

Evidence for widespread hydrated minerals on asteroid (101955) Bennu

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Early spectral data from the Origins, Spectral Interpretation, Resource Identification, and Security-Regolith Explorer (OSIRIS-REx) mission reveal evidence for abundant hydrated minerals on the surface of near-Earth asteroid (101955) Bennu in the form of a near-infrared absorption near 2.7 μm and thermal infrared spectral features that are most similar to those of aqueously altered CM-type carbonaceous chondrites. We observe these spectral features across the surface of Bennu, and there is no evidence of substantial rotational variability at the spatial scales of tens to hundreds of metres observed to date. In the visible and near-infrared (0.4 to 2.4 μm) Bennu's spectrum appears featureless and with a blue (negative) slope, confirming previous ground-based observations. Bennu may represent a class of objects that could have brought volatiles and organic chemistry to Earth.

The Origins, Spectral Interpretation, Resource Identification, and Security-Regolith Explorer (OSIRIS-REx) mission began its Approach phase to asteroid (101955) Bennu in August 2018. Before and just after arrival at Bennu on 3 December, the OSIRIS-REx Visible and InfraRed Spectrometer (OVIRS) and Thermal Emission Spectrometer (OTES) collected hyperspectral data of this B-type asteroid, which is thought to be related to the carbonaceous chondrite meteorites¹. The OVIRS instrument² is a hyperspectral, point spectrometer that measures the reflected and emitted energy of Bennu across the spectral region from 0.4 to 4.3 μm (25,000 to 2,300 cm^{-1}) with a circular, 4 mrad field of view (FOV). The OTES instrument³, the first thermal infrared spectrometer to visit an asteroid, is a hyperspectral, point spectrometer that measures the emitted radiance of Bennu across the spectral region from ~1,750 to 100 cm^{-1} (~5.71 to 100 μm) with a circular, 8 mrad FOV. The primary role of visible-to-infrared spectroscopy on the OSIRIS-REx mission is to characterize the mineralogy and chemistry of Bennu and aid in sample site selection⁴. The OTES radiance data also are used in conjunction with thermophysical models to determine properties of the surface, such as particle size and roughness, and to study the Yarkovsky effect⁵. The mineralogy and chemistry of the surface of Bennu provide information about the geological processes that have affected the asteroid, the potential for resource extraction and the accuracy of telescopic spectral

observations (with the final ground truth coming from measurements of the returned sample).

On five days between 2 and 9 November 2018, both spectrometers obtained whole-disk (sub-FOV) spectra of Bennu for 4.5 h, which is just over one full rotation period (~4.3 h). In December 2018, both instruments collected spatially resolved spectra of Bennu as 'ride-along' observations during imaging activities optimized for the PolyCam and MapCam imagers⁶.

Visible and near-infrared spectral characteristics

The ground-based, composite (0.4 to 2.4 μm) reflectance spectrum of Bennu shows a spectrally 'blue' (negative) continuum slope across the visible and near-infrared, characteristic of B-type asteroids¹. Clark et al.¹ did not find strong spectral absorptions in the Bennu telescopic data, and they identified CI- and CM-type carbonaceous chondrites as the most likely spectral matches, with a preference for a CM1-like composition. (Please note that throughout this Article we follow the standard convention of petrologic types for chondrites, such as CI1 and CM2, first introduced by Van Schmus and Wood⁷.) Thus, Bennu was predicted to have hydrated minerals, but no spectral features attributable to hydration were observed. The average OVIRS disk-integrated spectrum of Bennu compares very well with the telescopic data at these wavelengths, also having a negative slope and no clear absorption

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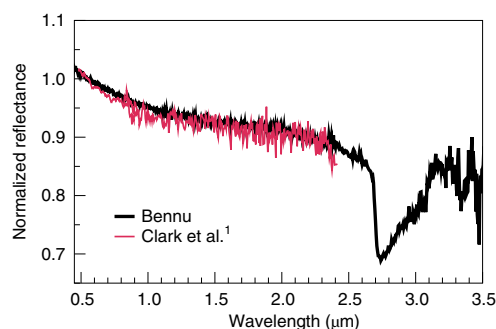


Fig. 1 | Average whole-disk, full-rotation OVIRS spectrum of Benu compared with the ground-based spectrum. The OVIRS radiance factor spectrum (black) and ground-based spectrum¹ (red) are normalized to a reflectance of 1.0 at 0.55 μm . The OVIRS data were acquired on DOY 306 (2 November 2018), and the FOV was approximately 40% filled during these observations.

features (Fig. 1). There is no variation in the spectra (above the noise) with rotational phase. Analysis of spatially resolved data is ongoing and will be used to confirm or refute ground-based observations of spectral slope changes⁸.

