



Cite this: *Chem. Commun.*, 2024, 60, 1444

Received 6th November 2023,
Accepted 22nd December 2023

DOI: 10.1039/d3cc05462d

rsc.li/chemcomm

Effective and sustainable depolymerization of Nylon 66 – a transamidation for the complete recycling of polyamides†

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The transamidation of polyamides with short primary amides is reported as an effective recycling technique. This novel depolymerization method is robust and only utilizes cheap and renewable reagents. The process requires a Nb₂O₅ catalyst, assisted by NH₃, and operates at relatively mild reaction conditions (i.e. 200 °C and 3 bar NH₃).

Polyamides (PAs) or nylon are an important class of nitrogen-containing polymers. With an annual production of more than 8 Mton, the PA market is mainly driven by the high demand for Nylon 6 (i.e. PA 6) and Nylon 66 (i.e. PA 66), with corresponding market shares of approximately 50% and 40%.¹ Polyamides are known for their strength, durability, thermal and chemical stability. These excellent mechanical properties make them versatile and desirable materials for a variety of applications. PAs can be found in everyday objects such as clothing or are used for high-end applications such as 3D printing.² The downside of PAs' durability is their high resilience towards degradation and depolymerization. This makes PAs notoriously difficult to recycle.

Over the past decades, many attempts were made to depolymerize PAs, e.g. via alcoholysis, ammonolysis, hydrolysis etc.^{3,4,6,8–10} In particular, the steam-assisted depolymerization of PAs proved to be very effective for lactam-based PAs such as Nylon 6. In supercritical liquid or with overheated steam molten Nylon 6 is depolymerized at temperatures of up to 420 °C.⁴ The generated caprolactam is immediately removed from the reaction mixture. Such a depolymerization was not successful for Nylon 66. During Nylon 66 hydrolysis, a reactive and unvolatile mixture of acids and amines is generated, which repolymerizes, thus leading to a Nylon 66 resin.⁵ In order to fully depolymerize Nylon 66, stoichiometric quantities of

mineral acids or alkali are required, trapping either the α , ω -dicarboxylic acids or α , ω -diamines in their salt form.^{4d,6}

To this date, the situation for Nylon 66 remains problematic. The lack of efficient recycling methods results in the predominant incineration of the material *via* waste-to-energy (WtE). While this method would be acceptable for plastics such as polyethylene (PE) or polypropylene (PP) this is not the case for nitrogen-containing plastics such as PA. Burning these polymers not only emits CO₂, but also toxic NO_x greenhouse gases (N₂O has a global warming potential that is 265 times higher than that of CO₂).⁷ The development of new and effective recycling methods is therefore essential to minimize potent greenhouse gas emission and to enable a circular economy.¹¹

In the past few years, several new PA recycling methods were proposed that utilize hydrogen (H₂) or ammonia (NH₃) as the splitting agent. However, each of these new processes suffers from significant disadvantages. For instance, the teams of Milstein and Schaub proposed homogeneous hydrogenolysis catalysts.⁸ While both groups were able to successfully split polyamides into α , ω -diamines and diols, or, aminoalcohols, the performance of their catalysts for Nylon 66 is remarkably low. Additionally, these specialized catalysts are expensive and easily poisoned by additives or impurities that might be present in actual plastic (waste). In the work of Stuyck *et al.* Nylon 66 was depolymerized *via* a sustainable and green ammonolysis process.⁹ Reactions were performed with a homogeneous Lewis acid catalyst and in a simple glycol solvent, e.g. biosourced waste glycerol. Unfortunately, the ammonolysis of PAs is an equilibrium reaction. As a result, this method is unable to split Nylon 66 completely into monomeric compounds and a fairly reactive product mixture is obtained. To deal with this drawback, Coeck *et al.* coupled the ammonolysis to an irreversible reaction.¹⁰ This approach worked excellently for long-chain polyamides (LCPA) to generate α , ω -diamines, α , ω -dinitriles or amidonitriles. However, these new recycling methods have not yet been proven effective for Nylon 66.

