

TEMPO Mission Overview and Status **Kelly Chance**

Smithsonian Astrophysical Observatory tempo.si.edu

July 28, 2016







Hourly Measurement of Pollution

60 minutes



- Currently on-schedule and on-budget
 - Passed System Requirements Review and Mission Definition Review in November 2013
 - Passed KDP-B April 2014
 - Most technical issues solved at the preliminary design level, following technical interchange meeting at Ball, April 2014
 - Passed PDR on July 31, 2014
 - Now in Phase C: KDP-C April 10, 2015
 - Instrument CDR June 2015
 - Ground Systems CDR May 2016
 - Test Readiness Review August 2016
- Select satellite host 2017+
 - TEMPO operating longitude and launch date are not known until after host selection
- Instrument delivery 08/2017 for launch 11/2018 or later, most likely in 2020 or 2021

Hourly atmospheric pollution from geostationary Earth orbit

PI: Kelly Chance, Smithsonian Astrophysical Observatory Instrument Development: Ball Aerospace Project Management: NASA LaRC Other Institutions: NASA GSFC, NOAA, EPA, NCAR, Harvard, UC Berkeley, St. Louis U, U Alabama Huntsville, U Nebraska, RT Solutions, Carr Astronautics International collaboration: Mexico, Canada, Cuba, Korea, U.K., ESA, Spain

Selected Nov. 2012 as NASA's first Earth Venture Instrument

- Instrument delivery May 2017
- NASA will arrange hosting on commercial geostationary communications satellite with launch expected NET 11/2018

Provides hourly daylight observations to capture rapidly varying emissions & chemistry important for air quality

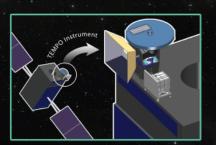
- UV/visible grating spectrometer to measure key elements in tropospheric ozone and aerosol pollution
- Distinguishes boundary layer from free tropospheric & stratospheric ozone

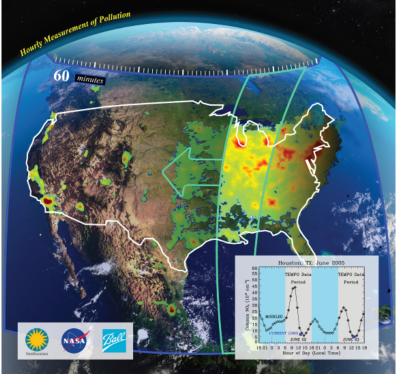
Aligned with Earth Science Decadal Survey recommendations

- Makes many of the GEO-CAPE atmosphere measurements
- Responds to the phased implementation recommendation of GEO-CAPE mission design team

vatory Tropospheric Emissions Monitoring of Pollution TEMPO's concurrent high temporal (hour)

TEMPO's concurrent high temporal (hourly) and spatial resolution measurements from geostationary orbit of tropospheric ozone, aerosols, their precursors, and clouds create a revolutionary dataset that provides understanding and improves prediction of air quality and climate forcing in Greater North America.





7/12/16 North American component of an international constellation for air quality observations

TEMPO instrument concept

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- Measurement technique
 - Imaging grating spectrometer measuring solar backscattered Earth radiance
 - Spectral band & resolution: 290-490 + 540-740 nm @ 0.6 nm FWHM, 0.2 nm sampling
 - 2 2-D, 2k×1k, detectors image the full spectral range for each geospatial scene

• Field of Regard (FOR) and duty cycle

- Mexico City/Yucatan, Cuba to the Canadian oil sands, Atlantic to Pacific
- Instrument slit aligned N/S and swept across the FOR in the E/W direction, producing a radiance map of Greater North America in one hour

Spatial resolution

- 2.1 km N/S × 4.7 km E/W native pixel resolution (9.8 km²)
- Co-add/cloud clear as needed for specific data products
- Standard data products and sampling rates
 - Most sampled hourly, including eXceL O₃ (troposphere, PBL)
 - NO₂, H₂CO, C₂H₂O₂, SO₂ sampled hourly (average results for \geq 3/day if needed)
 - Nominal spatial resolution 8.4 km N/S × 4.7 km E/W at center of domain (can often measure 2.1 km N/S × 4.7 km E/W)
 - Measurement requirements met up to 50° for SO₂, 70° SZA for other products

Baseline and threshold data products

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Species/Products	Required Precision	Temporal Revisit
0-2 km O ₃ (Selected Scenes) Baseline only	10 ppbv	2 hour
Tropospheric O ₃	10 ppbv	1 hour
Total O ₃	3%	1 hour
Tropospheric NO ₂	1.0×10^{15} molecules cm ⁻²	1 hour
Tropospheric H ₂ CO	1.0×10^{16} molecules cm ⁻²	3 hour
Tropospheric SO ₂	1.0×10^{16} molecules cm ⁻²	3 hour
Tropospheric C ₂ H ₂ O ₂	4.0×10^{14} molecules cm ⁻²	3 hour
Aerosol Optical Depth	0.10	1 hour

- Minimal set of products sufficient for constraining air quality
- Across Greater North America (GNA): 18°N to 58°N near 100°W, 67°W to 125°W near 42°N
- Data products at urban-regional spatial scales
 - Baseline ≤ 60 km² at center of Field Of Regard (FOR)
 - Threshold ≤ 300 km² at center of FOR
- Temporal scales to resolve diurnal changes in pollutant distributions
- Collected in cloud-free scenes
- Geolocation uncertainty of less than 4 km
- Mission duration, subject to instrument availability
 - Baseline 20 months
 - Threshold 12 months

TEMPO mission concept

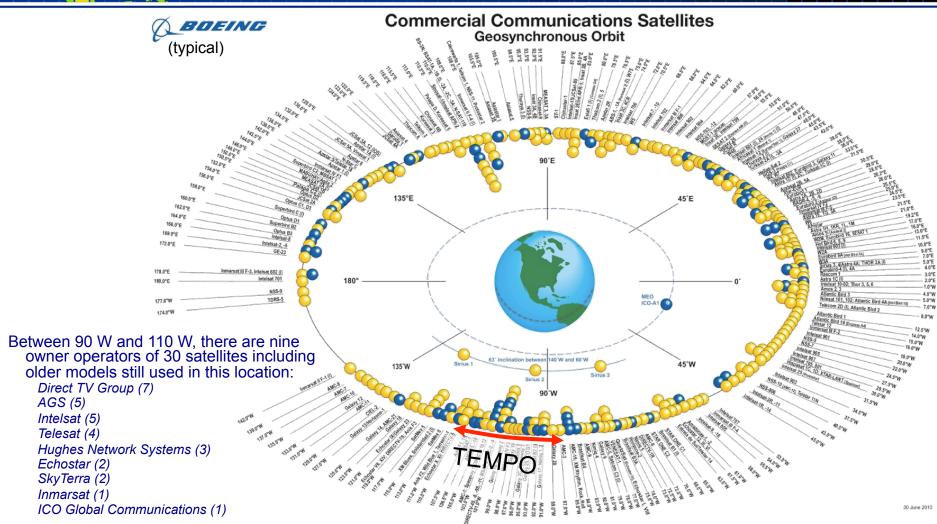
• Geostationary orbit, operating on a commercial telecom satellite

- NASA will arrange launch and hosting services (per Earth Venture Instrument scope)
 - 80-115° W acceptable latitude
 - Specifying satellite environment, accommodation
- Hourly measurement and telemetry duty cycle for at least ≤70° SZA

TEMPO is low risk with significant space heritage

- We proposed SCIAMACHY in 1985, as suggested by the late Dr. Dieter Perner
- All proposed TEMPO measurements have been made from low Earth orbit satellite instruments to the required precisions by SAO and Science Team members
- All TEMPO launch algorithms are implementations of currently operational algorithms
 - NASA TOMS-type O₃
 - SO₂, NO₂, H₂CO, C₂H₂O₂ from fitting with AMF-weighted cross sections
 - Absorbing Aerosol Index, UV aerosol, Rotational Raman scattering cloud
 - SAO eXceL profile/tropospheric/PBL O₃ for selected geographic targets
- Example higher-level products: Near-real-time pollution/AQ indices, UV index
- **TEMPO** research products will greatly extend science and applications
 - **Example research products:** BrO and IO from AMF-normalized cross sections; height-resolved SO₂; additional cloud/aerosol products; vegetation products; additional gases