A blue-sloped continuum could be explained in one or more ways; such a continuum has been observed in some CI and CM carbonaceous chondrites and, in CI meteorites, is attributed to the presence of fine-particulate magnetite and/or insoluble organic material; it is also commonly associated with larger-particle-size samples and possibly space weathering^{9–11}. Lauretta et al.¹² have identified a candidate magnetite feature at 0.55 μm (ref. 13) in the darkest materials imaged by the MapCam instrument; however, as of yet, no such feature has been observed in OVIRS spectra that would confirm this detection or its assignment to magnetite. Such a feature may become evident in the higher-spatial-resolution OVIRS data that will be collected later in the mission. Experimental space weathering of carbonaceous materials can result in reddening or bluing of the spectral slope^{11,14,15}; at present, we do not have sufficient information from OVIRS spectra to draw any conclusions about the nature or degree of space weathering on Benu as it relates to Benu's spectral slope or the presence of magnetite.

At longer wavelengths ($>2.4\mu\text{m}$), both disk-integrated and spatially resolved OVIRS spectra display an approximately 2.7 μm absorption feature. The 2.7 μm feature is apparent in all OVIRS spectra collected thus far and is similar to the feature observed in aqueously altered CM1 and CM2 carbonaceous chondrites^{16–19}. In analogue meteorites measured under appropriate conditions (Fig. 2), this absorption is due primarily to structural hydroxyl ions in hydrous clay minerals (typically poorly ordered to crystalline phyllosilicates of the kaolinite-serpentine group), which are common in CI and CM carbonaceous chondrites^{19–21}. Among carbonaceous chondrites, hydrated minerals also are a component of CR chondrites²². Adsorbed H_2O in CI/CM meteorite samples (commonly terrestrial in origin) exhibits a broad feature centred closer to 3.1 μm (ref. 19). Any potential H_2O feature in the OVIRS spectrum is weak and will be examined in greater detail using higher-spatial-resolution data.

The exact position of the approximately 2.7 μm band minimum in phyllosilicates shifts with mineral structure and composition^{19,23} and there is experimental evidence that its position may be altered by space weathering²⁴. The band centre in the OVIRS data is at 2.74 μm (± 0.01). Takir et al.¹⁹ have shown that CI and CM chondrites display three distinct types of spectra based on the position of this feature. In 'group 1' spectra, this feature ranges in position from 2.77 to 2.80 μm and is associated with petrologic subtypes between

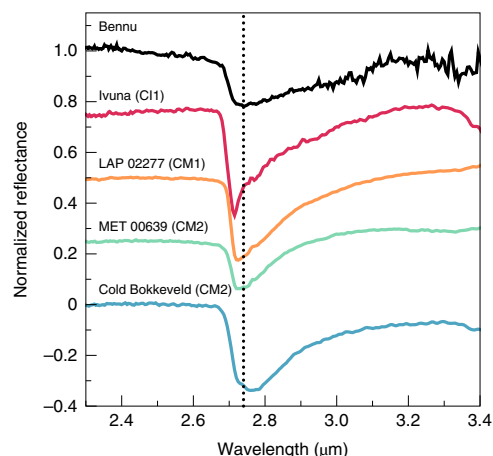


Fig. 2 | Average DOY 306 OVIRS spectrum between 2.3 and 3.5 μm compared with spectra of example carbonaceous chondrites. The carbonaceous chondrites were measured in vacuum after heating¹⁹ (see Methods for full meteorite names). The spectra are normalized to a reflectance of 1.0 at 2.4 μm and offset vertically for clarity. The dotted vertical line at 2.74 μm denotes the Benu band minimum position (see Methods).

CM2.3 and 2.6 (where decimal values indicate relative alteration within type 2, with smaller values representing greater alteration). The band centre for 'group 2' meteorites ranges from 2.76 to 2.78 μm and includes petrologic subtypes CM2.1 to 2.2, which are the most aqueously altered petrologic type-2 meteorites. Finally, 'group 3' meteorites are also CM2.1 to 2.2 but have a band centre at 2.72 μm . Ivuna, the only CI1 in the study, has a band centre at 2.71 μm . The OVIRS band centre lies between groups 2 and 3 and is consistent with meteorites having petrologic types of CM2.1–2.2. Meteorites with these petrologic types are among the most aqueously altered samples studied. Space weathering effects on asteroids in this spectral region do not always match predictions²⁵ but if solar wind irradiation is affecting this band in a manner consistent with experimental data on Murchison (CM2.5), the predicted effect would be to shift the band centre to slightly longer wavelengths (a maximum of 0.03 μm for Murchison) and introduce a concave shape²⁴. As seen in Fig. 2, the spectra of CI and CM1 and low petrologic subtype CM2 meteorites can display concave shapes in the absence of irradiation. The concavity of the Benu spectrum is visibly less than that observed in the analogue meteorites; therefore, we cannot uniquely ascertain whether or not the shape of the Benu spectrum in this region is indicative of space weathering.

Previous studies have identified four classes of so-called 3 μm band shapes among C-complex main belt asteroids, which includes the region of the 2.7 μm feature. These classes are named for their type examples: the asteroids Ceres, Pallas and Themis and the Jovian moon Europa^{26–29}. These classes correspond to different dominant surface materials. Benu's spectrum, with its smooth rise from 2.85 to $\sim 3.3\mu\text{m}$ and blue spectral slope, falls into the Pallas-like class, consistent with what is presumed to be a phyllosilicate-dominated composition.