In this work, we report a new and sustainable depolymerization method for PAs, optimized for Nylon 66 specifically. The

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† Electronic supplementary information (ESI) available. See DOI: <https://doi.org/10.1039/d3cc05462d>



Table 1 Variation of the reaction conditions for the transamidation of polyamides

Reaction scheme showing the transamidation of a polyamide (Nylon 66) with a primary amide (R-CONH₂) under heat (ΔT) and catalyst (NH₃) to produce two products: a monomer (1) and a dimer (2).

										Product yields ^b	
Substrate		Catalyst		Solvent (R-CONH ₂)		<i>P</i> _{NH₃}	Temp. (°C)	Time (h)			
									①	②	
1	Nylon 66	0.5 M ^a	—	—	Acetamide	10 g	3 bar	200	5	1% ^e	1%
2	Nylon 66	0.5 M ^a	Nb ₂ O ₅	0.1 g	Acetamide	10 g	—	200	5	4% ^e	4%
3	Nylon 66	0.5 M ^a	Nb ₂ O ₅	0.1 g	Acetamide	10 g	1 bar	200	5	7% ^e	7%
4	Nylon 66	0.5 M ^a	Nb ₂ O ₅	0.1 g	Acetamide	10 g	3 bar	200	5	12% ^e	13%
5	Nylon 66	0.5 M ^a	Nb ₂ O ₅	0.1 g	Acetamide	10 g	5 bar	200	5	9% ^e	8%
6	Nylon 66	0.5 M ^a	Nb ₂ O ₅	0.2 g	Acetamide	10 g	3 bar	200	16	65% ^e	23%
7	Nylon 66	0.5 M ^a	Nb ₂ O ₅	0.2 g	Acetamide	10 g	3 bar	225	16	94% ^e	5%
8	Nylon 66 + S ^c	0.5 M ^a	Nb ₂ O ₅	0.2 g	Acetamide	10 g	3 bar	200	16	69% ^e	20%
9	Nylon 66 + PP ^d	0.5 M ^a	Nb ₂ O ₅	0.2 g	Acetamide	10 g	3 bar	200	16	69% ^e	20%
10	Nylon 66	0.1 M ^a	Nb ₂ O ₅	0.2 g	Acetamide	10 g	3 bar	200	16	23% ^f	35% ^f
11	Nylon 66	0.1 M ^a	Nb ₂ O ₅	0.2 g	Acetamide	10 g	3 bar	200	72	98% ^e	2%
12	Nylon 66	0.5 M ^a	Nb ₂ O ₅	0.2 g	Acetamide	10 g	3 bar	200	72	93% ^e	7%
13	Nylon 66	1.25 M ^a	Nb ₂ O ₅	0.5 g	Acetamide	10 g	3 bar	200	72	89% ^e	9%
14	Nylon 66	0.5 M ^a	Nb ₂ O ₅	0.2 g	Propionamide	10 g	3 bar	200	72	92% ^e	8%

^a Amount of secondary amide bonds. ^b Leftover plastic not characterized. ^c S = *N*-butylbenzenesulfonamide, which was added to the mixture (0.022 g). ^d PP = polypropylene, which was added (0.25 g). ^e This value also corresponds to the adipamide yield, which is generated stoichiometrically. ^f Aside from the amides formed from Nylon 6, a yield of 5% caprolactam was obtained as well.

depolymerization involves a transamidation reaction with a short monofunctional amide (see top Table 1). The transamidation of PAs is a fairly unknown reaction. However, during melting or extrusion (≥ 250 °C) some sporadic, uncatalyzed transamidations occur.¹² In more recent years, spontaneous transamidation reactions at a high temperature allowed material scientists to generate “PA alloys”.¹³ However, such transamidations have never been performed catalytically or at a moderate temperature, nor was anyone ever able to effectively depolymerize PAs doing so.