Geostationary orbit opportunities of interest



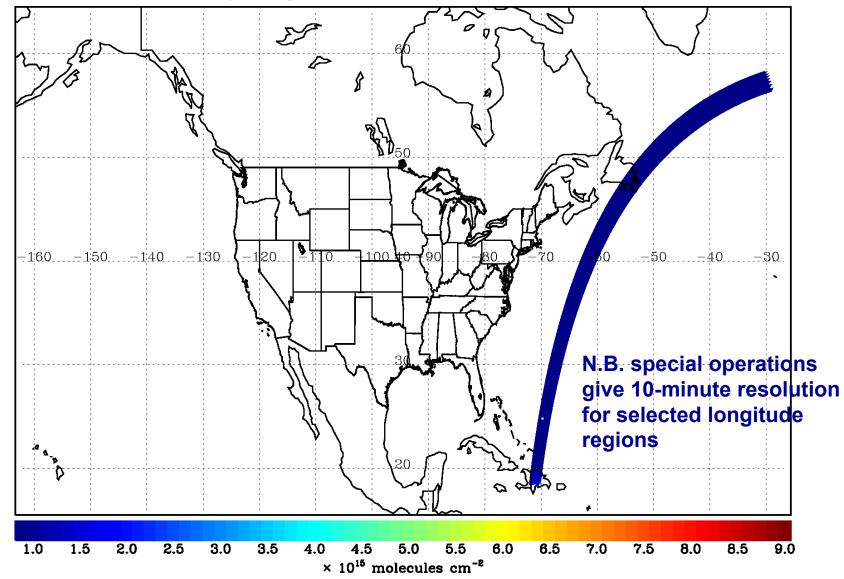
TEMPO can be located between 80 – 120 West

TEMPO hourly NO₂ sweep

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OMI NO₂ in April (2005-2008) over TEMPO FOR





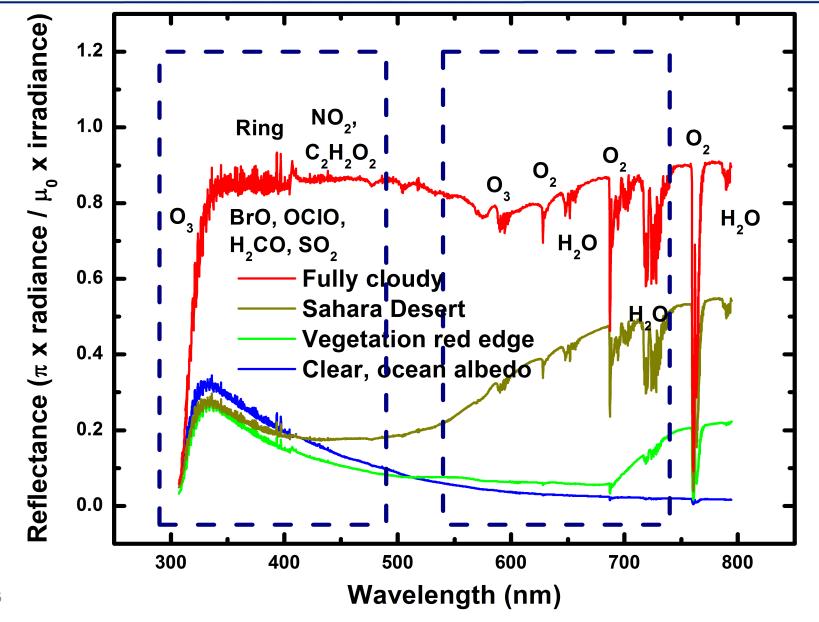
Air quality requirements from the GEO-CAPE Science Traceability Matrix

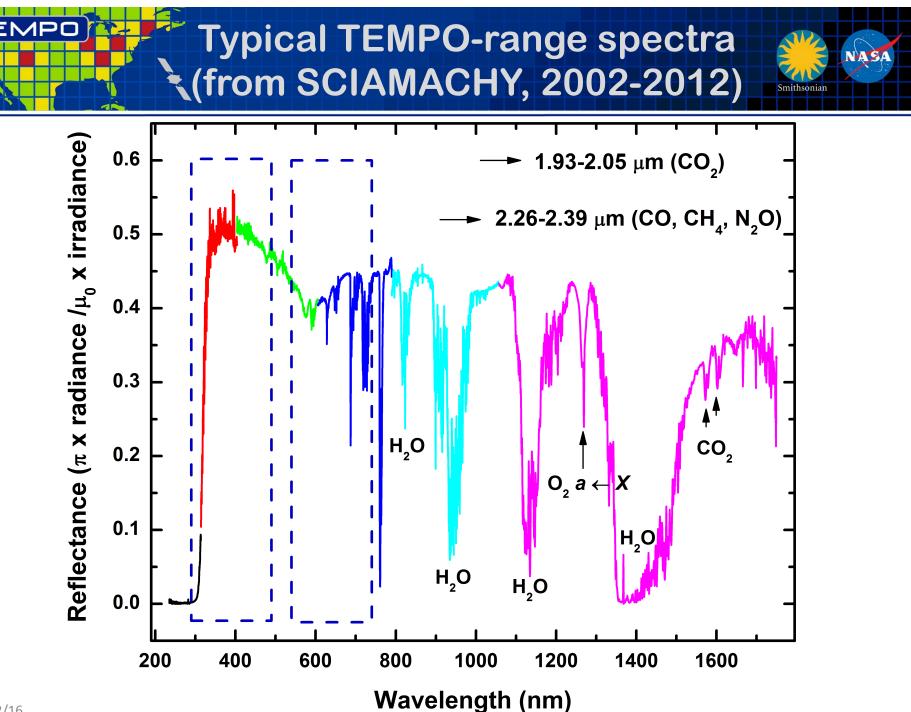
11-28-2011 DRAFT GEO-CAPE aerosol-atmospheres Science Traceability Matrix BASELINE and THRESHOLD Measurement Objectives Measurement Requirements Science Questions Measurement Rationale (color flag maps to Science Questions) (mapped to Measurement Objectives) Geostationary Observing Location: 100 W +/-10 Provides optimal view of North America. Baseline measurements¹ What are the O3, NO2, CO, SO2, HCHO, CH4, NH3, CHOCHO, temporal and Column measurements: [A to K] Continue the current state of practice in different temporal sampling frequencies, 4 km x 4 All the baseline and threshold species vertical: add temporal resolution spatial variations km product horizontal spatial resolution at the center of the domain; and AOD, AAOD, AI, aerosol optical of emissions of Cloud Camera 1 km x 1km horizontal spatial mprove retrieval accuracy, provide centroid height (AOCH), hourly for SZA<70 and 8 km dases and esolution, two spectral bands, baseline only diagnostics for gases and aerosol x 8 km product horizontal spatial resolution at the aerosols importan center of the domain. /ertical information: A to K for air quality and wo pieces of information in the 03.00 Threshold measurements1: Separate the lower-most troposphere climate? troposphere in daylight with (Baseline and CO hourly day and night; O3, NO2 hourly when from the free troposphere for O3, CO, sensitivity to the lowest 2 km Threshold) SZA<70; AOD hourly (SZA<50) ; at 8 km x 8 km product horizontal spatial resolution at the center of AOCH Detect aerosol plume height; improve How do physical ltitude (+/- 1km) he domain (baseline only) retrieval accuracy. chemical, and dynamical Product horizontal spatial resolution at the center of the domain. (nominally 100W, 35 N): A to H A Measure the threshold or baseline species or processes properties with the temporal and spatial km x 4 km (baseline) Gases determine resolution specified (see next column) to quantify Capture spatial/temporal variability; obtai km x 8 km (threshold) the underlying emissions, understand emission petter yields of products. tropospheric Aerosol processes, and track transport and chemical km x 8 km (baseline, threshold) properties composition and evolution of air pollutants 🚺 2, 3, 4, 5, 6) Over oper Inherently larger spatial scales, sufficient 6 km x 16 km (baseline only) air quality over B. Measure AOD, AAOD, and NH3 to quantify ocean to link to LEO observations scales ranging aerosol and nitrogen deposition to land and coastal regions [2, 4] Spectral region : 🛛 to H Typical use from urban to JV-Vis or UV-TIR 03 Provide multispectral retrieval information continental, Measure AOD AAOD and AOCH to relate SWIR. MWIR co in daylight diurnally to surface PM concentration, UV-B level and SO2 HCHO IV/ seasonally? visibility to aerosol column loading 🚺 2, 3, 4, 5 Retrieve gas species from their SWIR CH4 atmospheric spectral signatures (typical) **FIR** NH3 How does air Determine the instantaneous radiative forcings Obtain spectral-dependence of AOD for associated with ozone and aerosols on the pollution drive AOD, NO2, CHOCHO continental scale and relate them quantitative particle size and type information climate forcing to natural and anthropogenic emissions [3 5 6] Obtain spectral-dependence of AAOD for V-deep blue and how does aerosol type information G Observe pulses of CH4 emission from biogenic climate change JV-deep blue AL Provide absorbing aerosol information and anthropogenic releases; CO anthropogenic and wildfire emissions; AOD, AAOD, and AI from affect air quality /is-NIR AOCH Retrieve aerosol height 3 on a continental fires; AOD, AAOD, and AI from dust storms; SO2 and AOD from volcanic eruptions [1, 4, 6] Atmospheric measurements over Land/Coastal areas, baseline and threshold: 🛛 to K scale? Quantify the inflows and outflows of O3. CO. Time Typical pecies resolution SO2, and aerosols across continental boundaries 4. How can value to determine their impacts on surface air quality observations from two pieces of and on climate 2.3 sphere with space improve air 23 S74<70 2 km for surface Characterize aerosol particle size and type from quality forecasts spectral dependence measurements of AOD and AAOD [1, 2, 3, 4, 5, 6] forcing and assessments mass for societal 00 Acquire measurements to improve tical with benefit? representation of processes in air quality models daylight and improve data assimilation in forecast and irces AOD 5 How does assessment models [4] intercontinental Synthesize the GEO-CAPE measurements with nced/ 102 transport affect air information from in-situ and ground-based istry quality? remote sensing networks to construct an Additi K ements over Land/Coastal areas, baseline only: 🗖 enhanced observing system 1. 2. 3. 4. pecie 6. How do episodic Leverage GEO-CAPE observations into an integrated observing system including events, such as −сно[≠] geostationary satellites over Europe and Asia pected to peak at midday; try wild fires, dust together with LEO satellites and suborbital 502* outbreaks, and ms for assessing the hemispheric transport 1 2 3 4 5 61 volcanic eruptions, affect atmospheric Integrate observations from GEO-CAPE and composition and other platforms into models to improve H3 representation of processes in the models and to air quality? link the observed composition, deposition, and oso носн radiative forcing to the emissions from nemistry anthropogenic and natural sources [1, 2, 3, 4, 5, dust from nonsols: climate forcing Is near/above clouds and rly, SZA<70 snow/ice: aerosol events Determine plume height: large scale Hourly, SZA<70 Variable 1 km ransport, conversions from AOD to PM ean measurements: F.H.I.J.K baseline only, 16 km x 16 km 1/day Over open oceans, capture long-range transport of pollution, dust, and smoke into/out of North America 1/dav establish boundary conditions for North America AOD, AAOD, AI 1/dav AOD=Aerosol optical depth, AAOD=Aerosol at th, AI=Aerosol index. See next page for footnotes.