Spectra of Cb-type³⁰ asteroid (162173) Ryugu measured by the near-infrared spectrometer on the Japan Aerospace Exploration Agency-led Hayabusa2 mission exhibit a weak, narrow 2.72 μm hydroxyl band that does not vary spatially and is interpreted as indicating the presence of Mg-rich phyllosilicates³¹. The best meteorite analogues for the observed feature are thermally metamorphosed CI chondrites and shocked CM chondrites, suggesting that Ryugu has experienced more heating than Benu, although other

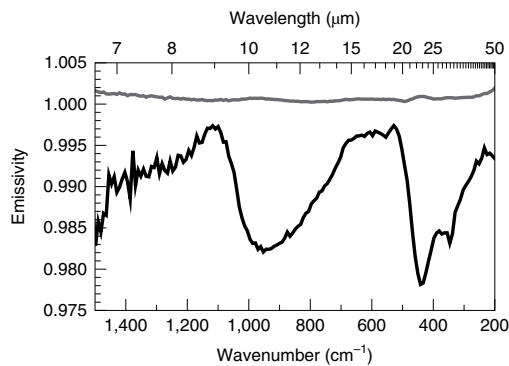


Fig. 3 | Average OTES spectrum of Benu between 1,500 and 200 cm^{-1} .

The Benu spectrum (black) represents slightly more than one full rotation of the asteroid as measured on DOY 347 (13 December 2018). The grey spectrum shows the standard deviation (offset +0.98).

interpretations are possible³¹. Regardless of the interpretation, it is clear that Ryugu differs from unheated or slightly heated, phyllosilicate-rich carbonaceous chondrites and from Benu.

There is not yet unambiguous evidence of organic features in the whole-disk or spatially resolved OVIRS spectra of Benu above the level of the noise in the data shown. The whole-disk observations filled only approximately 40% of the FOV, and the spatially resolved data were acquired at moderate phase angles (approximately $40\text{--}50^\circ$) on relatively hot (approximately 340 K) surfaces, which increases the contribution from thermal emission at the wavelengths where organic bands would be expected. Planned higher-spatial-resolution data on colder surfaces may yet reveal such signatures.

Thermal infrared spectral characteristics

Whole-disk emissivity spectra of Benu acquired in 2007 by the Infrared Spectrograph on the Spitzer Space Telescope have no discernible spectral features above the noise level of the data³² although a comparison is shown in ref. ³³. Comparable disk-integrated OTES observations require additional calibration because Benu does not fill the OTES FOV. However, spatially resolved (80 m per spot) OTES observations reveal thermal infrared spectra with a spectral contrast of approximately 2% that do not vary in shape with rotational phase above the level of the noise (Fig. 3).

The average thermal infrared spectrum of Benu exhibits a Christiansen feature (a peak on the high wavenumber/short wavelength side of the first major, usually silicate, absorption) position that is most similar to that of the CM1/2 and CM2 petrologic types. The spectrum also exhibits an absorption at the lowest wavenumbers (longest wavelengths) that is very similar to that observed in CI and CM carbonaceous chondrites (Fig. 4). Meteorites in the CI and CM groups are volumetrically dominated ($>55\text{ vol\%}$ (refs. ^{34,35})) by hydrated silicate minerals of the phyllosilicate group and are widely accepted to have been aqueously altered during their history within a parent body^{36–38}. Therefore, we can infer that Benu's surface is volumetrically dominated by phyllosilicates and represents aqueous alteration of the parent body.

It is notable that we have not yet observed a distinct Mg–OH feature near 625 cm^{-1} ($16\mu\text{m}$), as this feature is common to many meteorites of the CI and CM groups. The absence of this feature may be indicative of a non-Mg endmember (Fe-bearing) phyllosilicate composition, modest heating, disorder and/or a particle size effect. Although there is no 'smoking gun' match to Benu among the aqueously altered meteorites, spectra of Benu are distinctly dissimilar to carbonaceous meteorite groups that have either not undergone hydrothermal aqueous alteration or have experienced

alteration but are now 'dry' (for example, CO, CB, CV, CK³⁹) (Fig. 4 and Methods). Benu's spectral signature also is dissimilar to meteorites of the CR group, which may be aqueously altered but typically contain lesser amounts of phyllosilicates with abundant olivine and pyroxene³⁴ and have features that would be evident in the Benu spectrum (Fig. 4)^{39–41}.

OTES spectra of Benu also exhibit two features at 555 and 340 cm^{-1} that are probably attributable, at least in part, to magnetite (Fig. 5) and may support the proposed detection of a magnetite feature at $\sim 0.55\mu\text{m}$ in the darkest regions of the asteroid¹². Magnetite is believed to be a product of aqueous alteration and is present at abundances up to approximately 10% in CI chondrites. Magnetite abundance varies widely in CM chondrites, from approximately 0.3 to 8.4% depending on petrologic subtype^{34,35}. The abundance of magnetite on Benu has not yet been tightly constrained, but it is present at abundances of at least a few percent and its detection is consistent with our other observations that support an affinity with these meteorite groups.

