

Available online at www.sciencedirect.com



Urban Forestry & Urban Greening 4 (2006) 115-123



www.elsevier.de/ufug

Air pollution removal by urban trees and shrubs in the United States

David J. Nowak*, Daniel E. Crane, Jack C. Stevens

USDA Forest Service, Northeastern Research Station, 5 Moon Library, SUNY-ESF, Syracuse, NY 13210 USA

Abstract

A modeling study using hourly meteorological and pollution concentration data from across the coterminous United States demonstrates that urban trees remove large amounts of air pollution that consequently improve urban air quality. Pollution removal (O_3 , PM_{10} , NO_2 , SO_2 , CO) varied among cities with total annual air pollution removal by US urban trees estimated at 711,000 metric tons (\$3.8 billion value). Pollution removal is only one of various ways that urban trees affect air quality. Integrated studies of tree effects on air pollution reveal that management of urban tree canopy cover could be a viable strategy to improve air quality and help meet clean air standards. Published by Elsevier GmbH.

Keywords: Air quality; Urban forests; Urban forestry; Environmental quality

1. Introduction

Air pollution is a major environmental concern in most major cities across the world. An important focus of research has been on the role of urban vegetation in the formation and degradation of air pollutants in cities. Through the emission of volatile organic compounds (VOC), urban trees can contribute to the formation of ozone (O₃) (Chameides et al., 1988). However, more integrative studies are revealing that urban trees, particularly low VOC emitting species, can be a viable strategy to help reduce urban ozone levels (Cardelino and Chameides, 1990: Taha, 1996: Nowak et al., 2000). particularly through tree functions that reduce air temperatures (transpiration), remove air pollutants (dry deposition to plant surfaces), and reduce building energy and consequent power plant emissions (e.g., temperature reductions; tree shade). One study (Nowak et al., 2000) has concluded that for the US northeast

*Corresponding author. Tel.: +1 315 448 3212;

E-mail address: dnowak@fs.fed.us (D.J. Nowak).

1618-8667/\$ - see front matter Published by Elsevier GmbH. doi:10.1016/j.ufug.2006.01.007

coast, the physical effects of urban trees were more important than the chemical effects in terms of affecting ozone concentrations.

Nationally, urban trees and shrubs (hereafter referred to collectively as "trees") offer the ability to remove significant amounts of air pollutants and consequently improve environmental quality and human health. Trees remove gaseous air pollution primarily by uptake via leaf stomata, though some gases are removed by the plant surface. Once inside the leaf, gases diffuse into intercellular spaces and may be absorbed by water films to form acids or react with inner-leaf surfaces (Smith, 1990). Trees also remove pollution by intercepting airborne particles. Some particles can be absorbed into the tree, though most particles that are intercepted are retained on the plant surface. The intercepted particle often is resuspended to the atmosphere, washed off by rain, or dropped to the ground with leaf and twig fall. Consequently, vegetation is only a temporary retention site for many atmospheric particles.

To investigate the magnitude of air pollution removal by urban trees throughout the lower 48 United States, computer modeling of air pollution removal of carbon

fax: +13154483216.

monoxide (CO), nitrogen dioxide (NO₂), ozone, particulate matter less than $10 \,\mu\text{m}$ (PM₁₀) and sulfur dioxide (SO₂) was performed for 55 US cities and for the entire nation based on meteorological, pollution concentration, and urban tree cover data. Due to the need for various assumptions within the model, the model provides a first-order estimate of the magnitude of pollution removal by urban trees.

Methods

For each city, the downward pollutant flux (*F*; in $gm^{-2}s^{-1}$) was calculated as the product of the deposition velocity (V_d ; in ms^{-1}) and the pollutant concentration (*C*; in gm^{-3}) ($F = V_dC$). Deposition velocity was calculated as the inverse of the sum of the aerodynamic (R_a), quasi-laminar boundary layer (R_b) and canopy (R_c) resistances (Baldocchi et al., 1987). Hourly estimates of R_a and R_b were calculated using standard resistance formulas (Killus et al., 1984; Pederson et al., 1995; Nowak et al., 1998) and hourly weather data from nearby airports for 1994. R_a and R_b effects were relatively small compared to R_c effects.

Hourly canopy resistance values for O₃, SO₂, and NO₂ were calculated based on a modified hybrid of bigleaf and multilayer canopy deposition models (Baldocchi et al., 1987; Baldocchi, 1988). Canopy resistance (R_c) has three components: stomatal resistance (r_s) , mesophyll resistance (r_m) , and cuticular resistance (r_t) , such that: $1/R_c = 1/(r_s + r_m) + 1/r_t$. Mesophyll resistance was set to zero s m⁻¹ for SO₂ (Wesely, 1989) and 10 s m^{-1} for O₃ (Hosker and Lindberg, 1982). Mesophyll resistance was set to $100 \,\mathrm{s}\,\mathrm{m}^{-1}$ for NO₂ to account for the difference between transport of water and NO₂ in the leaf interior, and to bring the computed deposition velocities in the range typically exhibited for NO₂ (Lovett, 1994). Base cuticular resistances were set at 8000 sm^{-1} for SO₂, 10,000 sm⁻¹ for O₃, and $20,000 \,\mathrm{s}\,\mathrm{m}^{-1}$ for NO₂ to account for the typical variation in r_t exhibited among the pollutants (Lovett, 1994).

As removal of CO and particulate matter by vegetation are not directly related to photosynthesis/ transpiration, R_c for CO was set to a constant for in-leaf season (50,000 sm⁻¹) and leaf-off season (1,000,000 sm⁻¹) (Bidwell and Fraser, 1972). For particles, the median deposition velocity (Lovett, 1994) was set to 0.064 m s⁻¹ based on 50-percent resuspension rate (Zinke, 1967). The base Vd was adjusted according to in-leaf vs. leaf-off season parameters. To limit deposition velocities were set to zero during periods of precipitation.

