1 Inter-provincial Reliance for Improving Air Quality in China: A

2	Case Study on Black Carbon Aerosol
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14 ABSTRACT

15 Black carbon (BC) is of global concern because of its adverse effects on climate and human health. It can travel long distances via atmospheric movement, and can be geographically relocated 16 17 through trade. Here, we explored the integrated patterns of BC transport within 30 provinces in 18 China from the perspective of meteorology and inter-provincial trade using the Weather Research 19 and Forecasting with Chemistry (WRF/Chem) model and multi-region input-output analysis. In 20 general, cross-border BC transport, which accounts for more than 30% of the surface concentration, occurs mainly between neighboring provinces. Specifically, Hebei contributes 1.2 µg·m⁻³ BC 21 22 concentration in Tianjin. By contrast, trade typically drives virtual BC flows from developed 23 provinces to heavily industrial provinces, with the largest net flow from Beijing to Hebei (4.2 Gg). 24 Shanghai is most vulnerable to domestic consumption with an average inter-provincial consumption 25 influence efficiency of 1.5×10⁻⁴ (µg·m⁻³)/(billion Yuan·yr⁻¹). High efficiencies (~8×10⁻⁵ (µg·m⁻ ³)/(billion Yuan·yr⁻¹)) are also found from regions including Beijing, Jiangsu and Shanghai to 26 27 regions including Hebei, Shandong and Henan. The above source-receptor relationship indicates 28 two control zones—Huabei and Huadong control zones. Both mitigating end-of-pipe emissions and 29 rationalizing the demand for pollution-intense products are important within the two control zones 30 to reduce BC and other pollutants.

Keywords: black carbon, inter-provincial transport, source-receptor relationship, input-output
 analysis



39 INTRODUCTION

Black carbon (BC), which is generated by the incomplete combustion of 40 carbonaceous fuels,^{1,2} is an important combustion component of fine particulate matter 41 (PM_{2.5}).^{3, 4} Moreover, the scientific community has been increasingly concerned about 42 its adverse impact on climate change, air quality and human health.⁵⁻⁷ BC aerosols 43 influence climate both regionally and globally by absorbing solar radiation, which 44 reduces the atmospheric lapse rate and burns off cloud droplets.^{3, 7} BC level varies 45 consistently with carbon monoxide (CO), nitric oxide (NO) and other traffic-related 46 gaseous pollutants and occupies roughly a fixed proportion of particulate matter (PM) 47 concentration in summer and autumn.^{8,9} Additionally, pollution containing BC has been 48 proven to have a robust epidemiological association with many types of mortality, 49 particularly cardiovascular.^{10, 11} Thus, it is acknowledged that BC may serve as an 50 effective indicator of air quality and its health effects in helping to mitigate air pollution 51 including PM, CO and NO.^{4, 11, 12} Once emitted into the atmosphere, BC has a lifetime 52 of 2-10 days and can be transported long distances by atmospheric movement,¹³⁻¹⁶ 53 indicating its well-mixed condition in lower troposphere and regional, rather than local, 54 character.17 55

China has been the world's largest emitter of anthropogenic BC, organic matter 56 (OM) and other PM_{2.5} precursors.¹⁸⁻²⁰ In 2014, approximately 90% of the major cities 57 in China failed to meet the national air quality standard for PM_{2.5}.²¹ Emissions from the 58 industrial and transport sectors have been identified as the major sources of BC and 59 other combustion $PM_{2.5}$,²² which has led to the need for serious emissions control in 60 China. Recently, the "Law of the People's Republic of China on the Prevention and 61 Control of Atmospheric Pollution" has been revised to emphasize the national target of 62 63 air quality improvement from a concentration-based perspective and the supervision of pollution sources using emissions-based strategies.²³ This law also calls for 64 collaborative efforts across administrative boundaries for emissions control and air 65 quality improvement. Consequently, a quantitative understanding of the inter-66 provincial source-receptor relationship of air pollution transport and the underlying 67

68 economic drivers is of great importance.^{24, 25}

Previous studies have evaluated the possible sources of air pollution for a target 69 region.^{13, 26} For example, Guo et al. analyzed the observational data collected at 70 Changdao Island north of Shanghai and found that a significant amount of BC is 71 transported from Shandong and Jiangsu provinces.²⁷ Xue et al. utilized particulate 72 source apportionment technology (PSAT) in the CAMx model and found that 73 approximately 40% of ambient PM_{2.5} concentrations in Beijing, Shanghai and Jiangsu 74 are contributed by cross-boundary transport.²⁸ If the source-receptor relationship were 75 accepted by the relevant provincial governments, it might be the basis for promoting 76 multi-province cooperation on emissions control.²⁹ 77

Apart from observed atmospheric transport, domestic trade also affects the 78 distribution of emissions to a large extent and thus changes air pollution levels 79 geographically.^{30, 31} The production of traded products increases local emissions while 80 reducing the emissions in consuming regions. This trade-induced virtual emissions 81 transfer among provinces in China has been well documented for carbon dioxide (CO_2) , 82 83 sulfur dioxide (SO₂), primary PM_{2.5} and other atmospheric pollutants, which demonstrates that some developed provinces shift emissions to less-developed 84 provinces by importing products.^{25, 30} As with atmospheric transport, this virtual 85 transfer of emissions and the degradation of air quality via trade also lead to a source-86 receptor relationship. When considering the location disparity between consumers and 87 producers, the emissions generated in one province might be significantly driven by 88 final consumption in a different province. Thus, for purposes of cross-provincial action 89 on air pollution control, it is equally important to identify both the sources and the 90 91 drivers of air pollution.

In this study, we used BC as a proxy to establish the source-receptor relationship involving the atmospheric transport and trade-induced virtual transfer of primary pollution among 30 provinces in mainland China in 2007 (excluding only Tibet, where reliable data are unavailable). BC was chosen as a representative for cross-regional air pollutant mitigation because of its impact on environment, regional character and representative for other fine aerosol species. The model simulation of BC transport was

undertaken using the Weather Research and Forecasting with Chemistry (WRF/Chem) 98 model. An explicit tagging method was implemented in the WRF/Chem model to 99 efficiently track the pathways of BC transport.³² We also used multi-region input-output 100 (MRIO) analysis to examine the virtual transfer of BC emissions resulting from trading 101 goods and services.³³ By combining both physical and virtual transfers of BC emissions, 102 we quantified the direct and indirect inter-provincial linkages in terms of pollution 103 transport. This quantification leads to feasible suggestions on the priority of BC 104 105 reduction and the possibility of cooperative responsibility for pollution mitigation in China. 106

107 METHODOLOGY AND DATA

108 Emission inventory and data sources.

A production-based emissions inventory was developed by multiplying the energy 109 consumption data and BC emission factors.^{34, 35} Energy consumption data for 30 110 provinces in China (as listed in Table S1 in the supporting information) were derived 111 from provincial statistical yearbooks and energy balance tables from the 2008 Chinese 112 Statistical Yearbooks (Data were based on the investigation of year 2007) for each 113 province.³⁶ We aggregated the provincial BC emissions into 17 sectors (listed in Table 114 S2 in the supporting information) to conform with the Chinese MRIO Table.³⁷ Emission 115 factors for 8 types of energy (i.e., coal, coke, gasoline, kerosene, diesel, fuel oil, 116 liquefied petroleum gas and natural gas) were obtained from previous studies (listed in 117 Table S3 in the supporting information).^{34, 35} The derivation of production-based BC 118 emissions attributable to energy consumption for province *f* is expressed as: 119

$$C_{p}^{f} = \sum_{i=1}^{17} \sum_{m=1}^{8} E_{i,m}^{f} \times EF_{i,m}$$
(1)

where $E_{i,m}^{f}$ is the energy consumption of fuel *m* in sector *i*, province *f*; $EF_{i,m}$ is the emission factor of fuel *m* in sector *i*.

