Supplementary Material

Polycyclic aromatic hydrocarbons in atmospheric aerosols and air-sea exchange in the Mediterranean

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12 pages, 3 tables and 5 figures

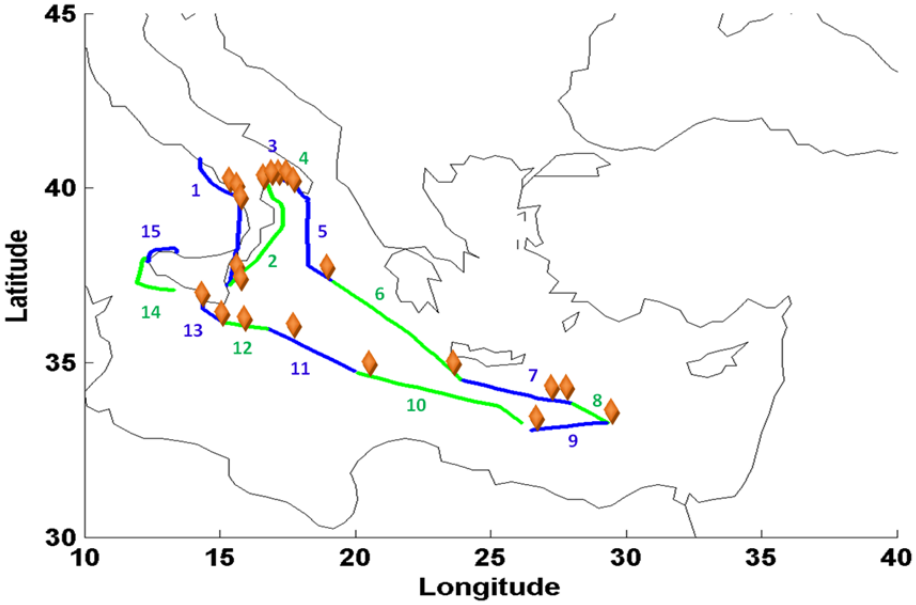
S1. Methodology

S1.1 Sampling

Table S1. Sampling times of high-volume air samples

Air sample	Start time (UTC)	Stop time (UTC)	Duration (h)
1	27.8.10 20:00	29.8.10 08:00	36
2	29.8.10 11:30	30.8.10 16:30	29
3	30.8.10 16:45	31.8.10 13:45	21
4	31.8.10 15:00	1.9.10 12:00	21
5	1.9.10 13:00	2.9.10 13:00	24
6	2.9.10 13:30	3.9.10 18:30	29
7	3.9.10 20:15	4.9.10 18:15	22
8	4.9.10 20:00	5.9.10 04:00	8
9	5.9.10 06:10	5.9.10 22:10	16
10	6.9.10 01:00	7.9.10 10:00	33
11	7.9.10 12:15	8.9.10 06:15	18
12	8.9.10 08:30	8.9.10 18:10	9.4
13	8.9.10 19:00	9.9.10 06:00	11
14	9.9.10 16:30	10.9.10 09:00	16.3
15	11.9.10 20:00	12.9.10 06:00	10

Figure S1. Spatial coverage of air (blue and green, numbers) and seawater samples (diamonds)



S1.2 Diffusive air-sea exchange flux calculation

A sensitivity analysis is done, to explore the influence of the variabilities of air and seawater temperatures and wind speed (expressed as their standard deviations) during individual sample duration on the air-sea exchange flux (Table S3).

Table S2. Sensitivity of the diffusive air-sea exchange flux, F_{aw}, calculated as $F_{aw} = | [100\times (F_{aw} + \sigma_i) - F_{aw}]/F_{aw} |$, with the standard deviations, σ_i, of i = wind speed (WS) or air or seawater temperature (T_a, T_w).

	FLT	PYR	RET
WS	146	148	185
T_a	1.3	0.19	4.1
T_w	13	9.5	4.1

S1.3 Two-box fugacity model

A non-steady state 2-box model is applied to test the hypothesis that seasonal depositional input of RET into the surface waters during the fire season (summer) triggers reversal of diffusive air-sea exchange.