TEMPO	Atmosph	Atmospheric measurements over Land/Coastal areas, baseline and threshold: [A to K]							
	Species	Time resolution	Typical value ²		Precision ²		Description		
	03	Hourly, SZA<70	9 x10 ¹⁸		2km-tropopause: 15 ppbv Stratosphere: 5% 0-2 km: 20ppbv 2km-tropopause: 20 ppbv		Observe O3 with two pieces of information in the troposphere with sensitivity to the lowest 2 km for surface AQ; also transport, climate forcing		
Infrared species	co	Hourly, day and 2 x10 ¹⁸ night		18			Track anthropogenic and biomass burning plumes; observe CO with two pieces of information in the vertical with sensitivity to the lowest 2 km in daylight		
	AOD	Hourly, SZA<70	0.1 –	1	0.05		Observe total aerosol; aerosol sources and transport; climate forcing		
	NO2	Hourly, SZA<70 6 x10 ¹⁵		15	1×10 ¹⁵		Distinguish background from enhanced/ polluted scenes; atmospheric chemistry		
	Addition	al atmospheri	c mea	suren	nents	over Land/	Coastal areas, baseline only: <mark>A to K</mark>		
	Species	Time resolution		Typic value	al 2	Precision ²	Description		
Ultraviolet/	нсно*	3/day, SZA	A<50 1×10 4 ×10		0 ¹⁶	1×10 ¹⁶	Observe biogenic VOC emissions, expected to peak at midday; chemistry		
visible	SO2*	3/day, SZA			16	1×10 ¹⁶	Identify major pollution and volcanic emissions; atmospheric chemistry		
	СН4	2/day			(10 ¹⁹ 20 ppbv		Observe anthropogenic and natural emissions sources		
species	NH3	2/day			2x10 ¹		6	0-2 km: 2ppbv	Observe agricultural emissions
(GOME,	сносно)* 2/day		2x10 ¹		2x10 ¹⁴		4×10 ¹⁴	Detect VOC emissions, aerosol formation, atmospheric chemistry
TEMPO,	AAOD	Hourly, SZ	A<70	0 – 0.05		0 – 0.05		0.02	Distinguish smoke and dust from non- UV absorbing aerosols; climate forcing
etc.)	AI	Hourly, SZ	A<70	-1 – +	-5	0.1	Detect aerosols near/above clouds and over snow/ice; aerosol events		
7/12/16	АОСН	Hourly, SZ	A<70	Varia	ble	1 km	Determine plume height; large scale transport, conversions from AOD to PM		

Typical TEMPO-range spectra (from ESA GOME-1)

NASA



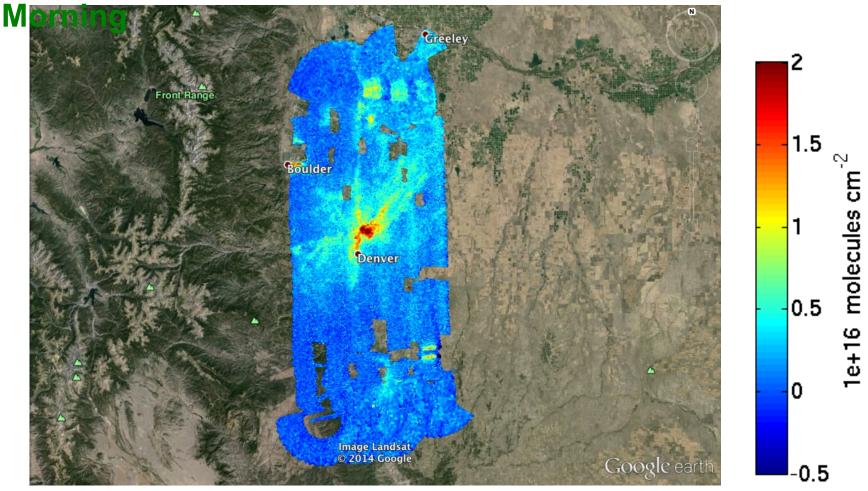


A full, minimally-redundant, set of polluting gases, plus aerosols and clouds is now measured to very high precision from satellites. Ultraviolet and visible spectroscopy of backscattered radiation provides O₃ (including profiles and tropospheric O_3), NO_2 (for NO_x), H_2CO and $C_2H_2O_2$ (for VOCs), SO_2 , H₂O, O₂-O₂, N₂ and O₂ Raman scattering, and halogen oxides (BrO, CIO, IO, OCIO). Satellite spectrometers we planned since 1985 began making these measurements in 1995.