Here, we considered only industrial BC emissions from all 17 aggregated economic sectors, because industrial emissions can "flow" in inter-provincial trade. According to the Chinese MRIO Table, every particular sector has a more or less monetary output to support non-local industry. By contrast, residential energy consumption cannot "flow" in trade, and was thus excluded in our analysis. We used the industrial BC emissions in WRF/Chem modeling for consistency with the MRIO analysis by mapping the emissions with high spatial resolution.³⁴ We also conducted additional simulations using revised all-source anthropogenic BC emission inventory from Wang et al. for model evaluation.³⁴

132 Model description and configuration.

WRF/Chem is a meteorological model that enables the simulation of atmospheric 133 phenomena across scales ranging from meters to thousands of kilometers.³⁸ 134 WRF/Chem includes chemical processes such as emissions, gas/aqueous phase 135 chemistry and dry/wet deposition.³⁹ WRF/Chem has been widely applied to simulate 136 the transport of BC and its radiative impact.^{38, 40, 41} Real meteorological data are used 137 as the initial and lateral boundary condition input for the WRF/Chem model to simulate 138 the physical transport of BC aerosols. Here we applied a data set from the NCEP FNL 139 Operational Model Global Tropospheric Analyses, which provide data every six hours 140 for the period from December 16st, 2006, to December 31st, 2007 for model simulation. 141 The first two weeks of the simulation were used for model spin-up. 142

143 To quantify source-receptor relationships among the 30 provinces, an explicit tagging technique was used in WRF/Chem to avoid modifying BC emissions.⁴² This 144 method differs from the traditional sensitivity approach to avoid reducing BC emissions 145 that may strongly disturb the local climate. Similar approaches have been previously 146 applied in global models to estimate the long-range transport of BC, OC and PM_{2.5} 147 between continental regions.^{13, 32, 43} In this tagging approach, two classes of BC tracers 148 are used for each "tagged" region. One is for hydrophobic BC, which represents freshly 149 emitted BC species, and the other is for hydrophilic BC, which represents aged BC and 150 151 has sufficient soluble coating to behave as cloud condensation nuclei (CCN). Therefore, 152 30 non-overlapping geographical regions were tagged individually with additional variables to track their transportation and transformation until deposition. Tagged BC 153

has the same physical and chemical properties as untagged BC, and the model thus accurately predicts the pathways of BC dispersion and its influence on surface concentration.

We use the WRF/Chem model to track the inter-provincial source-receptor relationships for BC in 2007 with a $0.2^{\circ} \times 0.2^{\circ}$ horizontal resolution. In general, the model agreed within a factor of 2 with the observations (As shown by Figure S1 in the supporting information - observational data were collected from published literatures.⁴⁴⁻ ⁵⁷ The spatial distribution of the data is shown in Figure S2). The output results were archived hourly and used to calculate the average surface concentrations for a province over a given period of time for analysis.

164 Multi-region Input-output (MRIO) Analysis.

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Originating from Leontief,⁵⁸ input-output analysis has been widely used to link 165 global and regional environmental issues with final consumption.^{31, 33} In the past decade, 166 environmental MRIO analysis has been developed to quantify emissions transfer via 167 inter-regional trade.^{33, 59, 60} Here, we used the Chinese MRIO Table from 2007 that was 168 developed by Liu et al. to quantify BC emissions embodied in traded products.³⁷ The 169 MRIO table consists of three parts. Part One is the intermediate input/output for 17 170 sectors in 30 provinces. Part Two consists of provincial final consumption (i.e., urban 171 172 household consumption, rural household consumption, government consumption and investment) and international export. Part Three consists of production-based BC 173 emissions for 17 sectors in 30 provinces. 174

For the entire system covering all provincial economies, we have the followingbalance of monetary flows:

177
$$X = AX$$

$$X = AX + Y \tag{2}$$

where X is a vector representing total monetary output for every province, A is a matrix with its elements defined as intermediate input to produce a unit output, and Y is a vector representing the total output of final consumption and international export in each province.



Consumption-based BC emissions can be obtained by introducing emission

183 intensity *EI*:

184
$$C_c = EI(I - A)^{-1}Y_c$$
 (3)

where *EI* is a vector with its elements defined as the direct BC emissions per unit of economic output, $(I - A)^{-1}$ is the Leontief inverse matrix and Y_c is the final consumption.

This basic formula can be further used to quantify emissions from the production of traded products. For instance, BC emissions embodied in the products exported from province f to province s can be calculated as

191
$$C_c^{fs} = EI^f (I - A)^{-1} Y_c^s$$
 (4)

where EI^f is a vector of BC emission intensity for province *f* but zero for all others and Y_c^s is the final consumption of province *s*.

194 **RESULTS**

195 Physical Transport of BC via Atmospheric Movement.

Figure 1 shows the major cross-boundary influence pattern of the area-weighted 196 annual mean surface BC concentration caused by industrial emissions. The annual 197 mean surface BC concentrations range from 0.025 µg·m⁻³ (Qinghai) to 5.7 µg·m⁻³ 198 (Shanghai). Shanghai and Tianjin (4.2 $\mu g \cdot m^{-3}$) have the highest BC concentration. 199 200 Major local sources of pollution for these two coastal megalopolises are traffic and transport sectors (as suggested by Figure S3), while emissions in their contiguous 201 provinces also exert considerable influence. Industry-dominant provinces including 202 Shandong (2.8 μ g·m⁻³), Henan (2.9 μ g·m⁻³) and Liaoning (2.0 μ g·m⁻³) also have heavy 203 BC concentrations. Moreover, provinces with heavier BC pollution are likely to be 204 205 located along or near the coastline.

Provincial BC concentrations are profoundly influenced by trans-boundary transport. The reciprocal effect between two contiguous provinces whose emissions share resemblances is generally comparable. It is particularly noticeable between Hebei and Shandong, Shandong and Henan, Jiangsu and Anhui, and Jiangsu and Shanghai, where approximately 10% of the BC concentrations in these provinces are contributed by one another. The northern provinces tend to be net contributors to the pollution load of the more southerly provinces in eastern China. Remarkably, Hebei is responsible for $0.59 \ \mu g \cdot m^{-3} \ (24\%)$ and $1.2 \ \mu g \cdot m^{-3} \ (28\%)$ surface BC concentrations in Beijing and Tianjin, respectively. It is also responsible for $0.13 \ \mu g \cdot m^{-3} \ (7\%)$ and $0.17 \ \mu g \cdot m^{-3} \ (6\%)$ of the BC concentration in Shanxi and Henan, respectively. Whereas Shanxi contributes $0.20 \ \mu g \cdot m^{-3} \ (22\%) BC$ in Shaaxi, and Henan contributes $0.23 \ \mu g \cdot m^{-3} \ (19\%)$ of the BC concentration in Hubei and $0.17 \ \mu g \cdot m^{-3} \ (8\%)$ in Anhui.