Each city was assumed to have a single-sided leaf area index within the canopy covered area of 6 and to be 10% coniferous (Nowak, 1994). Leaf area index value is total leaf area (m^2 : trees and large shrubs [minimum 1 in stem diameter]) divided by total canopy cover in city (m^2) and includes layering of canopies. Regional leaf-on and leaf-off dates were used to account for seasonal leaf area variation. Total tree canopy cover in each city was based on aerial photograph sampling (Nowak et al., 1996) or advanced very high resolution radiometer data (Dwyer et al., 2000; Nowak et al., 2001).

Hourly pollution concentration data (1994) from each city were obtained from the US Environmental Protection Agency (EPA). Missing hourly meteorological or pollution-concentration data were estimated using the monthly average for the specific hour. In some locations, an entire month of pollution-concentration data may be missing and are estimated based on interpolations from existing data. For example, O₃ concentrations may not be measured during winter months and existing O_3 concentration data are extrapolated to missing months based on the average national O₃ concentration monthly pattern. Data from 1994 were used due to available data sets with cloud cover information. To estimate percent air quality improvement due to dry deposition (Nowak et al., 2000), hourly boundary heights were used in conjunction with local deposition velocities for select cities with boundary layer height data. Daily morning and afternoon mixing heights from nearby stations were interpolated to produce hourly values using the EPA's PCRAMMIT program (US EPA, 1995). Minimum boundary-layer heights were set to 150 m during the night and 250 m during the day based on estimated minimum boundary-layer heights in cities. Hourly mixing heights (m) were used in conjunction with pollution concentrations ($\mu g m^{-3}$) to calculate the amount of pollution within the mixing layer ($\mu g m^{-2}$). This extrapolation from ground-layer concentration to total pollution within the boundary layer assumes a well-mixed boundary layer, which is common in the daytime (unstable conditions) (Colbeck and Harrison, 1985). Hourly percent air quality improvement was calculated as grams removed/(grams removed + grams in atmosphere), where grams in atmosphere = measured concentration $(gm^{-3}) \times$ boundary layer height (m) \times city area (m^2) .

To estimate pollution removal by all urban trees in the United States, national pollution concentration data (all EPA monitors) were combined with standardized local or regional pollution removal rates. Pollution removal rates (gm^{-2} of tree cover) standardized to the average pollutant concentration in the city (gm^{-2} per ppm or per μgm^{-3}). As flux rates are directly proportional to pollutant concentrations, standardized removal rates are used to account for concentration differences among urban areas.

For all urban areas in the United States outside of the 55 analyzed cities, local pollution monitoring data were

used to calculate the average pollution concentration in the urban area for each pollutant. Urban area boundaries are based on 1990 census definitions of urbanized areas (areas with population density ≥ 1000 people mi⁻²) and urban places (incorporated or unincorporated (census-defined) places with a population ≥ 2500) outside of urbanized areas. If pollutant monitors did not exist within the urban area, minimum state pollution concentration data were assigned to the urban area. Likewise, standardized pollution removal rates were assigned to each urban area based on data from the closest analyzed city within the same climate zone. All urban areas within a state were assigned to the dominant climate zone (cool temperate, Desert, Mediterranean, steppe, tropical, tundra, warm temperate) in the state, except for California and Texas where urban areas were individually assigned to one of multiple state climate zones.

For each urban area exclusive of the 55 analyzed cities, standardized pollution removal rates were multiplied by average pollutant concentration and total amount of tree cover to calculate total pollution removal for each pollutant in every urban area. Urban area pollution removal totals were combined to estimate the national total. Pollution removal value was estimated using national median externality values (Murray et al., 1994). Values were based on the median monetized dollar per ton externality values used in energy-decision-making from various studies. These values, in dollars per metric ton (t) are: $NO_2 = \$6752 t^{-1}$, $PM_{10} = \$4508 t^{-1}$, $SO_2 = \$1653 t^{-1}$, and $CO = \$959 t^{-1}$. Externality values for O_3 were set to equal the value for NO₂. Externality values can be considered the estimated cost of pollution to society that is not accounted for in the market price of the goods or services that produced the pollution.

Results and discussion

Total pollution removal and value varied among the cities from $11,100 \text{ ta}^{-1}$ (\$60.7 million a⁻¹) in Jacksonville, FL to 22 ta^{-1} (\$116,000 a⁻¹) in Bridgeport, CT (Table 1). Pollution removal values per unit canopy cover varied from $23.1 \text{ gm}^{-2} \text{ a}^{-1}$ in Los Angeles, CA to $6.2 \text{ gm}^{-2} \text{ a}^{-1}$ in Minneapolis, MN. The median pollution removal value per unit canopy cover was $10.8 \text{ gm}^{-2} \text{ a}^{-1}$.

Pollution removal values for each pollutant will vary among cities based on the amount of tree cover (increased tree cover leading to greater total removal), pollution concentration (increased concentration leading to greater downward flux and total removal), length of in-leaf season (increased growing season length leading to greater total removal), amount of precipitation (increased precipitation leading to reduced total removal via dry deposition), and other meteorological variables that affect tree transpiration and deposition velocities (factors leading to increased deposition velocities would lead to greater downward flux and total removal). All of these factors combine to affect total pollution removal and the standard pollution removal rate per unit tree cover.

Jacksonville's urban forest had the largest total removal, but had below median value of pollution removal per unit tree cover. Jacksonville's high total pollution removal value was due to its large city size (1965 km²) and relatively high estimated percent tree cover within the city (53%). Los Angeles had the highest pollution removal values per unit tree cover due to its relatively long in-leaf season, relatively low precipitation, and relatively high pollutant concentrations and deposition velocities. Minneapolis had the lowest pollution removal values per unit tree cover due, in part, to its relatively short in-leaf season.