The spectral slope of Benu from $1,500$ to $1,110\text{ cm}^{-1}$ (~ 6.6 to $9\mu\text{m}$) is relatively shallow and featureless—it does not clearly exhibit the spectral shapes and emissivity reductions in this region that are common to fine-particulate sample spectra and result from volume scattering (Fig. 4b). The region of silicate stretching bands ($\sim 1,100$ to 700 cm^{-1} ; ~ 9 to $14.3\mu\text{m}$) exhibits a broad, bowl-like shape that is not well reproduced by spectra equivalent to solid and coarse-particulate (for example, $>125\mu\text{m}$) meteorites or fine-particulate ($<125\mu\text{m}$) meteorites measured in vacuum with an induced thermal gradient (Fig. 4). Although there are similarities in the shape and breadth of the fine-particulate Orgueil (CI) chondrite spectrum and Benu in this region, there are distinct differences between these spectra at higher wavenumbers, so this feature shape might alternatively indicate an amorphous/disordered component rather than production of transparency features resulting from volume scattering.

Despite the lack of strong evidence for abundant, volume-scattering (fine) particulates at the approximately 80 m spatial scale of these observations, it is possible that these spectra represent a mixture of a small amount of fine ($<125\mu\text{m}$) and greater amount of coarse ($>125\mu\text{m}$) particulate materials, as well as the boulders that are present across the surface^{5,42}. The lack of variation in the spectra indicates that at these spatial scales, the distribution of particle sizes on the surface does not vary substantially. The thermal inertia of Benu is $350 \pm 20\text{ J m}^{-2}\text{ K}^{-1}\text{ s}^{-1/2}$, does not vary with rotational phase and indicates a mean particle size on the order of 0.5 to 5 cm (ref. ⁵). However, thermal inertia is not uniquely interpretable in terms of particle size, and the presence of numerous boulders for this relatively low value of thermal inertia could be interpreted as indicating that there also may be smaller particles present than the mean particle size estimate would suggest. Alternatively, it may be that the assumption about the thermal inertia of boulders on Benu is inaccurate and that their thermal inertia is lower than what is assumed for typical planetary materials⁵. The lack of rotational variability in thermal inertia is consistent with the lack of variability in the apparent particle size distribution from spectroscopy, despite their differing depth sensitivities.

OSIRIS-REx spectroscopic observations from visible to thermal infrared wavelengths are highly complementary and show that the pristine sample that will be returned from Benu has the potential to inform our understanding of water in the early Solar System and its origins on Earth. Benu's spectra indicate that the surface is consistent with and dominated volumetrically by some of the most aqueously altered CM chondrites. We cannot rule out the presence of a lesser component of CI material based on both the presence of magnetite and the visual variability among materials on the surface⁵.

The spectral datasets presented here are consistent with a surface having a range of particle sizes that does not vary spatially at