GeoTASO NO₂ Slant Column, 02 August 2014



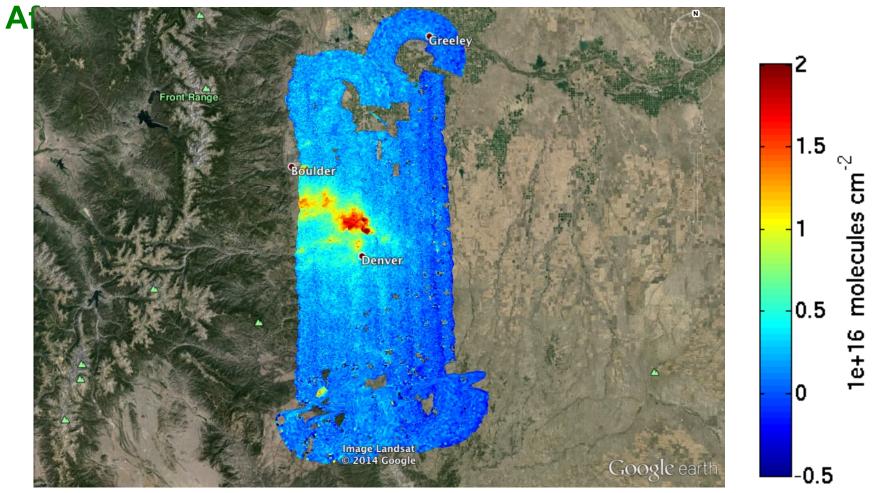
Co-added to approx. $500m \times 450m$

Morning vs. Afternoon

Preliminary data, C. Nowlan, SAO

TEMPO TEMPO measurements will capture the diurnal cycle of pollutant emissions

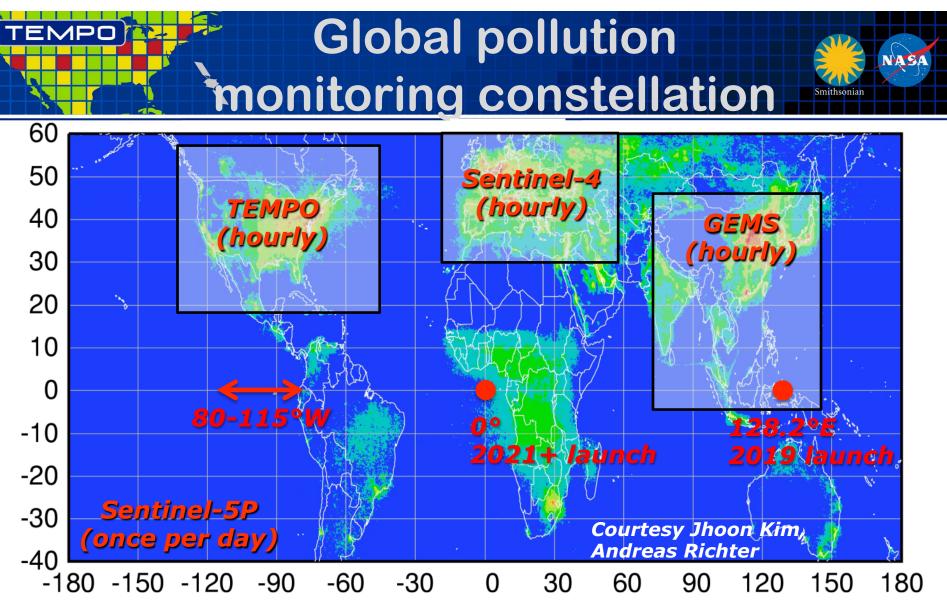
GeoTASO NO₂ Slant Column, 02 August 2014



Co-added to approx. $500m \times 450m$

Morning vs. Afternoon

Preliminary data, C. Nowlan, SAO



Policy-relevant science and environmental services enabled by common observations

• Improved emissions, at common confidence levels, over industrialized Northern Hemisphere

- · Improved air quality forecasts and assimilation systems
- Improved assessment, e.g., observations to support United Nations Convention on Long Range Transboundary Air Pollution



NO₂, SO₂, H₂CO, C₂H₂O₂ vertical columns

Direct fitting to TEMPO radiances

AMF-corrected reference spectra, Ring effect, etc.

DOAS option available to trade more speed for less accuracy, if necessary Research products could include H₂O, BrO, OCIO, IO

O₃ profiles, tropospheric O₃

eXceL optimal-estimation method developed @ SAO for GOME, OMI May be extended to SO₂, especially volcanic SO₂

TOMS-type ozone retrieval included for heritage

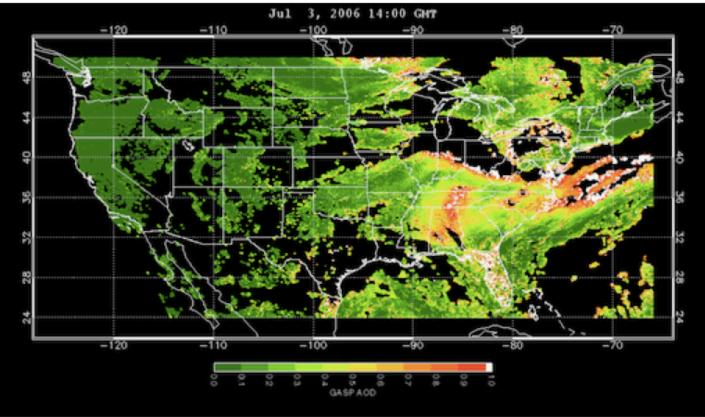
Aerosol products from OMI heritage: AOD, AAOD, Aerosol Index Advanced/improved products likely developed @ GSFC, U. Nebraska Cloud Products from OMI heritage: CF, CTP Advanced/improved products likely developed @ GSFC

UVB research product based on OMI heritage (FMI, GSFC)

Nighttime research products include city lights



TEMPO will use the EPA's Remote Sensing Information Gateway (RSIG) for subsetting, visualization, and product distribution – to make TEMPO YOUR instrument



See J. Szykman, tomorrow



What is an AQ index?"

"The Canadian Air Quality Health Index is a multipollutant index based on the sum of PM2.5, NO_2 , and O_3 , weighted by their contribution to mortality in daily time-series study across Canadian cities." [Cooper et al., 2012]

Cooper et al., for example, propose a satellite-based multipollutant index using the WHO Air Quality Guidelines (AQG):

$$SATMPI = \frac{PM_{2.5}}{AQG_{PM2.5}} \left[1 + \frac{NO_2}{AQG_{NO_2}} \right]$$

- Can we define different indices as appropriate to locations, seasons, times?
- Might they be formulated using RSIG?
- Might assimilation be included?

Cooper, M., R.V. Martin, A. van Donkelaar, L. Lamsal, M. Brauer, and J. Brook, A satellite-based multi-pollutant index of global air quality, *Env. Sci. and Tech.*, **46**, 8523-8524, 2012. 7/12/16



Volcanic **SO**₂ (column amount and plume altitude is a potential research product. Diurnal out-going **shortwave radiation and cloud forcing** is a potential research product.

Nighttime "**city lights**" products, which represent anthropogenic activities at the same spatial resolution as air quality products, may be produced twice per day (late evening and early morning) as a research product. Meeting TEMPO measurement requirements for NO₂ (visible) implies the sensitivity for city lights products over the CONUS within a 2-hour period at 2×4.5 km² to 1.1×10^{-8} W cm⁻² sr⁻¹ µm⁻¹.

Several additional first-measurement molecules are being studied.

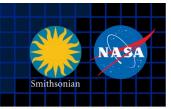
 H_2O will be produced at launch from the 7v vibrational polyad at 445 nm. Water vapor retrieved from the visible spectrum has good sensitivity to the planetary boundary layer, since the absorption is optically thin, and is available over both the land and ocean. The hourly coverage of TEMPO will greatly improve the knowledge of water vapor's diurnal cycle and make rapid variations in time readily observed.



Morning and evening higher-frequency scans The optimized data collection scan pattern during mornings and evenings provides multiple advantages for addressing TEMPO science questions. The increased frequency of scans coincides with peaks in vehicle miles traveled on each coast.

Biomass burning The unexplained variability in ozone production from fires is of particular interest. The suite of NO_2 , H_2CO , $C_2H_2O_2$, O_3 , and aerosol measurements from TEMPO is well suited to investigating how the chemical processing of primary fire emissions effects the secondary formation of VOCs and ozone. For particularly important fires it is possible to command special TEMPO observations at even shorter than hourly revisit time, probably as short as 10 minutes.

NO_x studies



Lightning NO_x Interpretation of satellite measurements of tropospheric NO₂ and O₃, and upper tropospheric HNO₃ lead to an overall estimate of 6 ± 2 Tg N y-1 from lightning [Martin et al., 2007]. TEMPO measurements, including tropospheric NO₂ and O₃, can be made for time periods and longitudinal bands selected to coincide with large thunderstorm activity, including outflow regions, with fairly short notice.