Average leaf-on daytime dry deposition velocities varied among the cities ranging from 0.44 to 0.29 cm s⁻¹ for NO₂, 0.40 to 0.71 cm s⁻¹ for O₃, and 0.38 to 0.69 cm s⁻¹ for SO₂. Deposition velocities did not vary for CO and PM₁₀ as deposition rates for these pollutants were not related to transpiration rates, but rates did vary based on leaf-off and leaf-on seasons. The deposition velocities for CO and PM₁₀ were based on literature averages and assumed to be constant. The highest deposition velocities occurred in San Jose, CA; the lowest in Phoenix, AZ.

Though urban trees remove tons of air pollutants annually, average percent air quality improvement in cities during the daytime of the vegetation in-leaf season were typically less than 1 percent (Table 2) and varied among pollutants based on local meteorological and pollution concentration conditions. Percent air quality improvement was typically greatest for particulate matter, ozone, and sulfur dioxide. Air quality improvement increases with increased percent tree cover and decreased mixing-layer heights. In urban areas with 100% tree cover (i.e., contiguous forest stands), average air quality improvements during the daytime of the inleaf season were around two percent for particulate matter, ozone, and sulfur dioxide. In some cities, shortterm air quality improvements (one hour) in areas with 100% tree cover are estimated to be as high as 16% for ozone and sulfur dioxide, 9% for nitrogen dioxide, 8% for particulate matter, and 0.03% for carbon monoxide (Table 2).

These estimates of air quality improvement due to pollution removal likely underestimate the total effect of the forest on reducing ground-level pollutants because they do not account for the effect of the forest canopy in preventing concentrations of upper air pollution from reaching ground-level air space. Measured differences in

City		O_3		PM_{10}		NO_2		SO_2		00		Total		
		(t)	$(\mathrm{gm^{-2}})$	(t)	$(g m^{-2})$	(t)	$(g m^{-2})$	(t)	$(\mathrm{g}\mathrm{m}^{-2})$	(t)	$(\mathrm{gm^{-2}})$	(t)	$(\mathrm{gm^{-2}})$	$(\$ \times 1000)$
Albany, NY ^a	Г	43	3.4	30	2.4	13	1.0	7	0.5	1	0.1	94	7.4	524
	¥ ⊦	12–54 78	1.0-4.3 2.4	12-47 84	0.9–3.7 7.6	6–15 37	0.5–1.2	4-10	0.3-0.8	na 11	na 0.2	35–128 244	2.7–10.1	183-698 1720
titut (anhtanhnatu	- X	7.6 26–123	0.8-3.9	33–132	1.0-4.1	ر 19–48	0.6-1.5	22–69	0.7 - 2.1	na	u.u na	244 111–382	3.5-12.0	501-1870
Atlanta, GA	Т	514	4.6	423	3.8	135	1.2	93	0.8	35	0.3	1200	10.7	6470
	R	106 - 610	0.9 - 5.4	165–661	1.5-5.9	61-156	0.5 - 1.4	40-137	0.4 - 1.2	na	na	407 - 1600	3.6-14.2	1970-8410
Austin, TX	ΗR	1150 295-1510	5.2 1 3-6 9	680 266–1060	3.1 1 7–4 8	436 180–543	2.0 0 8-2 5	374 154–596	1.7 0 7–2 7	138 113	0.6 na	2770 1030–3850	12.6 4 7–17 5	14500 4790–19800
Baltimore, MD	Τ	158	4.0	148	3.7	107	2.7	50	1.3	13	0.3	477	12.1	2550
	R	39–214	1.0 - 5.4	58-232	1.5 - 5.9	43-133	1.1 - 3.4	25-84	0.6 - 2.1	na	na	178-676	4.5–17.1	869–3540
Baton Rouge, LA	Т	508	4.8	369	3.5	133	1.3	97	0.9	50	0.5	1160	11.0	6200
	R	108-634	1.0-6.0	144-576	1.4-5.5	58-165	0.6 - 1.6	42–156	0.4 - 1.5	na	na	400-1580	3.8–15.0	1880-8300
Boston, MA		101	3.8	75	2.8	57	2.1	30 15 15	1.1	6	0.3	272	10.2 2 0 12 6	1470 402 1800
Bridgenort CT	¥ ⊢	2 4 –121 و	0.9-4.0 3 A	21110 7	1.1-4.4 2 0	24-00 1	C.2-4.0 7 1	10-4-0 ر	0.0-1./	па 1	Па 0 2	105-201 CC	0.01-0.0	492-1090 116
DINGCOUT, CI	- 2	0 0-0	0.8-3.9	,	2.0 1.1-4.4	+ C 4	0.7–1.8	- - 7	0.0	1 113	C.V BH	22 8-28	3.3-11.5	39–145
Buffalo, NY	H	72	3.7	41	2.1	20	1.0	29	1.5	ε	0.2	165	8.5	858
	R	18-85	0.9 - 4.3	16-64	0.8 - 3.3	9–23	0.5 - 1.2	12-42	0.6 - 2.1	na	na	58-216	3.0-11.1	274 - 1090
Charleston, WV	Τ	98	2.7	71	1.9	33	0.9	37	1.0	4	0.1	243	6.7	1270
	R	26-153	0.7 - 4.2	28-111	0.8 - 3.1	16-44	0.4 - 1.2	21–71	0.6 - 1.9	na	na	95–383	2.6 - 10.5	448–1950
Cincinnati, OH	F	213	3.0	241	3.3	126	1.8	85	1.2	19	0.3	683	9.5	3530
	R	51-278	0.7 - 3.9	94-376	1.3 - 5.2	52-155	0.7 - 2.2	37-131	0.5-1.8	na	na	253-959	3.5-13.3	1200-4860
Cleveland, OH	(- r	191	3.7	240	4.7	76	1.5 2	72	1.4	13	0.3	592	11.6	3010
	¥ F	40-227	0.9-4.0	04-370 272	1.8-7.4	34-89 01	0.7-1.7	31-107	0.0-2.1	na	na 0 4	218-811	4.3-15.9	1020-4010
Columna, SC		00L C7C	4.1 7 7 7 7	D/C	0.0	74 711 FA	0.100	1/1 70 701	CC 20	4 i	0. 1	0171	0.4	01/10
Columbus OH ^c	⊻⊦	143-728 448	2 7 2	147–587 357	1.2-4.6 2.0	4//II/ 211	0.4-0.9 1 8	167-8/	0.6-2.3	na v	na 0 0	464-17/0 1110	3.7-14.0 0.2	2120-8880 6210
Columbus, On	- 0	440 113 613	0.0 5 1	100 558	0.0 1 7 1 6	096 117	0.1 0	101 01	0.0	77	7.0 5.0	401 1570	3.2 13.0	0710 8640
Dallas, TX	4 F	1,160	4.7	905	3.6	296	1.2	110	0.4	124 124	0.5	2600	10.5	14200
	R	286 - 1420	1.2-5.7	354 - 1410	1.4-5.7	129–347	0.5 - 1.4	52-163	0.2 - 0.7	na	na	945–3470	3.8 - 14.0	4600 - 18700
Denver, CO	H	213	2.1	321	3.1	181	1.8	76	0.7	22	0.2	813	7.9	4250
	α θ	66 - 313 334	0.6 - 3.0	125-510 357	1.2-4.8	92-232 95	0.9-2.2	43–131 20	0.4-1.3	na	na	349–1210	3.4–11.6	1720–6180 2440
Detroit, MI	- 0	707 03	1.0	COF 001	0.4 12 52	00 C0	1.1 0 6 1 2	21 00	0.7	71	7.0	100 200	0./ 2.7 11 0	0440 1710 4550
FI Paso TX	4 ⊢	00-200 140	3.1	100-402 176	6.0-0.1	<i>ور</i> –۲4	0.1-0.0 0.4	06–10 47	c.1- 1 -0	19 19	11a 0 4	240-071 401	0.11-2.6	1210-4330 1960
	R	47-247	1.1 - 5.6	69–275	1.5-6.2	10-26	0.2 - 0.6	31–95	0.7 - 2.1	na	na	176-663	4.0 - 14.9	768-3260
Fresno, CA	Τ	221	5.1	292	6.7	69	1.6	28	0.6	13	0.3	622	14.3	3330
	R	66-390	1.5 - 8.9	114-456	2.6 - 10.4	31–97	0.7 - 2.2	16-58	0.4 - 1.3	na	na	239-1010	5.5-23.2	1210-5450
Honolulu, HI	Η¢	467	6.6 1 C T C	354 120 FF2	5.0	36 12 12	0.5	37	0.5	43	0.6	937	13.3	5090
Houston TY	⊻⊢	022-111	C./−0.1 4.5	200-851 817	2.0-1.2 7 7	13-42 356	0.2-0.0 2 0	217 217	0.2-0.8 1 8	na 73	na 0.4	319-1220 2340	4.2–1/.5 13.4	1330-0460 11900
TTOUSION, LY	- 2	164–968	0.9-5.5	319–1280	1.8-7.3	143–433	0.8-2.5	126-489	0.7 - 2.8	na	na na	826-3240	4.7–18.5	3790-16100
Indianapolis, IN	Τ	1280	4.5	958	3.4	298	1.0	317	1.1	62	0.2	2910	10.3	15500
•	R	313-1630	1.1-5.7	374-1500	1.3 - 5.3	147 - 364	0.5 - 1.3	133-471	0.5–1.7	na	na	1030-4020	3.6-14.2	5070-21000

