 °C,

when all urea is consumed. The produced amounts of biuret and triuret seem to be in the right order of magnitude, leading to correct predictions for cyanuric acid and ammelide. The predicted results again become even better for the 10 K/min ramp, when the first and the second decomposition step merge also in experiment.

7 Conclusions

At a first glance, there still seems to be too much uncertainty. The previous mechanism by Brack et al.¹³ was able to model the three decomposition steps of urea at the right temperature intervals. However, the proposed reaction scheme is advantageous in several ways:

First, it is stringently based on thermodynamics. Melting and solidification of substances are inherent to the thermodynamic data. The phase change during biuret decomposition can be explained by formation of an eutectic mixture. There is no need for



an extra matrix species.

Second, it is consistent with DSC experiments. From these, we can conclude that cyanuric acid does not decompose to isocyanic acid directly, but it sublimates instead. Furthermore we can also rule out direct decomposition of urea to gaseous ammonia and isocyanic acid.

Third, it scales correctly with crucible size and gradient of the temperature ramp. Thus, we have to distinguish between processes that are controlled by thermodynamic equilibrium and kinetically controlled surface reactions.

Last, but not least, the overall reaction scheme is much simpler. In the tradition of Occam's razor, it explains more effects with less assumptions compared to the Brack mechanism.

The reaction mechanism is not perfect yet. The main reason is the lack of data for the thermodynamic potentials of biuret and triuret. Eventually, a liquid triuret species should be added, because it should be part of the eutectic mixture. Currently, triuret decomposition is shifted to lower temperatures. This causes a coincidence of the first and second decomposition step for biuret.

For urea decomposition the results are in good agreement with the measurements by Wang et al.¹⁴ The dominant process for biuret and triuret formation is the self-combination of urea, which leads to the desorption of ammonia. However, this is a bit unsatisfying from a viewpoint of elementary reaction step mechanisms. According to the DSC signal, this reaction step cannot involve HNC(O). A more elegant explanation might be a route through an ionic reaction urea \rightarrow NH_4^+ + OCN^- as suggested by Kieke et al.⁴⁰

To summarize, the authors propose a new reaction mechanism for the urea decomposition, which is closer to the real processes than the previously published mechanisms. We want to emphasize the importance of thermodynamics in modeling of deposit formation in SCR systems.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

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Nomenclature

A	Cross-sectional area of the crucible (m^2)
A_k	Pre-exponential factor ($\text{mol}/\text{m}^3/\text{s}$ or $\text{mol}/\text{m}^2/\text{s}$)
a_i	Chemical activity coefficient (-)
a_{ni}	Coefficient of NASA polynomials
c_i	Concentration (mol/m^3)
c_i^\ominus	Reference concentration at normal pressure (mol/m^3)
$c_{p,i}$	Molar heat capacity ($\text{J}/\text{mol}/\text{K}$)
$E_{a,k}$	Activation energy (J/mol)
H	Enthalpy (J)

$H_{m,i}$	Molar enthalpy (J/mol)
h	Henry constant (-)
$K_{c,k}$	Equilibrium constant in terms of concentrations (variable units)
$K_{p,k}$	Equilibrium constant (-)
K_H	Henry's coefficient ($\text{mol}/\text{m}^3/\text{Pa}$)
k_W	Heat transfer coefficient ($\text{W}/\text{m}^2/\text{K}$)
M_i	Molar mass (kg/mol)
m	Mass (kg)
n_i	Molar amount (mol)
\dot{n}_{ik}	Molar rate of species S_i by reaction R_k (mol/s)
P_j	Phase
P^{DSC}	DSC signal (W)
p	Pressure (Pa)
p^\ominus	Standard pressure (101325 Pa)
p_i	Partial pressure (Pa)
p_i^{vap}	Vapour pressure (Pa)
R	Gas constant (8.31446 $\text{J}/\text{mol}/\text{K}$)
R_k	Reaction
r_k	Molar rate of reaction ($\text{mol}/\text{m}^3/\text{s}$ or $\text{mol}/\text{m}^2/\text{s}$)
S_i	Species
S^\ominus	Standard entropy at 298.15 K ($\text{J}/\text{mol}/\text{K}$)
$S_{m,i}$	Molar entropy ($\text{J}/\text{mol}/\text{K}$)
T	Temperature (K)
T^\ominus	Standard temperature (298.15 K)
t	Time (s)
V_j	Volume of phase P_j (m^3)
$V_{m,i}$	Molar Volume (m^3/mol)
X_i	Mol fraction (-)
Y_i	Mass fraction (-)
α_c	Accumulation factor for condensation (-)
β_k	Temperature exponent
$\Delta_{R_k} G^\ominus$	Reaction free enthalpy (J/mol)
$\Delta_f H^\ominus$	Standard enthalpy of formation at 298.15 K (J/mol)
$\Delta_{\text{sol}} H^\ominus$	Standard enthalpy of solution at 298.15 K (J/mol)
μ_i	Chemical potential of species
μ_i^0	Chemical potential of undiluted species
ν_{ik}	Stoichiometric coefficient
$\tilde{\nu}_{ik}$	Concentration exponent
ρ_i	Density (kg/m^3)