The model simulations for the period 2005-2010 are initialised by fire-related RET emissions into air for the East Mediterranean (28-45°N, 8-30°E). RET was in the region eventually also emitted to air from coal and crop residues combustion (Bi et al., 2008; Shen et al., 2012), and eventually emitted to seawater influenced by pulp or paper mill effluents (Leppänen and Oikari, 1999) or by diagenesis (Alexander et al., 1995) in the region. However, these sources of RET to air and seawater are neglected as expected to contribute insignificantly and to show less inter-annual variability. Moreover, advection of RET into the model domain e.g., from fires in the western Mediterranean is neglected for simplicity.

Temperature and wind speed data were taken from the Iraklion meteorological station (35°20'N / 25°11'E, 39 m a.s.l.), located close to the centre of the model domain. Wind speed data were extrapolated to 10 m above sea level assuming neutral conditions all the time (Stull, 1988). Input data are listed in Table S3. Only wind speeds of on-shore winds were considered representative, while periods (hourly data) of off-shore winds observed at Iraklion were rejected, leading to gaps in the time series of predicted F_{aw}. No experimental data for RET lifetime in seawater exist. Degradation rate in seawater is uncertain. It was derived from a model estimated halflife against hydrocarbon biodegradability in freshwater (56 days; BioHCwin; USEPA, 2009), which could be much longer for seawater. A factor of 10 is often applied to estimate lifetime in seawater from data in freshwater (EU, 1996).

Gaseous air and seawater concentrations and the air-sea exchange flux, F_{aw}, are output.

Two scenarios are considered, an 'Initially Estimated Parameter Set' (IEPS) representing mean values for environmental parameters, and an 'Upper Estimate Parameter Set' (UEPS) which represents realistic environmental conditions favouring seawater pollution (Table S3). UEPS considers lower estimates for the atmospheric and seawater mixing layers, the degradation rate in seawater (k_{OC}) and the export (settling) velocity in seawater (v_{exp}) and an upper estimate of the of fire-related $PM_{2.5}$ emission flux.

Table S3. Input parameters for the 2-box model, initially estimated parameter set (IEPS). For the upper estimate parameter set (UEPS) FRPM25 was replaced by FRPM25×5, KOC by KOC/100, VEXP by VEXP/10, HMIX by HMIX/2, and HMIXM by HMIXM/2.

	Parameter	Unit	Value adopted or	Reference

			mean (min-max)	
COH	OH concentration in air	molec cm^{-3}	$(0.5\text{-}2.2)\times10^{6}$ during day-time, 0 during nighttime	climatological data (Spivakovsky et al., 2000) temporally interpolated
DOC	Dissolved organic carbon	µM	61.5	Pujo-Pay et al., 2011
FRPM25	fire-related $PM_{2.5}$ emission flux	mg m^{-2} h^{-1}	3.02 $(0 - 496)\times10^{-7}$	Kaiser et al., 2012
FACEMPM25	Emission factor for $PM_{2.5}$	mg (kg fuel burnt)$^{-1}$	207	Schmidl et al., 2008, Andreae 1991
HENRY	Henry coefficient of RET	Pa m^{-3} mol^{-1}	2.3	USEPA, 2009
HMIX	Atmospheric mixing height	M	1000	Estimate
HMIXM	Mixing depth in ocean	M	40	d'Ortenzio et al., 2005
KDOC	Dissolved organic carbon/water partition coefficient	L g^{-1}	158.49	Karickhoff, 1981
KOC	1st order degradation rate coefficient in seawater	s^{-1}	1.0×10^{-7} $(4\times10^{-8}$ - $4\times10^{-7})$	USEPA, 2009, T dependence: EU, 1996
KOH	Gas-phase reaction rate coefficient with OH of RET	cm^3 $molec^{-1}$ s^{-1}	4.2×10^{-11}	Lammel et al., 2010a
KOW	Octanol/water partitioning coefficient of RET		2.24×10^{6}	USEPA, 2009
KOHEOR	Factor E/R in van 't Hoff equation for OH reaction of RET	K	-20.33	Calvert et al., 2002
KPOC	Particulate organic carbon/water partitioning coefficient	L g^{-1}	2240	assumed to be given by K_{ow}; Rowe et al., 2009
KS	Setschenow constant of RET	L mol^{-1}	0.38	Jonker and Muijs, 2010
POC	Concentration of particulate organic carbon in surface seawater	µM	3.08	Pujo-Pay et al., 2011
T_a	Air temperature	K	292.1 (275.2 - 311.2)	WMO, 2013
T_w	Surface seawater temperature	K	292.1 (27482 - 311.6)	estimated from T_a:

