Ammonia recovery from acidogenic fermentation effluents using a gas-permeable membrane

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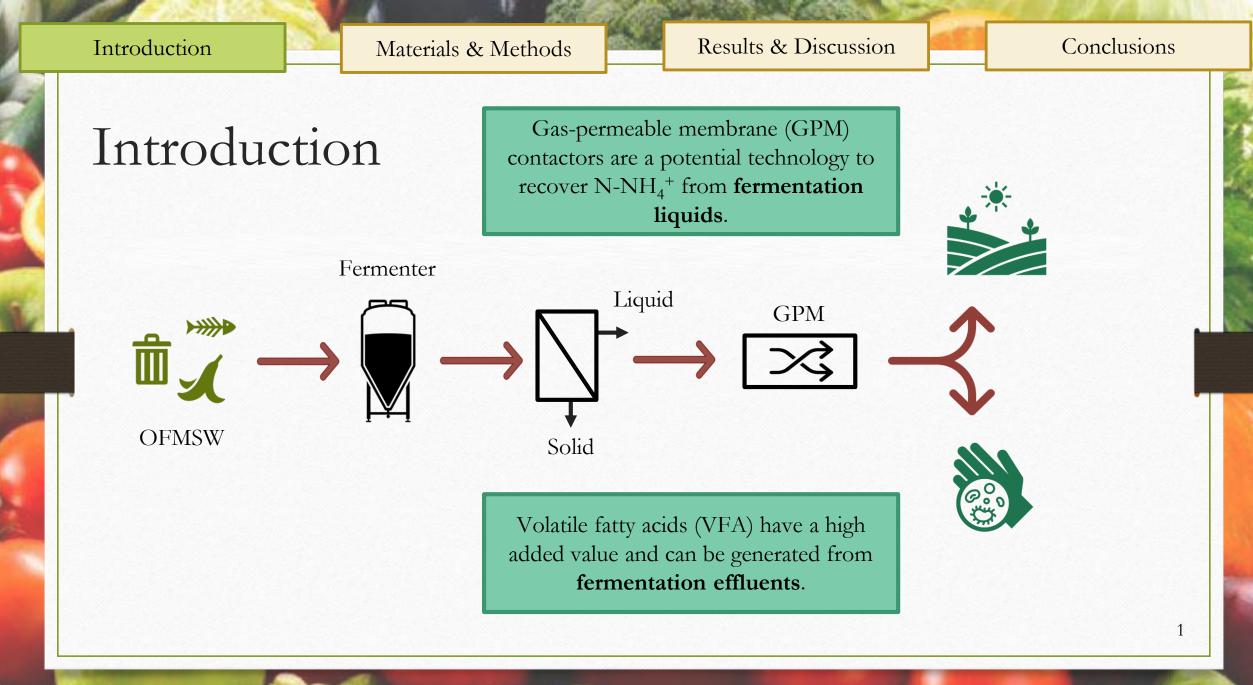
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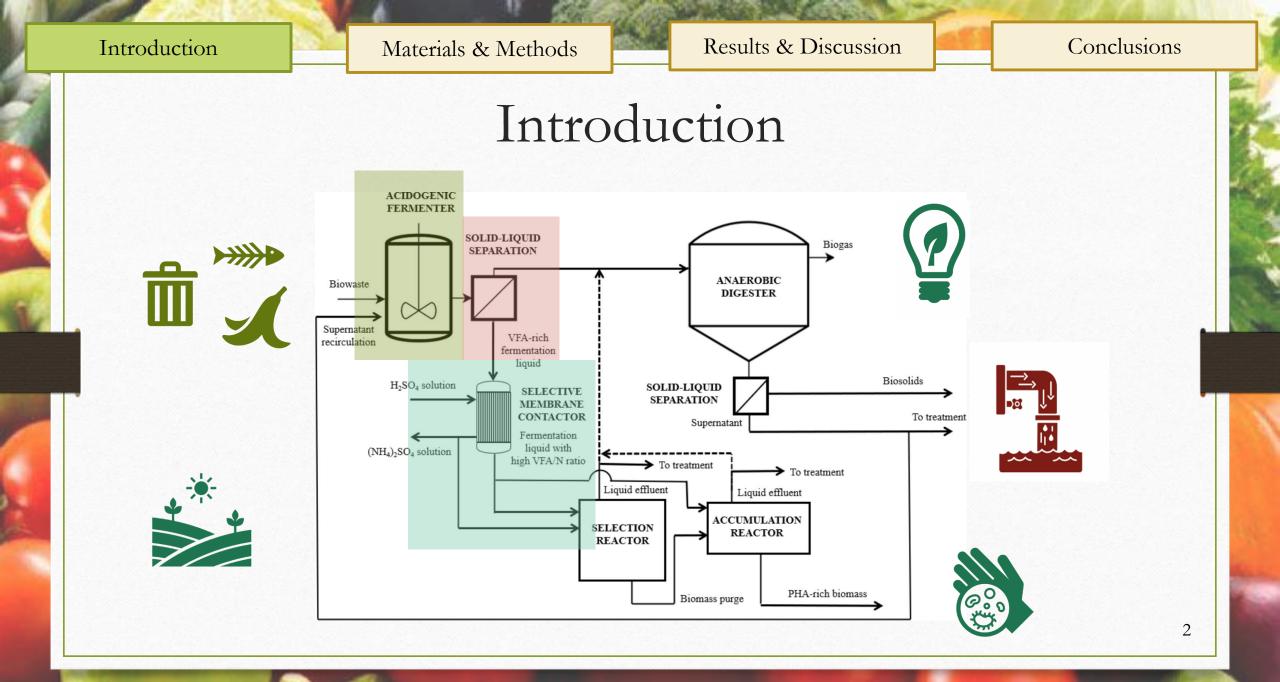


