Optical particle measurements are emerging as an important technique for understanding the ocean carbon cycle, including contributions to estimates of their downward flux, which sequesters carbon dioxide (CO$_2$) in the deep sea. Optical instruments can be used from ships or installed on autonomous platforms, delivering much greater spatial and temporal coverage of particles in the mesopelagic zone of the ocean than traditional techniques, such as sediment traps. Technologies to image particles have advanced greatly over the last two decades, but the quantitative translation of these immense datasets into biogeochemical properties remains a challenge. In particular, advances are needed to enable the optimal translation of imaged objects into carbon content and sinking velocities. In addition, different devices often measure different optical properties, leading to difficulties in comparing results. Here we provide a practical overview of the challenges and potential of using these instruments, as a step toward improvement and expansion of their applications.

**Keywords:** sinking particle fluxes, sinking velocities, carbon content, size, image processing, automated classification, *in situ* optical particle measurements, biological carbon pump
INTRODUCTION

Particulate Matter in the Ocean

Life and particulate organic matter in the ocean have fundamentally shaped our planet. On the most basic level, particulate organic matter can be defined as both living and non-living matter of biological origin with a size of ≥0.2 µm in diameter, including anything from a small bacterium (0.2 µm in size) to blue whales (20 m in size; see review by Blanchard et al., 2017). Organic matter plays a crucial role in regulating marine global biogeochemical cycles and events, from the Great Oxidation Event in Earth’s early history (Holland, 2006) to the sequestration of atmospheric carbon dioxide in the deep ocean (Volk and Hoffert, 1985; Heinze et al., 2015). Understanding the distribution, characteristics and dynamics of particulate matter in the ocean is hence fundamental in understanding and predicting the marine ecosystem, from food web dynamics to global biogeochemical cycles. In this review, we focus primarily on the particles and particle processes involved in the biological carbon pump within the mesopelagic zone (the region from below the productive layer to ~1,000 m depth) and on how these can be inferred from optical measurements. The technologies reviewed here can also be applied to other aquatic systems, and to problems other than the biological pump, such as the distribution and fate of microplastic in the ocean or the presence of specific organisms involved in harmful algal blooms.

The Biological Carbon Pump

The biological carbon pump describes the collection of biogeochemical processes associated with the production, sinking, and remineralization of organic carbon in the ocean (Volk and Hoffert, 1985; Giering and Humphreys, 2018). In brief, photosynthesis by microorganisms in the upper tens of meters of the water column fix inorganic carbon (any of the chemical species of dissolved carbon dioxide) into biomass. When this biomass sinks to the deep ocean, a portion of it fuels the metabolism of the organisms living there, including deep-sea fish and benthic organisms (Turner, 2015). Zooplankton play a critical role in shaping particle flux through ingestion and fragmentation of particles (e.g., Waite et al., 2000; Iversen and Poulsen, 2007; Poulsen and Iversen, 2008; Iversen et al., 2010; Giering et al., 2014; Svensen et al., 2014), production of fast-sinking fecal material (e.g., Turner, 2015; Iversen et al., 2017), and active vertical migration (e.g., Steinberg et al., 2000; Jónasdóttir et al., 2015; Kiko et al., 2017).

Besides the importance of “exported” organic carbon as a food source for deep ocean organisms, the biological carbon pump provides a valuable ecosystem function: Exported organic carbon transports an estimated 5–20 Gt C each year to the deep ocean (Henson et al., 2011), where some of it (~0.2–0.5 Gt C) (Guidi et al., 2015) is sequestered for several millennia. The biological carbon pump is hence of similar magnitude to current carbon emissions from fossil fuels (~10 Gt C year⁻¹). Any changes in its magnitude caused by a warming world may have direct implications for both deep-sea organisms and atmospheric carbon dioxide concentrations (Kwon et al., 2009; Passow and Carlson, 2012).

The magnitude and efficiency (amount of carbon sequestered relative to primary production) of the biological carbon pump, hence ocean carbon storage, is partly determined by the amount of organic matter exported and the rate at which it is remineralized (i.e., the rate with which sinking organic matter is reworked and respired in the mesopelagic zone region; Kwon et al., 2009; Iversen and Ploug, 2010; Reygondé et al., 2013). Especially particle size and composition are important parameters determining how fast a particle sinks (Ploug et al., 2008a; Iversen and Ploug, 2010), how much material it contains (Ploug et al., 2008b), and which organisms can find and utilize it (Kiorboe et al., 1999; Visser, 2001; Visser and Jackson, 2004).

Sinking particles can be phytoplankton, zooplankton, detritus, fecal pellets, or a mix of these (Simon et al., 2002; Turner, 2002, 2015). They range in size from a few micrometers to several centimeters, with particles of a diameter of >0.5 mm being referred to as “marine snow” (Allredge and Silver, 1988). In general, particles in a fluid are thought to sink once their densities are higher than the ambient fluid, i.e., when excess densities are larger than zero. Larger individual phytoplankton cells can thus contribute to sedimentary fluxes. For example, large diatom cells and diatom chains with a diameter of >5 µm have been shown to sink at rates up to several 10⁻¹ meters per day, though this is only possible owing to the heavy ballast of a silica frustule (Waite et al., 1997a; Miklasz and Denny, 2010). Both size and density affect particle sinking velocity; for example, for sinking velocities that follow Stokes’ Law, doubling the size of the particle increases the sinking speed by a factor of 4 (Moore and Villareal, 1996; Waite et al., 1997a). However, the highly porous nature of many marine particles means that they do not obey Stokes’ Law because small changes in particle density (i.e., compactness) can have a large impact on their sinking velocities (Iversen and Ploug, 2010). Large sinking particles are typically of two types: (1) aggregates formed from a number of primary particles, including phytoplankton, bacteria, fecal pellets, live protozoans and zooplankton and debris, and (2) zooplankton fecal pellets, which can dominate particle flux events and sink at velocities exceeding 1,000 m d⁻¹ (Turner, 2015).

Knowing the size, abundance, structure and composition (e.g., carbon content) of settling particles is important as these characteristics impose fundamental constraints on the biogeochemical cycling of carbon. For example, changes in climate are expected to facilitate a shift in species composition in a manner that alters the elemental composition of particulate matter, cell size and the trajectory of carbon through the food web, influencing the proportion of biomass exported to depth (Finkel et al., 2010). As such, any climate-induced change in the structure or function of phytoplankton communities is likely to alter the efficiency of the biological carbon pump, with feedbacks on the rate of climate change (Matear and Hirst, 1999; Le Quéré et al., 2007).

Autonomous Sampling on the Rise

The vastness of the ocean makes it difficult to accurately estimate the processes involved in the biological carbon pump. Remote sensing via satellites has only limited capabilities as it is restricted to the upper meters of the ocean. To date, our knowledge of
the biological carbon pump is based predominantly on data collected by sediment traps (e.g., Honjo, 1996; Francois et al., 2002; Buesseler et al., 2007; Honjo et al., 2008; Fischer et al., 2016), radioactive tracers such as Thorium-234 (e.g., Buesseler et al., 2006; Waite and Hill, 2008; Roca-Marti et al., 2017), and budgets of dissolved biogeochemical tracers such as nutrients (e.g., Schlitzer, 2002; Gehlen et al., 2006). Each of these methods has yielded important insights and has its strengths and weaknesses including limited spatio-temporal coverage and/or resolution or uncertain ocean circulation. Furthermore, carbon flux estimates from these diverse methods often differ widely, and the various spatio-temporal scales of the methods complicate efforts to compare their results (Boyd and Trull, 2007).