Table 1. (continued)														
City		03		PM_{10}		NO_2		SO_2		CO		Total		
		(t)	$(\mathrm{gm^{-2}})$	(t)	$(g m^{-2})$	(t)	(gm^{-2})	(t)	$(g m^{-2})$	(t)	$(\mathrm{gm^{-2}})$	(t)	(gm^{-2})	$($ \times 1000)$
Jacksonville, FL	μe	5210	5.0	3570	3.4	1137	1.1	931	0.9	295	0.3	11100	10.7	60800
Ionore City, NId	⊻⊦	11/0-62/0	1.1-6.0 2.6	1390–080 74	1.3-5.4	452-1380	0.4-1.3 7 7	355-1400 0	0.3–1.3 1 6	na 1	na 0.7	3670-14900 74	3.5-14.5 12.7	18100-/9400 275
JEISEY CILY, INJ	- 2	21 5–28	0.8-4.8	2 4 9–38	1.6-6.5	7–19	$\frac{2.7}{1.1-3.3}$	م 4–15	0.8-2.6	na 4	u./ na	74 29–103	5.0-17.8	131–515 131–515
Kansas City, MO	H	1150	4.0	940	3.3	209	0.7	236	0.8	82	0.3	2620	9.2	13900
	Ч	285 - 1400	1.0 - 4.9	367 - 1470	1.3 - 5.2	97–248	0.3 - 0.9	106 - 348	0.4 - 1.2	na	na	937–3550	3.3-12.5	4490–18400
Los Angeles, CA	Н	1260	6.9	1470	8.0	1150	6.3	105	0.6	228	1.2	4210	23.1	23300
	Ч	191-1290	1.0-7.1	573-2290	3.1–12.6	377-1350	2.1–7.4	42–161	0.2 - 0.9	na	na	1410-5320	7.7-29.2	6700-28600
Louisville, KY		189 16 350	4.3	153	3.5	60 77 75	1.4	84	1.9	12	0.3	498 170 712	11.3	2520 828 7510
Memphis. TN	4 ⊢	40-230 726	4.8	60–200 547	1.4-J.4 3.6	352	0.0-1./ 2.3	061-00 150	1.0-0.0 1.0	па 65	0.4	1/9-/12 1840	4.0-10.1 12.1	0100-020
	2	171–955	1.1 - 6.3	214-855	1.4-5.6	145-435	0.9-2.9	65-239	0.4 - 1.6	na	na.	655-2550	4.3 - 16.8	3260-13700
Miami, FL	Τ	104	5.5	88	4.6	33	1.7	7	0.4	10	0.5	243	12.7	1350
	Ч	24-121	1.3 - 6.4	34-137	1.8 - 7.2	12-40	0.6 - 2.1	3-11	0.2 - 0.6	na	na	84–320	4.4 - 10.8	414-1740
Milwaukee, WI	Ηı	157	3.3	105	2.2	55	1.2	23	0.5	6	0.2	350	7.4	1950
1. M. 1.	۲ ۲	39-180	0.8–3.8	41 - 165	0.9–3.5	25–63 11	0.5-1.3	11–32	0.2-0.7	na	na 0,7	125-449 177	2.6-9.5	645-2440 1040
Minneapous, MiN	- 2	8/ 25-107	$3.1 \\ 0.9 - 3.7$	30 12-47	0.4 - 1.7	44 20-51	0 7–1 8 0 7–1 8	$^{-10}$	$0.2 \\ 0.1 - 0.3$	x a	0.3 na	1/0 68-222	0.2 2 4-7 8	1040 370–1300
Nashville, TN	4 🛏	2190	3.4	2180	3.4	799	1.2	523	0.8	186	0.3	5870	9.1	31000
	Ч	577-3060	0.9 - 4.7	850-3400	1.3 - 5.3	372-976	0.6 - 1.5	237-851	0.4 - 1.3	na	na	2220-8470	3.5-13.1	10800 - 44200
New Orleans, LA	Н	467	4.8	471	4.8	162	1.6	66	1.0	63	0.6	1260	12.8	6600
	2	107-587	1.1 - 6.0	184–736	1.9-7.5	70-207	0.7 - 2.1	43-159	0.4 - 1.6	na	na	470–1750	4.8-17.8	2160-9000
New York, NY	H A	491	3.7	493 102 771	3.7	478	3.6	232	1.7	97	0.7	1790 706 2410	13.5	9240 2220 12200
Newark NI	⊻⊦	11/-623 30	0.9-4.7 3.6	193-//1 40	4 8 4 8	188268 73	1.4-4.5 7 7	0 0	0.8-2.7	na 3	na 03	/06-2410 105	5.3-18.2 12.5	3020-12200 556
	- 2	7-40	0.8-4.7	16-63	1.9–7.5	10-28	1.1 - 3.3	5-15	0.6-1.8	'na	u.o	39–148	4.7–17.6	193–767
Oklahoma City, OK	Г	1170	5.7	575	2.8	161	0.8	157	0.8	61	0.3	2120	10.4	11900
	Ч	284 - 1380	1.4 - 6.8	224-898	1.1 - 4.4	73-185	0.4 - 0.9	65-229	0.3 - 1.1	na	na	708–2760	3.5-13.6	3590-15100
Omaha, N E^e	Ηı	192	2.6	244	3.4	53	0.7	48	0.7	13	0.2	551	7.6	2850
Dhiladalnhia DA	⊻ ⊦	55-241 780	0.8–3.3 3 8	95-382 417	1.3-5.3 5 5	25-63 152	0.3-0.9	20-71 152	0.3-1.0 2.0	na 26	na 03	208-770 1040	2.9-10.6 13.7	1020-3900 5130
t miauvipina, t A	- 2	76-379	1.0-5.0	$\frac{1}{163}$ -651	2.2–8.6	67-184	0.9-2.4	132 74–240	1.0-3.2	na	u.v na	405-1480	5.4–19.6	1930-7160
Phoenix, AZ	H	404	4.3	670	7.2	219	2.3	21	0.2	86	0.9	1400	15.0	7350
	Ч	129-802	1.4 - 8.6	262 - 1050	2.8-11.2	102 - 336	1.1 - 3.6	14-49	0.1 - 0.5	na	na	593-2320	6.3-24.8	2850-12600
Pittsburgh, PA	Γ	147	3.6	153	3.8	65	1.6	96	2.4	10	0.2	470	11.6	2290
	Ч	34–184	0.8 - 4.5	60–238	1.5-5.9	29–76	0.7 - 1.9	45–141	1.1 - 3.5	na	na	177-650	4.4 - 16.0	778–3080
Portland, OR	Η¢	406 88 583	3.0 2.1 2.1	449 175 701	3.1	203 203	1.5	132	1.0	87	0.6	1280	9.2	6440 33.50 8330
Description D1	¥ ⊦	78C-86	0.7-4.3 2.7	10/-0/1	2.0-2.1 2.7	92-201	0.7–1.9 1.2	59-240 14	0.4-1.8	na 2	na C C	0/81-115 146	3.8-13.8 0.9	2250-9350 760
I TUVIUUIUU, INI	- 2	уо 13—63	0.9-4.2	22–86	7.7 1.5–5.8	8-20	0.6-1.3	7–20	0.5 - 1.4	r na	u.∠ na	53-193	3.6–13.0	255–987
Roanoke, VA	Г	134	4.6	118	4.0	29	1.0	32	1.1	٢	0.2	319	11.0	1690
	Ч	32-169	1.1 - 5.8	46 - 184	1.6 - 6.3	14-35	0.5 - 1.2	14-51	0.5 - 1.7	na	na	113-446	3.9–15.3	549–2300
Sacramento, CA	H	170	4.9	134	3.8	49	1.4	12	0.3	13	0.4	378	10.8	2110
	¥	42–255	1.2 - 7.3	53-210	1.5 - 6.0	23-70	0.6 - 2.0	6–22	0.2 - 0.6	na	na	136–569	3.9 - 16.3	698-3190