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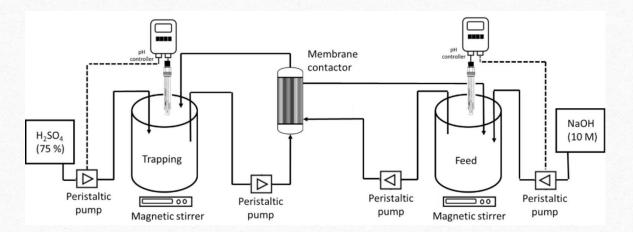


Results & Discussion

Conclusions

Materials and Methods

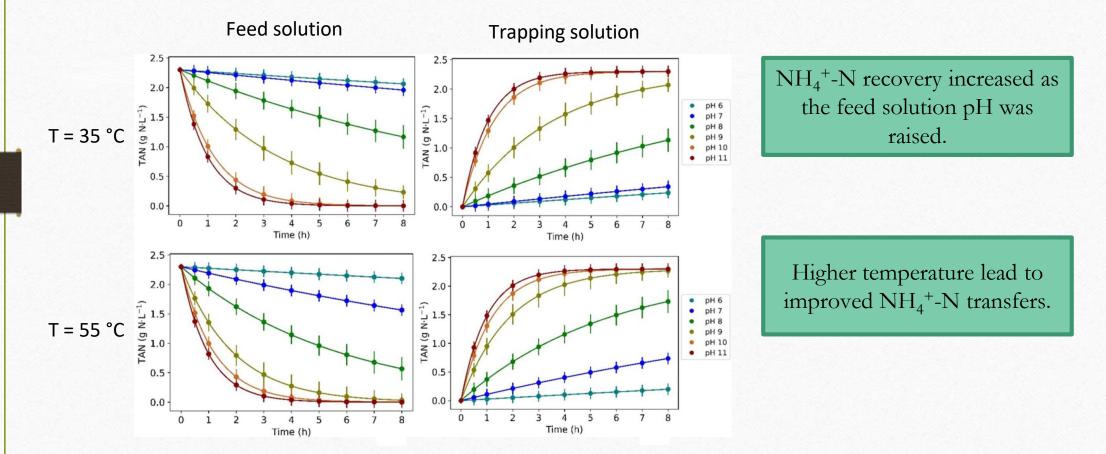
Ammonia removal took place in a nanoperforated hollow fibre membrane contactor that put in contact the feed solution and the trapping acidic solution.