218 Virtual Transfer of BC via Inter-provincial Trade.

Figure 2(a) shows the comparison of total production-based and consumption-219 based BC emissions in 2007 for 30 Chinese provinces. Total industrial BC emissions 220 amount to 894 Gg in China in 2007, which is consistent with previous studies.^{34, 61, 62} 221 From the production perspective, Shandong ranks first with emissions of 79.7 Gg, 222 followed by Henan (73.5 Gg), Shanxi (61.1 Gg) and Hebei (60.3 Gg). Provincial 223 consumption-based BC emissions present a different distribution pattern, with 741 Gg 224 (83%) emissions induced by domestic demand. This percentage is comparable to 225 previous results on primary PM_{2.5} and gaseous pollutants including SO₂ and NO_x.^{25, 63} 226 Except for Shandong (contributing 64.7 Gg emission), the southern provinces, 227 including Zhejiang (57.2 Gg), Jiangsu (55.0 Gg) and Guangdong (51.2 Gg), hold the 228 229 top positions. Remarkably, the Yangtze River Delta contributes 20% of the total embodied BC emissions, although its domain area is less than 2% of the total area. 230 Consumption-based emissions consist of emissions from four types of final 231 consumption. Investment is the dominant motor driving industrial BC emissions for 29 232 provinces (the exception is Xinjiang), contributing approximately 40%-70% of the 233 total consumption-based BC emissions. Urban household consumption is the second 234 largest driver of BC emissions, ranging from 15% in Shanxi to 43% in Tianjin. 235 Government consumption and rural household consumption account for the remaining 236 237 15%.

The difference between production-based and consumption-based BC emissions indicates that emissions are transferred via trade. Figure 2(b) illustrates net emissions

transfer through trade (only the largest fluxes between provinces are shown). 13 of 30 240 provinces are net emissions importers, and the other 17 provinces are net exporters. Net 241 importers are mainly industry-dominant provinces such as Hebei (23.2 Gg), Shanxi 242 (18.3 Gg), Liaoning (17.3 Gg) and Henan (15.7 Gg). Their industrial activities and 243 associated emissions enhanced by trade support consumption across the country, 244 particularly for a few developed provinces. Conversely, Zhejiang (24.1 Gg), Shanghai 245 (16.4 Gg), Beijing (14.5 Gg) and Guangdong (13.8 Gg) are major BC exporters. They 246 247 behave as exporters in trade with almost all other provinces, whereas the larger flows more often end up in Hebei and Henan. 248

The pattern of major flows is from southeastern China to the North China Plain 249 (NCP) geographically and from developed to less-developed provinces economically. 250 The demand-driven flows can be categorized into three types based on the economic 251 strength of emission exporter and importer, with the dominant pattern being from a 252 province with abundant capital to a province owning heavy industry (Figure 2b). First, 253 the largest BC emissions transfer occurs from Beijing to Hebei (with 4.2 Gg BC 254 255 emissions being relocated), followed by a flow of 4.1 Gg emissions from Zhejiang to Hebei. Second, shifts in emissions between contiguous provinces of comparable 256 economic strength are also noticeable. They occur noticeably within the Yangtze River 257 Delta and northeastern provinces, including Heilongjiang, Jilin and Liaoning, which 258 indicate intimate economic relationships between contiguous provinces. The typical 259 emission flows in this category are from Shanghai to Jiangsu (1.2 Gg) and from Jilin to 260 Liaoning (2.7 Gg). The third type of flow is from industrial provinces to resource-rich 261 but less-developed provinces, suggesting the need for inputting fundamental raw 262 materials for industrial activities. For example, 1.4 Gg of emissions are transported 263 from Hebei to Shanxi. In general, trade-induced emission flows across China occur 264 from south to north and coastal to inland, exhibiting a reversed source-receptor pattern 265 to BC dispersion via atmospheric transport. 266

Apart from revealing emission flows from a regional aspect, MRIO can also explore the sectors that undertake the transfer of BC emissions in trade.³¹ Figure 3 shows the sector-specific BC emission transfer embodied in inter-provincial trade. At the national level, nearly half of these trade-relevant emissions are caused by the production of intermediate products. This ratio is even higher in Shanxi, Gansu, Qinghai, Xinjiang and Yunnan, where more than 80% of the emissions are caused by the massive quantities of low-value-added raw materials and energy that are produced for export. By contrast, Zhejiang and Jiangsu have the highest proportion of intermediate goods from the import aspect because these provinces lack natural resources but are advanced in processing capabilities.

277 Moreover, final use accounts for the remaining 15%-55% of trade-embodied emissions, in which heavy industry (including the petro-chemical, nonmetallic mineral 278 products, and metallic mineral products) plays a dominant role in most provinces due 279 to its intensive energy consumption. In addition to heavy industry, exporting 280 agricultural products induces salient BC emissions in Hubei and Sichuan provinces; 281 mining and washing are responsible for most trade-relevant BC emissions in Shanxi 282 and Liaoning provinces. However, for Jiangsu and Zhejiang, light industry such as 283 textile and timber processing is key to generating BC emissions. With regard to 284 285 imported products, less-developed provinces tend to have a higher proportion of emission output in high-value products of light industry, whereas the provinces scarce 286 in energy and raw materials are likely to depend on products of mining and washing in 287 other provinces. 288

289 Surface BC Concentrations from a Production and Consumption Perspective.

By combining atmospheric BC transport and emission flow in trade, the source of 290 surface BC concentrations can be classified according to their on-site emission region 291 292 and the final consumer of the relevant products. With regard to whether BC is emitted locally or from other provinces, BC concentration can be classified as either local or 293 domestic production. Meanwhile, from a consumption-based perspective, BC 294 concentration can be referred to as local, domestic or foreign consumption. We specify 295 296 "domestic consumption" as the consumption from the other 29 provinces, and "foreign 297 consumption" as the internationally-exported products from the province where 298 production-based BC is generated. From an on-site emission perspective, 9 provinces

(i.e., Hubei, Anhui, Jiangxi, Hainan, Guangxi, Chongqing, Guizhou, Shaanxi and 299 Qinghai) have a share of domestic-production concentration (i.e., originated from the 300 other 29 provinces, namely the green, yellow plus light grey portions in Figure 4) over 301 50%. Others are typically within 30%-50%, with the lowest (3%) being Xinjiang. This 302 indicates the importance of local emissions, but trans-boundary transport is also a 303 noteworthy contributor. With regard to the key provinces for air pollution mitigation, 304 Beijing has a proportion of domestic-production BC surface concentration equal to 42%, 305 306 Tianjin 45%, Shanghai 22%, Jiangsu 39%, Zhejiang 41% and Guangdong 40%. BC concentration originating from other provinces but induced by local consumption (i.e., 307 the green portion in Figure 4) is very small (<3%) and mainly driven by neighboring 308 309 provinces.