The most direct method of measuring particle flux uses sediment traps, which collect sinking particles at a certain depth over a period of several days to months. The collected material is preserved in situ and available for biochemical analysis, including biomarkers. Sediment traps provide useful, quantitative and qualitative estimates of particle fluxes, but the small collection area of a single sediment trap (<1 m²) combined with the low number of traps that can feasibly be deployed complicates extrapolation to mesoscale and broader scale fluxes (Martin et al., 2011). Neutrally buoyant sediment traps, which were designed to overcome hydrodynamic biases and are considered the most accurate trap type (Buesseler et al., 2007), are also limited in temporal coverage to the length of a single oceanographic cruise, preventing the full characterization of seasonal and longer timescales. Full-year coverage is possible with moored sediment traps (e.g., Conte and Weber, 2014), but both spatial coverage and temporal resolution remain limited and questions of over- or under-collection due to hydrodynamic effects complicate interpretation (Yu et al., 2001). Moreover, particles are often pooled in sediment traps, making it hard to characterize the origin, size, and composition of the individual particles. An exception are gel traps, which are traps filled with a viscous, inert gel that slowly decelerates and isolates sinking particles, allowing investigation of individual particles (Jannasch et al., 1980; Waite and Nodder, 2001; Thiele et al., 2015; Flintrop et al., 2018).

Measurements of natural, particle-binding radioisotopes, including Thorium-234 and Polonium-210, can be used to estimate upper-mesopelagic particle fluxes on timescales of weeks to months (e.g., Buesseler, 1998; Le Moigne et al., 2013). These estimates do not, however, provide information about the nature of the particles responsible for the measured fluxes, and rely on assumptions regarding the conversion from radioisotopes to carbon that are difficult to evaluate (Waite and Hill, 2006). Moreover, radioisotopic estimates of particle flux are complicated by potential non-steady state dynamics and physical transport (Ceballos-Romero et al., 2018). Measurements are costly, as they need to be conducted during individual oceanographic cruises, and need to be calibrated with sediment traps.

Large-scale assessments of sinking particles in the marine carbon cycle focus on measuring dissolved biogeochemical tracers such as nutrients, oxygen or pH (e.g., Schlitzer, 2002; Gehlen et al., 2006; DeVries et al., 2012; Weber et al., 2016). These tracers reflect the net production and degradation of organic material combined with processes of particle transport and circulation. Major observational programmes that use dissolved tracers include GO-SHIP (Global Ocean Ship-Based Hydrographic Investigations Program, www.go-ship.org) and SOCCOM (Southern Ocean Carbon and Climate Observations and Modeling, https://soccom.princeton.edu), which uses biogeochemical sensors on profiling floats. The use of dissolved tracers has two advantages; the sensor techniques are relatively advanced, and estimated rates are integrated over space and time thus reducing observational needs. However, these approaches are unlikely to deliver any predictive understanding of how particle fluxes will respond to environmental change as they do not identify the processes that control the sinking and transformation of particles, nor the variability on interannual timescales. The largest global effort to characterize particle fluxes more directly was the Joint Global Ocean Flux Study (JGOFS), which carried out both surveys and process studies over more than a decade (e.g., Buesseler, 2001). It made major advances in connecting surface production to mesopelagic and deep particle fluxes, primarily via biogeochemical measurements, but also by including their correlation with bulk optical properties such as light transmission (e.g., Bishop, 1999; Gardner et al., 2006) and some early applications of photographic imaging techniques (e.g., Walsh and Gardner, 1992; Diercks and Asper, 1997; Bishop et al., 2002).

Recent developments in in situ optical devices for measuring particles offer the opportunity to advance the progress that began during JGOFS and other programmes. They are now much smaller, require much less power, and offer greater capabilities. Bulk optical property sensors, such as light transmission and scattering are available in multiple frequencies. Imaging systems have greater resolution and can be combined with image processing techniques for particle recognition to estimate particle type, size, and distribution. Optical devices for particle measurements can be used from ships (e.g., Herman, 2004; Davis et al., 2005; Picheral et al., 2010) or installed on remote platforms (e.g., moorings and Argo floats; Checkley et al., 2008; Rembauville et al., 2017). Several systems exist that can be deployed down to 6,000 m depth, and integration into standard CTD-rosettes allows for routine deployments as part of oceanographic surveys. Optical devices for measuring particles can provide high-resolution descriptions of particle abundances, sizes, and types (e.g., fecal pellet, diatom aggregate, mucous structures, zooplankton), which inform about particle origin and formation. The high-frequency spatial and temporal information collected by optical devices now allows inferring particle transformation mechanisms from observations on particle abundance and size-distributions at different water depths or from changes of their morphology over time. Imaging systems are also well-suited to investigate zooplankton-particle interactions, as they often allow for the simultaneous assessment of particles and zooplankton (e.g., Möller et al., 2012; Christiansen et al., 2018). Lastly, the use of optical devices for measuring particles has become increasingly attractive as they are continuing to become more affordable and technically more feasible.

The interpretation of optical measurements can be complicated as different optical devices generally measure distinct optical properties that are difficult to interpret and
compare. A lack of standardization in data analysis algorithms further impedes the direct comparison of different datasets. These issues are exacerbated during the translation of optical particle properties into flux estimates owing to a current lack of understanding of how particle optical properties such as size and type relate to particle sinking velocity and carbon content (Kriest, 2002; Guidi et al., 2008; Iversen et al., 2010; McDonnell and Buesseler, 2010; Le Moigne et al., 2013; Laurenceau-Cornec et al., 2015a; Nowald et al., 2015; Ramondenc et al., 2016).

Scope of Review

The advance of optical technology, autonomous ocean systems, and data processing power now promises a major leap in our understanding of the biological carbon pump. An important challenge now is to systematically improve the use of optical devices for measuring particles, which includes the comparison and inter-calibration of the outputs of available optical devices, as well as the collation and distribution of knowledge on how to efficiently convert optical information (abundance, size, and types of particles) into particle flux estimates. In this review, we provide an overview of the general issues that occur when trying to (1) interpret optical in situ measurements of particles in the mesopelagic zone and (2) convert these measurements into fluxes. We briefly present some of the common optical devices used for particle flux characterization in the mesopelagic zone and discuss their capabilities and limitations. A summary of currently commercially available optical devices for plankton research has been compiled by Lombard et al., (2019; Table 1), and most of these devices can be used for particle flux studies. The aim of this review is to give scientists the background needed to maximize the output of these optical devices for estimating particle flux and understanding particle dynamics.

OPTICAL IN SITU MEASUREMENTS OF PARTICLES

Unlike other measurements such as primary production, oxygen, salinity, or Chlorophyll a, for which there exist standard sampling and analysis protocols, there are currently no standards for optical particle sampling, data analyses and data deposition. Differences exist in data acquisition owing to the various optical devices and techniques, and data processing is often left to personal preferences, including image/signal analyses, classification and conversion algorithms. Hence, there is a great need for standardization to enable comparison of data collected by different instruments or analyzed by different scientists.

Data Acquisition: Particle Detection Methods

The size of a particle determines the detection method that is appropriate. For example, small (<2 µm) phytoplankton are much more abundant than large (>2 mm) zooplankton, so phytoplankton abundance can be measured using relatively small sample volumes. The small particle size makes detailed classification difficult as sufficient resolution of the shape at such small scales is technologically difficult. In situ characterization of small particles is therefore often restricted to estimates of abundance and biomass based on optical “bulk” properties such as transmission, backscatter, and fluorescence.

