

Solvometallurgical recovery of antimony from waste polyvinyl chloride plastic and co-extraction of organic additives

Jeroen Spooren ^{a,*}

^a. Waste Recycling Technologies, Materials & Chemistry Unit, Flemish Institute for Technological Research, VITO N.V., Boeretang 200, B-2400 Mol, Belgium.
jeroen.spooren@vito.be

Electronic supplementary information (ESI)

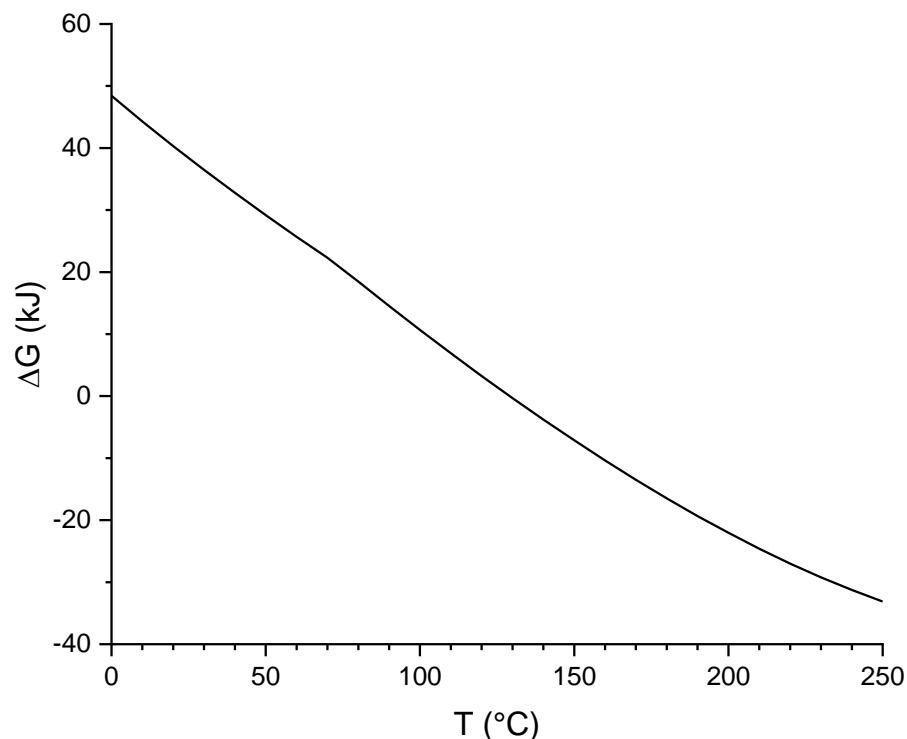


Figure S1 Gibbs free energy for the reaction $\text{Sb}_2\text{O}_5 + 3\text{HCl} \rightarrow 2\text{SbCl}_3 + 3\text{H}_2\text{O}$, as modelled by HSC Chemistry v8.0 software.



Figure S2 Photos of the PVC sample (left) as receive; (middle) cut in 1x1 cm flakes; (right) cryogenically milled to <4 mm.



Figure S3 Photo of FR1 after ethanol removal by distillation.

