Technical performance assessment of large scale process concepts for the recycling of end-of-life photovoltaic panels

Sz. Fogarasi*, F. Imre-Lucaci**, A.-M. Cormoş* and Á. Imre-Lucaci*

*Department of of Chemical Engineering, Faculty of Chemistry and Chemical Engineering, Babeş-Bolyai University, Cluj-Napoca, RO-400028, Romania

**Institute for Interdisciplinary Research in Bio-Nano-Sciences, Babeş-Bolyai University, Cluj Napoca, RO-400271, Romania

Presenting author e-mail: arpad.imre@ubbcluj.ro

Introduction

Photovoltaic power systems have emerged as one of the most promising alternatives to reduce the impact of the energy sector on climate change and to improve the energy security of countries all over the world [1]. Although photovoltaic panels (PhVP) have much larger useful lifespan than the majority of electronic and electrical equipment

they still become waste products after approximately 30 years.

According to literature data there are estimates for the accumulation of 1.7–8 million tons by 2030 and 60–78 million tons by 2050 of end-of-life PhVP (EOL PV) [2-4]. To manage such a large quantity of waste material goes beyond just enforcing policies, it needs a massive research and innovation effort to ensure the development and large scale deployment of ground-breaking, cost effective and eco-friendly EOL PV recycling technologies.

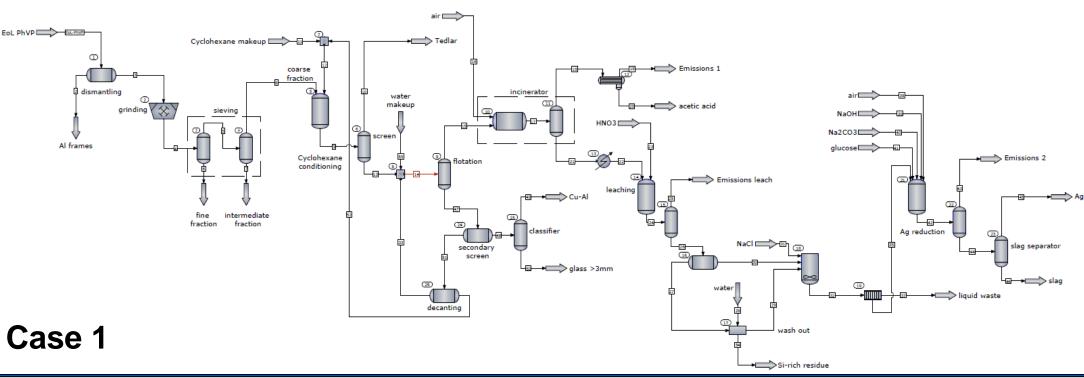
In attempt to provide a useful contribution to the field of EOL PV recycling, the current study presents novel large scale process concepts for the comprehensive treatment of EOL PV.

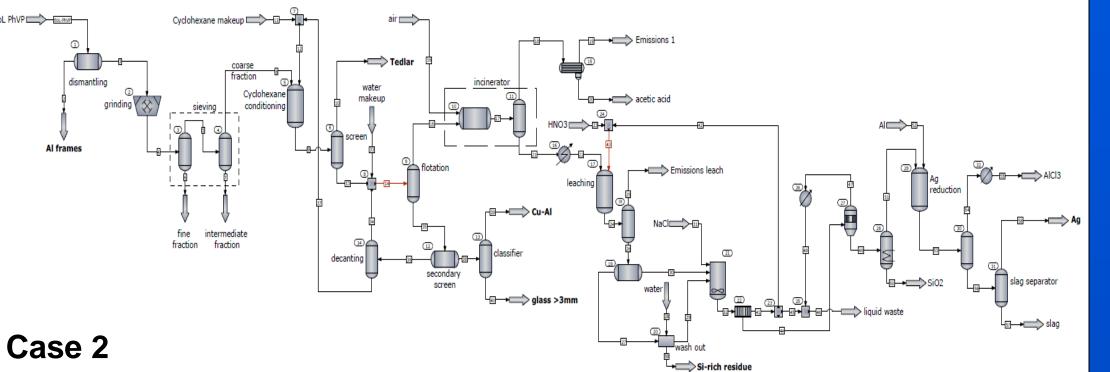
Results @ Discussions

The process of recycling the EOL PhVP is divided into the following subsystems (the PFDs developed in CHEMCAD are in Fig. 1):

- 1 Processing and mechanical separation
- 2 The processing of the coarse fraction
- 3 Ag and Si release by pyrolysis
- **4 Dissolution and purification**
- 5 Silver recovery by reducing AgCl to Ag considering two cases: Case 1 - the reducing system is glucose + NaOH + Na_2CO_3 Case 2 - the reducing agent is Al.

The mathematical models developed and presented in Fig.1 for the recovery process of metals from EOL PhVP were simulated in order to find the optimal technological option. For both cases, three different thermal integrations were considered: (I) - no thermal integration; (II a) - thermal integration only for the recycling process





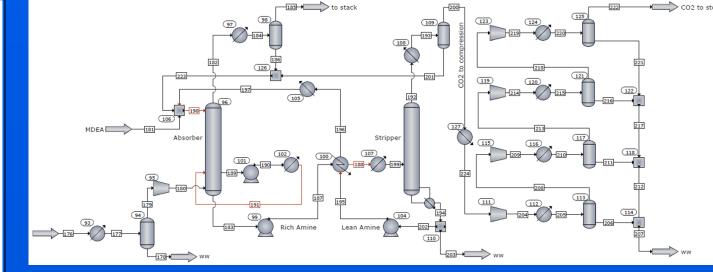


Fig. 2. PFD for the CO₂ capture process.