Large particles are rarer in abundance and contribute less to total biomass than small particles (Sheldon et al., 1972) and therefore require that a larger volume of water is measured. These particles have traditionally been collected with nets, bottles, or pumps, and identified visually based on shape or biochemically based on elements (e.g., diatoms via biogenic silica). The use of imaging systems that build on these classification methods is therefore convenient. Imaging can be based on photographic or holographic technology. Yet, whilst particle size and abundance can be retrieved relatively quickly from images, a more detailed classification still requires time-consuming manual identification. This step will become much faster in the coming years with the rapid advances in machine learning tools.

The decision when to use bulk water properties, like transmission, vs. imaging is fluid. The signal from an imaging system could be interpreted in a similar fashion to “bulk measurements” (i.e., looking at total frame properties rather than specific “regions of interest”), whilst anomalies in bulk signals (e.g., spikes in backscatter output) can be used to infer the size of particles (Briggs et al., 2011, 2013). “Hybrid” systems exist, such as the Laser Optical Particle Counter (LOPC; not commercially available anymore), which can combine several individual “one-pixel” photodetectors to generate two-dimensional information that can be used to investigate particle shape (Jackson and Checkley, 2011; Petrik et al., 2013).

Broadly, we can distinguish between four types of optical device: (a) single photodetectors, (b) simple photodetector arrays, (c) holographic systems, and (d) photographic systems. Box 1 explains the principles behind each of these types. Table 1 gives an overview of example devices for each type, highlighting the target range, classification level, typical sampling volumes, relevant threshold settings, and type of illumination.

Data Processing: Sizing

Once a suitable device has been chosen and optical particle measurements have been acquired, the next step is often to determine particle size. Individual particle measurements are especially useful for studying sinking particles because bulk measurements can be dominated by small, non-sinking particles. Nonetheless, bulk optical measurements can provide some amount of particle size information.

Single photodetectors do not provide direct particle size estimates, yet spike height (i.e., the high-frequency variability in the beam attenuation or backscattering signal) can be used as an indicator of particle size (Briggs et al., 2013; Box 1a). Further, the spectral slope of the beam attenuation coefficient (Boss et al., 2001) and the spectral slope of the backscattering coefficient (Slade and Boss, 2015) can be used to estimate mean particle diameter and the slope of the particle size distributions. Currently, in situ validation of these methods is either limited or conflicting (Reynolds et al., 2016), so they should be used with caution without further validation.
**BOX 1 | Basic principles of optical device types.**

(a) Single photodetector

A single photodetector measures a bulk optical property, such as optical backscattering or beam attenuation in a volume of water. These properties are empirically correlated with particle concentration in the ocean (e.g., Bishop, 1986; Reynolds et al., 2016) and, where organic particles dominate, particulate organic carbon (POC; e.g., Gardner et al., 2001; Cetinić et al., 2012). While a single photodetector cannot distinguish individual particles within its sample volume, information about particle size can in practice be extracted from a high-resolution time series or vertical profile. This interpretation is possible because a single large particle (> 150 μm) passing through a small sample volume causes a brief jump, or “spike,” in particle concentration (Figure 1, left panel). In the mesopelagic, such particles are generally rare enough relative to the sample volumes of commercial transmissometers that their optical signals can be completely separated from the background of smaller particles (see Figure 1, right panel). Then, either their numerical concentration (Rembauville et al., 2017) or POC concentration (Briggs et al., 2011) can be calculated. When combined with a sinking velocity estimate, the latter can be converted to an estimate of POC flux. Alternatively, spike height can be correlated with particle cross-sectional area, allowing estimates of particle size. This principle has been used to estimate mean particle diameter at high particle concentrations (Briggs et al., 2013), and it could also be applied to individual spikes at lower concentrations.

![Image](image.png)

**FIGURE 1 | Working steps to derive small and large particles from single photodetectors.** A median filter is fitted and assumed to be representative of small particles. Spikes are caused by large particles passing through the sampling frame.

(b) Simple photodetector array

Simple photodetector arrays use a similar principle to single photodetectors. The difference is that a number of photodetectors are arranged in a way that allows the extraction of additional particle properties. The most prominent example is the Laser Optical Plankton Recorder (LOPC; Herman, 2004). Thirty-five photodetectors (“photo-elements”) are arranged vertically and measure the absorbance of a laser sheet. As the instrument is towed through the water, particles passing through the light sheet block a portion of the light, and the receiving photo-elements register the change in voltage as digital size and transparency (Checkley et al., 2008).

Two types of particles are registered by the LOPC control software: single-element particles (SEPs) and multi-element particles (MEPs). SEPs are defined as particles occluding one or two photo elements, MEPs occlude three or more. For SEPs, only size information is recorded. For MEPs, in addition to their digital size, the occlusion of each photo-element is recorded providing information on shape and transparency.

The digital size of SEPs and MEPs is converted into equivalent spherical diameter (LOPC-ESD) using the manufacturer’s calibration with black spherical beads (Herman, 2004; Checkley et al., 2008; Gaardsted et al., 2010). The LOPC-ESD is thus the diameter of a particle equivalent to the diameter of a black sphere that would block the same amount of light, which means, e.g., that a large, transparent particle can have a relatively small LOPC-ESD. The LOPC per-se does not distinguish between particle types. However, for MEPs a separation based on transparency and/or shape can be done (Jackson and Checkley, 2011; Bacedov et al., 2013, 2014) based on the ratio of LOPC-ESD to occluded diameter (the width of all photo elements occluded); Jackson and Checkley, 2011; Trudnowska et al., 2014). A small LOPC-ESD: occluded diameter ratio means that particles are transparent and/or amorphous. To relate size to organisms, correlation relationships have been determined from organism concentrations collected with plankton nets simultaneously with LOPC observations (Gaardsted et al., 2010; Ohman et al., 2012; Marcolin et al., 2015). Another prominent instrument using a simple photodetector array is the LISST (Sequioa Scientific Inc.), which measures the angular distribution of forward-scattered laser light using concentric ring detectors (Gartner et al., 2001). Additional published methods exist that also use near-forward scattering for estimating particle size distributions (Twardowski et al., 2012). These are bulk property detectors, which can be processed to estimate the size distribution of equivalent spherical particles. This method depends on different assumptions than those applied to blocking of a beam, and thus the sizes are not directly comparable to those from the LOPC or other imaging approaches.

(c) Holographic system

Holographic systems record a digital hologram of the particles in a water sample. To do so, a sample volume is illuminated with a collimated laser. As the beam hits a particle, light is scattered and interferes with the incident light of the laser beam. The resulting interference pattern is recorded by a camera (e.g., a charge-coupled device) and can be used to reconstruct a holographic image of the particle (e.g., Brichard et al., 2013; Talapatra et al., 2013; Nayak et al., 2018).

While this approach sounds as if the hologram would provide information on the 3D structure of each particle, this is only partially true. Particles are holographically imaged only from one side with the structure of the “backside” of the particle remaining unknown. More importantly, the 3D information is largely disregarded during the data processing routine. Rather, the holographic information is used to precisely calculate the size and position of the particle within the sample volume: The digital hologram is reconstructed providing monochrome in-focus images of each particle corrected for its position in the z-axis (i.e., accounting for the distance between particle and camera). The reconstructed images can then be analyzed using image analysis programmes in the same way as used for images by photographic devices (Graham and Nimmo Smith, 2010; Davies et al., 2015).
For some devices, size is an explicit characteristic based on the assumptions of the method. For example, both the LISST-Holo and the LOPC report the equivalent spherical diameter (ESD) of detected objects without the user having to decide on the detection method. The advantage when size is provided as a standard output is that comparison of observations collected by different users and/or different devices of the same type are fairly straightforward. The user bias and limitations of the device and the exact definition of the size parameter. For example, the “ESD” provided by LISST-Holo is based on the pixels of the reconstructed particle. The area of the particle (in pixels) is used to find the diameter of a circle with the equivalent area (Box 2). The size of a particle is hence derived from a 2D image, similar to more traditional photographic imaging, with the advantage that all particles are in focus and size can be determined much more precisely (Graham and Nimmo Smith, 2010). For the LOPC, the light attenuated by a particle is converted into ESD using the manufacturer’s calibration with black spherical beads (Herman, 2004). The LOPC-ESD is thus the diameter of a black sphere that attenuates an equivalent fraction of light, which means that a large, transparent particle can have a relatively small LOPC-ESD.