To start, acetamide was chosen as the most ideal primary amide reagent because it can be cheaply biosourced (*e.g.* from acetic acid) and is readily biodegradable.¹⁴ In a blank reaction at 200 °C barely 1% of monomeric compounds was obtained (Table 1, entry 1), proving the need for a catalyst. Upon addition of *ortho*-Nb₂O₅, the yield of *N,N'*-hexamethylene bis(acetamide) (1) increased substantially (entries 2–5). This comes to no surprise; we previously reported the ability of Nb₂O₅ to activate secondary amide bonds.¹⁰ Ammonia (NH₃) was also introduced into the system, to assist in breaking down the polymer by ammonolysis. Under the applied conditions, virtually no amine products were detected. Thus, NH₃ merely acts as a co-catalyst. An optimum in NH₃ pressure was observed at 3 bar, with a yield of 12% *N,N'*-hexamethylene bis(acetamide) and 13% of the dimer (*i.e.* *N'*-(6-acetamidohexyl)adipamide (2)). Upon further addition of NH₃, the global catalytic performance decreases, most likely due to a poisoning effect of the high NH₃ concentration on the Nb₂O₅ surface.

Next, the reaction time was prolonged from 5 to 16 hours (entry 6). A spectacular increase in monomer yield (*i.e.* from 12% to 65% (1)) illustrates that the polymer is chopped up at random, gradually breaking it down into smaller pieces. By

slightly increasing the reaction temperature to 225 °C (entry 7), Nylon 66 was completely depolymerized into monomers (94% (1)) and dimers (5% (2)). Note that for every molecule of *N,N'*-hexamethylene bis(acetamide), a molecule of adipamide is generated as well. In addition to our desired products, we also detected very low quantities of pyrimidines and triazines, resulting from the trimerization of acetonitrile (*i.e.* dehydrated acetamide).¹⁵ This side reaction is caused by the Nb₂O₅ catalyst, which is known for its ability to (partially) dehydrate primary amides to nitriles.^{10,16} In order to maximize the solvent's reusability, we did not further increase the reaction temperature.

The system was also proven to be very robust (entries 8–10). The addition of other plastics (*e.g.* polypropylene, entry 9) or sulfur-containing plasticizers (*e.g.* *N*-butylbenzenesulfonamide, entry 8) did not hinder the reaction. In particular, the catalytic system's resilience towards plasticizers is a significant advance over current recycling techniques. Such sulfur-containing molecules typically poison metal-based catalysts almost immediately.^{8,10} In addition, we succeeded in breaking down Nylon 6 as well (entry 10). Therefore, our innovative depolymerization system has the potential to be a valuable approach for carpet recycling, where Nylon 66/Nylon 6/PP are commonly mixed and/or copolymerized.^{2,3a}

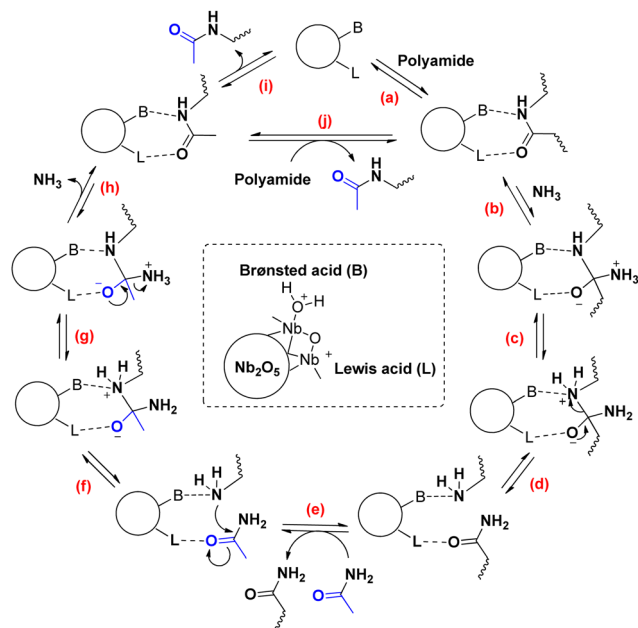
Next, reactions were performed with a long reaction time of 72 h and with a variable concentration of PA (entries 11–13). These experiments allow to identify the position of the thermodynamic equilibrium under each set of reaction conditions. In general, one could consider a single polymer string of Nylon 66 as a series of 1,6-diaminohexane (HMDA) molecules trapped in secondary amide bonds. It appears that the thermodynamic stability of the amide bond in HMDA-adipyl is similar to that of