From the consumption perspective, final demands within mainland China on 310 311 average induce 82% of provincial surface BC concentration across China. Domesticconsumption-induced concentration (i.e., the red plus yellow portions in Figure 4) is 312 greater than local-consumption-induced concentration (i.e., the blue plus green portions 313 in Figure 4) for all provinces, which indicates the profound influence of inter-regional 314 trade and trans-boundary transport on concentration. The proportion of BC 315 concentration generated locally but induced by domestic consumption ranges from 4% 316 in Hainan to 50% in Xinjiang and reflects the comparative industrial scales for 317 supporting local living standards or for securing economic growth via exports. This 318 319 proportion is above 30% for typical emission importers. Particularly, 43% and 41% of the concentration in Hebei and Liaoning, respectively, is contributed by the local 320 emissions induced by domestic consumption. Meanwhile, for emissions-exporting 321 provinces, such as Beijing, Zhejiang and Guangdong, this proportion falls to near 20%. 322 However, Shanghai has a higher percentage (40%) of local-domestic concentration 323 because of its considerable share of industry to support domestic consumption. 324

Source-Receptor Relationship from Combined Production and Consumption

326 Perspectives.

327 To combine the influence of atmospheric transport and inter-provincial trade on

surface BC concentrations and emissions, which is informative for judging the priority
of cooperative action for pollution mitigation, we introduce three source-receptor
indicators shown in Figure 5.

Figure 5(a) shows the emission-concentration relationship that is defined as the 331 annual averaged surface BC concentration over a receptor resulting from a unit of 332 emission in a source region (in $(\mu g \cdot m^{-3})/(Gg \cdot yr^{-1})$).²⁹ This atmospheric transport 333 efficiency is calculated by dividing the source-produced concentration in a receptor by 334 335 total annual emission in the source region. The colored chart shows a pattern in which a receptor is more sensitive to its own emissions than to domestic emissions and to 336 upwind contiguously located sources than to remote ones. Normalized BC 337 concentrations resulting from local emission range from 0.0047 ($\mu g \cdot m^{-3}$)/(Gg · yr⁻¹) in 338 Xinjiang to 0.20 ($\mu g \cdot m^{-3}$)/(Gg · yr⁻¹) in Shanghai, which is mainly determined by the 339 emission density. Non-local contributions to BC concentration from neighboring 340 provinces is typically 1-2 orders of magnitude smaller than local emissions but more 341 than 1-2 orders of magnitude larger than emissions from remote provinces. Provinces 342 343 within Jing-Jin-Ji and the Yangtze River Delta share a close relationship in BC concentration through atmospheric transport. For instance, 1 Gg annual emission in 344 Beijing and Hebei can increase the surface BC concentration in Tianjin by 0.022 and 345 $0.019 \,\mu \text{g} \cdot \text{m}^{-3}$, respectively. Similarly, the bilateral influence between Jiangsu, Shanghai 346 and Zhejiang ranges from 0.003 to 0.013 ($\mu g \cdot m^{-3}$)/(Gg · yr⁻¹). 347

Figure 5(b) shows the trade-induced consumption-emission relationship that is 348 defined as the production-based BC emissions of a receptor associated with a unit of 349 domestic consumption from a source (in tons of BC per billion Yuan), in which 350 consumption includes the sum of four final consumption.³⁰ This BC intensity is 351 calculated by dividing the source-induced BC emission in a receptor by total annual 352 final consumption in the source region. Unlike Figure 5(a) in which the pattern of 353 atmospheric transport exhibits a diagonal distribution, the consumption-emission graph 354 reflects a column-like distribution, which indicates that massive amounts of BC 355 imported from almost all other provinces via trade into some industry-dominant 356 provinces (e.g., Hebei, Henan, and Liaoning). Normalized production-based BC 357

emissions induced by local consumption range from 3.4 tons per billion Yuan in Tianjin 358 to 53.4 tons per billion Yuan in Shanxi. Generally, this value is higher in provinces with 359 massive energy consumption than in developed provinces with strict environmental 360 laws. However, those developed provinces relocate significant amounts of emissions to 361 other provinces. The proportion of emissions caused in other 29 provinces is 362 comparable to the proportion of local on-site emissions in developed provinces. Every 363 billion Yuan of consumption in Tianjin produces 5.6 tons of BC emissions in Hebei, 1.6 364 365 times its own local emission intensity. Similarly, Hebei and Henan receives 2.4 and 2.1 tons of BC emissions respectively for every billion Yuan of consumption in the Yangtze 366 River Delta. Heilongjiang and Jilin shift 7.2 and 8.4 tons of BC emissions, respectively, 367 to Liaoning for every one billion Yuan of consumption. 368

Figure 5(c) shows the consumption-concentration relationship, i.e., annual 369 averaged surface BC concentration in a receptor resulting from a unit of domestic 370 consumption in a source (in $(\mu g \cdot m^{-3})/(billion Yuan \cdot yr^{-1})$), which considers the joint 371 influence of trans-boundary transport and inter-regional trade on surface BC 372 373 concentration together. This consumption influence efficiency is calculated by concentration in a receptor caused by on-site emissions in 30 provinces that are induced 374 by a unit of annual consumption in a source. The pattern of the graph is a combination 375 of Figure 5(a) and (b), showing both aggregated groups along the diagonal and column-376 like distribution of eminent contributions. The bilateral influence of domestic 377 consumption within Jing-Jin-Ji and the Yangtze River Delta is more than 8×10^{-5} (µg·m⁻ 378 3)/(billion Yuan·yr⁻¹). Northeastern provinces also show intimate internal relationships 379 regarding both atmospheric transport and inter-provincial trade. Surprisingly, Tianjin 380 381 and Shanghai are the two provinces most vulnerable to consumption-based emissions in other provinces despite being net BC exporters in inter-provincial trade. The average 382 surface BC concentration resulting from a unit of consumption in a non-local source 383 region in Shanghai and Tianjin is $1.5 \times 10^{-4} (\mu g \cdot m^{-3})/(billion Yuan \cdot yr^{-1})$ and 1.4×10^{-4} 384 $(\mu g \cdot m^{-3})/(billion Yuan \cdot yr^{-1})$, respectively. This phenomenon can be attributed to the 385 386 considerably high BC concentration resulting from a unit of local-production emission and a prevailing proportion of local-production emission induced by domestic 387

consumption than by local consumption. Although they are densely urbanized 388 metropolises, Tianjin and Shanghai have a considerable scale of secondary industry, 389 which induces a massive emission import (see S3 in the supporting information). In 390 addition, the source-receptor relationship between consumption and concentration is 391 also noticeable within some emission exporters and importers in inter-provincial trade. 392 Per-billion Yuan annual consumption from a source province can lead to an 393 approximately $8 \times 10^{-5} \,\mu\text{g} \cdot \text{m}^{-3}$ increase in BC concentration in a receptor province. In 394 particular, BC concentrations in Liaoning and Hebei are largely affected by domestic-395 consumption emissions particularly from the Jing-Jin-Ji area and Shanxi, whereas the 396 concentrations in Anhui and Jiangsu are affected by emissions from the Yangtze Delta 397 area and Anhui. In Henan and Shandong, BC concentration is sensitive to consumption 398 in all the source provinces mentioned above. These provinces may show a close 399 relationship for air pollution control. 400

401 **DISCUSSION**

Using WRF/Chem modeling and environmental MRIO analysis, we quantified the source-receptor relationship of atmospheric transport and trade-induced geographical relocation of BC emissions. By combining the dual effects, we traced the influence from both producer and consumer perspectives on BC surface concentration for each province in China. The results can provide insights into collaborative efforts on air pollution control for policy makers.