The **acidogenic fermentation** of OFMSW was carried out in a 30 L reactor and lasted 8 days.

Feed solution	pH (-)	T (^o C)
	6.0	35
	7.0	35
	8.0	35
Synthetic	9.0	35
fermentation	10.0	35
liquid	11.0	35
(2.5 g N/L	6.0	55
3.0 g HAc/L)	7.0	55
	8.0	55
	9.0	55
	10.0	55
	11.0	55
OFMSW	9.0	35
fermentation	10.0	35
	9.0	55
liquid	10.0	55

Nitrogen recovery with synthetic solution



Nitrogen recovery with synthetic solution

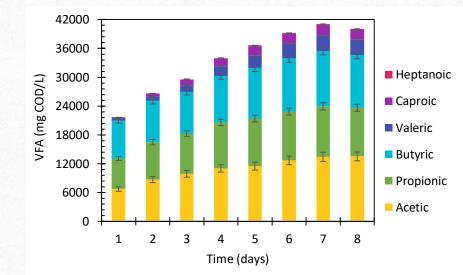
Exp.	Т	pН	TAN recovery at 8h	Average flux at 8h	K _m
	٥C	-	0⁄0	g N·day ⁻¹ ·m ⁻²	m·s ⁻¹
1A	35	6.0	8	2.5 ± 0.3	$(2.2 \pm 0.5) \cdot 10^{-8}$
1 B		7.0	16	4.3 ± 0.6	$(2.8 \pm 0.4) \cdot 10^{-8}$
1 C		8.0	65		$(9.5 \pm 1.3) \cdot 10^{-8}$
1D		9.0	91	62 ± 10	$(3.2 \pm 0.3) \cdot 10^{-7}$
1E		10.0	100	115 ± 16	$(9.2 \pm 0.4) \cdot 10^{-7}$
1F		11.0	100	138 ± 17	$(1.13 \pm 0.03) \cdot 10^{-6}$
2A	55	6.0	10	2.8 ± 0.3	$(1.3 \pm 0.3) \cdot 10^{-8}$
2B		7.0	36	10 ± 2	$(5.8 \pm 1.0) \cdot 10^{-8}$
2C		8.0	73	+ 32 ± 5	$(1.9 \pm 0.1) \cdot 10^{-7}$
2 D		9.0	99	84 ± 11	$(5.9 \pm 0.5) \cdot 10^{-7}$
2 E		10.0	100	121 ± 16	$(9.3 \pm 0.3) \cdot 10^{-7}$
2F		11.0	100	133 ± 16	$(1.15 \pm 0.04) \cdot 10^{-6}$

NaOH consumption decreased as the feed solution pH was raised.

 H_2SO_4 consumption decreased as the feed solution pH was raised (minimum at pH 10).

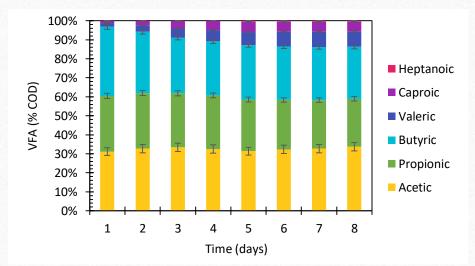
The best operation pH was determined to be 10.

Acidogenic fermentation



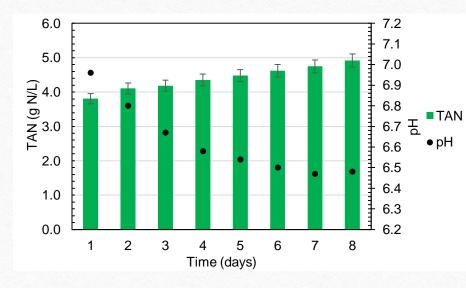
 $\begin{array}{c} {\rm VFA\ concentration} \\ {\rm reached\ 41\ g\ COD}_{\rm VFA}/{\rm L} \\ {\rm at\ the\ 7^{th}\ day.} \end{array}$