For imaging devices (holographic or photographic systems), particle detection and subsequent sizing rely heavily on background subtraction, threshold settings (Giering and Hosking, in review), edge detection, and segmentation algorithms. The ultimate choice is often left to the user, introducing operator bias on a very basic level before any further analysis is carried out. Next, the user needs to decide on a metric to report size. As mentioned above, ESD is often the preferred metric, though many others exist (equivalent circular diameter (ECD), equivalent circular perimeter diameter, Feret diameter, length and width). Note that many reported ESDs are based on ECDs as most $in situ$ systems work with particle area and not volume. Last, when ESD and ECD are calculated, a decision has to be made whether to include or exclude possible holes in the imaged particle, which can dramatically influence the final size estimate (Box 2).

Besides the difficulties in algorithm and metric choices, there are technical and practical issues for each of the different devices to capture the real size of a particle. The concept of
### TABLE 1 | Examples of instruments used for estimating particle flux.

| Device examples   | Name                                          | Target range | Information obtained | Classification level** | Automatic sizing? | Typical sampling volume (mL/image) (max) | Frames per second (max) | Type of illumination | Selected references |
|-------------------|-----------------------------------------------|--------------|----------------------|------------------------|-------------------|------------------------------------------|------------------------|----------------------|---------------------|
| Single photodetector | **OBS** Optical Backscatter Sensor** | –           | signal intensity     | 1 or 2*a   | N                | various; e.g., ~1 | various; e.g., ~1–4   | LED; various wavelengths | Briggs et al., 2011 |
|                   | **Transmissometer**                          | –           | signal intensity     | 1 or 2*a   | N                | various; e.g., ~5 | various; e.g., ~1–10 | LED; various wavelengths | Briggs et al., 2013 |
|                   | **Fluorescence**                             | –           | signal intensity     | 1 or 2*a   | N                | various; e.g., ~1 | various; e.g., ~1–4   | LED @ 470 nm (blue)    | Briggs et al., 2011 |
| Simple photodetector array | **LOPC** Laser Optical Plankton Counter | 100–4,000 (35,000) µm | diameter + transparency | 3   | Y                | NA                      | NA                     | 670nm (red) diode, focussed by a cylindrical lens producing a light sheet | Herman, 2004 |
|                   | **LISST Laser in Situ Scattering Transmissometer** | LISST-B: 1.25–250 µm; LISST-X: 2.5–500 µm; LISST-100: 5–200 µm | particle size distribution | 2   | Y                | –                      | 25                     | Solid state diode laser @ 670 nm (red) | Gartner et al., 2001 |
| Holographic system | **HoloSea** – (1.5) 20–2,000 µm | interference pattern to recognize size and shape | 4   | Y                | 0.1                    | 22                     | 405-nm (violet)            | http://4-deep.com/products/submersible-microscope/ |
|                   | **LISST** Laser in Situ Scattering Transmissometer | (4) 25–2,500 µm | interference pattern to recognize size and shape | 4   | Y                | 1.86                    | 20                     | Solid state diode laser @ 658 nm (red) | https://www.sequoiasci.com/product/lisst-holo/ |
| Photographic system | **VPR** Video Plankton Recorder >100 µm | Image, color | 5   | N                | 0.5–100 (depending on magnification settings) | 15–25            | strobe ring light synchronized with camera | Davis et al., 2005 |
|                   | **CPICS** Continuous Particle Imaging and Classification System | (0.9x) 200–10,000 µm | Image, color | 5   | N                | 0.33                    | 10                     | polarized light            | http://oceancubes.whoi.edu/instruments/cpic.html |
|                   | **UVP** Underwater Vision Profiler | 60–10,000 µm | Image, monochrome | 4   | N                | 1,020                   | 6                      | LED @ 625nm (red) in two glass cylinders either side of the imaging field | Picheral et al., 2010 |
|                   | **ISIS** In Situ Ichthyoplankton Imaging System | 60–130,000 µm | Image, monochrome | 4   | N                | 600                     | 3                      | back-illumination, 455 nm (blue) | https://www.planktonimaging.com/isis-optical-system |
|                   | **SIPPER** Shadowed Image Particle Profiling and Evaluation | >200 µm | Image, monochrome | 4   | N                | NA                      | NA                     | 635 nm (red) laser sheet | Samson et al., 2004 |
|                   | **LOKI** Lightframe On-Sight Keyspecies Investigation | 50–2,000,000 µm | Image, monochrome | 4   | N                | 2.6                     | 30                     | collimated laser, from the side; LED | Schmid et al., 2016 |

**Different magnifications available. Quoted details are for the magnification that is most suitable for marine snow.**

**Classification metric:**
1. Count only.
2. Size information.
3. Size + additional information (e.g., transparency) for rough grouping.
4. Black and white image or similar complex information for detailed grouping.
5. Color image, 3D or similar complex information for highest classification.

*a depending on processing.
size is in principle simple, however, in reality very complicated because of the complex shapes of marine particles, such as copepods with their legs and antennae, twisted diatom chains with their spines, radiolarians with delicate spikes, aggregates and feeding structures with complex shapes and empty cavities, or exopolymer particles, which are often undetected owing to their transparency. Not all imaging systems can resolve the necessary detail to capture such complex structures. Moreover, most imaging devices take a 2D image, which can substantially misrepresent the true size and shape of the original 3D particle.

For many devices with a relatively large depth of field, such as the Video Plankton Recorder, the z-position of a particle is unknown, meaning that the true size is unknown: A small particle close to the camera can have the same apparent size as a large particle further away (Figure 3). This imprecision is, however, likely averaged out over sufficient data provided that particles are illuminated consistently regardless of their z-position; i.e., small particles are as likely detected close to the camera than further away. For shadowgraph systems, the imaging of silhouettes using collimated light ensures that a particle is always imaged at the same size, regardless of how close it is to the camera (Cowen and Guigand, 2008; Ohman et al., 2019). Alternatively, a telecentric lens configuration can be used. In particle-rich environments, overlapping of objects might become an additional problem for devices with a large depth of field (Figure 3). Problems might also arise when only a small part of a larger particle is captured or illuminated, leading to a potential underestimation of particle size. Diffusion of light as it travels through the water from the particle to the sensor can cause a “halo” effect in which small particles appear larger than they really are. This effect can be corrected for with a size specific conversion factor between pixels and size (Picheral et al., 2010).

Estimating the absolute size of a particle using optical devices will likely always remain a challenge owing to the technical limitations and overall assumptions of the different methods. A big step forward to making data more comparable is increased transparency and standardization in data acquisition, analysis and description. We therefore recommend the following:
1. **Instrument.** The determination of size can be influenced by the type of device and/or its current configuration. Thus, to help end-users put data into context, we recommend that the instrument name and serial number are clearly reported. To reduce ambiguity, this should ideally be accompanied by an identifier from a standardized list, such as the SeaVox Device Catalog. Furthermore, relevant technical details should be explicitly mentioned, such as illumination type and frequency, sampling volume and frequency, and relevant calibrations.