The source-receptor relationship of physical BC transport among provinces is 408 largely determined by both the amount of BC emissions and the direction of prevailing 409 410 winds. Depending on locations, more than 20% of surface BC concentration may originate from a neighboring province, particularly for provinces such as Tianjin and 411 412 Hubei, which are located downwind contiguously of major BC source provinces (such as Hebei and Henan). By contrast, provinces such as Hebei, Shanxi, Henan, Zhejiang 413 414 and Guangdong transport substantial BC pollution to their neighbor provinces. Notably, in inland China, where the north wind is dominant particularly in autumn and winter 415 416 because of the influence of the Siberian High, BC transport typically occurs more

usually from north to south. In southeastern coastal China, however, where airflow is 417 driven by the Hawaiian High, BC transport occurs mainly northwestward during the 418 summer. Combining these two factors, the proportion of surface BC concentration 419 caused by the other provinces varies from 3% to 71%. Provinces such as Hubei, Shaanxi, 420 and Hainan are vulnerable to non-local emissions because of their relatively small scale 421 of industry compared with their neighbors with massive industrial production. Beijing, 422 Shandong, Jiangsu and other provinces with on-site emissions comparable to those of 423 424 their neighbors are also sensitive to domestic-production emissions. Several provinces, such as Xinjiang, Liaoning and Shanghai, have a percentage above 75% of the local-425 production surface BC concentration owing to their location and the total amount of 426 BC emissions, which indicates a local-production dominant situation. 427

Unlike atmospheric transport, which is driven by natural forces, domestic trade 428 relocates BC in a different way. Beijing, Tianjin, Guangdong and the Yangtze River 429 Delta are more likely to outsource BC emissions via inter-provincial trade to industrial 430 provinces including Hebei, Henan, Shanxi and Liaoning. For developed provinces such 431 432 as Beijing and Shanghai, consumption-based BC emissions can be double the production-based BC emissions, whereas in industry-dominant provinces such as Hebei 433 and Shanxi, net BC emissions transferred via trade amount to approximately 30% of 434 their production-based emissions. Surface BC concentration generated by local 435 emission but induced by domestic consumption can account for more than 30% of the 436 total concentration in these provinces. In addition, three northeastern provinces show 437 tight economic connections, with Liaoning playing the major role of emissions importer. 438 This imbalance in inter-provincial trade may be due largely to the enormous disparity 439 in wealth and economic structure among provinces.⁶⁴ Although emission transport via 440 trade is bilateral, developed provinces are more likely to import low value-added 441 commodities in the heavy industry, mining and washing, and agricultural sectors from 442 less-developed provinces according to their mainstay industry, while exporting 443 444 technology-containing commodities from light industry.

445 Combining these two aspects, it is reasonable to say that the patterns of 446 atmospheric transport and inter-provincial trade are the opposite of one another.

Emission flows from Beijing and Tianjin to Hebei are transported in reverse to 447 influence the local BC concentration. Similar patterns can also be observed from 448 Shanghai to Jiangsu and Zhejiang. Moreover, consumption in developed provinces 449 located along the southeastern coastline increases BC concentration and exerts adverse 450 influence on air quality in those emission input industry-dominant provinces, mainly 451 on NCP via trade. These phenomena may serve as a major motivation for the close 452 cooperation between provinces on air pollution control as advocated in the "Law of the 453 454 People's Republic of China on the Prevention and Control of Atmospheric Pollution." By introducing advanced technology from developed provinces to industry-dominant 455 provinces and by taking cross-regional governance into consideration, the supportive 456 provinces benefit in that their surface pollutant concentration caused by domestic-457 production emissions is reduced. Meanwhile, cooperation contributes to reducing 458 surface BC concentration in the emission source, which may compensate for the 459 pollutant transferred via inter-provincial trade. Overall air quality in China can also be 460 enhanced because downwind provinces suffer less from trans-boundary emissions. 461 462 Generally, neighboring provinces have a more intimate relationship concerning both trans-boundary transport and inter-provincial trade, which indicates their optimal 463 prospects for joint efforts on mitigating air pollution. Two partly overlapping control 464 zones for collaborative action on air pollution mitigation with prior concern are 465 promoted according to our study. The Huabei control zone, led by the Jing-Jin-Ji area, 466 together with Shandong, Shanxi, Liaoning and Henan has a close relationship with 467 respect to both trans-boundary transport and virtual transfer of emissions. Liaoning is 468 also intimate with Jilin and Heilongjiang and may implement multilateral supervision 469 470 action. Meanwhile, for Shandong and Henan, cooperation with the Yangtze River Delta 471 (i.e., Jiangsu, Shanghai and Zhejiang) and Anhui can also achieve enhanced efficiency in emission mitigation. These 6 provinces comprise the Huadong control zone. 472

473 ASSOCIATED CONTENT

474 Supporting Information. Tables S1-S4, Figures S1-S3 include provinces and sectors
 475 information, emission factors of BC for 8 fuel types, sectoral distribution of provincial

production-based BC emissions and the evaluation of model simulation with
observations. Supporting data contain simplified Chinese MRIO Table (2007) for 30
provinces and 17 sectors. This material is available free of charge via the Internet at
http://pubs.acs.org.

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- 488 Notes
- 489 The authors declare no competing financial interest.

490 **REFERENCE**

- (1) Wu, J.; Jiang, W. M.; Fu, C. B.; Su, B. K.; Liu, H. N.; Tang, J. P., Simulation of the
 radiative effect of black carbon aerosols and the regional climate responses over
 China. *Adv. Atmos. Sci.* 2004, *21*, (4), 637-649.
- Kondo, Y.; Oshima, N.; Kajino, M.; Mikami, R.; Moteki, N.; Takegawa, N. et al.,
 Emissions of black carbon in East Asia estimated from observations at a remote
 site in the East China Sea. *J. Geophys. Res. Atmos.* 2011, *116*, D16201.
- (3) Bond, T. C.; Doherty, S. J.; Fahey, D. W.; Forster, P. M.; Berntsen, T.; DeAngelo,
 B. J. et al., Bounding the role of black carbon in the climate system: A scientific
 assessment. J. Geophys. Res. Atmos. 2013, 118, (11), 5380-5552.
- (4) Lewandowska, A.; Falkowska, L.; Murawiec, D.; Pryputniewicz, D.; Burska, D.;
 Beldowska, M., Elemental and organic carbon in aerosols over urbanized coastal
 region (southern Baltic Sea, Gdynia). *Sci. Total Environ.* 2010, *408*, (20), 4761-
- 503 4769.

