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

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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 + Na₂CO₃ 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 and (II b) - thermal integration include also the CO₂ caption process.

The obtained results are presented in Tables 1-6.

Table 1. Production rate and recovery yield for the main products of the end-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.

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Process type	Raw material		H ₂ O	HNO ₃	NaCl	NaOH	Na ₂ CO ₃	C ₆ H ₁₂ O ₆	aer	C ₆ H ₁₂	TOTAL
Cons.4 Consu	Congumntion	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
Coop 2	Canaumontian	kg/h	919	210	2.5	-	-	-	16000	203.1	-
Case 2 Cons	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 different subsystems of the spent PhVP recycling process.

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Process type	Subsystem	1	2	3	4	5	TOTAL			
Case 1	TOTAL, (kg/h)	10203	266	16000	2392	9.6				
	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 5. Overall thermal energy balance for the recycling process of end-of-life PhVP.

Process type		Davamatava			Thermal	energy	Equivalent CH ₄ , kg CH ₄ /h		
		Parameters	Generated, MJ/h			Consumed, MJ/h	Generated	Consumed	
	,	T, °C	600	400	600	60			
	ı	Q, MJ/h	-1.4	-56570	-1	333	1131	6.66	
Case 1	II o	T, °C		400					
Case I	II a	Q, MJ/h		-56365			1127		
	II b	T, °C		400					
		Q, MJ/h		-46487			930		
		T, °C	40	400	500	60			
	1	Q, MJ/h	-491	-58109	-7	631	1162	12.63	
Case 2	II a	T, °C		400					
Case 2		Q, MJ/h		-57577			1152		
	Пb	T, °C		400					
		Q, MJ/h		-47699			954		

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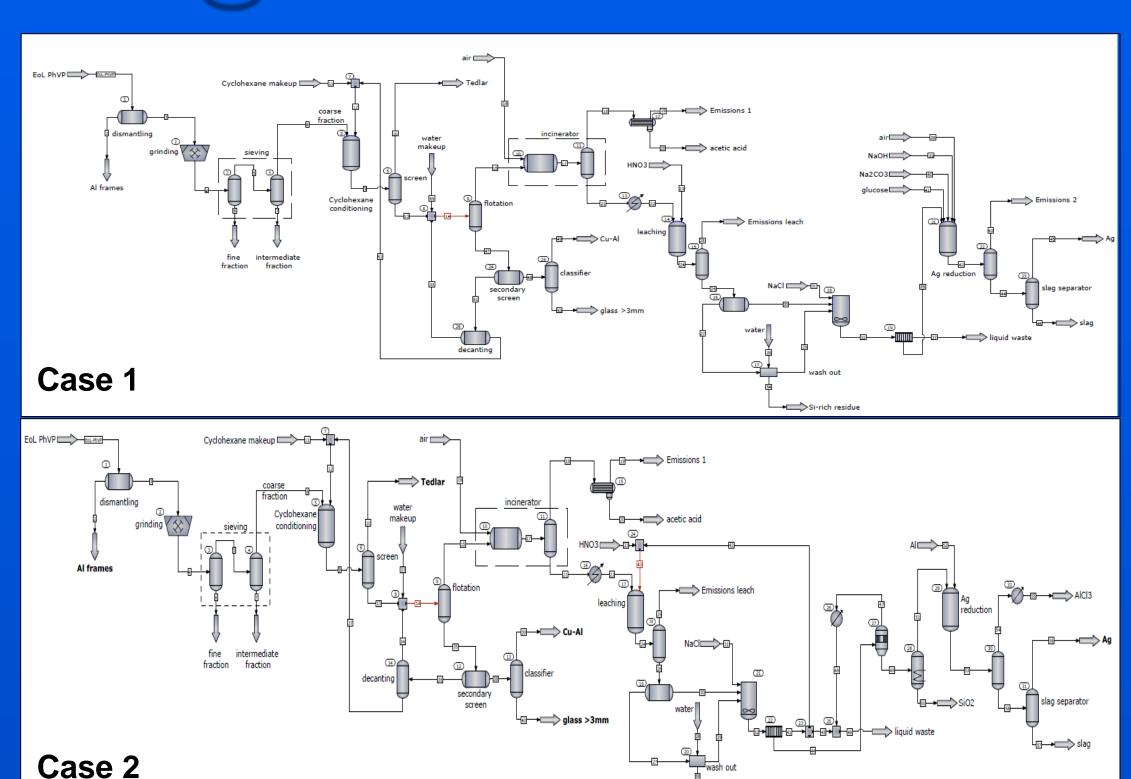