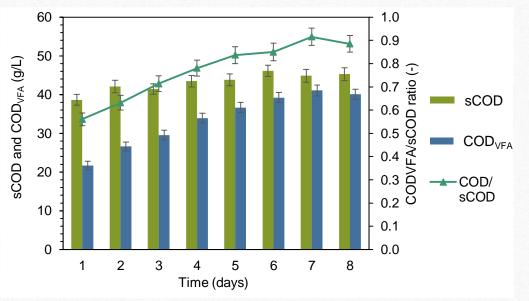




Acidogenic fermentation

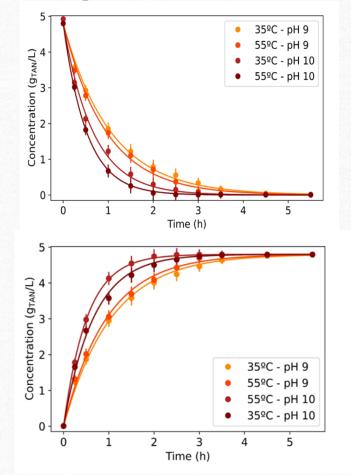
 $N-NH_4^+$ concentration reached 4.9 g N/L.



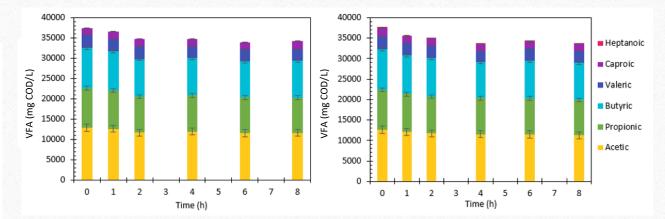


COD_{VFA} increased due to the acidogenic fermentation.

Nitrogen recovery with fermentation liquid



The GPM allowed a complete recovery of $N-NH_4^+$ with a minimum loss of VFA.



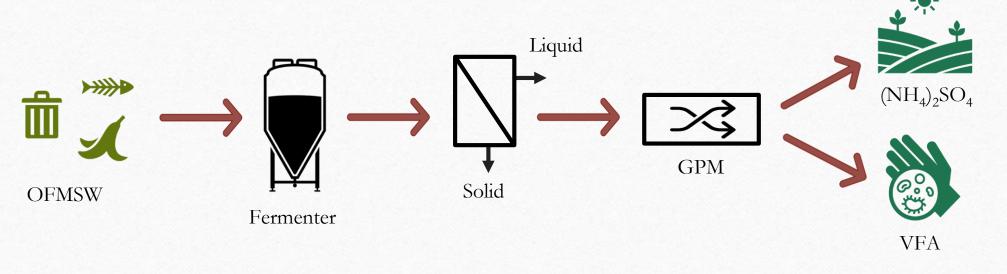
Nitrogen recovery with fermentation liquid

Exp.	Т	pН	TAN recovery at 8h	Average flux at 8h	K _m	Synthetic K _m
	٥C	-	0⁄0	g N∙day⁻¹∙m⁻²	m·s ⁻¹	m·s ⁻¹
3A	35	9.0	100	78.4 ± 11.0	$(2.8 \pm 6.0) \cdot 10^{-7}$	$(3.2 \pm 0.3) \cdot 10^{-7}$
3B		10.0	100	120.1 ± 14.4	$(5.0 \pm 9.2) \cdot 10^{-7}$	$(9.2 \pm 0.4) \cdot 10^{-7}$
3C	55	9.0	100	91.0 ± 12.7	$(3.3 \pm 6.3) \cdot 10^{-7}$	$(5.9 \pm 0.5) \cdot 10^{-7}$
3D		10.0	100	138.8 ± 16.7	$(5.4 \pm 5.7) \cdot 10^{-7}$	$(9.3 \pm 0.3) \cdot 10^{-7}$

N-NH₄⁺ transfer decreased when we operated with fermentation liquid.

The K_m improve from pH 9 to 10 were lower in fermentation liquid.

- The **N-NH₄⁺ can be completely recovered** from fermentation effluents at 35 and 55 °C and pH values above 9 efficiently.
- The process produce a (NH₄)₂SO₄ solution suitable as fertiliser and a rich VFA liquid free of N-NH₄⁺.



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Thank you for your attention













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