- (5) IPCC, Climate Change 2007: The Physical Science Basis. Contribution of
 Working Group I to the Fourth Assessment, Report of the Intergovernmental Panel
 on Climate Change. *Cambridge University Press, Cambridge, United Kingdom, and New York, NY, USA* 2007.
- 508 (6) EPA, Effects of Black Carbon. *http://www.epa.gov/blackcarbon/effects.html*.
- (7) Ramanathan, V.; Carmichael, G., Global and regional climate changes due to black
 carbon. *Nat. Geosci.* 2008, *1*, (4), 221-227.
- (8) Reche, C.; Querol, X.; Alastuey, A.; Viana, M.; Pey, J.; Moreno, T. et al., New
 considerations for PM, Black Carbon and particle number concentration for air
 quality monitoring across different European cities. *Atmos. Chem. Phys.* 2011, *11*,
 (13), 6207-6227.
- Gong, W.; Zhang, T. H.; Zhu, Z. M.; Ma, Y. Y.; Ma, X.; Wang, W., Characteristics
 of PM_{1.0}, PM_{2.5}, and PM₁₀, and Their Relation to Black Carbon in Wuhan, Central
 China. *Atmosphere* 2015, *6*, (9), 1377-1387.
- (10) Geng, F. H.; Hua, J.; Mu, Z.; Peng, L.; Xu, X. H.; Chen, R. J.; Kan, H. D.,
 Differentiating the associations of black carbon and fine particle with daily
 mortality in a Chinese city. *Environ. Res.* 2013, *120*, 27-32.
- (11) Janssen, N. A.; Hoek, G.; Simic-Lawson, M.; Fischer, P.; van Bree, L.; ten Brink,
 H. et al., Black carbon as an additional indicator of the adverse health effects of
 airborne particles compared with PM₁₀ and PM_{2.5}. *Environ. Health Persp.* 2011, *119*, (12), 1691-1699.
- (12) Invernizzi, G.; Ruprecht, A.; Mazza, R.; De Marco, C.; Mocnik, G.; Sioutas, C.;
 Westerdahl, D., Measurement of black carbon concentration as an indicator of air
 quality benefits of traffic restriction policies within the ecopass zone in Milan,
 Italy. *Atmos. Environ.* 2011, 45, (21), 3522-3527.
- (13) Wang, H. L.; Rasch, P. J.; Easter, R. C.; Singh, B.; Zhang, R. D.; Ma, P. L.; Qian,
 Y.; Ghan, S. J.; Beagley, N., Using an explicit emission tagging method in global
 modeling of source-receptor relationships for black carbon in the Arctic:
 Variations, sources, and transport pathways. *J. Geophys. Res. Atmos.* 2014, *119*,
 (22), 12888-12909.

- (14) Liu, J. F.; Fan, S. M.; Horowitz, L. W.; Levy, H., Evaluation of factors controlling
 long-range transport of black carbon to the Arctic. *J. Geophys. Res. Atmos.* 2011, *116*, D04307.
- (15) Saikawa, E.; Naik, V.; Horowitz, L. W.; Liu, J.; Mauzerall, D. L., Present and
 potential future contributions of sulfate, black and organic carbon aerosols from
 China to global air quality, premature mortality and, radiative forcing. *Atmos. Environ.* 2009, 43, (17), 2814-2822.
- (16) Sharma, S.; Ishizawa, M.; Chan, D.; Lavoue, D.; Andrews, E.; Eleftheriadis, K.;
 Maksyutov, S., 16-year simulation of Arctic black carbon: Transport, source
 contribution, and sensitivity analysis on deposition. *J. Geophys. Res. Atmos.* 2013, *118*, (2), 943-964.
- 545 (17) Nunes, T. V.; Pio, C. A., Carbonaceous aerosols in Industrial and Coastal
 546 Atmospheres. *Atmos. Environ., Part A* 1993, 27, (8), 1339-1346.
- 547 (18) Wang, R.; Tao, S.; Shen, H. Z.; Huang, Y.; Chen, H.; Balkanski, Y. et al., Trend in
 548 Global Black Carbon Emissions from 1960 to 2007. *Environ. Sci. Technol.* 2014,
 549 48, (12), 6780-6787.
- (19) Bond, T. C.; Streets, D. G.; Yarber, K. F.; Nelson, S. M.; Woo, J. H.; Klimont, Z.,
 A technology-based global inventory of black and organic carbon emissions from
 combustion. *J. Geophys. Res. Atmos.* 2004, *109*, D14203.
- (20) Huang, Y.; Shen, H. Z.; Chen, H.; Wang, R.; Zhang, Y. Y.; Su, S. et al.,
 Quantification of Global Primary Emissions of PM_{2.5}, PM₁₀, and TSP from
 Combustion and Industrial Process Sources. *Environ. Sci. Technol.* 2014, *48*, (23),
 13834-13843.
- 557 (21) Ministry of Environmental Protection of the People's Republic of China, The Ministry of Environmental Production Anounced the Air Quality of Major Areas 558 74 and Cities in Year 2014 (in Chinese). available 559 at: 560 http://www.zhb.gov.cn/gkml/hbb/qt/201502/t20150202_295333.htm 2015.
- (22) Pui, D. Y. H.; Chen, S. C.; Zuo, Z. L., PM_{2.5} in China: Measurements, sources,
 visibility and health effects, and mitigation. *Particuology* 2014, *13*, 1-26.
- 563 (23) Ministry of Environmental Protection of the People's Republic of China, Law of

- 564the People's Republic of China on the Prevention and Control of Atmospheric565Pollution(inChinese).availableat:566http://www.zhb.gov.cn/ztbd/rdzl/gwv/wj/201509/t20150906309355.htm2015.
- 567 (24) Jiang, C.; Wang, H.; Zhao, T.; Li, T.; Che, H., Modeling study of PM_{2.5} pollutant
 568 transport across cities in China's Jing-Jin-Ji region during a severe haze episode in
 569 December 2013. *Atmos. Chem. Phys.* 2015, *15*, (10), 5803-5814.
- (25) Zhao, H. Y.; Zhang, Q.; Guan, D. B.; Davis, S. J.; Liu, Z.; Huo, H.; Lin, J. T.; Liu,
 W. D.; He, K. B., Assessment of China's virtual air pollution transport embodied
 in trade by using a consumption-based emission inventory. *Atmos. Chem. Phys.*
- **2015**, *15*, (10), 5443-5456.
- (26) Ma, P. L.; Gattiker, J. R.; Liu, X. H.; Rasch, P. J., A novel approach for determining
 source-receptor relationships in model simulations: a case study of black carbon
 transport in northern hemisphere winter. *Environ. Res. Lett.* 2013, *8*, (2).
- 577 (27) Guo, Q. F.; Hu, M.; Guo, S.; Wu, Z. J.; Hu, W. W.; Peng, J. F. et al., The
 578 identification of source regions of black carbon at a receptor site off the eastern
 579 coast of China. *Atmos. Environ.* 2015, *100*, 78-84.
- (28) Xue W.B.; F. F.; Wang J.N.; Tang G.Q.; Lei Y.; Yang J.T.; Wang Y.S., Numerical
 study on the characteristics of regional transport of PM_{2.5} in China (in Chinese). *Chin. Environ. Sci.* 2014, *34*, (6), 1361-1368.
- (29) Liu, J. F.; Mauzerall, D. L., Potential influence of inter-continental transport of
 sulfate aerosols on air quality. *Environ. Res. Lett.* 2007, 2, (4).
- (30) Guo, J. e.; Zhang, Z.; Meng, L., China's provincial CO₂ emissions embodied in
 international and interprovincial trade. *Energy Policy* 2012, *42*, 486-497.
- (31) Zhang, B.; Chen, Z. M.; Xia, X. H.; Xu, X. Y.; Chen, Y. B., The impact of domestic
 trade on China's regional energy uses: A multi-regional inout-output modeling. *Energy Policy* 2013, 63, 1169-1181.
- (32) Liu, J. F.; Mauzerall, D. L.; Horowitz, L. W.; Ginoux, P.; Fiore, A. M., Evaluating
 inter-continental transport of fine aerosols: (1) Methodology, global aerosol
 distribution and optical depth. *Atmos. Environ.* 2009, *43*, (28), 4327-4338.
- 593 (33) Davis, S. J.; Caldeira, K., Consumption-based accounting of CO₂ emissions. *Proc.*