Considering the importance of CO_2 capture in the industrial sector, the recycling plant of EOL PhVP was coupled with a post-combustion CO_2 capture based on amine absorption process (the PFD developed in CHEMCAD is presented in Fig. 2).

and (II b) - thermal integration include also the CO₂ caption process. The obtained results are presented in Tables 1-6.

Fig. 1. Process flow diagrams of the EOL PhVP recycling plant.

Table 1. Production rate and recovery yield for the main products of theend-of-life PhVP recycling process.

Process type	Product	Glass	Tedlar	Si	Ag	AI	Cu
	Production rate, (kg/h)	6950	430	330.20	4.45	1000	108
Case 1 and Case 2	Recovery yield, (%)	97	98	89.23	89.01	98.83	99.99

Table 2. Specific consumption of raw materials in kg/kg PhVP to obtain the main products of the recycling process of end-of-life PhVP.

Process type	Raw m	H ₂ O	HNO ₃	NaCl	NaOH	Na ₂ CO ₃	$C_6H_{12}O_6$	aer	C ₆ H ₁₂	TOTAL	
0	Concurrention	kg/h	919	1713	26.39	2.42	1.17	1.12	16004	203.1	-
Case 1	Consumption	kg/kg PhVP	0.09	0.17	~ 0				2.75	0.02	3.03
00000	o D Concurrention	kg/h	919	210	2.5	-	-	-	16000	203.1	-
Case 2	Consumption	kg/kg PhVP	0.09	0.02	~ 0	-	-	-	2.75	0.02	2.88

Table 3. Specific consumption of raw materials in kg/kg product for differentsubsystems of the spent PhVP recycling process.

Process type	Subsystem	1	2	3	4	5	TOTAL
000004	TOTAL, (kg/h)	10203	266	16000	2392	9.6	
Case 1	W, (kg/kg)	2.91	0.05	0.91	4.81	0.52	9.2
Case2	TOTAL, (kg/h)	10203	266.1	16000	864.9	0.38	
	W, (kg/kg)	2.91	0.05	0.91	1.74	0.06	5.67

Table 4. Overall thermal energy balance for the recycling process of end-of-life PhVP.

Thermal energy	Equivalent CH_4 , kg CH_4 /h
inemai energy	

Table 5. Thermal energy balance for different subsystems of the end-of-life PhVP recycling process for the two non-thermally integrated case studies.

Process		Parameters of	Thermal energy				TOTAL	Specific	Equivalent
	Subsystem	energy flows	G	enerate	d	Consumed	consumption,	consumption,	consumption,
type		energy nows	1	2	3	1	MJ/h	kJ/kg	kg CH₄/h
	1	T, °C				35			
		Q, MJ/h				68	68	6.80	1.36
	2	T, °C				34			
	Z	Q, MJ/h				61	61	12.05	1.21
Case 1	3	T, °C	400	400	400				
Case I	3	Q, MJ/h	-4900	-51581	-89				
	4	T, °C	0			60			
		Q, MJ/h	0			204	204	411	4.08
	5	T, °C	600						
		Q, MJ/h	-1.40						
	1	T, °C				35			
	I.	Q, MJ/h				39	39	3.90	0.78
	2	T, °C				34			
	Z	Q, MJ/h				61	61	12.04	1.21
Case 2	3	T, °C	400	400	400				
Case 2	3	Q, MJ/h	-4900	-53120	-89				
	4	T, °C	40			60			
	4	Q, MJ/h	-491			532	532	1070	10.64
	5	T, °C	500						
	5	Q, MJ/h	-7						

Table 6. Total and specific CO2 emissions, respectivelly energy consumptionof the CO2 capture process for the recycling of end-of-life PhVP.

		No ca	apture	With c	Capture energy	
Process	type	Total, (kg CO ₂ /h)	kg CO₂/t PhVP	Total, (kg CO ₂ /h)	kg CO₂/t PhVP	consumption CO ₂ , (GJ/h)
	I	2989	298.92	149	14.95	9.94
Case 1	lla	2971	297.09	149	14.85	9.88
Case 2	I	3006	300.55	150	15.03	9.99
	lla	2971	297.07	149	14.85	9.88

Process	туре	Parameters	Generated, MJ/h			Consumed, MJ/h	Generated	Consumed
		T, °C	600	400	600	60		
	1	Q, MJ/h	-1.4	-56570	-1	333	1131	6.66
Case 1		T, °C		400				
Case 1	lla	Q, MJ/h		-56365			1127	
	ll b	T, °C		400				
		Q, MJ/h		-46487			930	
		T, °C	40	400	500	60		
		Q, MJ/h	-491	-58109	-7	631	1162	12.63
	ll a	T, °C		400				
Case 2		Q, MJ/h		-57577			1152	
	ШЬ	T, °C		400				
	llb	Q, MJ/h		-47699			954	

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Conclusions

- It was found that in the best operating conditions the average recovery rate of critical materials is over 89 % and the purity of the obtained compounds was more than 99 % which makes them suitable for new PhVP production.
- The results proved that in both case studies the pyrolysis and combustion of organic materials provides the necessary amount of energy for the thermal integration of the process.
- As an overall conclusion it can be stated that both case studies remain net thermal energy producers with the capacity to provide an equivalent of over 900 kg CH₄/h even in this situation with CO₂ capture.

Acknowledgements

This work was supported by a grant of the Ministry of Research, Innovation and Digitization of the Romanian Government, CNCS/CCCDI - UEFISCDI, project code COFUND-LEAP-RE-RESTART, within PNCDI III.



This project has received funding from the European Union's Horizon 2020 Research and Innovation Program under Grant Agreement 963530.