2. **Image resolution.** The final estimated size of particles is partly determined by the resolution of the image. A higher resolution allows not only the detection of smaller particles but also the description of more complex shapes. The resulting size estimate may vary significantly. A clear description of the image resolution, pixel size, and particle detection criteria (i.e., minimum and maximum particle size, see also Box 1d) will help to compare datasets.

3. **Image/signal processing.** All details of image or signal processing should be reported and the code made available (e.g., via GitHub). Image and signal processing steps may include background subtraction, noise reduction, dilation and erosion techniques, object recognition, and segmentation.

4. **Thresholding and edge detection** (for holographic and photographic systems). The most appropriate threshold (e.g., for black and white images the gray-scale value for the transformation of the image into a black-white binary field representing background and particle, which are used to calculate particle statistics) or edge detection algorithm should be determined. This could be done by using calibration beads or real aggregates and plankton of known size. Uncertainties should be clearly stated in the methods. Alternatively, a sensitivity analysis using a range of thresholds or algorithms and their effect on estimated particle size should be carried out.

5. **Size metric.** The metric used for describing size should be clearly stated. We recommend using ESD/ECD as this is the most widely used metric (Box 2).

6. **Data deposition and sharing.** (For a detailed discussion, see section data deposition and sharing). Particles imaged at high enough resolution to allow identification (generally ∼30 × 30 px) should be saved as separate "vignettes" (images of individual particles extracted from the frame) and made publicly available to allow future image-based analyses. A unique identifier or hashtag could be assigned to the particle. In addition, a file in text-format containing the measurements on all individual particles (e.g., the parameters given by image analyses programmes such as imageJ, Matlab’s Imaging Processing Tool Box, or the plugins for Python’s image analyses) should be provided.

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**Data Processing: Classification**

The size and abundance of particles is useful information and sufficient for many applications (e.g., identification of changes in particle populations with depth and its possible links to disaggregation and flux attenuation; Stemmann et al., 2004a,b; Jouandet et al., 2011; Kiko et al., 2017). To fully understand ecosystem processes, however, the identity of the particles is key: e.g., whether it is an amorphous aggregate or an individual zooplankton. Several devices have been designed specifically for quantification and identification of zooplankton and marine snow [e.g., the Underwater Vision Profiler (Picheral et al., 2010) or the Video Plankton Recorder (Davis et al., 1992, 2005)], often targeting the mesozooplankton size range (∼0.2–2 mm). For these devices, a visual classification that is based on morphological features is very fruitful, and taxonomic guides can often be used to identify zooplankton to a fairly high taxonomic level (sometimes down to species). For single photodetectors or simple photodetector arrays, classification is much harder as information on particle type is very limited. However, with a combination of different devices (for example different wavelengths of backscatter, or backscatter combined with fluorescence measurements), some level of classification can be achieved, e.g., the chlorophyll fluorescence to red light backscatter ratio can be used to estimate the relative abundances of algal vs. non-algal particles (Iversen et al., 2010; Barbieux et al., 2018; Schallenberg et al., 2019).

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1. [https://www.bodc.ac.uk/resources/vocabularies/vocabulary_search/L22/](https://www.bodc.ac.uk/resources/vocabularies/vocabulary_search/L22/)
Methods Based on Visual Classification

While underwater imaging of marine particles has recently become widely used, image analysis software tools have lagged behind hardware developments (Hu and Davis, 2005). The large amount of visual data emerging from in situ plankton samplers, benchtop systems, and cabled underwater observatories require automatic procedures. Besides saving time, automation avoids bias and errors inherent to the fatiguing process of manually classifying vast amounts of images (Culverhouse et al., 2003). Several automated methods for the analysis and classification of plankton and particle images have been developed, but their limited accuracy (around 80% on a realistic number of classes in recent attempts; Zheng et al., 2017) still requires significant manual verification to obtain accurate counts and identification or to further identify behavioral modes like trophodynamic plankton—particle interactions (Möller et al., 2012).

Image processing typically proceeds along the following steps. (1) When necessary, focus detection is used to restrict the effort to in-focus particles by eliminating out-of-focus frames. (2) Objects (i.e., “regions of interest”) are isolated from full frames using binarization, segmentation, and/or connectivity routines. For underwater image analysis, this step may be challenging due to variable illumination, scales, and orientation of objects and their non-rigid deformation (Py et al., 2016). Also, large organisms may exceed the sampling frame (and therefore be cut) or obstruct other, smaller objects. (3) Once detected, regions of interest are processed individuately to compute size and extract morphological features used for automatic classification. These regions of interest are often saved as separate files (“vignettes”) for classification techniques that require an image as input.

The classification process then starts with the manual labeling of a set of images of objects by human operators, to provide examples on which machine learning classifiers can be trained. This training (or learning) set has to be as representative of the full data set as possible. Larger training set sizes usually significantly improve the performance of the classifier, although it saturates at some point (Gorsky et al., 2010; Ellen et al., 2015). Therefore, this step is time-consuming.

A wide variety of morphological features and classifiers have been trained to sort particle images automatically into plankton taxa or particle categories. Their full review is beyond the scope of this paper; Benfield et al. (2007) wrote a good synthesis of the field and a more recent list of papers is in the introduction by Zheng et al. (2017). Briefly, the morphological features can be global descriptors of the object (such as the area, the average gray level, etc.; Grosjean et al., 2004; Sosik and Olson, 2007) or the concatenation of local shape and texture characteristics, such as Fourier descriptors (Tang et al., 1998) or Histograms of Oriented Gradients (Bi et al., 2015). The former is more immediately interpretable with respect to the overall characteristics of the object (big vs. small, dark vs. light, etc.), in particular for ecologists; the latter is more common in the image analysis domain because it often yields better classification results. Once the set of features is chosen, the difference between the various classifiers (Support Vector Machines, Random Forests, Artificial Neural Networks, etc.) is usually small (Grosjean et al., 2004). Rather, accuracy is gained by introducing richer input images (such as color images), by combining different types of features (such as shape and texture; Hu and Davis, 2006), or by combining classifiers into ensemble models (Ellen et al., 2015; Zheng et al., 2017).

Finally, recently developed Convolutional Neural Networks (CNNs) have scored higher than any other technique on major image classification challenges (Krizhevsky et al., 2012). CNNs learn both the extraction of relevant features directly from the images and their classification. Thanks, in part, to an online machine learning competition that prompted the interest of the computer vision community and provided a standardized test dataset (https://www.kaggle.com/c/datasciencetbowl, 2015), CNNs are increasingly used to classify marine particle images (e.g., Py et al., 2016; Luo et al., 2018). Training a CNN is computationally intensive and requires a large number of training examples. Fortunately, this constraint can be partly alleviated by kick-starting the process using a model pre-trained on another dataset because the features they extract are often quite generic (transfer learning; Orenstein and Beijbom, 2017; Lumini and Nanni, 2019), by guiding human operators to grow the training set with only problematic images (active learning; Bochinski et al., 2019), or by post-processing the features output by the network to learn new classes after the training step (low-shot learning; Schröder et al., 2019).

While these techniques hold promise, the classification of marine particle images is a challenging task because the image quality is often suboptimal for smaller particles, the range of sizes of particles is huge, and a few classes are much more numerous than others, which makes it difficult to tune the classification performance on the rarer, often interesting, classes. Moreover, novel classes that occur in the sample but do not occur in the learning set will also be misclassified.

Methods Based on Optical Properties

For single photodetectors or simple photodetector arrays that do not capture particle images, or for particle images that are small relative to image resolution, a combination of optical properties and/or a size parameter can provide information useful for very broad particle classification.