- 594 *Natl. Acad. Sci. USA* **2010**, *107*, (12), 5687-5692.
- (34) Wang, R.; Tao, S.; Wang, W. T.; Liu, J. F.; Shen, H. Z.; Shen, G. F. et al., Black
 Carbon Emissions in China from 1949 to 2050. *Environ. Sci. Technol.* 2012, 46,
 (14), 7595-7603.
- (35) Meng, J.; Liu, J.; Guo, S.; Li, J.; Li, Z.; Tao, S., Trend and driving forces of
 Beijing's black carbon emissions from sectoral perspectives. *J. Clean Prod.* 2016, *112*, 1272-1281.
- 601 (36) CESY, China Energy Statistical Yearbook 2008. *China Statistics Press, Beijing*602 2008.
- (37) Liu, W. D.; Chen, J.; Tang, Z.P.; Liu, H.G.; Han, D.; Li, F.Y., Compliment theory
 and practice of China's interregional input–output table for 30 regions in 2007 (in
 Chinese). *China Statistics Press, Beijing, P.R.China* 2012a.
- (38) Nordmann, S.; Cheng, Y. F.; Carmichael, G. R.; Yu, M.; van der Gon, H. A. C. D.;
 Zhang, Q. et al., Atmospheric black carbon and warming effects influenced by the
 source and absorption enhancement in central Europe. *Atmos. Chem. Phys.* 2014, *14*, (23), 12683-12699.
- (39) Matsui, H.; Koike, M.; Kondo, Y.; Moteki, N.; Fast, J. D.; Zaveri, R. A.,
 Development and validation of a black carbon mixing state resolved threedimensional model: Aging processes and radiative impact. *J. Geophys. Res. Atmos.* **2013**, *118*, (5), 2304-2326.
- (40) Gao, Y.; Zhao, C.; Liu, X.; Zhang, M.; Leung, L. R., WRF-Chem simulations of
 aerosols and anthropogenic aerosol radiative forcing in East Asia. *Atmos. Environ.*2014, 92, 250-266.
- (41) Tao, W.; Liu, J.; Ban-Weiss, G. A.; Hauglustaine, D. A.; Zhang, L.; Zhang, Q.;
 Cheng, Y.; Yu, Y.; and Tao, S., Effects of urban land expansion on the regional
 meteorology and air quality of eastern China. *Atmos. Chem. Phys.* 2015, *15*, 85978614.
- (42) Rasch, P. J.; Barth, M. C.; Kiehl, J. T.; Schwartz, S. E.; Benkovitz, C. M., A
 description of the global sulfur cycle and its controlling processes in the National
 Center for Atmospheric Research Community Climate Model, Version 3. J.

- 624 Geophys. Res. Atmos. 2000, 105, (D1), 1367-1385.
- (43) J. Zhang; J. Liu; S. Tao; and G. A. Ban-Weiss, Long-range transport of black
 carbon to the Pacific Ocean and its dependence on aging timescale. *Atmos. Chem. Phys.* 2015, *15*, 11521-11535.
- (44) Cao, J. J.; Lee, S. C.; Ho, K. F.; Zhang, X. Y.; Zou, S. C.; Fung, K.; Chow, J. C.;
 Watson, J. G., Characteristics of carbonaceous aerosol in Pearl River Delta Region,
 China during 2001 winter period. *Atmos. Environ.* 2003, *37*, (11), 1451-1460.
- (45) Sun, Y. L.; Zhuang, G. S.; Ying, W.; Han, L. H.; Guo, J. H.; Mo, D.; Zhang, W. J.;
 Wang, Z. F.; Hao, Z. P., The air-borne particulate pollution in Beijing concentration, composition, distribution and sources. *Atmos. Environ.* 2004, *38*,
- 634 (35), 5991-6004.
- (46) Duan, F. K.; He, K. B.; Ma, Y. L.; Yang, F. M.; Yu, X. C.; Cadle, S. H.; Chan, T.;
 Mulawa, P. A., Concentration and chemical characteristics of PM_{2.5} in Beijing,
 China: 2001-2002. *Sci. Total Environ.* 2006, *355*, (1-3), 264-275.
- (47) Cao, J. J.; Lee, S. C.; Chow, J. C.; Watson, J. G.; Ho, K. F.; Zhang, R. J. et al.,
 Spatial and seasonal distributions of carbonaceous aerosols over China. J. *Geophys. Res. Atmos.* 2007, 112, (D22).
- (48) Qu, W. J.; Zhang, X. Y.; Arimoto, R.; Wang, D.; Wang, Y. Q.; Yan, L. W.; Li, Y.,
 Chemical composition of the background aerosol at two sites in southwestern and
 northwestern China: potential influences of regional transport. *Tellus B* 2008, *60*,
 (4), 657-673.
- (49) Zhang, X. Y.; Wang, Y. Q.; Zhang, X. C.; Guo, W.; Gong, S. L., Carbonaceous
 aerosol composition over various regions of China during 2006. *J. Geophys. Res. Atmos.* 2008, *113*, D14111.
- (50) Huang, H.; Ho, K. F.; Lee, S. C.; Tsang, P. K.; Ho, S. S. H.; Zou, C. W.; Zou, S.
 C.; Cao, J. J.; Xu, H. M., Characteristics of carbonaceous aerosol in PM_{2.5}: Pearl
 Delta River Region, China. *Atmos. Res.* 2012, *104*, 227-236.
- 651 (51) Zhao, P. S.; Dong, F.; He, D.; Zhao, X. J.; Zhang, X. L.; Zhang, W. Z.; Yao, Q.;
- Liu, H. Y., Characteristics of concentrations and chemical compositions for $PM_{2.5}$
- 653 in the region of Beijing, Tianjin, and Hebei, China. *Atmos. Chem. Phys.* 2013, 13,

654 (9), 4631-4644.

