

Senate Standing Committee on Economics
ANSWERS TO QUESTIONS ON NOTICE
Innovation, Industry, Science and Research Portfolio
Budget Estimates Hearing
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AGENCY/DEPARTMENT: INNOVATION, INDUSTRY, SCIENCE AND RESEARCH – CHIEF SCIENTIST

TOPIC: Climate Change –human action

REFERENCE: Question on Notice (Hansard 1 June 2009, E 99)

QUESTION No. BI-87

Senator ABETZ—The largest portion of the change in the CO₂ levels is due to human action. Are you able to assist the committee as to what the body of scientific knowledge is? You may need to take that on notice. I could accept that you would not necessarily have this figure in your back pocket, but if you do it would be great. You say that the largest portion of the change is due to human action. What proportion would you say is due to human action? Would you put it at 51 per cent or 80 per cent? Can we quantify it in any way?

Prof. Sackett—It can be quantified. As in most good science, it has a range. In fact, what science does is not only tell one a best estimate but give some uncertainty around that, or rather a degree of certainty around the range in which those numbers are believed to lie. I would not like to state that in a forum such as this. I would like to take that on notice.

ANSWER

The proportion of the change in atmospheric CO₂ (carbon dioxide) concentration that is due to human (anthropogenic) action depends on the timescale over which one measures the change. A large body of scientific evidence indicates that nearly all of the long-term increase of more than 35% (107ppm, from 280 to 387ppm) in atmospheric CO₂ concentrations since 1800 is due to anthropogenic causes. Natural causes over this period contribute short-term variability on time scales of several months to a few years with transient amplitudes of about 10 ppm or less. These conclusions rest on data arising from ice core analysis, direct atmospheric CO₂ measurements, careful accounting of the net amount of CO₂ released into the air by humans, and isotopic ‘‘fingerprint’’ analysis of atmospheric and marine CO₂. In sum, at any given moment in the present, about 90-99% of the industrial era *change* of 107ppm in the atmospheric CO₂ concentration is due to human activity. The variability of this percentage is due to short-term, oscillatory natural causes. The much larger, steady long-term increasing trend is due to human activities. I will now summarize the body of evidence that supports these conclusions.

The baseline data: Atmospheric CO₂ over millennia during the pre-industrial era

Over a period stretching over most of the Holocene from more than 11,000 years ago to the year 1750, CO₂ concentrations have been stable at 280 ppm, with variations of 20ppm or less (Etheridge *et al.* 1996; Etheridge *et al.* 1998; Indermühle *et al.*, 1999, Sabine *et al.* 2004a). The primary source of atmospheric CO₂ concentration data during this period comes from

analysis of the composition of air enclosed in bubbles in ice cores from Greenland and Antarctica. The measured variation of no more than 20 ppm (ie, 7% of pre-industrial levels, or about 5% of current levels) during this time frame is judged to be largely due to non-human sources.

The data: Atmospheric CO₂ over the whole of the Industrial era

Since the beginning of the 19th century, carbon dioxide in the atmosphere has risen substantially. The primary source of data for CO₂ concentrations during this period comes from ice core analysis and, since the late 1950s, also from direct measurements of atmospheric CO₂, most notably from carefully calibrated observing stations at Mauna Loa, Hawai'i (Keeling *et al.*, 1976; Thoning *et al.*, 1989; Conway *et al.*, 1994) and Cape Grim Tasmania (Francey 2005). These records are in agreement, and show that by 1958 atmospheric CO₂ had risen to 315ppm, and then rose even more sharply to the present value of 387ppm. It is this change of 107ppm over the past 200 years that is attributed almost entirely to human activity, a change more than five times the variability in atmospheric CO₂ from the end of the last ice age to the dawn of the industrial age.

The accounting: Net anthropogenic contribution to atmospheric CO₂

The main evidence that human activities are responsible for the increased CO₂ during the industrial era comes from an accounting of the actual amount of carbon dioxide released due to humans. Tallying the increased CO₂ from industrial processes, the amount of fossil fuel extracted and combusted (Marland & Rotty 1984; Marland *et al.* 2006; Raupach *et al.* 2007), and the reduction of CO₂ sinks caused primarily by land clearing (Houghton 2003), indicates that humans have produced far more CO₂ than now remains in the atmosphere (Sabine *et al.* 2004a; Denman *et al.*, IPCC 2007, Ch 7; Canadell *et al.* 2007; Raupach *et al.* 2008). The total amount of carbon released from 1850 to 2007 that is attributed to these activities (Canadell *et al.* 2007) is enough to have raised the atmospheric concentration of CO₂ to nearly 500 ppm if all the carbon had remained as CO₂ in the atmosphere. Concentrations have not reached that level because the ocean and the terrestrial biosphere have the capacity to absorb some of the CO₂ (Sabine *et al.*, 2004b). Humans have produced CO₂ *faster* than the ocean and biosphere can absorb it; the remainder explains the observed increase in the atmosphere.