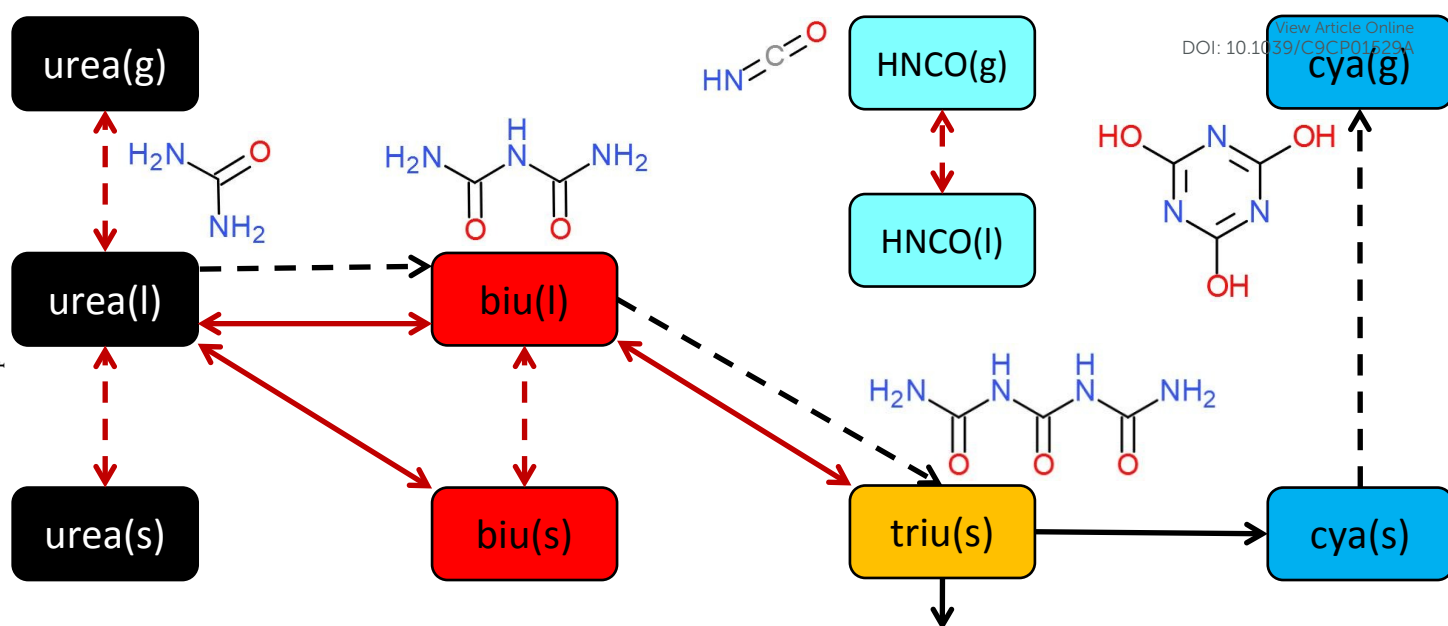
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A new reaction mechanism of urea decomposition is proposed after a strict treatment of thermodynamics in all phases.