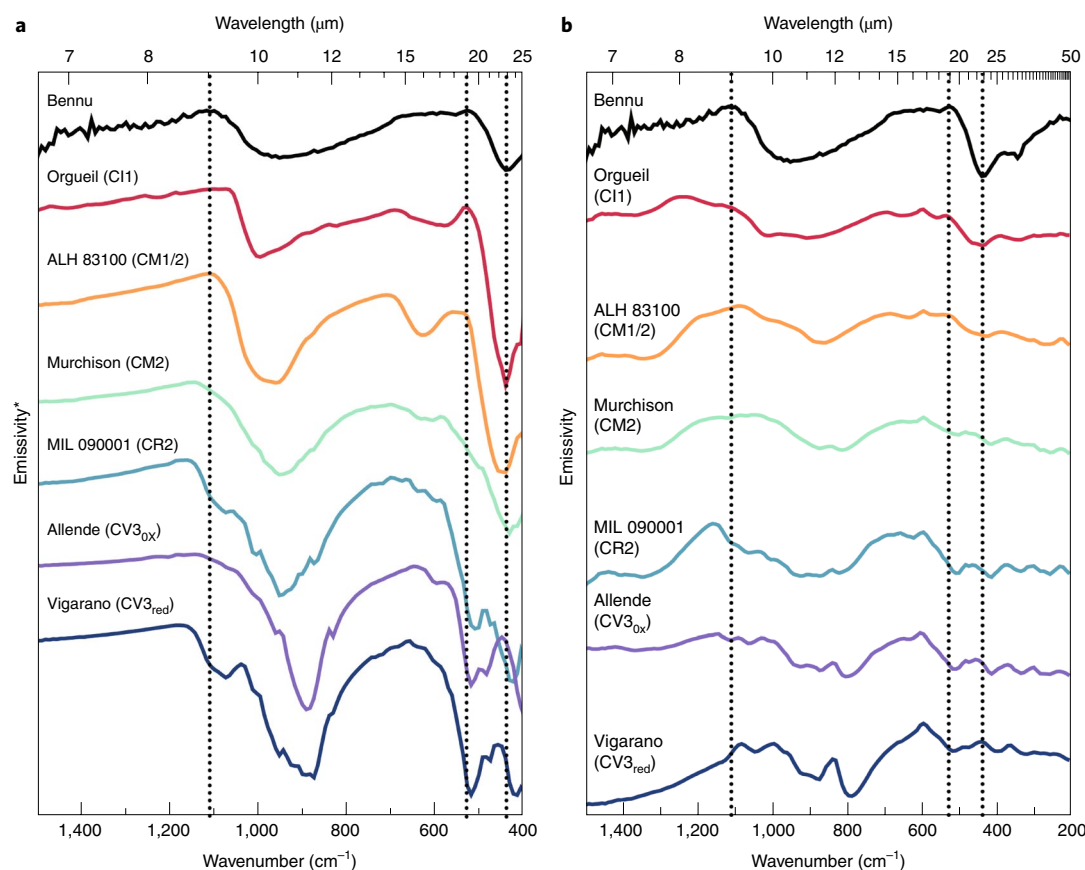


Fig. 4 | Average OTES spectrum of Benu compared with spectra of whole-rock and fine-particulate carbonaceous chondrite meteorites. a, Comparison with whole-rock samples, where the asterisk in the y-axis label indicates that laboratory spectra were measured in reflectance and converted to emissivity (see Methods). **b**, Comparison with fine-particulate (<125 μm) samples. All spectra have been scaled and offset for comparison (see Methods). Dotted vertical lines at 1,110 and 530 cm^{-1} indicate the positions of diagnostic peaks in the Benu spectrum, and the dotted vertical line at 440 cm^{-1} denotes a diagnostic absorption.

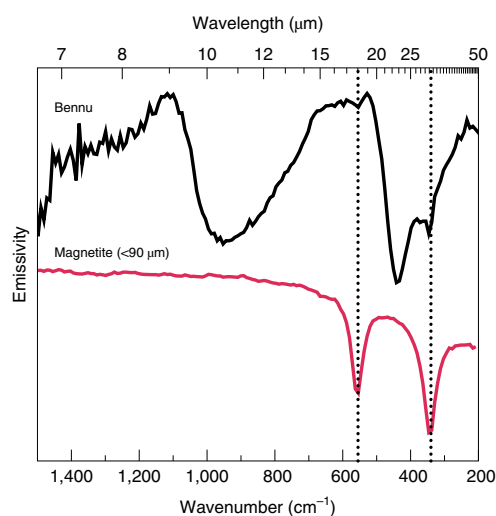


Fig. 5 | Average OTES Benu spectrum compared with a spectrum of pure, fine-particulate (<90 μm) magnetite. Spectra have been scaled and offset for comparison. Dotted vertical lines at 555 and 340 cm^{-1} indicate the positions of diagnostic absorptions in both spectra.

scales down to 80 m as evidenced by the lack of variation in the spectral reflectance and emissivity. Other observed properties may help explain the apparent spatial uniformity of the spectral

signatures at relatively large scales if there are compositional variations present among the mobile materials, but material movement leads to homogenization of their distribution. The lack of rotational and spatial variation in particle size distribution may reflect surficial redistribution processes rather than compositional uniformity, given the observed variations in albedo⁵. Redistribution processes are supported by the geopotential at Benu's surface, which reveals that disturbed material moves towards the equator and/or escapes⁴³. In addition, analysis of the geological characteristics of Benu's surface indicates that it is an old rubble pile but has experienced recent dynamical and geological processes⁴². With these and future, higher-spatial-resolution spectral observations, we will be able to: (1) provide vital context for analysis of the returned sample; (2) address the history and degree of aqueous alteration experienced by Benu's parent body based on details of mineral distribution, abundance and composition (for example, Mg/Fe proportions in phyllosilicates and abundance of magnetite); and (3) constrain the presence or absence of organics.

Methods

OVIRS instrument, calibration and data processing. The OVIRS design is derived from the New Horizons Lisa Hardaway Infrared Mapping Spectrometer (LEISA) portion of the Ralph instrument⁴⁴ with an extended wavelength and simplified optics. The spectrometer uses five linear variable filters to collect the spectrum. Details of the various operating modes (for example, super pixel summing) are described elsewhere². To measure compositional spectral features with >5% absorption depth at spatial resolutions of 5 to 50 m, OVIRS meets a performance requirement of a signal-to-noise ratio of >50 across the entire

spectral range assuming an asteroid surface albedo of approximately 3–4% at a solar range of 1.2 au and 300 K thermal radiation. To characterize and map variations in space weathering on surfaces with an albedo of >1%, OVIRS meets an accuracy requirement of 2.5% with a precision of 2%. OVIRS calibrations and performance assessments were performed on the ground and in-flight during the OSIRIS-REx Earth encounter in September 2017 (ref. 45).