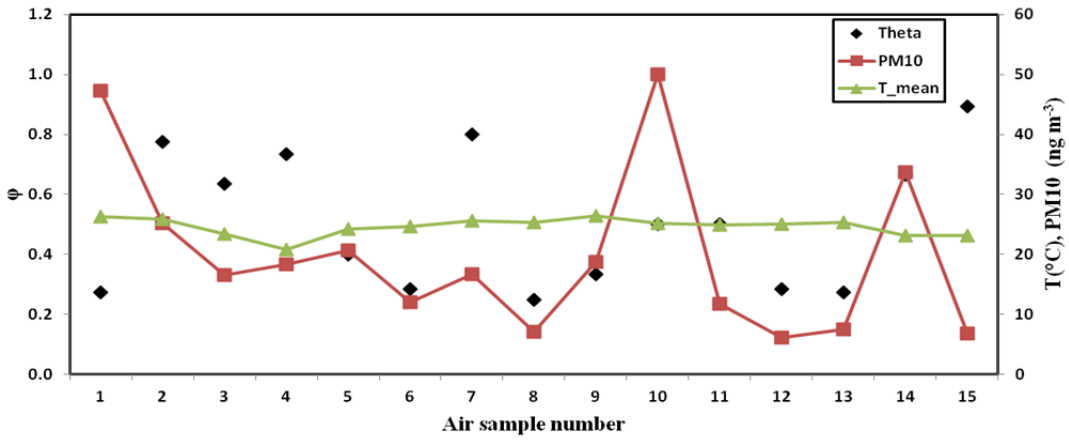
				$T_w = T_a + 0.4$ K during day-time, $T_w = T_a - 0.4$ K during nighttime
THETA	Particulate mass fraction of RET in air		0.05 (0.02 - 0.14)	Lammel et al., 2010a, T dependence: Lammel et al., 2010b
VEXP	Export (settling) velocity of particle-sorbed molecule in seawater	m s^{-1}	8×10^{-6}	Schwarzenbach et al., 2003
VDEPP	Deposition velocity of particle-sorbed molecule in air	m s^{-1}	6.5×10^{-5}	Franklin et al., 2000
WS	Wind speed	m s^{-1}	6.1 (0.6-30.7)	WMO, 2013

S2. Results

S2.1 Gas-particle partitioning

Fig. S2. Time series of particulate mass fraction (Theta), air temperature T (°C) and PM_{10} concentration (ng m^{-3}) for (a) BAA, (b) TRI, (c) CHR and (d) BBF.

a.