Soil NO_x Jaeglé et al. [2005] estimate 2.5 - 4.5 TgN y⁻¹ are emitted globally from nitrogen-fertilized soils, still highly uncertain. The US a posteriori estimate for 2000 is 0.86 ± 1.7 TgN y⁻¹. For Central America it is 1.5 ± 1.6 TgN y⁻¹. They note an underestimate of NO release by nitrogen-fertilized croplands as well as an underestimate of rain-induced emissions from semiarid soils.

TEMPO is able to follow the temporal evolution of emissions from croplands after fertilizer application and from rain-induced emissions from semi-arid soils. Higher than hourly time resolution over selected regions may be accomplished by special observations. Improved constraints on soil NO_x emissions may also improve estimated of lightning NO_x emissions [Martin *et al.* 2000].



Halogens



BrO will be produced at launch, assuming stratospheric AMFs. Scientific studies will correct retrievals for tropospheric content. **IO** was first measured from by SAO space using SCIAMACHY spectra [Saiz-Lopez *et al.*, 2007]. It will be produced as a scientific product, particularly for coastal studies, assuming AMFs appropriate to lower tropospheric loading.

The atmospheric chemistry of halogen oxides over the ocean, and in particular in coastal regions, can play important roles in ozone destruction, oxidizing capacity, and dimethylsulfide oxidation to form cloud-condensation nuclei [Saiz-Lopez and von Glasow, 2012]. The budgets and distribution of reactive halogens along the coastal areas of North America are poorly known. Therefore, providing a measure of the budgets and diurnal evolution of coastal halogen oxides is necessary to understand their role in atmospheric photochemistry of coastal regions. Previous ground-based observations have shown enhanced levels (at a few pptv) of halogen oxides over coastal locations with respect to their background concentrations over the remote marine boundary layer [Simpson et al., 2015]. Previous global satellite instruments lacked the sensitivity and spatial resolution to detect the presence of active halogen chemistry over mid-latitude coastal areas. TEMPO observations together with atmospheric models will allow examination of the processes linking ocean halogen emissions and their potential impact on the oxidizing capacity of coastal environments of North America.

TEMPO also performs hourly measurements one of the world's largest salt lakes: the Great Salt Lake in Utah. Measurements over Salt Lake City show the highest concentrations of BrO over the globe. Hourly measurement at a high spatial resolution can improve understanding of BrO production in salt lakes. $\frac{7/12}{16}$

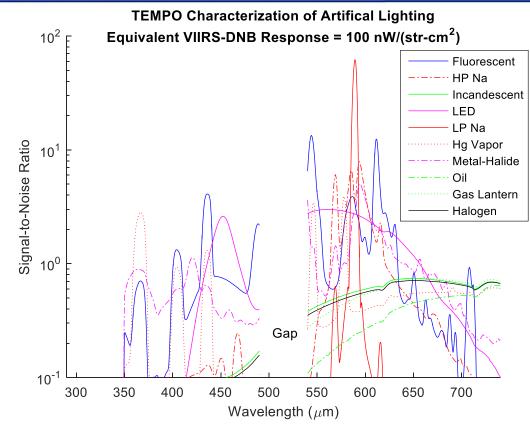
Spectral indicators

Fluorescence and other spectral indicators Solar-induced fluorescence (SIF) from chlorophyll over both land and ocean will be measured. In terrestrial vegetation, chlorophyll fluorescence is emitted at red to far-red wavelengths (~650-800 nm) with two broad peaks near 685 and 740 nm, known as the red and far-red emission features. Oceanic SIF is emitted exclusively in the red feature. SIF measurements have been used for studies of tropical dynamics, primary productivity, the length of carbon uptake period, and drought responses, while ocean measurements have been used to detect red tides and to conduct studies on the physiology, phenology, and productivity of phytoplankton. TEMPO can retrieve both red and far-red SIF by utilizing the property that SIF fills in solar Fraunhofer and atmospheric absorption lines in backscattered spectra normalized by a reference (*e.g.*, the solar spectrum) that does not contain SIF.

TEMPO will also be capable of measuring **spectral indices developed for estimating foliage pigment contents and concentrations**. Spectral approaches for estimating pigment contents apply generally to leaves and not the full canopy. A single spectrally invariant parameter, the Directional Area Scattering Factor (DASF), relates canopy-measured spectral indices to pigment concentrations at the leaf scale.

UVB TEMPO measurements of daily UV exposures build upon heritage from OMI and TROPOMI measurements. Hourly cloud measurements from TEMPO allow taking into account diurnal cloud variability, which has not been previously possible. The OMI UV algorithm is based on the TOMS UV algorithm. The specific product is the downward spectral irradiance at the ground (in W m⁻² nm⁻¹) and the erythemally weighted irradiance (in W m⁻²).

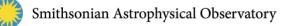




SNRs of various artificial lighting types normalized to a common VIIRS-DNB response with a 10s dwell, no co-addition, and no spatial binning, assuming a dark current of 2800 e⁻ s⁻¹. City lights over all of greater North America can either be observed piecemeal over several days or in a single scan near the winter solstice.









Default Launch Data Products

Product	Algorithm	Hourly Coverage @ ≤ 4.5×8 km²
O ₃	TOMS-Vn	15 - 50.25°N, 60 - 130°W
O ₃	XL optimal estimation	Selected urban areas and burning regions
NO ₂	Direct fitting, AMF (λ)	15 - 50.25°N, 60 - 130°W
SO ₂	Direct fitting, AMF (λ)	15 - 50.25°N, 60 - 130°W
H ₂ CO	Direct fitting, AMF (λ)	15 - 50.25°N, 60 - 130°W
$C_2H_2O_2$	Direct fitting, AMF (λ)	15 - 50.25°N, 60 - 130°W
Aerosol OD and SSA	AERUV	15 - 50.25°N, 60 - 130°W
Cloud pressure and fraction	CLDRR	15 - 50.25°N, 60 - 130°W
UBV and Eryth. dose	UVB	15 - 50.25°N, 60 - 130°W
AQ indices	L3-L4 based	15 - 50.25°N, 60 - 130°W



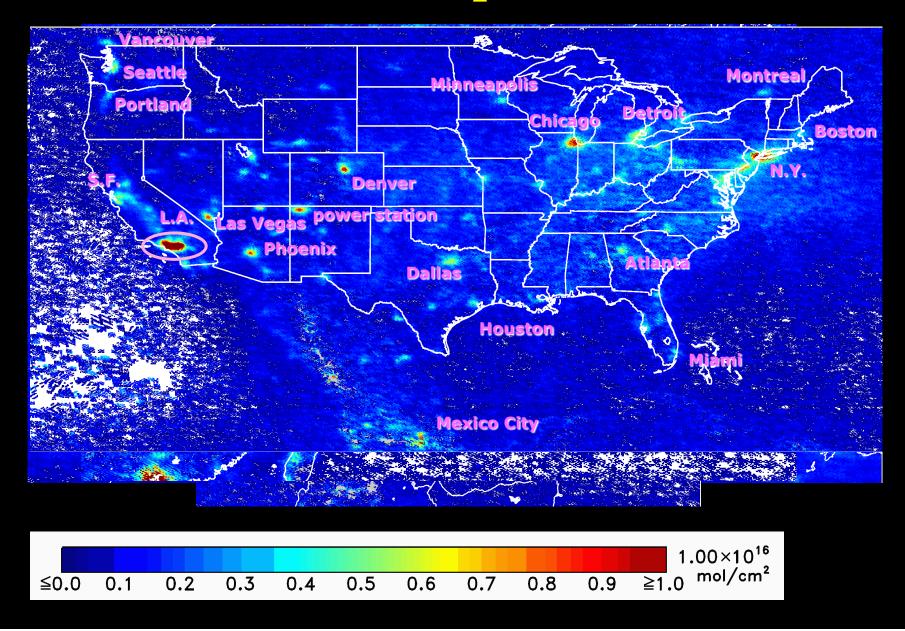


Secondary and Improved Data Products

Product	Algorithm	Hourly Coverage @ ≤ 4.5×8 km²
O ₃	XL optimal estimation	15 - 50.25°N, 60 - 130°W (or, extended regions)
BrO	Direct fitting, AMF (λ)	15 - 50.25°N, 60 - 130°W
H ₂ O	Direct fitting, AMF (λ)	15 - 50.25°N, 60 - 130°W
Aerosols	AERUV+	15 - 50.25°N, 60 - 130°W
Clouds	CLDRR+	15 - 50.25°N, 60 - 130°W
SO ₂	Height-resolved	15 - 50.25°N, 60 - 130°W