HMDA-acetyl. In other words, the results depicted in entries 11–13 nearly perfectly match the situation in which each amino group of a molecule HMDA does not differentiate between a condensation reaction with adipamide or acetamide (*i.e.* $R_1-NH_2 + NH_2-C(=O)-R_2$ to $R_1-NH-C(=O)-R_2$). For example, in entry 12, the reaction contains 169 mmol carbonyl groups from acetamide and 5 mmol from adipamide (ratio of 97:3). If no differentiation occurs, one would expect a *N,N'*-hexamethylene bis(acetamide) yield of 94% ($= 0.97 \times 0.97$), which matches the experimental value of 93% very well. It is worth noting that even at a high polymer concentration, *i.e.* 12.5 wt% Nylon 66 (entry 13), the plastic can still be depolymerized completely into mainly monomeric compounds (*i.e.* 89% yield) with the dimer as the secondary side product (9% yield). This is the consequence of the large molar excess of the small molecule acetamide *versus* adipamide (*i.e.* 169 mmol acetamide *versus* 6.25 mmol adipamide).

Finally, as an alternative to acetamide, a transamidation reaction was performed with propionamide, which poses fewer health risks for mammals (*e.g.* humans; entry 14).¹⁴ Although the final monomer yield is slightly lower than with acetamide, a final yield of 92% *N,N'*-hexamethylene bis-(propionamide) was obtained with 8% of the dimer (*i.e.* *N*¹-(6-propionamido-hexyl)adipamide) as the only side product.

Based on prior knowledge and our experimental results, we propose a reaction mechanism for the ammonia assisted transamidation of PAs (Scheme 1).¹⁰ Initially, the carbonyl group of the secondary amide (*i.e.* PA) needs the Lewis acid site (L) on the catalytic surface to be reactive (a). The positive effect of NH_3 on the reaction rate is in agreement with the next step, *i.e.* a nucleophilic attack of NH_3 on the surface-adsorbed amide to form a primary amide and a free amine group (b)(c).

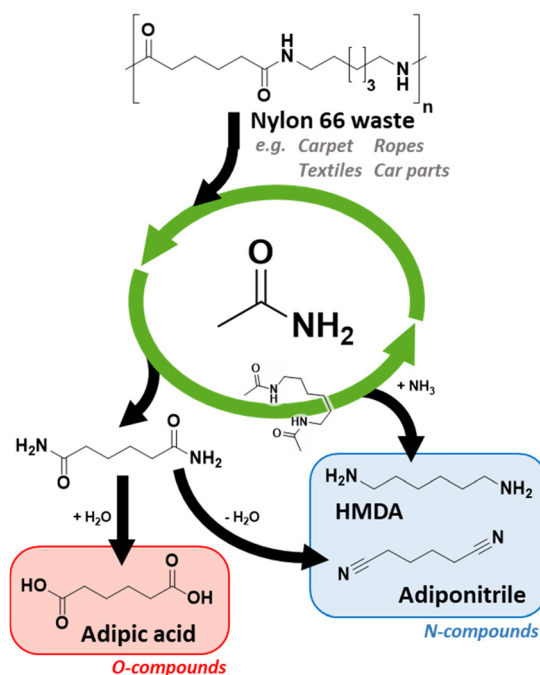


Scheme 1 Proposed reaction mechanism for the ammonia assisted transamidation of polyamides with acetamide on a Nb_2O_5 catalyst.

During this reaction, a neighboring (Brønsted) acid group facilitates the leaving of the amine group (d). If the surface-adsorbed primary amide end-group is displaced by an acetamide molecule from the solvent (e), the ammonolysis reaction can be reversed. First, the free amine chain end attacks the now activated carbonyl group of acetamide (f). The resulting intermediate releases a molecule of NH_3 , yielding the final transamidation product (g)(h). After desorption of the product, a new secondary amide molecule can coordinate to the catalytic surface (i)(j). All elementary steps in this mechanism are reversible. Nevertheless, the PA is depolymerized completely as a consequence of the large molar excess of the small molecule acetamide *versus* adipamide.