Natural causes of small, short-term variations in atmospheric CO₂

On short time frames ranging from seasons to a few years, natural causes have affected the modern atmospheric concentration of CO₂. Over a single year, northern hemisphere observing stations show significant, regular and cyclic variability of 5 to 7 ppm due to seasonal plant growth (Keeling *et al.*, 1976; Thoning *et al.*, 1989; Conway *et al.*, 1994); changes in the Southern hemisphere are less (Francey 2005). Over 3-5 year time periods, consequences arising from volcanoes and major ocean upwellings (notably ENSO) can cause interannual variability of about 4 ppm and 1 ppm, respectively (Bacastow 1976; Jones and Cox 2001; Jones *et al.*, 2001; Prentice *et al.*, 2001, IPCC 2001, Ch 3; Lintner 2002; Denman *et al.*, IPCC 2007). These changes are small, short-lived, and oscillatory compared to the large, long-trend rise of 107 ppm observed over the last 200 years.

An anthropogenic fingerprint: The decreasing ratio of ¹³C to ¹²C:

Human activities are also shown to be the cause of recent increases in atmospheric CO₂ concentrations by isotopic analysis of oceanic and atmospheric carbon dioxide. Since the last ice age, the pre-industrial age atmosphere has had a relatively constant ratio between two types of carbon, the isotopes ¹²C and ¹³C, varying slightly over a few millennia (Indermühle *et al.*, 1999). CO₂ produced from burning fossil fuels or land clearing, however, has a distinct isotopic fingerprint. Plants have a preference for lighter isotopes (¹²C vs ¹³C) when building their tissue (Park and Epstein 1960), resulting in a lower ¹³C/¹²C ratio than the atmosphere. Since fossil fuels are ultimately derived from ancient plants, fossil fuels also have low ¹³C/¹²C. As CO₂ from these materials is released into, and mixes with, the atmosphere, the average ¹³C/¹²C ratio of the atmosphere decreases (Prentice *et al.*, IPCC 2001), as has been directly measured (Keeling *et al.* 2005). This is consistent with the decreasing ¹³C/¹²C ratio measured from CO₂ trapped in ice cores for the period since 1850 (Francey 1999). Furthermore, ¹³C/¹²C analysis indicates that this ratio is also decreasing in the ocean. The atmosphere has an even lower ¹³C/¹²C ratio (fingerprint) than the ocean, indicating that ¹³C-depleted carbon from fossil fuels and deforestation is passing from the atmosphere into the ocean (Quay *et al.*, 1992; Quay *et al.*, 2003).

References

(The following references are a small part of the supporting body of evidence.)

Bacastow R (1976) Modulation of atmospheric carbon dioxide by the Southern Oscillation, *Nature*, 261, 116-118.

Canadell JG, Le Quere C, Raupach MR, Field CB, Buitenhuis ET, Ciais P, Conway TJ, Gillett NP, Houghton RA, Marland G (2007) Contributions to accelerating atmospheric CO₂ growth from economic activity, carbon intensity, and efficiency of natural sinks. *Proceedings of the National Academy of Sciences (PNAS)* **104**:18866-18870

Conway TJ, Tans PP, Waterman LS, Thoning KW, Kitzis DR, Masarie KA and Zhang N, 1994: Evidence for interannual variability of the carbon cycle from the National Oceanic and Atmospheric Administration/Climate Monitoring and Diagnostics Laboratory global air sampling network, *Journal of Geophysical Research*, **99**, 22831-22855

Denman, K.L., G. Brasseur, A. Chidthaisong, P. Ciais, P.M. Cox, R.E. Dickinson, D. Hauglustaine, C. Heinze, E. Holland, D. Jacob, U. Lohmann, S Ramachandran, P.L. da Silva Dias, S.C. Wofsy and X. Zhang, (2007): Couplings between changes in the climate system and biogeochemistry, in: *Climate Change 2007: The Physical Science Basis. Contribution of Working Group I to the Fourth Assessment Report of the Intergovernmental Panel on Climate Change* [Solomon, S., D. Qin, M. Manning, Z. Chen, M. Marquis, K.B. Averyt, M.Tignor and H.L. Miller (eds.)]. Cambridge University Press, Cambridge, United Kingdom and New York, NY, USA.

Etheridge DM, Steele LP, Langenfelds RL, Francey RJ, Barnola JM, Morgan VI (1996) Natural and anthropogenic changes in atmospheric CO₂ over the last 1000 years from air in Antarctic ice and firn. *J. Geophys. Res. Atmos.* **101**:4115-4128