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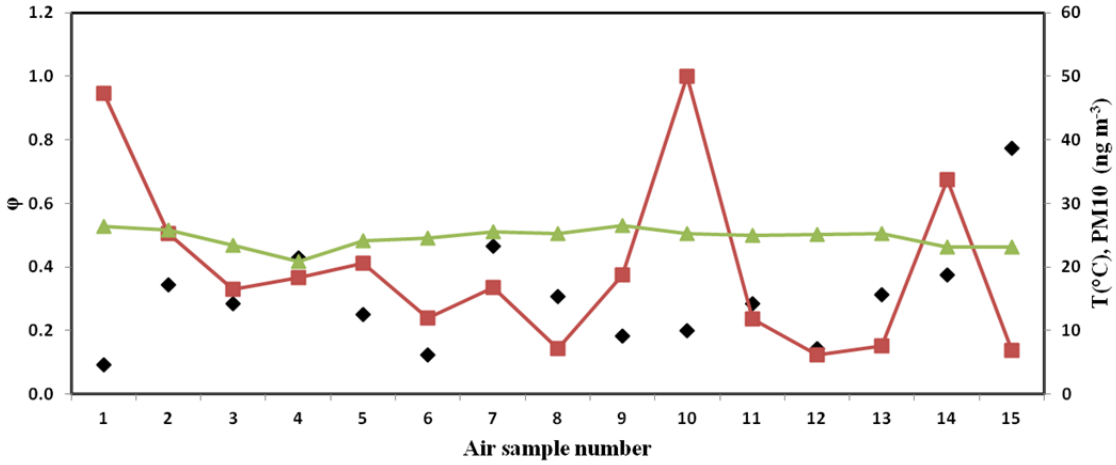

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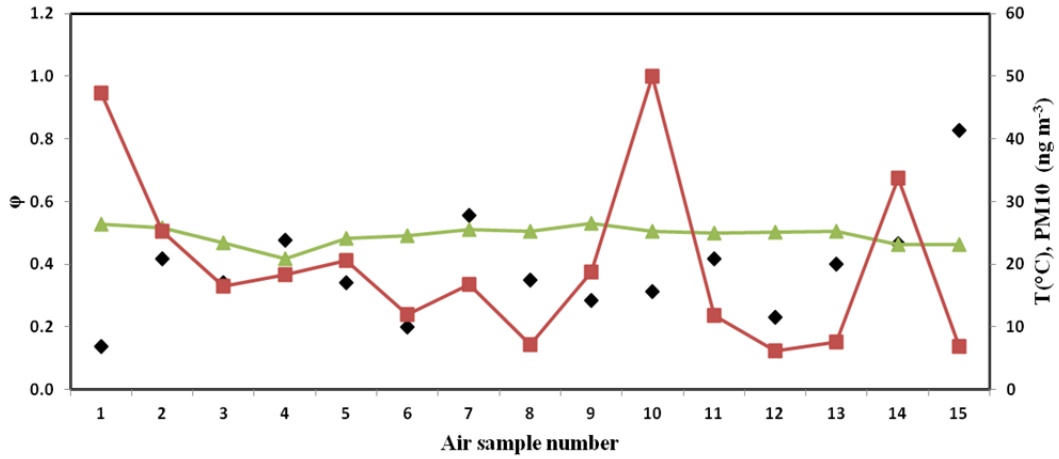
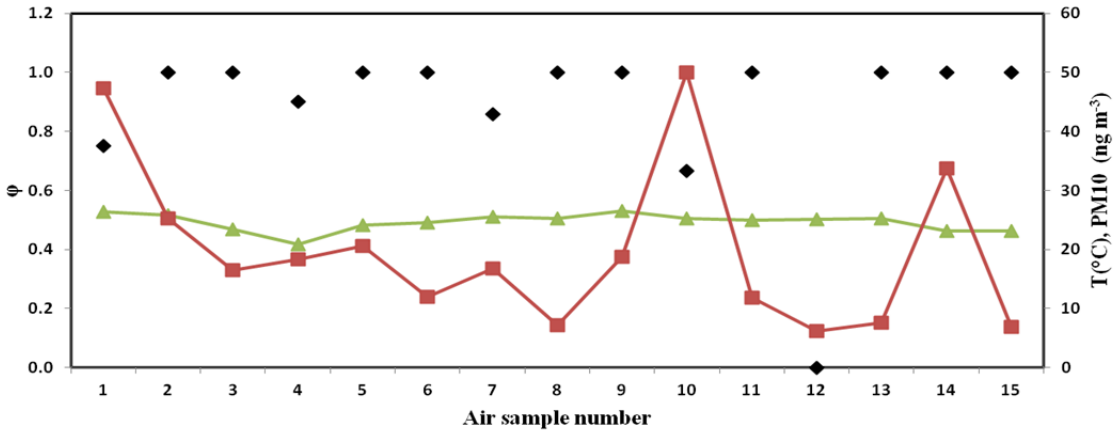


d.