For example, the ratio of backscattering to beam attenuation is related to the ratio of organic to inorganic matter (Jamet et al., 2018) and to the refractive index, which can indicate plankton—particle interactions (Twardowski et al., 2001). The ratio of chlorophyll fluorescence to optical backscattering can inform about the contribution of phytoplankton to particulate matter; and bulk birefringence can be a proxy for suspended CaCO₃ concentrations (Guay and Bishop, 2002).

Another example is the combination of the spike signals detected by several single photodetectors, such as backscattering and fluorescence sensors. These sensors can identify individual particles larger than ~150 μm by the brief spikes they induce when passing through the sensor’s sampling frame. The ratio of fluorescence spikes to backscattering spikes in a population of particles can distinguish aggregates of phytoplankton from other large particles (Briggs et al., 2011). The ratio of spikes of different wavelengths of backscattering, or beam attenuation spikes...
to backscattering spikes, likely contains further information, although this information has not yet been investigated.

The combination of size and transparency can inform whether the particle is an aggregate or zooplankton (e.g., Petrik et al., 2013; see also Box 1b). Devices measuring fluorescence can distinguish particles containing chlorophyll (phytoplankton) and/or phycocerythrin (cyanobacteria) from other particles, and cameras measuring birefringence have been used to distinguish particles containing calcium carbonate (Bishop et al., 2016). For particle images, particle brightness (or light attenuation for light field images) and color are optical properties that can help distinguish particle types (Wilson et al., 2008).

Data Deposition and Sharing

As a next step, processed data should be deposited and made freely available for future research. Currently, there is no standardized procedure to deposit data on optical particle measurements, and there is a clear lack of agreed metadata, particle descriptors, data formats, classification criteria, and accessibility. Any community agreement should consider the FAIR Guiding Principles (Wilkinson et al., 2016), a set of standards to improve the findability, accessibility, interoperability, and reusability of data. As discussed above, we need increased transparency and standardization for data sharing, comparison and future data interpretation. Embedding well-structured metadata and data provenance information in data workflows are fundamental to ensuring user trust in data and any data products generated (Buck et al., 2019). As discussed in section Data Processing: Sizing, using common standards such as controlled vocabularies to annotate data help reduce ambiguity and facilitate interoperability. Many journals require datasets to be cited with Digital Object Identifiers to support scientific results. In addition, persistent identifiers (PIDs) are well-established in the academic community to improve transparency, and there are international efforts to use PIDs to identify “real-world” instruments. Such tools could help a user to relate back to the manufacturer’s calibration or configurations of a device to put particle size data into context. Several marine observational programmes use agreed formats that are machine-readable and enriched with common standards to facilitate data sharing, automation and comparison within a community.

For “live” planktonic particles a start point is the widely-adopted Darwin Core format used by the Ocean Biogeographic Information System (OBIS) for biogeographic data (Nakamura et al., 2017), but it lacks many descriptors necessary for it to be directly applied to the wide range of particles in the ocean. The Argo Climate Forecast (CF)-NetCDF is more flexible. However, NetCDF requires complex software and some level of expertise to access it. The World Wide Web Consortium’s (W3C) “CSV on the Web” offers the same benefits as NetCDF but in a simpler format that may be more accessible to the biogeochemical community, requiring standard software such as Microsoft Excel. Ideally, all raw images (i.e., full frames) should be saved. However, raw images require considerable storage capacity and often contain mostly empty space. Instead, a common practice is to save segmented individual particle images (“vignettes”). Vignettes should be saved to the highest resolution to facilitate re-analysis and avoid deterioration if compressed. Vignettes should be saved on a global databank or distributed database, allowing others to carry out their routines on the raw particle images, particularly machine learning and image processing for classification. While a similar approach has been started (e.g., EcoTaxa, https://ecotaxa.obs-vlfr.fr; Picheral et al., 2017), a current big limitation is the required storage capacity, especially with particle imaging becoming ever more popular. EcoTaxa alone already hosts >80 million vignettes (as of May 2019).

In addition, funding bodies that require data deposition often prefer only the particle descriptors of each image in text-format as it is less memory intensive. When describing images in text-format, there is substantial information loss. For example, particles are often described in terms of size and particle type only. Besides the inconsistencies and ambiguities of sizing (see Section Data Processing: Sizing), potentially crucial information (e.g., color, shape, and texture) are lost. We therefore recommend depositing raw images and/or vignettes whenever possible. Finally, a unique identifier or hashtag could also be assigned to each particle image to allow tracking of information regarding this particle. e.g., different scientists might carry out different image processing steps or assign different classifications to the same particle depending on their data analysis procedures or classification scheme. Furthermore, more detailed analysis on single particles, including sinking velocity measurements and carbon content, might be available for some particles. If a unique identifier is assigned, identification and detailed information could be harvested and used for further in-depth analyses and meta-analyses.

Any software and image analysis codes used to process images should be assigned a PID and cited in accompanying documentation to improve transparency to users. Image analysis codes should also be made available using code-hosting facilities (such as GitHub).

FROM IMAGE TO PARTICLE FLUX

Optical particle measurements and classification are only the first steps to understanding particle dynamics in the ocean and the biological pump. The next steps involve the conversion of this information into flux estimates. Particle fluxes are typically calculated as

\[ F = c \times w_{avg}, \quad \text{or} \]
\[ F = \Sigma (m_{part} \times w_{part}) / V_{sample} \]

where \( F \) is the matter flux (in mg m\(^{-2}\) d\(^{-1}\)), \( c \) is matter concentration (typically in mg m\(^{-3}\)), \( w_{avg} \) and \( w_{part} \) are the bulk and individual particle sinking velocity, respectively (in m d\(^{-1}\)), and \( m_{part} \) is the matter content of an individual particle (in mg C) with \( \Sigma (m_{part} \times w_{part}) \) being the sum of all particles within a known sampling volume (\( V_{sample} \) in m\(^{-3}\)).
These equations can be further expanded to explicitly separate the composition of particles from their masses and numerical abundance to represent the fluxes of specific components, e.g., for the particulate organic carbon (POC) flux:

\[ F = \frac{\sum_part(n V_{part} \rho_i w_{part})}{V_{sample}} \]  

(3)

in which \( n \) is the number of particles in that class (\( n = 1 \) for individuals), \( V_{part} \) the particle volume (typically in mm\(^3\)), and \( \rho_i \) is the density of the constituent \( i \) in the particle (e.g., mg C mm\(^{-3}\)). The sum is then taken over all particles and normalized to the sampled volume.

Optical devices for particle measurements provide great direct estimates of particle numbers (\( n \)) and good estimates for particle volume (\( V_{part} \)), but they are not able to provide direct information for chemical particle composition (i.e., \( \rho_i \)), with \( i \) being POC or any other component of interest such as carbonate or silica minerals. Hence, all optical devices require an estimation of the approximate elemental particle composition in order to estimate POC concentrations and, ultimately, flux.

Particle sinking velocities (\( w \)) can be estimated using a range of methods (Section Sinking Velocity); however, most optical devices are not capable of providing sinking velocities and rely on additional data or assumptions. Two bulk optical methods bypass the need for information on sinking velocities by quantifying particle fluxes more directly either by optically measuring particle accumulation on a horizontal surface ("optical sediment trap"; Bishop, 2004; Estapa et al., 2013; Bourne et al., 2019) or by tracking the accumulation of integrated particle concentrations below a depth threshold (Dall’Olmo and Mork, 2014).