In summary, we have developed a very robust recycling method that only uses renewable and/or low-cost materials and reagents. Nylon 66 could be depolymerized into easily processable monomers at relatively high concentrations, *i.e.* 12.5 wt% of Nylon 66 (in acetamide). After a transamidation reaction with acetamide, two distinct monomers are obtained: *N,N'*-hexamethylene bis(acetamide) and adipamide (Scheme 2). These monomers differ greatly in chemical properties, facilitating an easy separation. Adipamide's limited solubility in most organic solvents allows a separation *via* crystallization. In addition, the boiling point of these monomers is very different, allowing classical isolation methods such as distillation.

After isolation, *N,N'*-hexamethylene bis(acetamide) can be reconverted to HMDA *via*, *e.g.*, ammonolytic reactive distillation.^{10a} In doing so, acetamide is liberated, closing the loop for this primary amide molecule. Alternatively, one can utilize the homogeneous amide hydrogenation catalysts developed by the teams of Milstein and Schaub.⁸



Scheme 2 Complete recycling of Nylon 66 waste based on a transamidation depolymerization technique.



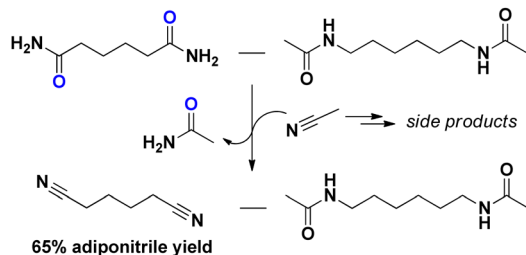


Fig. 1 Transfer dehydration of a 1–1 molar mixture of *N,N'*-hexamethylene bis(acetamide) and adipamide to acetamide. Reaction conditions: Adipamide (1 mmol), *N,N'*-hexamethylene bis(acetamide) (1 mmol), 200 °C, Nb₂O₅ (0.2 g), acetonitrile (10 mL), 16 hours.

The resulting overall reaction for such a process would be as follows:

Nylon 66 (repeating unit) + 4 H₂ + 2 acetamide → HMDA + 2 ethanol [+ adipamide].

Although in this case acetamide could not be recycled, the overall reaction could still be economically feasible.

Considering adipamide, this molecule is a well-known intermediate in the PA industry. One of the first (industrial) synthesis methods of adiponitrile was *via* the dehydration of adipamide/ammonium adipate (under a stream of NH₃).¹⁷ If adiponitrile is the desired end product, a (complete) separation of *N,N'*-hexamethylene bis(acetamide) and adipamide might be superfluous. As a proof of concept, we performed a niobia catalyzed transfer-dehydration with acetonitrile as the solvent and sacrificial water scavenger. Although severe solvent degradation was observed, we obtained a 65% adiponitrile yield without major degradation of the *N,N'*-hexamethylene bis(acetamide) monomer (Fig. 1).

Alternatively, adipamide could be hydrolyzed with water to the original adipic acid. For this reaction, Nb₂O₅ would be an excellent catalyst as well.¹⁸ Hence, the developed transamidation process could potentially serve as the foundation for the closed-loop recycling of Nylon 66.

Conflicts of interest

There are no conflicts to declare.