S2.2 Air-sea exchange

S.2.2.1 FLT and PYR

FLT, PYR and dimethylphenanthrenes were near equilibrium or net volatilisational in coastal waters of the southeastern Mediterranean and FLT in the Black Sea, and also FLT (FR = f_a/f_w = 0.1) and PYR (FR = 0.3) in two pairs of samples collected in May 2007 in the open southeastern Mediterranean Sea, in the same regions than our samples No. 8 and 10 (Castro-Jiménez et al., 2012). In this study (June-July 2006 and May 2007), for FLT and PYR mean deposition fluxes F_{aw} = -5.87 (-11.42 – -1.11) ng m^{-2} d^{-1} and F_{aw} = -7.29 (-12.18 – -2.46) ng m^{-2} d^{-1}, respectively, were derived in the ISS, while for the SEM for FLT and PYR mean volatilisation fluxes F_{aw} = 14.33 (4.54 – 41.04) ng m^{-2} d^{-1} and F_{aw} = 15.90 (-0.56 – +62.58) ng m^{-2} d^{-1}, respectively, were derived. In our study (August-September 2010), in the ISS (paired air and water samples No. 11-13) we obtain F_{aw} = 3.14 (-38.79 – +29.16) ng m^{-2} d^{-1} and F_{aw} = 60.95 (-14.21 – +195.0) ng m^{-2} d^{-1} for FLT and PYR, respectively, and for the SEM (paired air and water samples No. 7-10) F_{aw} = - 47.36 (-169.23 – +24.45) ng m^{-2} d^{-1} and F_{aw} = -53.64 (-243.67 – +62.74) ng m^{-2} d^{-1} for FLT and PYR, respectively. This comparison shows opposite findings. However, spatial variability was very high during the 2010 cruise: In the ISS different signs of flux are indicated for the various sampling sites. In the SEM the mean fluxes derived from the paired air and water samples No. 7 and 9-10 (neglect of samples No. 8), F_{aw} = 7.48 (-2.62 - 24.45) and F_{aw} = 42.64 (17.73-62.74) for FLT and PYR, respectively, is close to the 2006-07 findings. C_a in sample No. 8 were very high, apparently because the air mass had passed over an industrial area in western Turkey (Izmir; see also SM Fig. S5a). This caused a correspondingly high deposition flux, similar to the mean annual fluxes derived for 2001-02 at Finokalia, Crete, with then lower C_w (i.e., F_{aw} = -240 ng m^{-2} d^{-1} and F_{aw} = -187 ng m^{-2} d^{-1} for FLT and PYR, respectively; Tsapakis et al., 2006). In conclusion, considering spatial and temporal variabilities and different seasons (spring vs. summer) no trend, in particular no reversal of air-sea exchange is indicated by these two data sets, 3 years apart.