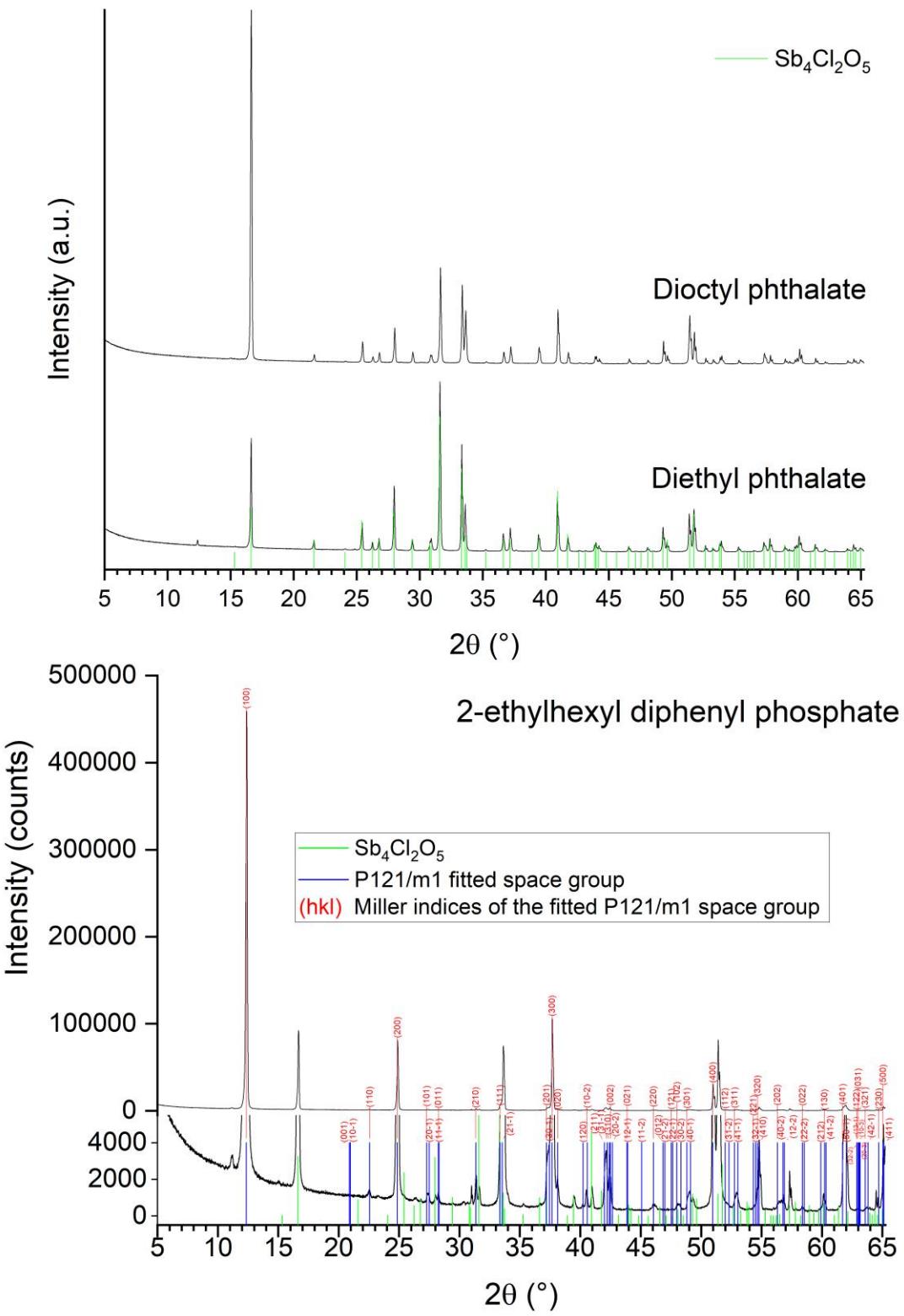


Figure S4 XRD diffractograms of the precipitates formed by hydrolysis precipitation of an acidic SbCl_3 solution in the presence of (top) phthalates and (bottom) 2-ethylhexyl diphenyl phosphate. In the bottom graph the peak positions of a fitted space group (P121/m1, $a = 8.691(4)$ Å, $b = 5.477(3)$ Å, $c = 5.160(3)$ Å, $\beta = 106.856(8)^\circ$) are plotted, accompanied by their respective Miller indices.