The observing sequence on day of year (DOY) 306 consisted of pointing OVIRS at Bennu for 4.5 h while scanning in a slight ‘up and down’ pattern but keeping Bennu within the FOV at all times to obtain whole-disk measurements. The phase angle during these observations was approximately 5.2°. The spectrum shown in Figs. 1 and 2 is the average of 17,061 radiance factor (observed radiance over solar irradiance or flux, I/F) spectra where Bennu filled approximately 40% of the FOV; the excursions in the spectra are representative of the point-to-point scatter in the data. The OVIRS calibrated radiance spectra were obtained according to methods described by refs. 245. In brief, OVIRS raw data were converted from counts per second to absolute radiance units using an automated calibration pipeline. First, the closest space view was identified to create an average background file. The background subtracted counts were converted to physical units using radiometric and out-of-band coefficients derived from ground testing and in-flight calibration activities. The full calibration approach is described in more detail elsewhere⁴⁵. Slight adjustments were made to the previously derived radiometric and out-of-band coefficients to adjust the response in a few spectral regions based on the Bennu Approach data to ensure filter overlap regions aligned. Calibrated radiances were then resampled onto a common wavelength axis by removing outlying noise spikes more than 1.8 standard deviations from the mean and a performing a weighted average on the remaining spectral points in each wavelength bin. The common wavelength axis had a spectral sampling of 2 nm from 0.4 to 2.4 μm and 5 nm from 2.4 to 4.3 μm . Data were then converted to radiance factor (I/F) by dividing by the solar spectrum scaled for Bennu’s distance.

The OVIRS disk-integrated data shown in this work are not photometrically corrected. The geometric albedo of Bennu (0.044 ± 0.002) as determined from imaging results is given by ref. 46. The geometric albedo of asteroids (extrapolated to 0° phase) is known to be higher than the values measured in laboratory settings at 30° phase, where for Bennu’s phase function, this scale factor is approximately 2. If we apply this scaling factor to meteorite albedo values presented in Fig. 4 of ref. 1, CI and CM chondrite values are most comparable to the geometric albedo of the hemispherically integrated observation of Bennu and meteorites of the CK, CO, CV, CR and CH groups are not consistent. However, because there is evidence in higher-resolution imaging of materials on Bennu’s surface having considerably higher albedos⁵, we are not prepared to assert that any compositions are ruled out by the global geometric albedo value.

Analysis of OVIRS spectra beyond approximately 2 μm requires removal of the contribution to the signal from thermal emission. We tested two methods for removing this ‘thermal tail’, with both giving similar results; we show the spectrum obtained by the first method. The first approach to computing the thermal contribution to the total radiance uses a smooth-surface thermophysical model assuming a spherical asteroid. The thermal portion of the measured flux was estimated assuming that the spectrum of Bennu is flat from 2.2 to 4.0 μm . The thermal model was run while varying thermal inertia and asteroid size to fit the thermal portion of the measured flux for each OVIRS spectrum. The reflected radiance was computed as a straightforward subtraction of the model thermal radiance from the total measured radiance. In this approach, all the uncertainty and any remaining calibration artefacts are assumed to reside in the reflected radiance. Because the absolute uncertainties remain unchanged but the radiance itself is decreased substantially at wavelengths with significant thermal contribution, the relative uncertainties at these longer wavelengths increase, leading to an apparent increase in noise at longer wavelengths in the subtracted spectrum. For purposes of searching for potential spectral features, we also computed total model radiance by adding the thermal model radiance to a model reflected radiance (computed by scaling the solar spectrum to OVIRS radiance at 2.2 μm), then divided the measured OVIRS spectra by the model total spectra (Figs. 1 and 2).

In the second method, the thermal contribution to the total radiance was computed using the OSIRIS-REx thermal model described in ref. 5. The computation was performed independently for each OVIRS spectrum for the instantaneous spacecraft distance and rotation phase of Bennu, using the shape of Bennu derived from OSIRIS-REx images⁴⁷. We used the v13 shape model at the lowest (12 m) resolution. The disk-integrated thermal models are not affected by the small changes in the newer version (v20) of the shape model. For some rotation phases, the model thermal radiance does not perfectly match the OVIRS measurements due to remaining imperfections in the shape model. We therefore scaled the model thermal radiance to the average measured radiance (averaged from 3.5 to 4.0 μm) of each spectrum before subtracting from the total radiance. For scaling purposes, the measured thermal radiance was estimated assuming that the reflectance of Bennu is flat from 2.2 to 4.0 μm . The reflected radiance was computed as a straightforward subtraction of the model thermal radiance from the total measured radiance. In this approach, all the uncertainty and any remaining calibration artefacts are assumed to reside in the reflected radiance as described above.

Determination of the 2.7 μm band centre was calculated (after the correction for thermal emission) using two methods that give virtually the same result to

within the uncertainty of the measured spectrum. The first method is to fit a sixth-order polynomial to the measured spectrum between 2.65 and 2.85 μm and find the minimum of that fit; this is the same method used by ref. 19, although those authors did not report the wavelength range over which they did their fitting. This fit was calculated for both versions of the average thermal-radiance-removed Bennu spectrum. The derived minima vary by a single channel between the two spectra, being fit at 2.74 and 2.745 μm . Because the thermal emission correction can influence the position of this band, and these spectra represent a whole-disk measurement with variable temperatures and phase angles, this result suggests our uncertainty is relatively small (on the order of the channel-to-channel uncertainty).