S2.2.2 RET

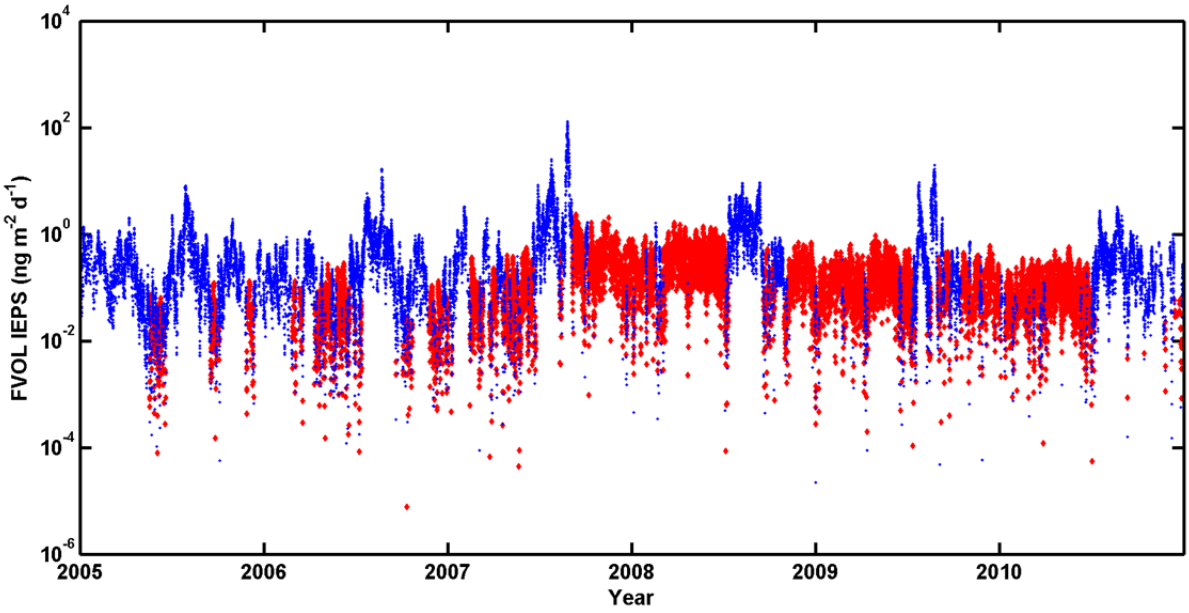
Under UEPS (Fig. S4), for 6 out of 12 observed (i.e., fugacity ratio-derived) F_{aw} (all > 0) agreement within one order of magnitude is found (underpredicting), the wrong sign (F_{aw} < 0) is predicted for 2 such cases (31.8.2010 and 2.9.2010) and no prediction was possible for 4 such cases. Note that because of a high frequency of nocturnal off-shore winds at the coastal

station from where wind speed data were adopted from (land breeze at Iraklion, Crete), data gaps in the simulated time series of F_{aw} occur more often during night-time than during day time (visible in Fig. S4). Because of the diurnal variation of temperature and wind speed these data gaps are often corresponding with maxima rather than minima of predicted F_{aw}. Underprediction could be due to neglected emissions to air and seawater in the region other than fire related (no or little seasonality) or neglect of advection into the model region (similar seasonality as captured emissions). Therefore, also the amplitude of the high frequency (daily) fluctuations could be underestimated. On the other hand, F_{aw} derived from observed concentrations C_a and C_w is uncertain, too. The biggest contribution is expected to be caused by sampling air and water not simultaneously (but combining short seawater sampling intervals with 10-20 h air sampling periods, often starting or ending when seawater samples were collected).

A sensitivity analysis (section S1.2) was performed to quantify the uncertainty of the calculated flux, F_{aw}, accounting for the variabilities of wind speed, air and seawater temperatures during sampling periods (Table S2). F_{aw} is found most sensitive to wind speed, changes on average for all the samples about 160% when adding or subtracting one SD of wind speed (hourly data) from the mean. The flux is much less sensitive to variation of the air and seawater temperatures, leading to changes of approximately 2 and 9%, respectively, when adding or subtracting one SD from the mean. While the sensitivity of F_{aw} to wind speed would be even higher when based on higher time-resolved data, hourly data appear appropriate considering mixing times of surface waters. This sensitivity to input uncertainties may explain part of the underestimate, but not up to one order of magnitude.