Both chemical particle composition and sinking velocity can vary greatly. POC content can be as low as \( \sim 1% \) by weight in particle fluxes dominated by lithogenic and biogenic minerals (e.g., Armstrong et al., 2002; Klaas and Archer, 2002) and as high as 40% in aggregates (e.g., Allardredge, 1998). Sinking velocities typically range from 1 to 1,000 m d\(^{-1}\) (e.g., Kriest and Evans, 1999; Turner, 2002; Laurenceau-Cornec et al., 2015a). Hence, the biggest bottleneck associated with translating optical particle measurements into accurate flux estimates is the uncertainty in these two parameters. We explore some approaches to measure particle composition and sinking velocity in the following sections.

**Organic Matter Content and Concentrations**

The approaches for estimating bulk matter concentrations (\( c \)) and matter content of individual particles (\( \rho_i \times V_{part} \)) differ in that the former is an estimate based on the entire sample whereas the latter is specific to objects of interest.

Bulk estimates are based on the entire particle field and how they affect the optical properties of the water volume they are in. In simple terms, the more light there is, the more light will be scattered and absorbed by the particles (i.e., less light will be transmitted through the water). Devices that measure scatter and transmission in one way or another can therefore be used to infer bulk matter concentrations. Empirical data have shown that both particulate backscattering (\( b_{bp} \)) and attenuation (\( c_p \)) are good proxies for the concentration of POC in pelagic environments (Gardner et al., 1993; Bishop, 2009; Bishop and Wood, 2009), though the relationships are dependent on the local particle populations and potentially on methodological differences such as sensor calibrations (Cetinici et al., 2012). To use such bulk proxies, it is therefore preferable to take POC samples along with the optical property measurements to derive a site-specific relationship between the two parameters. In addition, natural variability in lithogenic matter concentrations (e.g., sediment load) can strongly affect the relationship between POC and \( b_{bp} \) or \( c_p \) (e.g., Reynolds et al., 2016), so these optical proxies cannot be used where sediments dominate the optical signal.

For individual particles, determination of the organic matter content becomes more complex. The typical work process is as follows. The optical device images a volume of water, the individual particles are sized (section Data Processing: Sizing), and the size estimate is converted into POC content using empirical relationships. As imaged particles are rarely also captured and brought to the surface for elemental analysis, most studies rely on published size-to-POC conversion equations. The most commonly applied equation is the one by Alldredge (1998). Alldredge (1998) photographed particles (>0.5 mm in diameter) in situ from the surface ocean in the Santa Barbara Basin, pooled these according to size and type (three size classes and four types), and analyzed them for POC, particulate organic nitrogen and dry mass. Organic matter content of particles clearly increased with increasing size (\( p < 0.001, R^2 = 0.86, n = 25 \)), though larger aggregates contained less matter per unit volume than smaller aggregates (Alldredge, 1998). Later laboratory experiments have shown that the size-to-POC conversion for aggregates is strongly dependent on the phytoplankton community composition (Ploug et al., 2008a; Iversen and Ploug, 2010) and can vary substantially from the size-to-POC conversion for zooplankton fecal pellets (Gonzalez and Smetacek, 1994). These observations suggest that there is substantial variability in the size-to-POC conversions depending on plankton community structure, particle type, season, location and likely depth (Kriest and Evans, 1999; Kriest, 2002; Iversen et al., 2010).

On the most basic level, the uncertainties associated with converting a single particle image into POC concentrations are not different from the problems encountered for bulk analysis. The main difference is that the sample size of large particles imaged using holographic or photographic devices is relatively small compared to the sample size imaged by e.g., backscatter sensors. Backscatter sensors integrate over a much larger particle population size, which can therefore be related to bulk POC measurements with more certainty. A similar bulk approach could be applied to large particles to reduce uncertainties in the conversion. Waite (unpublished) explored this possibility by filtering large volumes (50–100 L) of seawater onto a mesh of known mesh size (in this case 20 \( \mu \)m), rinsing the material off and measuring its POC content. The POC content of these particles related reasonably well to the particle biovolume measured using an Underwater Vision Profiler, providing a site-specific size-to-POC conversion (Figure 4). While gravitational filtration or pumping (e.g., as in situ stand-alone pumping systems) could
cause particles to break up and change characteristics, they are relatively easy to make, not very time demanding and not very prone to subjective bias. This "bulk approach" for larger particles might thus be more feasible at sea than studies on individual particles, and could be adopted for quasi-global measurements.

There is clearly a need for more data on size-to-POC relationships and how they vary in space and time. We therefore highly recommend a concerted effort to collect information on size-to-POC relationships. These measurements are relatively tedious to collect and often only yield a few data points, which is frequently deemed insufficient for data interpretation purposes. However, a joined effort and data portal that collates all these individual measurements would soon produce a respectable database that could form the basis for more accurate and appropriate size-to-POC conversion equations.

Sinking Velocity
The range of recorded sinking velocities of particles in the oceans spans from negative (i.e., particles float toward the surface; Azetsu-Scott and Passow, 2004; Acuña et al., 2010) to several km per day (e.g., salp fecal pellets; Iversen et al., 2017). When considering the sinking velocity of an individual particle, a first approximation can be obtained from Stoke's law (originally derived for spherical, non-porous particles and laminar flow) combined with White's approximation (White, 1991), which suggest that sinking velocity increases linearly with excess density (the difference from the water density) and the square of particle diameter (i.e., linearly with the particle area). Building on these expectations, many studies have tried to relate sinking velocity primarily to size, which has been shown to be a useful predictor for particles generated in controlled environments (e.g., roller tanks; Gårdes et al., 2011; Iversen and Ploug, 2013; Iversen and Robert, 2015). However, strong relationships were only observed when all particles were generated using the same water/plankton community (Iversen et al., 2010). When particles were made by different plankton communities, size alone was a bad predictor (e.g., Diercks and Asper, 1997) strongly supporting notions that particle densities and shapes vary widely depending on the source material (Iversen et al., 2010).

Packaging and porosity contribute appreciably to determining sinking velocities. On the one hand, adding ballasting materials, such as diatom frustules, to aggregates may lead to an increase in sinking velocities owing to the increase in excess density. On the other hand, the addition of ballasting mineral particles to marine particle populations frequently leads to smaller more densely packed aggregates that sink slower because of their smaller size (Hamm, 2002; Passow et al., 2014). Mucous-rich particles have been shown to float despite relatively large sizes (Azetsu-Scott and Passow, 2004; Bochdansky et al., 2016), whereas oil- or plastic-containing aggregates have been shown to sink rapidly despite the presence of substances with an excess density smaller than seawater (Long et al., 2015; Passow et al., 2019). In natural environments, particles are formed through different mechanisms, by different organisms, and under varying environmental conditions that affect aggregation (e.g., salinity, pH, minerals), ballasting (e.g., dust deposition, sediment load; Iversen et al., 2010; Iversen and Robert, 2015; van der Jagt et al., 2018) and sinking behavior (e.g., viscosity; Taucher et al., 2014). A universal conversion of size-to-sinking velocity is hence impracticable (Jouandet et al., 2011).

 Nonetheless, estimates of size-to-sinking-velocity relationships are powerful when determined site-specifically for either distinct types of particles or large particle populations, which negates the effect of individual outliers. To do so, a sufficient number of particles needs to be imaged and their sinking velocities measured for each location and—if possible—for each particle class. Unfortunately, measuring the sinking velocity of individual particles directly remains challenging, and many studies rely on indirectly approximated sinking velocities from bulk measurements. The following sections explore the most commonly used methods for determining sinking velocities for individual particles and from bulk measurements.