× .														
City		O ₃		PM_{10}		NO_2		SO_2		CO		Total		
		(t)	$(g m^{-2})$	(t)	(gm^{-2})	(t)	(gm^{-2})	(t)	$(\mathrm{g}\mathrm{m}^{-2})$	(t)	$(g m^{-2})$	(t)	$(\mathrm{gm^{-2}})$	$($ \times 1000)$
Salt Lake City, UT	Г	219	3.0	381	5.2	116	1.6	37	0.5	22	0.3	774	10.5	4060
ì	К	68-347	0.9 - 4.7	149–595	2.0 - 8.1	60-152	0.8 - 2.1	20-71	0.3 - 1.0	na	na	318-1190	4.3 - 16.2	1590-6190
San Diego, CA	F	549	7.6	401	5.6	202	2.8	56	0.8	48	0.7	1260	17.4	7020
	Ч	105 - 603	1.5 - 8.4	157-627	2.2-8.7	72–243	1.0 - 3.4	21 - 84	0.3 - 1.2	na	na	403 - 1610	5.6 - 22.2	1980-8730
San Francisco, CA	H	80	3.5	107	4.7	63	2.7	12	0.5	15	0.7	276	12.1	1480
	Ч	19 - 90	0.8 - 3.9	42 - 166	1.8 - 7.3	22-75	1.0 - 3.3	4-17	0.2 - 0.8	na	na	103 - 364	4.5-15.9	491 - 1910
San Jose, CA ^f	μ	305	4.6	243	3.7	188	2.8	28	0.4	34	0.5	798	12.0	4500
	Ч	55-312	0.8 - 4.7	95 - 380	1.4 - 5.7	67 - 207	1.0 - 3.1	10 - 38	0.1 - 0.6	na	na	261 - 972	3.9 - 14.6	1300 - 5320
Seattle, WA ^g	μ	105	3.3	98	3.1	47	1.5	48	1.5	21	0.6	319	10.0	1570
	Ч	23-136	0.7 - 4.3	38 - 153	1.2 - 4.8	21 - 60	0.7 - 1.9	20-78	0.7 - 2.5	na	na	123-448	3.9 - 14.1	525-2160
St. Louis, MO	μ	115	3.3	133	3.8	53	1.5	63	1.8	6	0.3	373	10.7	1840
	Ч	29 - 150	0.8 - 4.3	52-207	1.5 - 6.0	24-65	0.7 - 1.9	30 - 103	0.9 - 3.0	na	na	144-534	4.1 - 11.5	650-2560
Tampa, FL	H	156	5.8	123	4.5	29	1.1	64	2.4	13	0.5	385	14.2	1920
	Ч	34 - 184	1.3 - 6.8	48-192	1.8 - 7.1	11 - 35	0.4 - 1.3	23 - 94	0.9 - 3.5	na	na	129–518	4.8 - 19.2	569-2510
Tucson, AZ	H	252	4.6	213	3.8	70	1.3	15	0.3	21	0.4	572	10.3	3190
	Ч	75–398	1.3 - 7.2	83–333	1.5 - 6.0	37–95	0.7 - 1.7	8^{-30}	0.1 - 0.5	na	na	224-877	4.0 - 15.8	1160 - 4900
Tulsa, OK	H	266	5.3	182	3.6	46	0.9	58	1.2	18	0.4	569	11.4	3040
	Ч	63-322	1.3 - 6.5	71-284	1.4 - 5.7	21 - 56	0.4 - 1.1	25-83	0.5 - 1.7	na	na	196 - 762	3.9 - 15.3	940-4000
Virginia Beach-Norfolk, VA	H	1660	5.8	794	2.8	428	1.5	330	1.2	71	0.2	3280	11.5	18300
	Ч	420 - 2030	1.5 - 7.1	310 - 1240	1.1 - 4.3	184 - 516	0.6 - 1.8	152 - 503	0.5 - 1.8	na	na	1140 - 4360	4.0 - 15.3	5790-23700
Washington, DC	μ	192	3.9	161	3.3	100	2.0	81	1.6	24	0.5	558	11.3	2850
	Ч	46-251	0.9 - 5.1	63-252	1.3-5.1	42–119	0.9 - 2.4	37-125	0.7 - 2.5	na	na	212-771	4.3–15.6	963–3860
K 46-251 0.9-5.1 0.9-5.1 1.3-5.1 42-119 0.9-24 37-125 0.7-2.5 na na 212-771 4.3-15.6 963-3860	¥	40-21	0.9–0.1	03-252	1.3-5.1	42-119	0.9-2.4	37-125	0.7–7.5	na	na	1//-717	4.3-15.6	