- (52) Pathak, R. K.; Wang, T.; Ho, K. F.; Lee, S. C., Characteristics of summertime PM_{2.5}
 organic and elemental carbon in four major Chinese cities: Implications of high
 acidity for water-soluble organic carbon (WSOC). *Atmos. Environ.* 2011, 45, (2),
 318-325.
- (53) Wang, G.; Li, J.; Cheng, C.; Hu, S.; Xie, M.; Gao, S.; Zhou, B.; Dai, W.; Cao, J.;
 An, Z., Observation of atmospheric aerosols at Mt. Hua and Mt. Tai in central and
 east China during spring 2009-Part 1: EC, OC and inorganic ions. *Atmos. Chem. Phys.* 2011, *11*, (9), 4221-4235.
- (54) Duan, J. C.; Tan, J. H.; Cheng, D. X.; Bi, X. H.; Deng, W. J.; Sheng, G. Y.; Fu, J.
 M.; Wong, M. H., Sources and characteristics of carbonaceous aerosol in two
 largest cities in Pearl River Delta Region, China. *Atmos. Environ.* 2007, *41*, (14),
 2895-2903.
- (55) Feng, Y. L.; Chen, Y. J.; Guo, H.; Zhi, G. R.; Xiong, S. C.; Li, J.; Sheng, G. Y.; Fu,
 J. M., Characteristics of organic and elemental carbon in PM_{2.5} samples in
 Shanghai, China. *Atmos. Res.* 2009, *92*, (4), 434-442.
- (56) Li, X. X.; Shen, Z.X.; Cao, J.J.; Liu, S.X.; Zhu, C.S.; Zhang, T., Distribution of
 carbonaceous aerosol during Spring 2005 over the Horqin sandland in northeastern
 China. *China Particuology* 2006, *4*, 316-322.
- (57) Zhang, F. W.; Zhao, J. P.; Chen, J. S.; Xu, Y.; Xu, L. L., Pollution characteristics
 of organic and elemental carbon in PM_{2.5} in Xiamen, China. *J. Environ. Sci.-China* **2011**, *23*, (8), 1342-1349.
- (58) Leontief, W., Environmental Repercussions and the Economic Structure: An InputOutput Approach. *Rev. Econ. Stat.* 1970, *52*, 262-271.
- (59) Lenzen, M.; Moran, D.; Kanemoto, K.; Foran, B.; Lobefaro, L.; Geschke, A.,
 International trade drives biodiversity threats in developing nations. *Nature* 2012,
 486, (7401), 109-112.
- (60) Wiedmann, T. O.; Schandl, H.; Lenzen, M.; Moran, D.; Suh, S.; West, J.;
 Kanemoto, K., The material footprint of nations. *Proc. Natl. Acad. Sci. USA* 2015, *112*, (20), 6271-6276.

- (61) Lei, Y.; Zhang, Q.; He, K. B.; Streets, D. G., Primary anthropogenic aerosol
 emission trends for China, 1990–2005. *Atmos. Chem. Phys.* 2011, *11*, (3), 931-954.
- (62) Zhang, N.; Qin, Y.; Xie, S. D., Spatial distribution of black carbon emissions in
 China. *Chinese Sci. Bull.* 2013, 58, (31), 3830-3839.
- (63) Meng, J.; Liu, J. F.; Xu, Y.; Tao, S., Tracing Primary PM_{2.5} emissions via Chinese
 supply chains. *Environ. Res. Lett.* 2015, *10*, (5).
- 690 (64) Feng, K. S.; Davis, S. J.; Sun, L. X.; Li, X.; Guan, D. B.; Liu, W. D.; Liu, Z.;
- Hubacek, K., Outsourcing CO₂ within China. *Proc. Natl. Acad. Sci. USA* 2013, *110*, (28), 11654-11659.
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Figure 1. Largest surface BC concentration contribution via atmospheric transport within 30 provinces. The colors in the map indicate annual mean surface BC concentration. Arrows on the map reflect a typical contribution above $0.1 \ \mu g \cdot m^{-3}$. The thickness of the arrows indicates the relative magnitude of the absolute inter-provincial contribution of surface concentration.





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Figure 2. (a) A comparison between production-based and consumption-based BC emissions. 704 705 Consumption-based BC emissions are categorized into 4 types based on final consumption; (b) Largest net fluxes in "traded" BC emissions among 30 provinces. Color in the map indicates total 706 707 net emission budget (emission imports minus exports) via trade. Red indicates an emission importer, 708 i.e., more BC is emitted due to the inter-provincial trade. Blue indicates an emission exporter. The 709 arrows reflect typical cross-border net emission flows above 1 Gg in inter-provincial trade. The thickness of the arrows indicates the relative magnitude of the net BC emissions transferred between 710 711 provinces.



Figure 3. Sectoral BC emissions embodied in exported and imported products via inter-provincial
trade for 30 provinces; the 17 sectors in MRIO are sorted into 8 for clearer presentation (listed in
Table S2). Intermediate products (dark gray) are embodied BC emissions used by industry.





719 Figure 4. The contribution of surface BC concentration in 30 provinces from both production and 720 consumption perspectives. Blue (Local-Local), red (Local-Domestic), dark grey (Local-Foreign) 721 columns indicate the percentage of surface concentration in a province that is contributed by its own 722 on-site emissions but induced by consumptions from its own province, the rest 29 provinces, and foreign countries, respectively. Comparably, green (Domestic-Local), yellow (Domestic-Domestic), 723 724 and light grey (Domestic-Foreign) columns, respectively, indicate the percentage that is contributed 725 by the emissions released in the other 29 provinces but induced by local, domestic and foreign 726 consumptions. The dashed line at 50% marks the comparison between surface concentrations 727 contributed by local and non-local on-site emissions. 728



Figure 5. (a) Source-receptor relationship between provincial emission and BC concentration (area-730 731 weighted at the surface) among 30 provinces (unit: $(\mu g \cdot m^{-3})/(Gg \cdot yr^{-1})$); (b) Source-receptor 732 relationship between provincial final consumption and on-site emission (unit: tons per billion Yuan); 733 (c) Source-receptor relationship between provincial final consumption and surface BC concentration (unit: $(\mu g \cdot m^{-3})/(billion Yuan \cdot yr^{-1})$). Province abbreviations (or see Table S1 in the 734 735 supporting information) are: HL-Heilongjiang, JL-Jilin, LN-Liaoning, BJ-Beijing, TJ-Tianjin, HB-736 Hebei, SD-Shandong, SX-Shanxi, HE-Henan, UB-Hubei, HU-Hunan, AH-Anhui, JX-Jiangxi, JS-737 Jiangsu, SH-Shanghai, ZJ-Zhejiang, FJ-Fujian, GD-Guangdong, HA-Hainan, GX-Guangxi, CQ-738 Chongqing, SC-Sichuan, GZ-Guizhou, YN-Yunnan, NM-Neimeng, SA-Shaanxi, GS-Gansu, QH-739 Qinghai, NX-Ningxia, SJ-Xinjiang.