- Etheridge DM, Steele LP, Langenfelds RL, Francey RJ, Barnola JM, Morgan VI (1998) Historical CO₂ records from the Law Dome DE08, DE08-2, and DSS ice cores. *Trends: A Compendium of Data on Global Change, Carbon Dioxide Information Analysis Center*, Oak Ridge National Laboratory, U.S. Department of Energy, Oak Ridge, Tennessee, USA
- Francey RJ, Allison CE, Etheridge DM, Trudinger CM, Enting IG, Leuenberger M, Langenfelds RL, Michel E, and Steele LP (1999) A 1000-year high precision record of d¹³C in atmospheric CO₂, *Tellus*, Ser. B, 51, 170– 193
- Francey RJ (2005) Recent record growth in atmospheric CO₂ levels. *Environmental Chemistry* 2:3-5
- Houghton RA (2003) Revised estimates of the annual net flux of carbon to the atmosphere from changes in land use and land management 1850-2000. *Tellus* 55B:378-390
- Indermühle A, Stocker TF, Joss F, Fischer H, Smith HJ, Wahlen M, Deck B, Mastroianni D, Tschumi J, Blunier T, Meyer R, and Stauffer B (1999), Holocene carbon-cycle dynamics based on CO₂ trapped in ice at Taylor Dome, Antarctica. *Nature*, 398, 121-126.
- Jones, CD and Cox, PM (2001) Modeling the volcanic signal in the atmospheric CO₂ record, *Global Biogeochemical Cycles*, 15, 453-465.
- Jones CD, Collins M, Cox, PM, and Spall, SA (2001) The carbon cycle response to ENSO: a coupled climate-carbon cycle model study, *Journal of Climate*, 14, 4113–4129.
- Keeling CD, Bacastow RB, Bainbridge AE, Ekdahl CA, Guenther PR, and Waterman LS (1976) Atmospheric carbon dioxide variations at Mauna Loa Observatory, Hawaii, *Tellus*, vol. 28, 538-551.
- Keeling CD, Bollenbacher AF, and Whorf TP (2005) Monthly atmospheric ¹³C/¹²C isotopic ratios for 10 SIO stations. In: *Trends: A Compendium of Data on Global Change. Carbon Dioxide Information Analysis Center*, Oak Ridge National Laboratory, U.S. Department of Energy, Oak Ridge, TN
- Lintner BR (2002) Characterizing global CO₂ interannual variability with empirical orthogonal function/principal component (EOF/PC) analysis. *Geophysical Research Letters*, 29, 19, 1921-1924
- Marland G, Rotty RM (1984) Carbon dioxide emissions from fossil fuels: A procedure for estimation and results for 1950-82. *Tellus Ser. B* 36:232-261
- Marland G, Boden TA, Andres RJ (2006) Global, regional, and national CO₂ emissions. *Trends: A Compendium of Data on Global Change, Carbon Dioxide Information Analysis Center*, Oak Ridge National Laboratory, U.S. Department of Energy, Oak Ridge, Tennessee, USA
- Park R and Epstein S (1960) Carbon isotope fractionation during photosynthesis, *Geochimica et Cosmochimica Acta*, 21, 110-126

- Prentice IC, Farquhar GD, Fasham MJR, Goulden ML, Heimann M, Jaramillo VJ, Khashgi HS, Le Quéré C, Scholes RJ, Wallace DWR (2001) Climate Change 2001: IPCC Third Assessment, The Carbon Cycle and Atmospheric Carbon Dioxide in: *The Scientific Basis. Contribution of Working Group I to the Third Assessment Report of the Intergovernmental Panel on Climate Change*, Cambridge University Press, Cambridge, United Kingdom and New York
- Quay PD, Tilbrook B, and Wong CS (1992) Oceanic Uptake of Fossil Fuel CO₂: Carbon-13 Evidence, *Science*, Vol. 256. no. 5053, 74 – 79
- Quay P, Sonnerup R, Westby T, Stutsman J, and McNichol A (2003) Changes in the ¹³C/¹²C of dissolved inorganic carbon in the ocean as a tracer of anthropogenic CO₂ uptake, *Global Biogeochemical Cycles*, **17**, 1, 1004-1023
- Raupach MR, Marland G, Ciais P, Le Quere C, Canadell JG, Klepper G, Field CB (2007) Global and regional drivers of accelerating CO₂ emissions. *Proceedings of the National Academy of Science* **104**:10288-10293
- Raupach MR, Canadell JG, Le Quere C (2008) Anthropogenic and biophysical contributions to increasing atmospheric CO₂ growth rate and airborne fraction. *Biogeosciences* **5**:1601-1613
- Sabine CL, Heimann M, Artaxo P, Bakker DCE, Chen C-TA, Field CB, Gruber N, Le Quere C, Prinn RG, Richey JD, Romero Lankao P, Sathaye JA, Valentini R (2004a) Current status and past trends of the global carbon cycle. In: Field CB, Raupach MR (eds) *The Global Carbon Cycle: Integrating Humans, Climate, and the Natural World*. SCOPE 62, Island Press, Washington, 17-44
- Sabine CL, Feely RA, Gruber N, Key RM, Lee K, Bullister JL, Wanninkhof R, Wong CS, Wallace DWR, Tilbrook B, Millero FJ, Peng T-H, Kozyr A, Ono T., Rios AF (2004b) The Oceanic Sink for Anthropogenic CO₂, *Science* **16**, Vol. 305. no. 5682, pp. 367 – 371
- Thoning KW, Tans PP, and Komhyr WD (1989) Atmospheric carbon dioxide at Mauna Loa Observatory 2. Analysis of the NOAA GMCC data, 1974-1985, *J. Geophys. Research*, vol. 94, 8549-8565.