5, ^aNO₂ data from Buffalo, NY ^bSO₂ data from El Paso, TX ^cNO₂ data from Cincinnati, OH ^dWeather data from Newark, NJ ^cNO₂ data from Newark, NJ ^cNO₂ data from Sans City, MO ^fWeather data from San Francisco, CA ^sNO₂ data from Portland, OR

City	%tree cover	% air quality ir	nprovement			
		СО	NO ₂	O ₃	PM ₁₀	SO ₂
Atlanta, GA	32.9	0.002	0.5	0.7	0.7	0.7
		(0.001 - 0.009)	(0.1 - 2.5)	(0.1 - 4.4)	(0.3 - 2.8)	(0.1 - 4.3)
Boston, MA	21.2	0.002	0.4	0.6	0.6	0.5
		(0.000 - 0.006)	(0.0 - 1.8)	(0.1 - 3.4)	(0.1 - 1.8)	(0.1 - 3.4)
Dallas, TX	28.0	0.002	0.4	0.6	0.6	0.6
		(0.001 - 0.008)	(0.1 - 2.2)	(0.1 - 3.9)	(0.2 - 2.4)	(0.1 - 3.8)
Denver, CO	26.0	0.001	0.2	0.3	0.4	0.3
		(0.000 - 0.007)	(0.0 - 1.5)	(0.0 - 2.1)	(0.1 - 2.2)	(0.0 - 2.0)
Milwaukee, WI	19.1	0.001	0.3	0.4	0.4	0.4
		(0.000 - 0.005)	(0.0 - 1.5)	(0.1 - 2.7)	(0.1 - 1.6)	(0.0 - 2.7)
New York, NY	16.6	0.001	0.3	0.4	0.5	0.4
		(0.000 - 0.005)	(0.0 - 1.4)	(0.1 - 2.6)	(0.1 - 1.4)	(0.1 - 2.6)
Portland, OR	42.0	0.003	0.6	0.8	1.0	0.7
		(0.001 - 0.012)	(0.1 - 2.7)	(0.1 - 3.7)	(0.3 - 3.5)	(0.1 - 4.0)
San Diego, CA	8.6	0.001	0.2	0.3	0.3	0.3
		(0.000 - 0.002)	(0.0 - 0.7)	(0.0 - 1.4)	(0.1 - 0.7)	(0.0 - 1.4)
Tampa, FL	9.6	0.001	0.2	0.2	0.2	0.2
• ·		(0.000 - 0.003)	(0.0 - 0.8)	(0.0 - 1.4)	(0.1 - 0.8)	(0.0 - 1.4)
Tucson, AZ	13.7	0.001	0.1	0.1	0.2	0.1
		(0.000 - 0.004)	(0.0 - 1.0)	(0.0 - 1.7)	(0.1 - 1.2)	(0.0 - 1.7)
Washington, DC	31.1	0.002	0.4	0.6	0.7	0.6
6		(0.001 - 0.009)	(0.2 - 2.3)	(0.1–3.9)	(0.2 - 2.6)	(0.1–3.9)