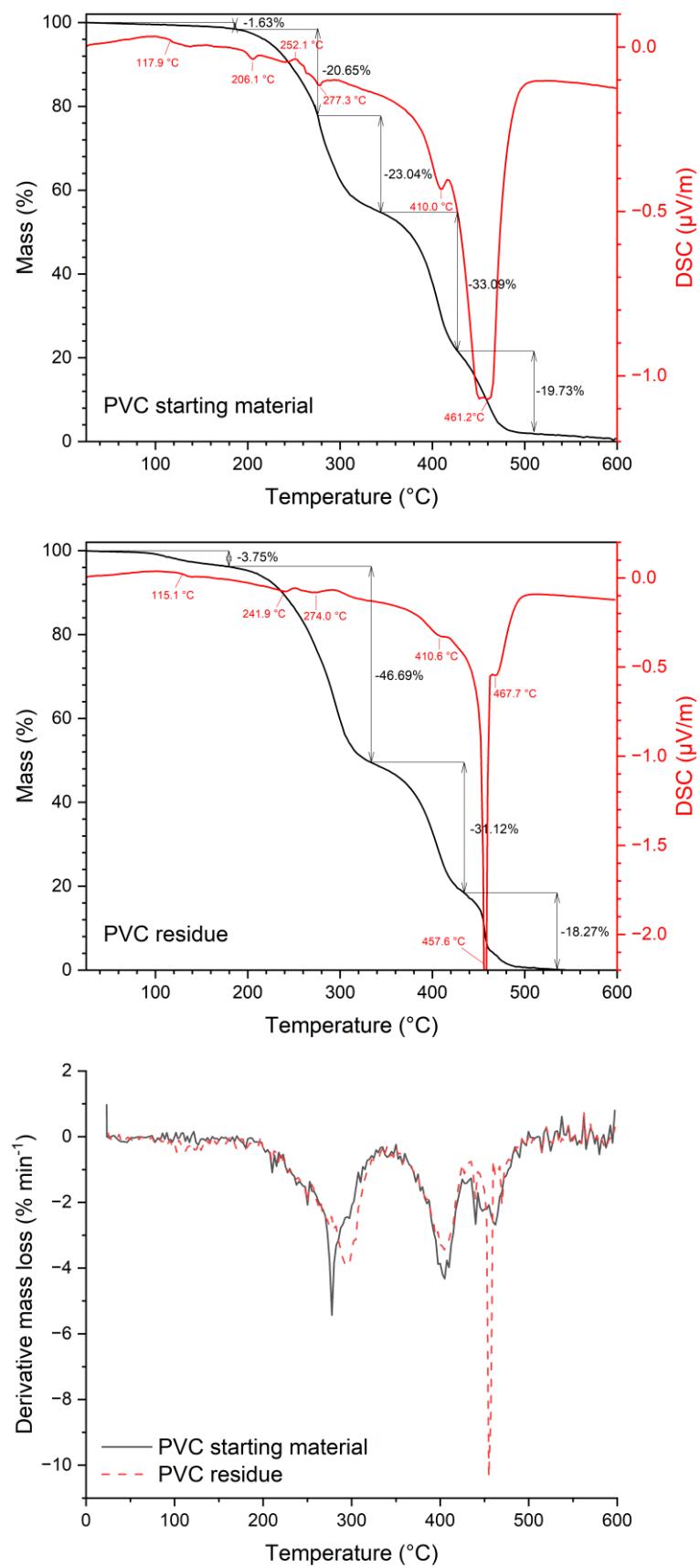


Figure S5 DSC and TGA curves of the original cryogenically milled PVC sample (top) and the residue obtained after leaching in 4 M ethanolic HCl solution at 80 °C for 4 h (bottom). The derivative mass loss of the two samples is shown in the graph at the bottom.

Table S1 Average pH of 2 M, 4 M and 6 M HCl aqueous and ethanolic solutions after leaching reactions at 50 °C and 80 °C. Standard deviations are given as error.

[HCl] (M)	Average pH aqueous solution	Average pH ethanolic solution
2	-0.095 ± 0.149	-0.422 ± 0.151
4	-0.682 ± 0.108	-1.029 ± 0.027
6	-0.848 ± 0.288	-1.272 ± 0.018