Fig. 1. Process flow diagrams of the EOL PhVP recycling plant.

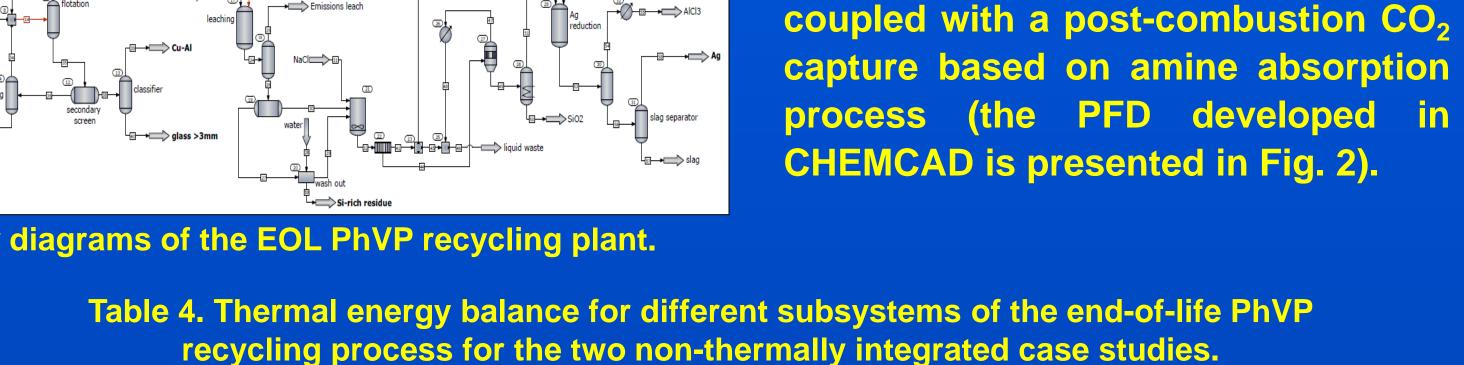


Fig. 2. PFD for the CO₂ capture process.

Considering the importance of CO₂

capture in the industrial sector, the

recycling plant of EOL PhVP was

Thermal energy **TOTAL Specific** Equivalent **Parameters of Process Subsystem** Generated Consumed consumption, consumption, consumption, energy flows type MJ/h kg CH₄/h kJ/kg T, °C Q, MJ/h 6.80 1.36 68 68 T, °C Q, MJ/h 12.05 1.21 61 T, °C 400 Case ' Q, MJ/h -4900 -51581 -89 T, °C 204 204 411 Q, MJ/h 4.08 T, °C 600 Q, MJ/h -1.40 T, °C Q, MJ/h 3.90 39 0.78 T, °C Q, MJ/h 61 12.04 1.21 400 T, °C Case 2 Q, MJ/h -4900 -53120 -89 T, °C 60 40 532 1070 Q, MJ/h 532 10.64 -491 T, °C 500 Q, MJ/h

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

		No c	apture	With o	Capture energy	
Process	type	Total, (kg CO ₂ /h)	kg CO ₂ /t PhVP	Total, (kg CO ₂ /h)	kg CO ₂ /t PhVP	consumption CO ₂ , (GJ/h)
04	ı	2989	298.92	149	14.95	9.94
Case 1	lla	2971	297.09	149	14.85	9.88
0000	ı	3006	300.55	150	15.03	9.99
Case 2	lla	2971	297.07	149	14.85	9.88

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.

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