Table 2. Estimated percent air quality improvement in selected US cities due to air pollution removal by urban trees

Estimates are given for actual tree cover conditions in city for ozone (O₃), particulate matter less than $10 \,\mu m$ (PM₁₀), nitrogen dioxide (NO₂), sulfur dioxide (SO₂), and carbon monoxide (CO) based on local boundary layer height and pollution removal estimates. Bounds of total tree removal of O₃, NO₂, SO₂, and PM₁₀ were estimated using the typical range of published in-leaf dry deposition velocities (Lovett, 1994)

 O_3 concentration between above- and below-forest canopies in California's San Bernardino Mountains have exceeded 50 ppb (40-percent improvement) (Bytnerowicz et al., 1999). Under normal daytime conditions, atmospheric turbulence mixes the atmosphere such that pollutant concentrations are relatively consistent with height (Colbeck and Harrison, 1985). Forest canopies can limit the mixing of upper air with groundlevel air, leading to significant below-canopy air quality improvements. However, where there are numerous pollutant sources below the canopy (e.g., automobiles), the forest canopy could have the inverse effect by minimizing the dispersion of the pollutants away at ground level.

The greatest effect of urban trees on ozone, sulfur dioxide, and nitrogen dioxide is during the daytime of the in-leaf season when trees are transpiring water. Particulate matter removal occurs both day and night and throughout the year as particles are intercepted by leaf and bark surfaces. Carbon monoxide removal also occurs both day and night of the in-leaf season, but at much lower rates than for the other pollutants.

Urban areas are estimated to occupy 3.5% of lower 48 states with an average canopy cover of 27%. Urban tree cover varies by region within the United States with

cities developed in forest areas averaging 34.4% tree cover, cities in grassland areas: 17.8%, and cities in deserts: 9.3% (Dwyer et al., 2000; Nowak et al., 2001). Total pollution air removal (5 pollutants) by urban trees in coterminous United States is estimated at 711,000 t, with an annual value of \$3.8 billion (Table 3).

Though the estimates given in this paper are only for a 1-year period (1994), analysis of changes in meteorology and pollution concentration on pollution removal by urban trees over a 5-year period in Chicago (1991–1995) reveals that annual removal estimates were within 10% of the 5-year average removal rate. Estimates of pollution removal may be conservative as some of the deposition-modeling algorithms are based on homogenous canopies. As part of the urban tree canopy is heterogeneous with small patches or individual trees, this mixed canopy effect would tend to increase pollutant deposition. Also, aerodynamic resistance estimates may be conservative and lead to a slight underestimate of pollution deposition.

Though the average percent air quality improvement due to trees is relatively low (<1%), the improvement is for multiple pollutants and the actual magnitude of pollution removal can be significant (typically hundreds to thousands of metric tons of pollutants per city per

Pollutant	Removal (t)	Value ($\$ \times 10^6$)
03	305,100	2,060
	(75,000-390,200)	(506-2635)
PM_{10}	214,900	969
	(84,000-335,800)	(378–1514)
NO_2	97,800	660
-	(42,800-119,100)	(289-804)
SO_2	70,900	117
-	(32,200-111,100)	(53–184)
CO	22,600	22
	na	Na
Total	711,300	3828
	(256,600-978,800)	(1,249–5158)

Table 3. Air pollution removal and value for all urban treesin the coterminous United States

Estimates are given for ozone (O₃), particulate matter less than $10 \,\mu\text{m}$ (PM₁₀), nitrogen dioxide (NO₂), sulfur dioxide (SO₂), and carbon monoxide (CO). The monetary value of pollution removal by trees is estimated using the median externality values for the United States for each pollutant (Murray et al., 1994). Externality values for O₃ were set to equal the value for NO₂. Bounds of total tree removal of O₃, NO₂, SO₂, and PM₁₀ were estimated using the typical range of published inleaf dry deposition velocities (Lovett, 1994).

year). Percent air quality improvement estimates are likely conservative and can be increased through programs to increase canopy cover within cities. Air pollution removal is also only one aspect of how urban trees affect air quality. Ozone studies that integrate temperature, deposition and emission effects of trees are revealing that urban trees can have significant effects on reducing ozone concentrations (Cardelino and Chameides, 1990; Taha, 1996; Nowak et al., 2000). Based in part on these findings, the US Environmental Protection Agency has introduced urban tree cover as a potential emerging measure to help meet air quality standards (US EPA, 2004). So even though the percent air quality improvement from pollution removal by trees may be relatively small, the total effect of trees on air pollution can produce impacts that are significant enough to warrant consideration of tree cover management as a means to improve air quality.

Conclusion

Through pollution removal and other tree functions (e.g., air temperature reductions), urban trees can help improve air quality for many different air pollutants in cities, and consequently can help improve human health. While the existing percent air quality improvements due to pollution removal by urban trees are modest, they can be improved by increasing urban tree canopy cover. The combined total effects of trees on air pollutants are significant enough that urban tree management could provide a viable means to improve air quality and help meet clean air standards in the United States.