The second method for determining the 2.7 μm band position fits a smoothing spline function to the spectrum between 2.69 and 2.85 μm for the DOY 306 (2 November 2018) average spectrum; the best fit is obtained using a smoothing value of 0.999999. The first derivative is then calculated, and the inflection point is used to determine the position of the band, which is 2.74 ± 0.007 . Applying the same analytical approach to all of the spectra acquired on DOY 306, we obtain the same answer, to within the uncertainty of the data. Based on the consistency of the results obtained by these two methods and their estimated uncertainties, we conservatively identify the feature as being located at 2.74 ± 0.01 μm .

OTES calibration and data processing. The OTES instrument⁸ is a Michelson interferometer with heritage from the Mars Exploration Rovers Mini-Thermal Emission Spectrometer and Mars Global Surveyor Thermal Emission Spectrometer^{48,49}. Spectral sampling is 8.66 cm^{-1} across the entire spectrum. To confidently identify spectral features with a >5% band depth and achieve a 1.5% total emitted radiance accuracy requirement, OTES meets a signal-to-noise ratio of 320 at a reference temperature of 325 K and has a single-spectrum radiometric precision of $\leq 2.2 \times 10^{-8} \text{ W cm}^{-2} \text{ sr}^{-1} / \text{cm}^{-1}$ between 1,350 and 300 cm^{-1} . The absolute integrated radiance error is <1% for scene temperatures ranging from 150 to 380 K.

Observing sequences designed to obtain whole-disk OTES spectra consisted of pointing OTES at Bennu for 4.5 h (a little longer than one full rotation of the asteroid) without scanning. However, the standard calibration of OTES data depends on the scene and the calibration targets all filling the FOV; when the scene (Bennu) fills only a portion of the FOV, wavelength-dependent, off-axis modulation of energy through the interferometer results in an apparent low signal at short wavelengths. Correcting this effect requires a substantially more complex calibration approach, which is under consideration. As such, we show here spectra acquired on DOY 347 (13 December 2018) when the FOV was fully filled and the standard calibration approach is appropriate for the observations. The average phase angle during these observations was approximately 45.5°. The DOY 347 observations cover the equator and southern (relative to the plane of the ecliptic) hemisphere and are equally representative of observations in the northern hemisphere collected during Preliminary Survey sequences on other days.

The calibration of OTES data generally consists of an automated processing pipeline that transforms OTES raw interferograms into voltage spectra and then into absolute radiance units⁵. More specifically, the measured voltage spectrum is the difference between the radiance of the scene, foreoptics and the detector times the instrument response function; the radiance of the detector and the instrument response function are unknowns, but can be determined by periodically observing space and an internal calibration target, at which point it becomes possible to solve for the scene radiance and account for temperature fluctuations of the instrument (detector) that result from the instrument heater cycling during the observations. After the acquisition of Earth observations in September 2017, an adjustment was made to the calibration pipeline to account for slopes in the interferograms that occur during the transition between cold space and a hot target (for example, Earth or Bennu). This slope results from the time constant associated with the DC-correction electronics (which is longer than the 2 s integration) and, if uncorrected, results in high-frequency ‘ringing’ in the spectra. In addition, many of the ‘ride-along’ observation sequences in the Approach and Preliminary Survey⁴ that were designed for imaging did not include periodic views of space, instead measuring space only at the start and end of sequences that lasted on the order of 4.5 h. As a result, an alternative calibration approach was developed to account for instrument (detector) temperature fluctuations during these sequences; this involves using a look-up table that correlates in-flight measurements of the temperature measured by a thermistor adjacent to the detector to the detector radiance.

The afternoon local time of the DOY 347 observations (approximately 15:00–15:30) results in viewing surfaces having different temperatures (for example, sunlit and shadowed) thus requiring an emissivity–temperature separation that allows for the fitting of multiple temperatures. We fit the OTES calibrated radiances using a non-negative linear least squares algorithm⁵⁰ that takes as input a suite of Planck functions with temperatures between 150 and 380 K. The mixture of Planck functions that provides the best fit to the measured radiance is divided into the measured Bennu radiances to obtain emissivity, where the maximum emissivity is assumed to be 0.97 based on reflectance measurements of relevant carbonaceous chondrite meteorites³⁹. The Bennu spectrum shown in Figs. 3 and 4 is the average of 974 spectra with spatial resolutions of approximately 80–90 m per spot collected on DOY 347.

Meteorite samples. The meteorites shown in Fig. 2 are Ivuna (CI1), LaPaz Icefield (LAP) 02277 (CM1), Meteorite Hills (MET) 00639 (CM2) and Cold Bokkeveld (CM2)³⁹. Cold Bokkeveld may have been very mildly and briefly heated based on Raman spectroscopy of the insoluble organic material, but the evidence is ambiguous^{41,53} and there is no mineralogical evidence of heating that would change our interpretation of the observed 2.71 μm feature (where mineralogy is the property to which the laboratory and remote-sensing measurements shown here are sensitive). Because meteorites have interacted with the Earth's environment, even if briefly, they are prone to mineralogical and chemical alteration, including the adsorption and absorption of terrestrial water (which can be recognized through oxygen isotope analysis). The spectra shown in Fig. 2 were measured under vacuum after the samples were heated to between 400 and 475 K, which drives out adsorbed and absorbed terrestrial water. The laboratory spectra have been resampled to the OVIRS spectral sampling. See ref. ¹⁹ for details of sample preparation, characterization and measurement.

