



Volcanological applications of SO₂ cameras

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ABSTRACT

Ground-based volcanic gas and ash imaging has the potential to revolutionise the way in which volcanoes are monitored and studied. The ability to track and quantify volcanic emissions in space and time with unprecedented fidelity opens the door to integration with geophysical measurements, allowing breakthroughs in our understanding of the physical processes driving volcanic activity. In May 2013 a European Science Foundation funded Plume Imaging workshop was conducted in Stromboli, Italy, with the objective of bringing the ground-based volcanic plume imaging community together in order to examine the state of the art, and move towards a 'best-practice' for volcanic ash and gas imaging techniques. A particular focus was the development of SO₂ imaging systems, or SO₂ cameras, with six teams deploying and testing various designs of ultraviolet and infrared-based imaging systems capable of imaging SO₂. One conclusion of the workshop was that the term 'SO₂ camera' should be applied to any SO₂ imaging system, regardless of wavelength of radiation used.

This Special Issue on Volcanic Plume Imaging is the direct result of the Stromboli workshop, and together the papers presented here represent the state of the art of ground-based volcano plume imaging science and technology. In this work, we examine in detail the volcanological applications of the SO₂ camera, reviewing previous works and placing the new research contained in this Special Issue in context. The development of the SO₂ camera, and future developments extending imaging to other volcanic gases, is one of the most exciting and novel research frontiers in volcanology today.

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1. Introduction

In May 2013 a European Science Foundation-funded Plume Imaging workshop was conducted on Stromboli, Italy, with the objective of bringing the ground-based volcanic plume imaging community together in order to examine the state of the art, and move towards a 'best-practice' for volcanic ash and gas imaging techniques of volcanic ash and gas imaging techniques. A particular focus was the development of SO₂ imaging systems, or SO₂ cameras, with six teams deploying and testing various designs of ultraviolet and infrared-based imaging systems capable of imaging SO₂ (Kern et al., 2015a, b). This special issue is a direct product of that workshop, and represents the state of the art in volcanic gas and ash imaging research.

The recent development of gas and ash imaging systems has opened up new frontiers in volcanology. The ability to detect rapid changes in gas and ash emissions allows a whole new series of phenomena to be examined over a range of scales, with a particular focus on explosive volcanism and the dynamics of passive degassing. Volcanologists may now combine comprehensive studies of gas emissions with seismic signals, producing unprecedented new constraints on the underlying

physical processes that drive both geochemical and geophysical observations. Compared to earlier plume scanning technologies (e.g. Edmonds et al., 2003; Burton et al., 2007; Burton et al., 2009; Salerno et al., 2009; Galle et al., 2010) SO₂ cameras allow more precise and much faster measurement of gas fluxes.

Clearly, however, direct interdisciplinary comparisons also place great demands on the quality of data produced by each discipline. When used to make truly quantitative inferences on a physical process we require data that is both accurate, in order to e.g. compare with mass fluxes of lava, and precise, in order to distinguish real variations in flux from uncertainties. Furthermore, we begin a natural progression from short-term experimental field studies by research teams towards permanent and automatic data acquisition and analysis by volcano observatories. These demands require a rapid maturation of methodology, and the ongoing efforts of the community to ensure comparability of standards.

In this paper we examine the role of the SO₂ camera in volcanology and discuss what steps need to be made to fully realise the potential of the approach, highlighting how the papers in this special issue and previously published SO₂ camera research have contributed to this goal. A detailed review of SO₂ camera technologies is presented by Platt et al., 2015, and therefore in this work we focus on three main subjects: (i) Quantification of gas emission rates, (ii) volcanic processes which

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can be investigated with the SO₂ camera, and (iii) implementation of SO₂ cameras in monitoring networks. Platt et al. (2015) highlight that whilst the majority of SO₂ cameras developed to date are based on ultraviolet spectroscopy, infrared spectroscopy offers tangible advantages, particularly in terms of night-time viewing. One conclusion of the workshop was the term ‘SO₂ camera’ which should be applied to any SO₂ imaging system, regardless of wavelength of radiation used.

2. Quantification of gas emission rates

The majority of SO₂ camera systems used to date (and 1-D plume scanning systems) are based on ultraviolet absorption spectroscopy using scattered sunlight as a source of radiation. This approach has been a staple of volcano observatories looking to monitor SO₂ emissions from active volcanoes since the 1980s. As we discuss in more detail in Section 4, volcano observatory personnel are generally focussed on the interpretation of signals generated by their instruments, and tend to trust the results obtained by them. However, frequently it is overlooked that direct trace gas column measurements can be misleading unless appropriate corrections are applied (e.g. Lübcke et al., 2013). Light scattering below volcanic plumes can dilute the SO₂ signal measured at the ground. This may be quite a significant problem, particularly when the distance between the instrument and the plume is large or the plume is ash-rich; unfortunately SO₂ fluxes during eruptions are of great interest to volcano observatories. A theoretical treatment of these multiple scattering issues is described by Kern et al. (2013). Campion et al. (2015) use the brightness of the volcanic edifice to determine how much ‘light dilution’ is occurring, and use this to correct for the effect. Burton and Sawyer (2013) present an intensity-based spectrum analysis approach to solve the issue. Also, care has to be taken that the SO₂ camera correctly calibrated, Lübcke et al. (2013) describe possible pitfalls in calibration and how problems can be avoided. More work is required to fully understand and resolve this challenging problem.

