

# Dermal in vitro bioavailability of HMW PAHs in Gasworks soils

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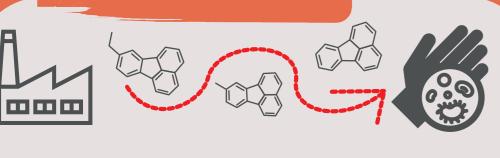
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### **BACKGROUND**



This NERC and industry funded PhD research project has investigated the relationships between the in vitro dermal bioavailability of different high molecular weight polycyclic aromatic hydrocarbons (HMW PAHs) and the bulk soil organic matter (OM) from highly contaminated gasworks sites. Both parent and alkylated HMW PAHs were investigated.

### **RESEARCH QUESTIONS**



Are there differences in the dermal bioavailability between different parent and alkylated PAHs?



Which soil bulk organic matter (OM) properties determined by **Pyrolysis** Rock-Eval(6) have influence on the dermal bioavailability of both parent and alkylated PAHs?

## **METHODS**



Dermal in vitro experiments were conducted in triplicate for each time step for each soil sample. Dermal matrices (soil, membrane and receptor solution) were dried, weighed, and processed using accelerated solvent extraction (ASE)

and solid phase extraction (SPE). Quantification of **PAHs** was determined by GC-MS/MS.

### **Experiment Conditions:**

- Soil surface area: 8.55 cm<sup>2</sup>
- Soil moisture content: 25%
- Temperature: 32 °C
- Time Steps: 1, 10, 24 h
- 5 GW soils and 1 CRM soil

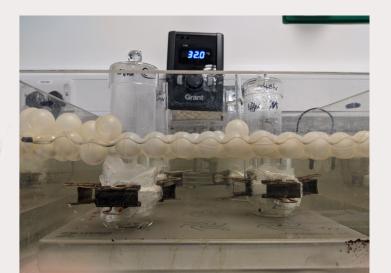
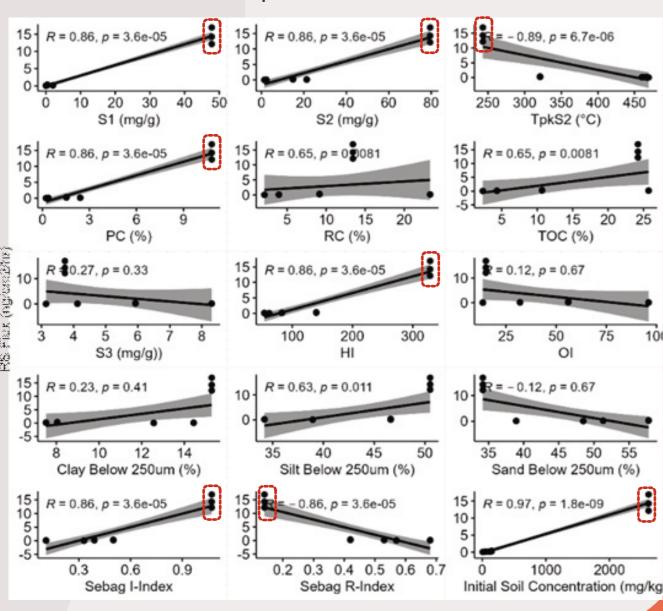


Figure: Photo of dermal in vitro experiment in lab.

### **CORRELATIONS WITH ROCK-EVAL(6) PYROLYSIS**

Rock-Eval(6) Pyrolysis (RE) is capable of measuring the release of hydrocarbons with increasing temperature. Key findings:

- Common correlation trends for RS fluxes and RE parameters for 12-13 HMW PAHs for the 5 MGP soils.
- Increasing amounts of labile OM (cracked at lower temperatures) increased dermal fluxes. E.g. High S1, S2, TpkS2, HI and I-index.
- Correlations predominantly driven by highly contaminated E1.5 (highlighted in red) - creating uncertainties with predictions.



### **SAMPLES MGP Soil Samples Pilot study ANALYSED** with CRM BCR-E1.5 H16 **E2.7 A11** 524 ∑EPA16 HMW PAHs 523 230 83.7 700 5,490 986 (mg/kg) 267 10,823 1,079 1,934 ∑PAH51 (mg/kg) 171 831 **Applied** soil x3 all x3 all x3 all x3 all x3 all x3 all timesteps timesteps timesteps timesteps timesteps timesteps **Synthetic** membrane x1 1-h & 10-h x3 at 24-h x3 all timesteps **Synthetic** receptor solution x3 24-h x3 all x3 all x3 all x3 all x3 all (RS) timesteps timesteps detected not timesteps timesteps timesteps quantified) Simplified dermal diffusion cell Measured all Measured all triplicate RS at Limited triplicate all timesteps samples samples quantified for all timesteps

### **CONCLUSION**

Some of the key findings found in this research are:



 HMW PAHs measured different dermal bioavavilbilities in different soils - suggests soil impact.