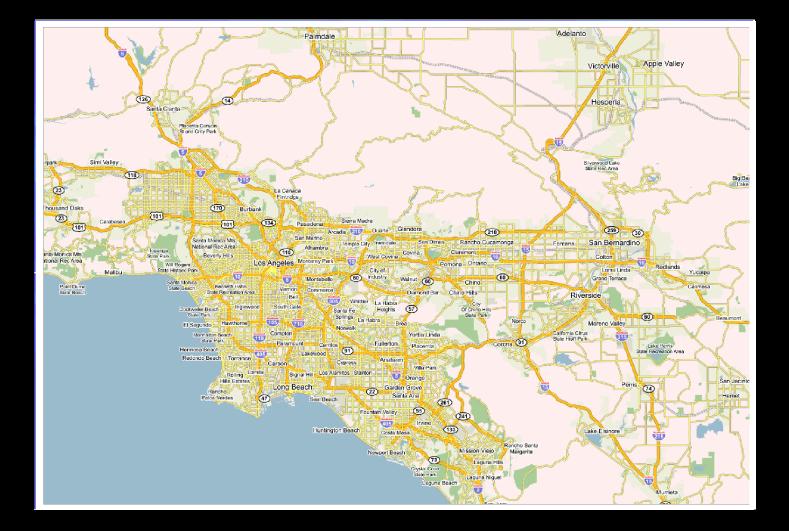


 NO_2





NO₂ - Los Angeles





A day in the life

NASA

Smithsonian

 AM
 PM
 PM
 AM
 AM

 6:00
 7:00
 8:00
 9:00
 10:00
 11:00
 12:00
 1:00
 2:00
 3:00
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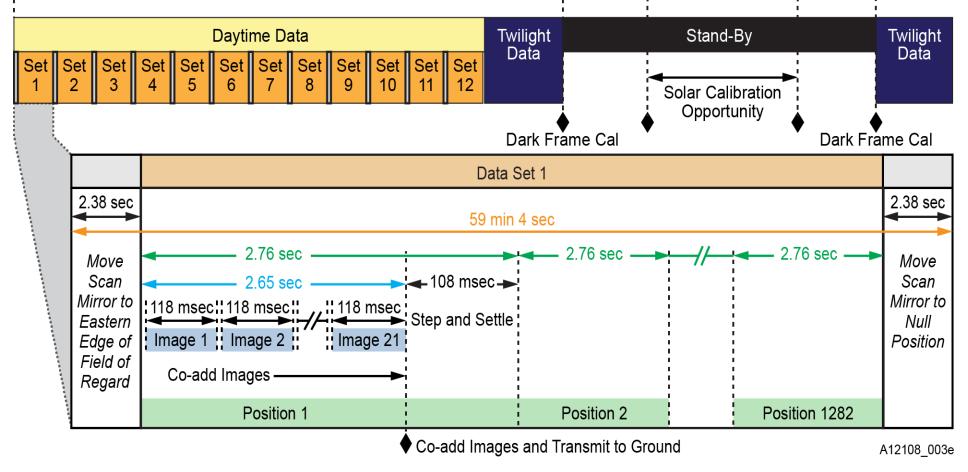


Figure 7. Nominal daily operations for TEMPO instrument.

Air quality and health to subsolin

TEMPO's hourly measurements allow better understanding of the complex chemistry and dynamics that drive air quality on short timescales. The density of TEMPO data is ideally suited for data assimilation into chemical models for both air quality forecasting and for better constraints on emissions that lead to air quality exceedances. Planning is underway to combine TEMPO with regional air quality models to **improve EPA air quality indices** and to directly supply the public with near real time pollution reports and forecasts through website and mobile **applications**. As a case study, an OSSE for the Intermountain West was performed to explore the potential of geostationary ozone measurements from TEMPO to improve monitoring of ozone exceedances and the role of background ozone in causing these exceedances (Zoogman et al. 2014).



Clouds The launch cloud algorithm is be based on the rotational Raman scattering (RRS) cloud algorithm that was developed for OMI by GSFC. Retrieved cloud pressures from OMCLDRR are not at the geometrical center of the cloud, but rather at the optical centroid pressure (OCP) of the cloud. **Additional** cloud products are possible using the O_2 - O_2 collision complex and/ or the $O_2 B$ band.

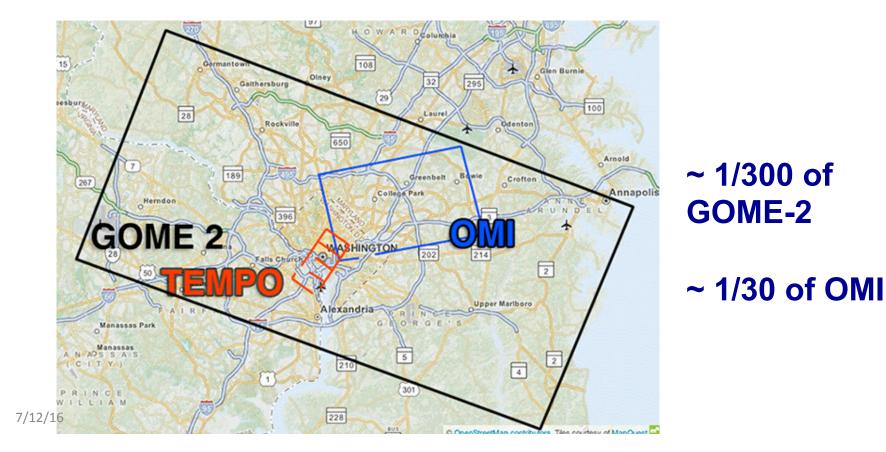
Aerosols TEMPO's launch algorithm for retrieving aerosols will be based upon the OMI aerosol algorithm that uses the sensitivity of near-UV observations to particle absorption to retrieve Absorbing Aerosol Index (AAI), aerosol optical depth (AOD) and single scattering albedo (SSA). TEMPO may be used together with the advanced baseline imager (ABI) instruments on the NOAA GOES-R and GOES-S satellites for aerosol retrievals, reducing AOD and fine mode AOD uncertainties from 30% to 10% and from 40% to 20%.

Science studies

NA SA

mithsonian

- Spatial resolution: allow tracking pollution at sub-urban scale
 - GEO at 100°W: 2.1 km N/S × 4.7 km E/W = 9.8 km² (native) at center of FOR (36.5°N, 100°W)
 - Full resolution for NO₂, HCHO, total O₃ products
 - Co-add 4 N/S pixels for O_3 profile product: 8.4 km N/S × 4.7 km E/W



TEMPO

TEMPO Science Team, U.S.

NASA

Team Member	Institution	Role	Responsibility		
K. Chance	SAO	PI	Overall science development; Level 1b, H ₂ CO, C ₂ H ₂ O ₂		
X. Liu	SAO	Deputy PI	Science development, data processing; O_3 profile, tropospheric O_3		
J. Al-Saadi	LaRC	Deputy PS	Project science development		
J. Carr	Carr Astronautics	Co-I	INR Modeling and algorithm		
M. Chin	GSFC	Co-I	Aerosol science		
R. Cohen	U.C. Berkeley	Co-I	NO ₂ validation, atmospheric chemistry modeling, process studies		
D. Edwards	NCAR	Co-I	VOC science, synergy with carbon monoxide measurements		
J. Fishman	St. Louis U.	Co-I	AQ impact on agriculture and the biosphere		
D. Flittner	LaRC	Project Scientist	Overall project development; STM; instrument cal./char.		
J. Herman	UMBC	Co-I	Validation (PANDORA measurements)		
D. Jacob	Harvard	Co-I	Science requirements, atmospheric modeling, process studies		
S. Janz	GSFC	Co-I	Instrument calibration and characterization		
J. Joiner	GSFC	Co-I	Cloud, total O ₃ , TOA shortwave flux research product		
N. Krotkov	GSFC	Co-I	NO ₂ , SO ₂ , UVB		
M. Newchurch	U. Alabama Huntsville	Co-I	Validation (O ₃ sondes, O ₃ lidar)		
R.B. Pierce	NOAA/NESDIS	Co-I	AQ modeling, data assimilation		
R. Spurr	RT Solutions, Inc.	Co-I	Radiative transfer modeling for algorithm development		
R. Suleiman	SAO	Co-I, Data Mgr.	Managing science data processing, BrO, H ₂ O, and L3 products		
J. Szykman	EPA	Co-I	AIRNow AQI development, validation (PANDORA measurements)		
O. Torres	GSFC	Co-I	UV aerosol product, Al		
J. Wang	U. Nebraska	Co-I	Synergy w/GOES-R ABI, aerosol research products		
J. Leitch 7/12/16	Ball Aerospace	Collaborator	Aircraft validation, instrument calibration and characterization		
D. Neil	LaRC	Collaborator	GEO-CAPE mission design team member		