Fig. S3. Model predicted diffusive air-sea exchange flux, F_{aw}, of RET (ng m^{-2} d^{-1}; downward in blue and upward in red) using the initially estimated parameter set (IEPS) for the Eastern Mediterranean (28-45°N/8-30°E) 1.1.2005-31.12.2010, hourly means. Data filtered against

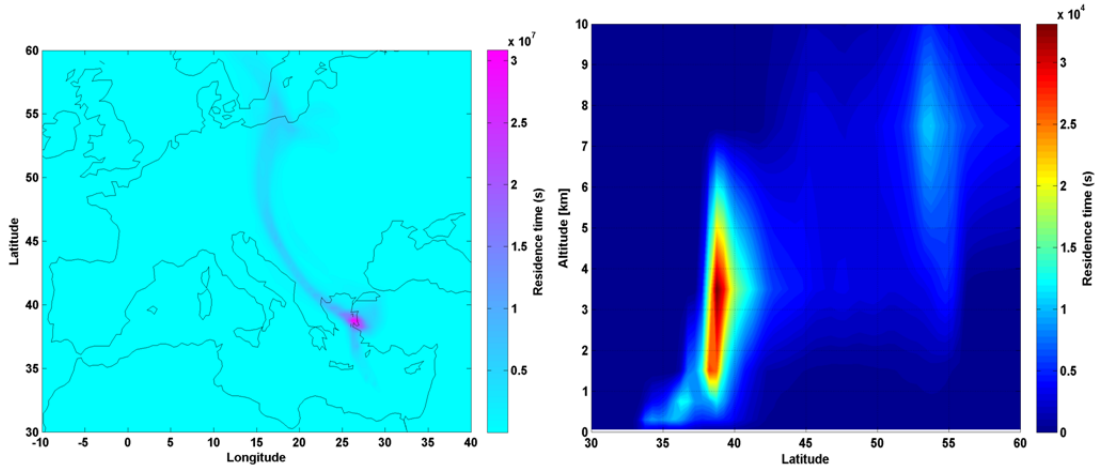
off-shore winds (see main text). (Same as Fig. 4, but IEPS)



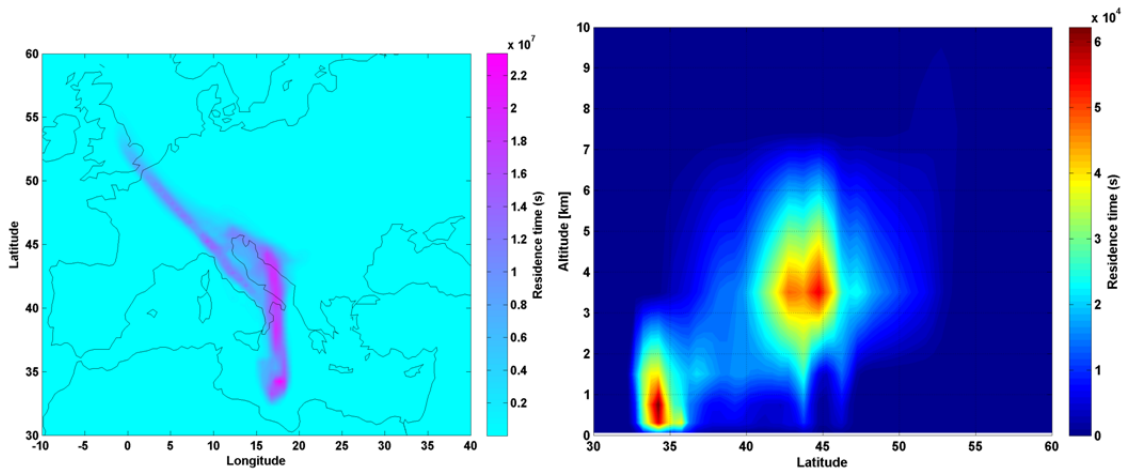
S2.3 Long-range transport

Fig. S5. Residence time distribution (left: latitude vs. longitude, right: latitude vs. altitude) of particles in backward simulations corresponding to (a) maximum and (b) minimum atmospheric PAH concentrations.

a)



b)



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