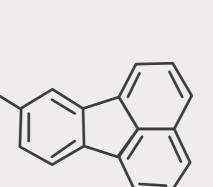
- 4-ring HMW PAHs measured highest dermal absorptions.
- HMW PAHs can remain for long periods in the skin (especially with increased ring size).
  - Alkylated C1-Fla/Pyr breakthrough into membrane and RS at longer timesteps.
  - RE could potetially help with esimating dermal bioavailbilities risks with HMW PAHs, but future work needed on a larger soil dataset.

## **SAMPLE H16 STUDY**

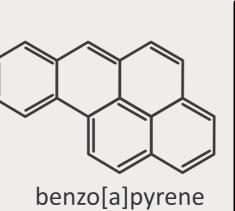
Dermal fluxes (pentration rate) for 27 parent & alkylated HMW PAH at each timestep were calculated. Examples in plots on the right. Key findings:

- Majority of available PAHs within membrane, membrane flux >> RS flux.
- Increase ring size or alkylation decreases dermal fluxes.
- The only alkylated HMW PAH compound detected and third highest flux at longer timesteps was C1fluoranthrene/pyrene (C1-Fla/Pyr).
- HMW PAHs fluxes fall below previous studies and current HHRAs guidance with BaP. (UK dermal absoprtion fraction (ABS<sub>d</sub>) 0.13, H16 ABS<sub>d</sub>was 0.0086 for 24-h).

fluoranthene (Fla) soil conc.147 mg/kg



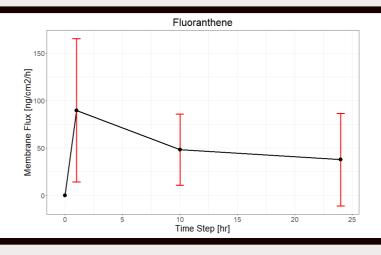
C1-Fla/Pyr soil conc.78 mg/kg

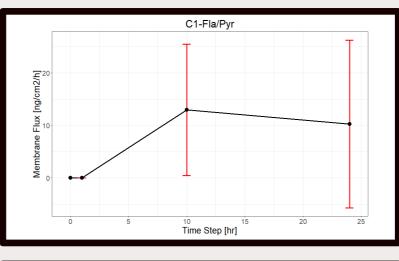


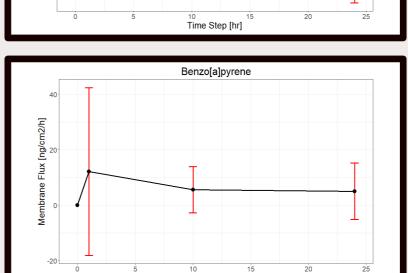
(BaP)

soil conc.150 mg/kg

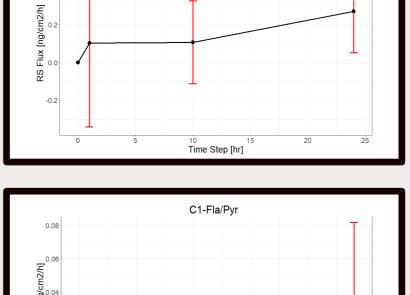
### **Membrane**

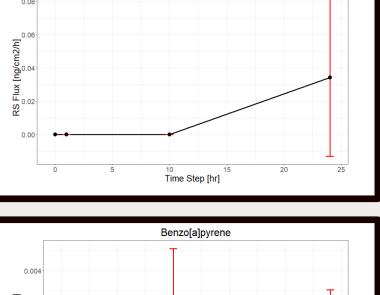


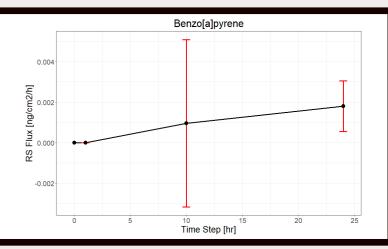




# **Receptor Solution**







\*Note y-axis flux (ng/cm2/hr) different scales, error bars 95% CI.