A further major challenge in derivation of gas emission rates from SO₂ camera data is the conversion of sequences of images of SO₂ slant column amounts into a time series of SO₂ flux. This was dealt within the first papers (e.g. Mori and Burton, 2006) by defining a line orthogonal to the plume advection direction and integrating the SO₂ slant column amounts along that line, before multiplying by a plume velocity derived from the cross-correlation lag between parallel lines. This approach works well when the plume movement is quite uniform, which is a good approximation when the SO₂ camera is far from the plume. However, one of the strengths of the SO₂ camera is its spatial and temporal resolutions, which make it very often attractive to deploy it close to a source of gas, where the plume direction is far from uniform. In this case it is non-trivial to determine a flux, as there is no single velocity for the gas. Peters et al. (2015) address this problem using motion detection algorithms to find the overall flux of gas passing through a user-defined line or curve, allowing accurate determination of gas fluxes even close to the gas source. Kern et al. (2015a, b) report a very similar approach, used to help automate SO₂ flux calculations on Kilauea. Validation of SO₂ fluxes measured with the SO₂ camera can also be obtained using a calibrated emission source, and such a test is reported by Smekens et al. (2015).

A key element in the evolution of the SO₂ camera is the intercomparison of the diverse approaches that have developed since the inception of the technique. The May 2013 plume imaging workshop on Stromboli provided the ideal vehicle for such an intercomparison, and the results from this are reported by Kern et al. (2015a, b).

3. Volcanic and degassing processes which can be investigated with the SO₂ camera

Magmatic degassing is one of the main drivers of volcanic processes, coupling with crystallisation and viscosity variations to determine the style and intensity of volcanic activity (Sparks, 2003). The magnitude

of gas release is related to the mass of magma ascending in the volcanic system, allowing inferences on magma dynamics from mass balance considerations. In the following we review how SO₂ cameras can improve our understanding of volcanic processes.

3.1. SO₂ imagery and geophysical signals associated with volcanic activity

One of the greatest strengths of the SO₂ camera is its ability to measure fast changes in degassing rates at source, thanks to both, a potentially fast frame rate (cameras can collect up to 18 frames per second in ideal conditions) and the fact that each two-dimensional image records the history of degassing captured within the frame, like a ticker-tape timer. This ability makes the camera ideal for investigating the rates of degassing during explosions (see Section 3.2, below) and for studying the links between degassing and seismic and acoustic signals. Quantitative inversion of very long period (VLP) seismic events allows the volume change associated with the passage of a gas slug to be estimated (Chouet et al., 2003; Zuccarello et al., 2013). This information can be directly compared with results from the SO₂ camera, in order to test our understanding of the physical processes which link gas slug ascent with VLP events. Volcanic tremor, an almost ubiquitous seismic signal on active volcanoes, is linked with magma and gas flow (e.g. Tamburello et al., 2013; Zuccarello et al., 2013), and therefore comparisons with gas flux data can deepen our understanding of the processes driving this seismic signal as well. We summarise the work performed to date comparing SO₂ imagery with geophysical parameters below, and in Table 1.

SO₂ is typically a minor gas component in magmatic degassing, which is instead dominated by H₂O and CO₂. VLP signals are generated by magmatic gas, not just the SO₂ component, and therefore the composition of the explosion gas must be known in order to convert from SO₂ masses measured with the SO₂ camera to total gas masses. In the future, technological developments may permit multiple gases to be imaged with a single instrument (see Platt et al., 2015).

Dalton et al. (2010) were the first to study the relationship between acoustic signals and explosive gas release measured with an SO₂ camera at Pacaya volcano, Guatemala. They found that gas masses derived from gas and acoustic datasets agreed to the same order of magnitude, and that the infrasound explosion records correlate with small pulses in degassing, but that longer-term degassing trends were probably controlled by a deeper process.

The first SO₂ camera comparisons with VLP signals were conducted by Kazahaya et al. (2011) and Nadeau et al. (2011), on Asama (Japan) and Fuego (Guatemala) volcanoes, respectively. Both authors found a linear correlation between SO₂ amounts released in each explosion and VLP amplitude. Furthermore, Nadeau et al. (2011) found a clear correlation between volcanic tremor and gas emission rate, and suggested that decreases in gas emission rate may be the result of rheological stiffening in the upper conduit of Fuego.

Tamburello et al. (2012) performed SO₂ camera measurements on Stromboli, finding again a linear relationship between the VLP amplitude and explosion gas mass (2009) and extended their investigation to include thermal signals, which were also well-correlated with gas emissions. SO₂ camera investigations of the degassing rate from Etna, Italy, show a correlation with volcanic tremor amplitude during passive degassing (Tamburello et al., 2013).

Lopez et al. (2013) used an infrared imaging system, NicAIR, to measure SO₂ and ash emission rates at Karymsky volcano, Kamchatka, which produced a wide range of degassing and explosive behaviours during their observation period. The NicAIR SO₂ camera was used to quantify pulsatory magmatic degassing, but in this case no clear correlation with infrasound pressure or thermal variations were detected, probably due to the temporal delay between gas emission at the vent and SO₂ detection downwind.

In what is probably the most detailed study of the relationship between SO₂ degassing rates measured with an SO₂ camera and geophysical signals during explosive volcanic activity, Waite et al. (2013)

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