More than 80 substances from different chemical families were identified (Figure 1) and about 45 were quantified (Table 2).

**CONCLUSIONS**

Results indicate that VOCs concentrations in different sites were low. High concentrations were found in the recycling materials classification zone and in the waste unloading area. The most significant compounds are limonene (0.2-620 µg/m³), ethanol (2-1673 µg/m³), toluene (3-211 µg/m³), acetone (4-182 µg/m³), tetrachloroethylene (0.2-169 µg/m³), octanal (1.5-90 µg/m³) and n-decane (0.4-40 µg/m³), but all them were below known effect level.

**REFERENCES**

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**INTRODUCTION**

Volatile organic compounds (VOCs) released by waste treatment systems may lead to major impacts such as toxic air levels, discomfort incidents and annoying odours. The main objectives of this work were to identify and quantify selected VOCs, emitted during waste collection and processing, to control exposure hazards. An analytical method, previously developed and validated in our laboratory, for the determination of VOCs within a wide range of volatility and polarity and with variable concentrations was used.

**MATERIALS AND METHODS**

**Sampling**

Custom packed glass multi-sorbent cartridge tubes (Carbotrap 20/40, 70 mg; Carbobap X 40/60, 100 mg and Carboxen 569 20/45, 90 mg) were connected to an air collector pump sampler for the retention of VOCs.

Dynamic personal samples were taken in four sites of a waste collection centre: recycled materials classification room, waste unloading area, evaporation zone and osmosis plant. Besides, an ambiental sample of the facilities was taken.

**Desorption and analysis**

Analysis of VOCs was performed by Automatic Thermal Desorption (ATD) coupled with capillary Gas Chromatography (GC)/ Mass Spectrometry Detector (MSD), using a Perkin Elmer ATD 400 (Perkin Elmer, Boston, Massachusetts, USA) and a Thermo Quest Trace 2000 GC (ThermoQuest, San Jose, California, USA) fitted with a Thermo Quest Trace Finnigan MSD (Figure 1). VOCs standards were prepared in methanol and injected at 30°C on the multisorbent-tubes under an inert Helium gas flow (100 ml min⁻¹) using a conventional gas chromatograph packed column injector. The instrumental settings and operating conditions are shown in Table 1.

**RESULTS**

More than 80 substances from different chemical families were identified (Figure 1) and about 45 were quantified (Table 2).