MEDIUM-LONG TERM STORAGE CONDITION OF CAMBRIDGE FILTER PADS (CFPs) FOR NICOTINE DOSIMETRY



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GRAPHYCAL ABSTRACT

INTRODUCT

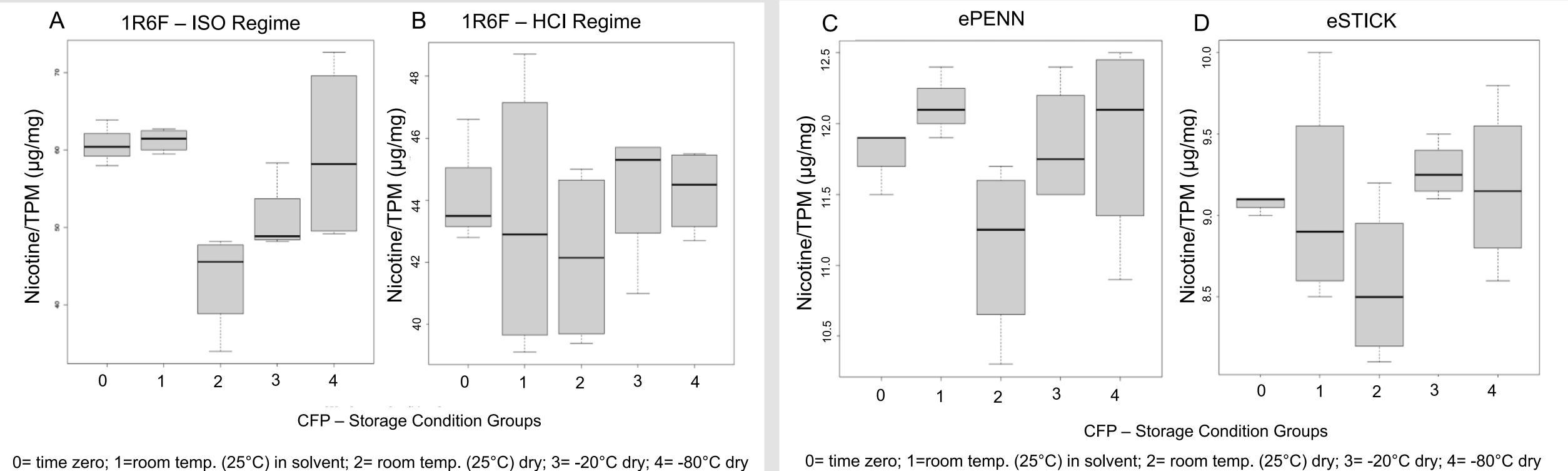
Nicotine is an alkaloid extracted from tobacco leaves. It is a dibasic compound with pyridine and pyrrolidine rings and a pKa of 8.5. Nicotine is water soluble and separates preferentially by organic solvents depending on the solution pH. Its degradation mechanisms include photolysis, thermolysis, oxidation, and hydrolysis.

Since the rising numbers of comparison between studies cigarettes and electronic nicotine delivery products (ENDs) in order to a better evaluation of health human effects, it would be necessary to establish a storing protocol for CFPs containing nicotine post exposure.

The aim of the present study was to medium-long term the assess storage condition of Cambridge Filter Pads (CFPs) for nicotine dosimetry.

The study shows that different exposure regimes and different products can affect the stability of nicotine in CFPs and storage of samples at -80 °C prevents loss of nicotine. These conditions are the best recommended in order to ensure proper harmonization of exposure tests in multicentric studies and nicotine analysis over time.

Under ISO regime, we observed a significant difference of mean value between CFP groups stored at room temperature and at temperature of -20°C (p<0.05). Moreover, the data shows a significant difference of variance within CFP groups stored at room temperature. For Vype ePen, a significant difference (p<0.05) was observed for the mean value of the CFPs group stored in solvent, higher than the mean value of CFPs of the control group.



Smoke/vapor was generated by a reference tobacco cigarette (1R6F) and Electronic Nicotine Delivery Systems (ENDS) under different exposure regimes (ISO, HCI and CRM81) and collected in CFPs. Data about nicotine was normalized by Total Particular Matter. One sample t-test and Kruskal-**Wallis** test were performed to assess statistical differences between CFPs analyzed at time zero (control) group) and the others stored under different conditions (solvent, room temperature, -20°C and -80°C) and analyzed after 30 days.

References

- CORESTA In Vitro Toxicology Task Force, 2004;
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CONCLUSION

RESULTS

METHODOLOGY

Adamson J, et al., 2017. Nicotine Quantification In Vitro : A Consistent Dosimetry Marker for e-Cigarette Aerosol and Cigarette Smoke Generation . Appl. Vitr. Toxicol. 3, 14–27. https://doi.org/10.1089/aivt.2016.0025 Pietro Zuccarello, et. Al, Nicotine dosimetry and stability in Cambridge Filter PADs (CFPs) following different smoking regimen protocols and condition storage.bioRxiv 2020.09.09.289371;

0= time zero; 1=room temp. (25°C) in solvent; 2= room temp. (25°C) dry; 3= -20°C dry; 4= -80°C dry