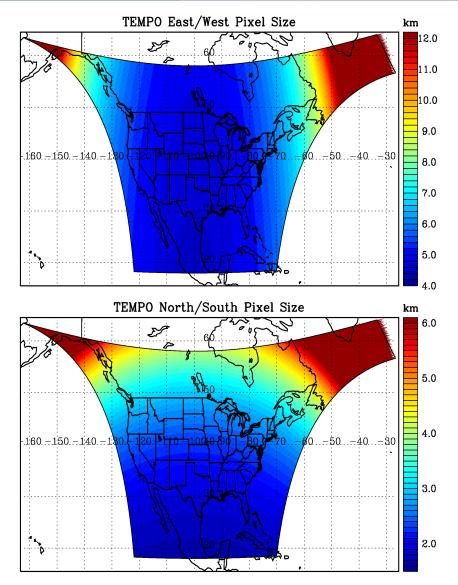
TEMPO Science Team, non-U.S.

NASA

Team Member	Institution	Role	Responsibility
Randall Martin	Dalhousie U.	Collaborator	Atmospheric modeling, air mass factors, AQI development
Chris McLinden	Environment Canada	Collaborator	Canadian air quality coordination
Michel Grutter de la Mora	UNAM, Mexico	Collaborator	Mexican air quality coordination
Gabriel Vazquez	UNAM, Mexico	Collaborator	Mexican air quality, algorithm physics
Amparo Martinez	INECC, Mexico	Collaborator	Mexican environmental pollution and health
J. Victor Hugo Paramo Figeuroa	INECC, Mexico	Collaborator	Mexican environmental pollution and health
Brian Kerridge	Rutherford Appleton Laboratory, UK	Collaborator	Ozone profiling studies, algorithm development
Paul Palmer	Edinburgh U., UK	Collaborator	Atmospheric modeling, process studies
Alfonso Saiz-Lopez	CSIC, Spain	Collaborator	Atmospheric modeling, process studies
Juan Carlos Antuña Marrero	GOAC, Cuba	Collaborator	Cuban Science team lead, Cuban air quality
Osvaldo Cuesta	GOAC, Cuba	Collaborator	TEMPO validation, Cuban air quality
René Estevan Arredondo	GOAC, Cuba	Collaborator	TEMPO validation, Cuban air quality
J. Kim	Yonsei U.		Korean GEMS, CEOS constellation of GEO pollution monitoring
C.T. McElroy			CSA PHEOS, CEOS constellation of GEO pollution monitoring
B. Veihelmann			ESA Sentinel-4, CEOS constellation of GEO pollution monitoring
J.P. Veefkind	KNMI		ESA Sentinel-5P (TROPOMI)

EMPO

TEMPO footprint (GEO at 100° W)



Location	N/S (km)	E/W (km)	GSA (km²)
36.5°N, 100°W	2.11	4.65	9.8
Washington, DC	2.37	5.36	11.9
Seattle	2.99	5.46	14.9
Los Angeles	2.09	5.04	10.2
Boston	2.71	5.90	14.1
Miami	1.83	5.04	9.0
Mexico City	1.65	4.54	7.5
Canadian oil sands	3.94	5.05	19.2

NASA

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Assumes 2000 N/S pixels

For GEO at 80°W, pixel size at 36.5°N, 100°W is 2.2 km × 5.2 km.

Low Earth orbit:

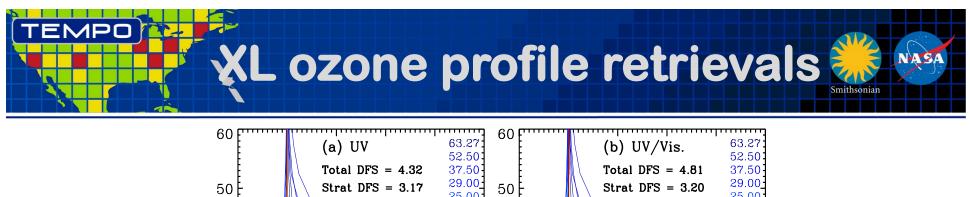
Sun-synchronous nadir heritage

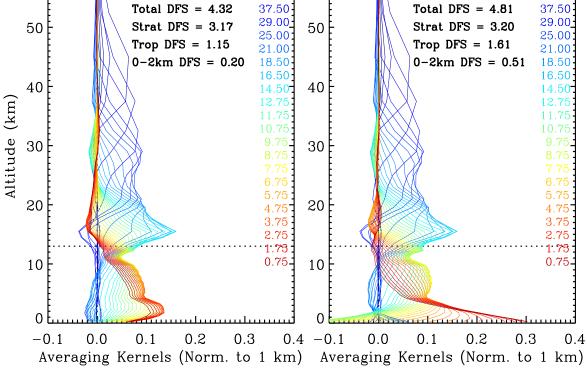
NA SA

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Instrument	Detectors	Spectral Coverage [nm]	Spectral Res. [nm]	Ground Pixel Size [km ²]	Global Coverage
GOME-1 (1995-2011)	Linear Arrays	240-790	0.2-0.4	40×320 (40×80 zoom)	3 days
SCIAMACHY (2002-2012)	Linear Arrays	240-2380	0.2-1.5	30×30/60/90 30×120/240	6 days
OMI (2004)	2-D CCD	270-500	0.42-0.63	13×24 - 42×162	daily
GOME-2a,b (2006, 2012)	Linear Arrays	240-790	0.24-0.53	40×80 (40×10 zoom)	near-daily
OMPS-1 (2011)	2-D CCDs	250-380	0.42-1.0	50×50	daily

Previous experience (since 1985 at SAO and MPI) Scientific and operational measurements of pollutants O₃, NO₂, SO₂, H₂CO, C₂H₂O₂ ^{7/12/16}(& CO, CH₄, BrO, OCIO, CIO, IO, H₂O, O₂-O₂, Raman, aerosol,)





Retrieval averaging kernels based on iterative nonlinear retrievals from synthetic TEMPO radiances with the signal to noise ratio (SNR) estimated using the TEMPO SNR model at instrument critical design review in June 2015 for (a) UV (290-345 nm) retrievals and (b) UV/ Visible (290-345 nm, 540-650 nm) retrievals for clear-sky condition and vegetation surface with solar zenith angle 25°, viewing zenith angle 45° and relative azimuthal angle 86°. DFS is degrees of freedom for signal, the trace of the averaging kernel matrix, which is an indicator of the averaging kernel matrix, which is an indicator of pieces of independent information in the solution.

- 1. What are the temporal and spatial variations of **emissions** of gases and aerosols important for air quality and climate?
- 2. How do physical, chemical, and dynamical **processes** determine tropospheric composition and air quality over scales ranging from urban to continental, diurnally to seasonally?
- 3. How does air pollution drive **climate** forcing and how does climate change affect air quality on a continental scale?
- 4. How can observations from space improve **air quality forecasts and assessments** for societal benefit?
- 5. How does **intercontinental transport** affect air quality?
- 6. How do **episodic events**, such as wild fires, dust outbreaks, and volcanic eruptions, affect atmospheric composition and air quality?



Table D.2-3 TEMPO STM¹ clearly links science questions with instrument and investigaton requirements.