The meteorites shown in Fig. 4 are Orgueil (CI1), Allan Hills (ALH) 83100 (CM1/2), Murchison (CM2), Miller Range (MIL) 090001 (CR2), Allende (CV3_{un}), and Vigarano (CV3_{un}). All of these spectra were acquired as part of the development of the OSIRIS-REx spectral library for the analysis of OTES data and have been resampled to the OTES spectral sampling. The spectral acquisition methods are described below. The text indicates that thermal infrared spectra of meteorite groups CO, CB and CK do not resemble the OTES spectrum of Bennu; spectra of these groups are contained in the research collection of V.E.H. and are not shown here but have been shown elsewhere³⁹.

Laboratory spectroscopy. The meteorite spectra shown in Fig. 4a were measured by V.E.H. in reflectance on uncoated thin sections using a Thermo Scientific Nicolet iN10 microscope at Southwest Research Institute in Boulder, CO. The microscope is equipped with a KBr beamsplitter and a nitrogen-cooled, extended-range mercury-cadmium-telluride detector and measures spectra from 4,000 to 400 cm^{-1} ; the optical geometry of this microscope is such that the spectra are equivalent to emission spectra according to Kirchhoff's law⁵³. The spectra have been scaled by differing amounts to minimize spectral contrast variations and simplify the comparison of spectral shapes. These spectra are appropriate for comparison with OTES emissivity spectra of coarse particulates and solids that do not exhibit volume scattering and are not susceptible to thermal gradients^{54,55}.

Figure 4b shows fine-particulate (<125 μm) versions of the same meteorite samples measured by K.L.D.H. in a simulated asteroid environment at Oxford University; the sample preparation, characterization and spectral measurements are described in detail in ref. ³⁶. The <90 μm magnetite spectrum in Fig. 5 was measured under the same conditions and its spectrum is virtually identical to magnetite spectra measured as coarse materials and under ambient conditions. All of these spectra are appropriate for comparison with OTES emissivity spectra of dominantly fine particulates that exhibit volume scattering and are potentially susceptible to the development of thermal gradients.

Data availability

The data that support the plots within this paper and other findings of this study are available from the corresponding author upon reasonable request. Raw and calibrated spectral data will be available via the Planetary Data System (PDS) (<https://sbn.psi.edu/pds/resource/orex>). Data are delivered to the PDS according to the OSIRIS-REx Data Management Plan available in the OSIRIS-REx PDS archive. Higher-level products will be available in the PDS one year after departure from the asteroid. Laboratory spectral data are deposited in the spectral library hosted by Arizona State University (<http://speclib.mars.asu.edu/>).

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Author contributions

V.E.H. is the spectral analysis working group lead, the OTES deputy instrument scientist, and wrote this manuscript. A.A.S. is the spectral analysis working group deputy, the OVIRS deputy instrument scientist, and led the calibration of the OVIRS data and production of the disk-integrated average spectrum. P.R.C. is the OTES instrument scientist and led the calibration of OTES data. D.C.R. is the OVIRS instrument scientist. B.E.C. is the OSIRIS-REx Mission Asteroid Scientist. M.A.B., H.H.K., R.D.H. and A.P. contributed to the analysis of the OVIRS 2.7 μm band. N.E.B. hosts the laboratory that made the simulated asteroid environment spectral measurements. W.V.B. is the mission instrument scientist and contributed to ensuring the mission plan enables the instruments to meet their observation requirements. J.R.B., E.A.C., S.F., C.L., J.-Y.L., F.M., S.A.S., C.A.T. and X.-D.Z. contributed to the development of science pipeline software. H.C.C. Jr is the mission sample scientist and helped guide the selection and acquisition of the meteorite samples used in this work. K.L.D.H. measured the samples shown in Fig. 4b. J.P.E. and B.R. contributed to the subtraction of thermal emission from OVIRS spectra. H.L.E. is the deputy principal investigator for the OSIRIS-REx mission. C.W.H. contributed to the data processing and analysis of OTES spectra. E.S.H. contributed to the development of science pipeline software and provided manual processing of some of the data shown in this manuscript. L.P.K. and T.J.M. helped guide the selection and acquisition of the meteorite samples used in this work. L.F.L. contributed to extensive discussions about the laboratory measurements. M.C.N. is the science team chief and contributed the resampled solar spectrum used in the calibration of OVIRS data. D.L.S. contributed to the preparation and characterization of meteorite samples used in this work. D.S.L. is the OSIRIS-REx principal investigator and the entire OSIRIS-REx Team made the Bennu encounter possible.

Competing interests

The authors declare no competing interests.

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