Notes and references

- (a) Statista. Market volume of polyamide industry worldwide from 2017 to 2018 with a forecast until 2025, <https://www.statista.com/>; (b) Grand View Research. Global Nylon Market Size, Share & Trends Report, 2030, <https://www.grandviewresearch.com/>.
- B. Herzog, M. I. Kohan, S. A. Mestemacher, R. U. Pagilagan, K. Redmond and R. Sarbandi, *Ullmann's Encyclopedia of Industrial Chemistry*, Wiley, 2020, vol. 14, pp. 1–47.
- (a) C. Mihut, D. K. Captain, F. Gadala-Maria and M. D. Amiridis, *Polym. Eng. Sci.*, 2001, **41**, 1457–1470; (b) A. Kamimura, K. Ikeda, S. Suzuki, K. Kato, Y. Akinari, T. Sugimoto, K. Kashiwagi, K. Kaiso, H. Matsumoto and M. Yoshimoto, *ChemSusChem*, 2014, **7**, 2473–2477.
- (a) K. Kaiso, T. Sugimoto, K. Kashiwagi and A. Kamimura, *Chem. Lett.*, 2011, **40**, 370–371; (b) K. Kaiso and T. Sugimoto, *European Pat.*, EP1975156A, 2008; (c) S. Sifniades, A. B. Levy and J. A. J. Hendrix, *US Pat.*, US5932724, 1999; (d) E. F. Moran, *World Pat.*, WO1994008942A, 1994.
- Carpet America Recovery Effort, CARE 2016 Annual Report, <https://carpetrecovery.org/>.
- (a) M. G. Peter, *US Pat.*, US3069465A, 1962; (b) D. D. Davis and E. D. Wilhoit, *World Pat.*, WO9700846A, 1997.
- S. L. Konopa, J. A. Mulholland, M. J. Realff and P. M. Lemieux, *J. Air Waste Manage. Assoc.*, 2008, **58**, 1070–1076.
- (a) A. Kumar, N. von Wolff, M. Rauch, Y.-Q. Zou, G. Shmul, Y. Ben-David, G. Leitus, L. Avram and D. Milstein, *J. Am. Chem. Soc.*, 2020, **142**, 14267–14275; (b) W. Zhou, P. Neumann, M. Al Batal, F. Rominger, A. S. K. Hashmi and T. Schaub, *ChemSusChem*, 2021, **14**, 4176–4180.
- W. Stuyck, K. Janssens, M. Denayer, F. De Schouwer, R. Coeck, K. V. Bernaerts, J. Vekeman, F. De Proft and D. E. De Vos, *Green Chem.*, 2022, **24**, 6923–6930.
- (a) R. Coeck, A. De Bruyne, T. Borremans, W. Stuyck and D. E. De Vos, *ACS Sustain. Chem. Eng.*, 2022, **10**, 3048–3056; (b) R. Coeck, M. Houbrechts and D. E. De Vos, *Chem. Sci.*, 2023, **14**, 7944–7955.
- (a) J. H. Clark, T. J. Farmer, L. Herrero-Davila and J. Sherwood, *Green Chem.*, 2016, **18**, 3914–3934; (b) W. R. Stahel, *Nature*, 2016, **531**, 435–438; (c) T. Keijer, V. Bakker and J. C. Sloopweg, *Nat. Chem.*, 2019, **11**, 190–195; (d) J.-P. Lange, *Green Chem.*, 2002, **4**, 546–550.
- J. G. Ramos and M. A. Lozano, *An. R. Soc. Esp. Fis. Quim., Ser. B*, 1960, **56B**, 833–840.
- (a) L. Wang, X. Dong, Y. Gao, M. Huang, C. C. Han, S. Zhu and D. Wang, *Polymer*, 2015, **59**, 16–25; (b) K. L. L. Eersels, G. Groeninckx, Y. Mengerink and S. van der Waals, *Macromolecules*, 1996, **29**, 6744–6749.
- G. L. Kennedy and R. D. Short, *CRC Crit. Rev. Toxicol.*, 1986, **17**, 129–182.
- (a) X. Tao, T. Liu, H. Tao, R. Liu and Y. Qian, *J. Mol. Catal. A: Chem.*, 2003, **201**, 155–160; (b) J. H. Forsberg, V. T. Spaziano, S. P. Klump and K. M. Sanders, *J. Heterocycl. Chem.*, 1988, **25**, 767–770.
- R. Coeck, J. Meeprasert, G. Li, T. Altantzis, S. Bals, E. A. Pidko and D. E. De Vos, *ACS Catal.*, 2021, **11**, 7672–7684.
- K. Weissmermel and H.-J. Arpe, *Industrial Organic Chemistry*, Wiley, Weinheim, Germany, 2003, vol. 11, pp. 239–266.
- S. M. A. H. Siddiki, M. N. Rashed, A. S. Touchy, M. A. R. Jamil, Y. Jing, T. Toyao, Z. Maeno and K. Shimizu, *Catal. Sci. Technol.*, 2021, **11**, 1949–1960.