Acknowledgments

This work was supported by funds through the USDA Forest Service's RPA Assessment Staff, and State and Private Forestry's, Urban and Community Forestry Program. We thank D. Baldocchi, M. Ibarra, E.L. Maxwell, and M.H. Noble for assistance with model development and data processing.

References

- Baldocchi, D., 1988. A multi-layer model for estimating sulfur dioxide deposition to a deciduous oak forest canopy. Atmospheric Environment 22, 869–884.
- Baldocchi, D.D., Hicks, B.B., Camara, P., 1987. A canopy stomatal resistance model for gaseous deposition to vegetated surfaces. Atmospheric Environment 21, 91–101.
- Bidwell, R.G.S., Fraser, D.E., 1972. Carbon monoxide uptake and metabolism by leaves. Canadian Journal of Botany 50, 1435–1439.
- Bytnerowicz, A., Fenn, M.E., Miller, P.R., Arbaugh, M.J., 1999. Wet and dry pollutant deposition to the mixed conifer forest. In: Miller, P.R., McBride, J.R. (Eds.), Oxidant Air Pollution Impacts in the Montane Forests of Southern California: A Case Study of the San Bernardino Mountains. Springer, New York, pp. 235–269.
- Cardelino, C.A., Chameides, W.L., 1990. Natural hydrocarbons, urbanization, and urban ozone. Journal of Geophysical Research 95 (D9), 13,971–13,979.
- Chameides, W.L., Lindsay, R.W., Richardson, J., Kiang, C.S., 1988. The role of biogenic hydrocarbons in urban photochemical smog: Atlanta as a case study. Science 241, 1473.
- Colbeck, I., Harrison, R.M., 1985. Dry deposition of ozone: some measurements of deposition velocity and of vertical profiles to 100 metres. Atmospheric Environment 19 (11), 1807–1818.
- Dwyer, J.F., Nowak, D.J., Noble, M.H., Sisinni, S.M., 2000. Assessing our nation's urban forests: connecting people with ecosystems in the 21st century. USDA Forest Service General Technical Report PNW-460, Portland, OR.
- Hosker, R.P., Lindberg, S.E., 1982. Review: atmospheric deposition and plant assimilation of gases and particles. Atmospheric Environment 16 (5), 889–910.
- Killus, J.P., Meyer, J.P., Durran, D.R., Anderson, G.E., Jerskey, T.N., Reynolds, S.D., Ames, J., 1984. Continued Research in Mesoscale Air Pollution Simulation Modeling, Vol. V: refinements in numerical analysis, transport, chemistry, and pollutant removal, EPA/600/3-84/095a. US EPA, Research Triangle Park, NC.
- Lovett, G.M., 1994. Atmospheric deposition of nutrients and pollutants in North America: an ecological perspective. Ecological Applications 4, 629–650.
- Murray, F.J., Marsh, L., Bradford, P.A., 1994. New York State Energy Plan, vol. II: Issue Reports. New York State Energy Office, Albany, NY.

- Nowak, D.J., 1994. Air pollution removal by Chicago's urban forest. In: McPherson, E.G., Nowak, D.J., Rowntree, R.A. (Eds.), Chicago's Urban Forest Ecosystem: Results of the Chicago Urban Forest Climate Project. USDA Forest Service General Technical Report NE-186, Radnor, PA, pp. 63–81.
- Nowak, D.J., Rowntree, R.A., McPherson, E.G., Sisinni, S.M., Kerkmann, E., Stevens, J.C., 1996. Measuring and analyzing urban tree cover. Landscape and Urban Planning 36, 49–57.
- Nowak, D.J., McHale, P.J., Ibarra, M., Crane, D., Stevens, J., Luley, C., 1998. Modeling the effects of urban vegetation on air pollution. In: Gryning, S.E., Chaumerliac, N. (Eds.), Air Pollution Modeling and its Application XII. Plenum Press, New York, pp. 399–407.
- Nowak, D.J., Civerolo, K.L., Rao, S.T., Sistla, G., Luley, C.J., Crane, D.E., 2000. A modeling study of the impact of urban trees on ozone. Atmospheric Environment 34, 1610–1613.
- Nowak, D.J., Noble, M.H., Sisinni, S.M., Dwyer, J.F., 2001. Assessing the US urban forest resource. Journal of Forestry 99 (3), 37–42.
- Pederson, J.R., Massman, W.J., Mahrt, L., Delany, A., Oncley, S., den Hartog, G., Neumann, H.H., Mickle, R.E., Shaw, R.H., Paw, UKT., Grantz, D.A., MacPherson,

J.I., Desjardins, R., Schuepp, P.H., Pearson, R., Arcado, T.E., 1995. California ozone deposition experiment: methods, results, and opportunities. Atmospheric Environment 29 (21), 3115–3132.

- Smith, W.H., 1990. Air Pollution and Forests. Springer, New York.
- Taha, H., 1996. Modeling impacts of increased urban vegetation on ozone air quality in the South Coast Air Basin. Atmospheric Environment 30 (20), 3423–3430.
- US Environmental Protection Agency (US EPA), 2004. Incorporating Emerging and Measures in a State Implementation Plan (SIP). US Environmental Protection Agency, Research Triangle Park, NC. http://www.epa. gov/ttn/oarpg/t1/memoranda/evm_ievm_g.pdf (last accessed: October 2005).
- US Environmental Protection Agency (US EPA), 1995. PCRAMMIT User's Guide. US Environmental Protection Agency, Research Triangle Park, NC.
- Wesely, M.L., 1989. Parameterization for surface resistance to gaseous dry deposition in regional-scale numerical models. Atmospheric Environment 23, 1293–1304.
- Zinke, P.J., 1967. Forest interception studies in the United States. In: Sopper, W.E., Lull, H.W. (Eds.), Forest Hydrology. Pergamon Press, Oxford, pp. 137–161.