Foliage from balsam fir, Mt. Mansfield, VT; collected 6/10 and 6/11, Mike Aucott, NJDEP, Trenton, NJ and Rutgers U., Dept. Env. Sci., New Goal was to replicate findings of Hartmut Frank and others of high le trichloroacetic acid in foliage of apparently stressed conifers. Ide trichloroacetic acid may be exerting some herbicidal effect.

Foliage collected from trees at 5 to 10 meter hight, put in sealed gl refrigerated until analyzed 6/13

Then, analyzed for dichloroacetic acid (dcaa) and trichloroacetic aci in the manner of Frank, et al., Environ. Sci. and Pollut. Res., 1, 4-Frank, et al., J. High Resol. Chromatog., 18, 83-88, 1995.

Low levels of tcaa were found in the foliage of previous year's needl correlation with altitude was noted. High findings of tcaa of Frank, were not replicated.

Raw data, notes., etc. appear below...

				micrograms/liter					
elevationwashed grams			dcaa		tcaa	analyte valu- blank va			
feet	or un w	'danalyzed	vial :	#blank	blank	dcaa	tcaa	dcaa	tcaa
3850	u	1.31	3	0.652	0.274	0.132	1.17	-0.5	0.9
3620	u	1.67	4	0.652	0.274	-0.080	1.02	-0.7	0.75
3050	u	1.53	5	0.652	0.274	0.824	1.59	0.17	1.31
3050	W	1.45	6	0.652	0.274	-0.176	1.16	-0.8	0.89
2590	u	1.66	7	0.652	0.274	0.218	1.12	-0.4	0.84
2050	W	1.8	9	0.652	0.274	-0.246	0.72	-0.8	0.44
2050	u	1.71	8	0.652	0.274	72.926	277.	72.2	277.

plain water blank tcaa reading was 0.448, external standard indicated so blank assumed to be 0.274

plain water dcaa blank was -.125 and external standard indicated 1.43 so blank assumed to be 0.652

quantity in analyte was extracted into 9.0 ml water from grams of nee so ppb val is determined by multiplying (analyte-blank val) by 9/1000 dividing this quan by grams analyzed, this will give ug/g, or ppm, so 1000 to get ppb

anomalous value is assumed to be inaccurate; possible explanation is this value was from unwashed sample collected moring of 6/11, after 1 and other sample were either washed (from the 2050 elev.) or collecte before, under dry conditions, that this quan represents quan of tcaa else eluting at same time) that was in the moisture clinging to the n

4 samples of these needles, which had been placed in sealed container at -10 degrees F, were washed 6/29, and the wash analyzed the results showed very modest levels of toaa and essentially ND for so whatever caused the anomolous high reading was either gone, or not associated with the needles in the first place. The latter se the more logical explanation.

data from foliage washings (15 ml. wash water):

					ug/I				
foliage wt			tcaa	blank	tcaa	tcaa in wash			
	grams	vial #	(est	imated))val	per g foliage	(calc'd	as	ppb)
	3.86		7	0.27	0.532	1.018			
	6.7		8	0.27	0.512	0.541			
	11.82		9	0.27	1.112	1.068			
	13.31	1	0	0.27	0.504	0.263			

avg 0.723

so, looks like the amount of toaa on the surface of needles w. can be by washing amounts to about 0.7 ppb (by wt) of the total; this is cons with the pair of samples from 3050 feet above, where the washed sampl about 1.2 ppb less toaa than the unwashed sample

also of potential interest is the amount of water which fir foliage a hold. I took dry foliage samples, and spritzed water on with an atom the type used for windex. The average of three readings indicates th of needles can hold 0.6 grams of water.

Brunswick, NJ vels of a is that ass vials, d (tcaa) 14, 1994 and es. No et al ppb, foliage dcaa tcaa -3.57 6.183 -3.94 4.068 1.008 7.752 -5.14 5.548 -2.35 4.597 -4.49 2.245 380.3 1459. this value almost certainly result of error. 0.1 dles shown , and then multiply by that, since ight rain, d day (or something eedles dcaa, ems

removed
istent
e had

ppears able to
izer of
at 1 gram