				_			
Science Objective	Science Measurement	t Requirement		Instrument Function Requirements			Investigation Requirements
	Observables	Physical Pa	rameters	Parameter	Req.	Predicted	
What are the -High temporal resolution		Relevant absorption bands		Spectral Range	290-690 nm	290-690 nm	
measurements to capture changes				Spectral Resolution	0.6 nm	0.6 nm	1-year mission
in pollutant gas distributions.		for aero	sols	Spectral Sampling	0.2 nm	0.2 nm	lifetime (minimum)
- High spatial resolution measurements that sense urban	suitable for retrievals of O ₃ , NO ₂ , H ₂ CO, SO ₂ and C ₂ H ₂ O ₂ at spatial	Baseline Trac	e gas column	• •	s noted, hourly (@ 8×4.5 km ²	On-orbit Calibration
scale pollutant gases across GNA		Species	Precision	Band	Signal to No	oise (hourly)	
and surrounding areas.	atmospheric chemistry models.			O::Vis (546-648 nm)	958	1254	FOR
Magaurament of major elements							encompasses
	Multispectral data in suitable O ₃					1000	CONUS and
including multispectral	absorption bands to provide vertical			423-451 nm	1233	1910	adjacent areas
measurements to improve sensing	distribution information.						
		SO ₂ # (3/day)	10.0	305-345 nm	1297	1820	GEO Longitude:
	Spectral radiance measurements with	C ₂ H ₂ O ₂ # (2/day)	0.40	433-457 nm	1350	2331	Preferred: 100W Acceptable:
	suitable quality (SNR) to provide Baseline Aerosol/Cloud properties			sol/Cloud properties hourly	urly @ 8×4.5 km²		75W – 137W
		Property	Precision	Band	Signal t	to Noise	GEO Bus Pointing:
spatial resolution for quantifying		AOD#	0.05		1000		Control <0.1°
			0.03	354, 388 nm		1596	Knowledge <0.04°
loading.							
Determine the instantaneous	reflectance spectrum for solar zenith			346-354 nm	600	1608	Provide near-real-
	angles <70°.	CTP#	100 mb				time products to user communities
O_3 and aerosols on the continental		Solar irradianc	e snectrally			within 2 hrs to	
scale.							enable assimilation
				λ-independent	< 3%	2.0%	into chemical
- Integrate observations from TEMPO and other platforms into				Spectral Accuracy	<0.02 nm	<0.02 nm	models (NOAA & EPA) and use by
models to improve representation	No additional observable			Polarization Factor			smart-phone
	requirements	requirem	nents	Coolection Accuracy			applications
ayatem.							Archive and
- Quantify the flow of pollutants							distribute TEMPO science data
across continental boundaries;		No additional physical		GSD E/W*	4.0 km		products
Join a global observing system.	requirements require		Tents	MTF: N/S×E/W	0.3×0.3	0.50×0.46	
	 High temporal resolution measurements to capture changes in pollutant gas distributions. High spatial resolution measurements that sense urban scale pollutant gases across GNA and surrounding areas. Measurement of major elements in tropospheric O₃ chemistry cycle, including multispectral measurements to improve sensing of lower-tropospheric O₃, with precision to clearly distinguish pollutants from background levels Observe aerosol optical properties with high temporal and spatial resolution for quantifying and tracking evolution of aerosol loading. Determine the instantaneous radiative forcings associated with O₃ and aerosols on the continental scale. Integrate observations from TEMPO and other platforms into models to improve representation of processes in the models and construct an enhanced observing system. Quantify the flow of pollutants across continental boundaries; 	Science Ubjective Observables -High temporal resolution measurements to capture changes in pollutant gas distributions. Spatially imaged & spectrally resolved, solar backscattered earth radiance, spanning spectral windows suitable for retrievals of O ₃ , NO ₂ , H ₂ CO, SO ₂ and C ₂ H ₂ O ₂ at spatial scales comparable to regional atmospheric O ₃ chemistry cycle, including multispectral measurements to improve sensing of lower-tropospheric O ₃ , with precision to clearly distinguish pollutants from background levels Multispectral data in suitable O ₃ absorption bands to provide vertical distribution information. - Observe aerosol optical properties with high temporal and spatial resolution for quartifying and tracking evolution of aerosol loading. Spetially imaged, wavelength dependence of atmospheric reflectance spectrum for solar zenith angles <70°.	ObservablesPhysical Pa-High temporal resolution measurements to capture changes in pollutant gas distributions.Spatially imaged & spectrally resolved, solar backscattered earth radiance, spanning spectral windows suitable for retrievals of O ₅ , NO ₂ , H ₂ CO, SO ₂ and C ₂ H ₂ O ₂ at spatial scales comparable to regional atmospheric O ₅ chemistry cycle, including multispectral measurements to improve sensing of lower-tropospheric O ₃ , with prolutants from background levelsMultispectral data in suitable O ₃ absorption bands to provide vertical distribution information.Baseline Trac O ₃ :0-2 km- Measurements to improve sensing of lower-tropospheric O ₃ , with prolutants from background levelsMultispectral data in suitable O ₃ absorption bands to provide vertical distribution information.O ₃ :* Total NO ₂ # H ₂ CO* (3/day)- Observe aerosol optical properties with high temporal and spatial resolution of quantifying and tracking evolution of aerosol loading.Spectral radiance measurements with suitable quality (SNR) to provide multiple measurements over daylight hours for solar zenith angles <70°.	Science Objective Observables Physical Parameters I-High temporal resolution measurements to capture changes in pollutant gas distributions. Spatially imaged & spectrally resolved, solar backscattered earth radiance, spanning spectral windows suitable for retrievals of O ₂ , NO ₂ , H ₂ CO, SO ₂ and C ₂ H ₂ O ₂ at spatial scales comparable to regional atmospheric C ₂ , themistry models. Relevant absorption bands for trace gase & windows for aerosols - Measurement of major elements in tropospheric O ₂ themistry cycle, including multispectral measurements to improve sensing of lower-tropospheric O ₂ , with pollutants from background levels Multispectral data in suitable O ₂ absorption bands to provide vertical distribution information. Baseline Trace gas column - Observe aerosol optical properties with high temporal and spatial resolution for quantifying and tracking evolution of aerosol loading. Multispectral radiance measurements wird suitable Q ₂ at spatial resolution for quantifying and tracking evolution of aerosol loading. Spectral radiance measurements were daylight hours for solar zenith angle < 70°.	Science Ubjective Observables Physical Parameters Parameter High temporal resolution measurements to capture changes in pollutant gas distributions. - High spatial resolution measurements that sense urban scale pollutant gases across GNA and surounding areas. - Measurement of major elements in tropospheric O ₂ , termistry cycle, including multispectral measurements to inprove sensing of lower-tropospheric O ₂ , with progerise with high temporal and spatial resolution for quantifying and tracking evolution of aerosol bading. Multispectral data in suitable O ₂ absorption bands to provide vertical distribution information. Relevant absorption bands for aerosols Spectral Resolution Spectral Sampling Observe aerosol optical properties with high temporal and spatial resolution for quantifying and tracking evolution of aerosol 	Science Objective Observables Physical Parameters Parameter Req. -High temporal resolution measurements to capture changes in pollutant gase distributions. - High spatial resolution measurements that sense urban scale pollutant gases across GNA and surrounding multispectral and surrounding multispectral intropospheric O, chemistry cycle, including multispectral measurements to improve sensing of ower-tropospheric O, with precision to clearly distinguish pollutant from background levels Spectral Range 200-690 nm Spectral Range Spectral Range Spect	Science Ubjective Observables Physical Parameters Parameter Reg. Predicted -High temporal resolution measurements to capture change in pollutant gas distibutions. Spatially imaged & spectrally resolved, solar backscattered earth radiance, spanning spectral windows suitable for treievals of 0, No., the Q, solar distibution, measurements that sense uban cale pollutant gass across GNA and surrounding areas. Spatially imaged & spectral windows in topospheric 0, with intuding multispectral atmospheric c chemistry models. Relevant absorption bands for aerosols Spectral Range 290-690 nm 290-690 nm 290-690 nm 290-690 nm 290-690 nm 200-690 nm 200 nm 200-690 nm 200 nm 200-690 nm 200-690 nm 200 nm 200 nm 200-690 nm 200 nm 200 nm 200 nm 200 nm 200 nm 200 nm

¹FT=**Free Tropo**sphere (2km-tropopause), SOC=Stratospheric Ozone Column, AOD=Aerosol optical depth, AAOD=Aerosol absorption optical depth, AI=Aerosol index, CF=Cloud Fraction & CTP=Cloud Top Pressure, Albedo=Radiance/Irradiance, FOR=Field Of Regard, IFOV=Instantaneous Field Of View, GSD=Ground Sample Distance. *Projected to 36.5°N,100°W from GEO 100°W. # Threshold Products at 8×9km² and 80-minute intervals instead of hourly.