Individual Particles
The majority of data on individual particle sinking velocities has been generated ex-situ. A big advantage of ex situ sinking velocity measurements is that the particles can be retrieved after the measurement and analyzed for elemental composition. It is thus possible to generate data on size, sinking velocity and carbon content for individual particles, allowing the calculation of carbon flux (by that particle) with relatively high certainty. Box 3 lists some of the methods used to measure the sinking velocity of individual particles ex situ.

Briefly, particles for ex situ measurements are either generated in roller tanks or collected and resuspended. The first approach is problematic as these particles may—as described above—not reflect natural particles and particle behaviors (Ploug, 2001; Prairie et al., 2015). Its advantages are that it allows the testing of
targeted interactions and effects on sinking velocities. The second approach can be tricky as collection and handling procedures may alter the particles (i.e., fragmentation, aggregation, change of shape, porosity, etc.). To measure sinking velocities, individual particles are introduced into an experimental container, such as a tank or settling column, and their sinking velocity is measured either traditionally with a stop-watch (Box 3a) or using time-lapse photography (i.e., video analysis; Box 3b). Another method is the use of a flow chamber, where the particle is kept in suspension by an upward flow of water (Box 3c; Ploug and Jørgensen, 1999; Peterson et al., 2005; Iversen and Ploug, 2010). The flow speed thus reflects the sinking velocity of the particle. This method further allows simultaneous imaging of the particle as well as additional measurements such as oxygen gradients within the particle (Ploug and Grossart, 1999; Belcher et al., 2016a,b).

In situ sinking velocities of individual particles have been measured by divers (Allredge and Gotschalk, 1988) or using time-lapse camera systems taking rapid sequential photos of particles as they settle through the water column. To negate lateral advection and internal waves, some camera systems image particles as they settle through a column (Diercks and Asper, 1997; Cartwright et al., 2013), other systems are installed on neutrally-buoyant drifting platforms in a Lagrangian fashion (Pilskaln et al., 1998, Lampitt and Iversen, unpublished) or as moored platforms making seasonal measurements of settling velocities on individual aggregates in the deep ocean (Diercks et al., 2018; Iversen, unpublished). Images are often taken in bursts every few hours to conservatively use battery power and data storage, and the capture rate during these bursts needs to be sufficiently fast to accurately track fast-sinking particles (Diercks et al., 2018). The advantage of these methods is that they provide the most accurate sinking velocity estimates of natural particles to date as they minimize any manipulation and preserve in situ conditions. Moreover, they can collect data on a large number of particles spanning a relatively large particle size range (depending on the camera specifications), and their data can thus be used to construct both particle size spectra as well as sinking velocity spectra and investigate the relationship between size, shape, and sinking velocity. On the flip side, construction and deployment of time-lapse camera systems are relatively complicated and expensive. A noteworthy methodology to calculate sinking velocity from such in situ image time series is particle image velocimetry (PIV; e.g., Steinbuck et al., 2010; Smith and Friedrichs, 2015), which has also been used to observe filtration rates of larvaceans in situ (Katiia et al., 2017).

**Bulk Measurements**

Approaches to estimate the average sinking velocity for the entire particle population or a particle size class are plentiful with a great variety of methodologies (Box 4). These methods are mostly based on in situ observations. In situ approaches often use an observed change in characteristic (e.g., Chlorophyll a) over time and depth (Box 4a) or modified sediment traps that allow the collection of differential settling particles (Box 4b). When combined with conventional sediment traps or gel traps, in situ observations of particle size spectra can be used to infer sinking velocities of individual particle size classes (Box 4c) (Guidi et al., 2008; McDonnell and Buesseler, 2010) and to estimate mass fluxes (Guidi et al., 2008; Iversen et al., 2010; McDonnell and Buesseler, 2012; Nowald et al., 2015). Recently, the use of naturally occurring radionuclide pairs ($^{210}$Po/$^{210}$Pb and $^{234}$Th/$^{238}$U) has been added as a tool to estimate average sinking velocities (Box 4d) (Villa-Alfageme et al., 2014, 2016). The “SETCOL method” is a relatively easy and fast ex situ method that can provide sinking velocity estimates for selected particle groups (e.g., specific species, types or size classes) (Box 4e; Bienfang, 1981).

**MODELS AS A TOOL**

Optical devices for particle measurements provide a wealth of information, which can be used far beyond simple particle classification and carbon flux estimates. For example, particle size distributions can reveal trophic positions (Zhou, 2006; Basedow et al., 2016) and, when combined with models, particle interactions such as aggregation and disaggregation ([O’Brien et al., 2004; Stemmann et al., 2004b; Karakas et al., 2009].

Models can play two essential scientific roles. (1) They synthesize our best understanding of a process. Our understanding can then be tested and improved by comparing the quantitative predictions by models with observations and experiments. (2) Models can be used to estimate hard-to-measure parameters. For example, despite successful measurements of aggregation/disaggregation parameters in the laboratory (Waite et al., 1997b; O’Brien et al., 2004; Iversen and Ploug, 2013), to date, we have no means of observing in situ particle aggregation and disaggregation rates. A comparison of modeled and observed particle size distributions, however, has been successfully used to infer these rates between particle size classes (Stemmann et al., 2004a,b; Karakas et al., 2009]. To use models with any degree of accuracy or precision requires a careful assembly of additional data beyond just the observed size distributions, such as phytoplankton community composition and size distributions in sediment traps (Jackson et al., 2005).

Models of marine particle size distributions commonly describe particle interactions in terms of coagulation theory (Burd and Jackson, 2009), which provides the basic framework for calculating the rates of collisions between particles. These models require estimates for a suite of biophysical parameters that will vary between sites and models (Table 2). For example, a simple model that assumes a single source of particle requires relatively easy-to-get auxiliary data (e.g., primary production; Jackson, 1990), whereas a complex model, e.g., simulating coagulation between different types of particles such as diatoms and fecal pellets, will require complex auxiliary data (e.g., production rates for each type of particle; Jackson, 2001; Jokulsdottir and Archer, 2016). Moreover, as coagulation depends on physical processes and particle geometry, estimates of turbulent dissipation rates and particle fractal dimensions may also be needed. Some of these parameters (such as turbulent dissipation rates) can be estimated from detailed measurements of physical water-column, properties whereas particle-based
parameters such as fractal dimensions can be obtained from particle image analysis (e.g., Kilps et al., 1994). These complex auxiliary data are largely contextual data that provide the model with the correct biophysical description to estimate the particle size distribution and make meaningful comparisons between model results and observations. Models can be made without such contextual data with the missing parameters considered as fitting parameters, a situation that becomes more and more unsatisfactory as the number of missing parameters increases.

Although both modeled and observed particle size distributions can be calculated using different widths of the size bins, it is useful to know in advance of the model calculations what these size bins are. This is particularly critical because of the different rationales for choosing bin sizes in models and data; bin sizes in models are often chosen to improve computational speed, whereas bin sizes in data are often chosen to optimize signal over noise. In addition, models and observations should use the same definition of particle size (i.e., the modeler should be aware of how the size was calculated from the images; Jackson et al., 1997).

Estimates of the uncertainties of the observed particle size distribution can help when using the model as a tool to help interpret the observations. This is because model results tend to be smooth functions of particle size, often approximated by power laws over specific size ranges (Burd and Jackson, 2002), whereas observed distributions show varying degrees of noise (Jackson et al., 1997). Significant differences between the modeled and observed distributions may suggest the presence of certain processes, such as size-specific grazing rates, and departures from a smooth power-law distribution may indicate the occurrence of significant disaggregation (Burd and Jackson, 2002). Contextual data such as turbulent dissipation rates and zooplankton abundances may help to identify the possible causes of differences between modeled and observed distributions.

A good example of how models and particle image data can be used together to infer both rates and processes is provided by

**BOX 3 | Ex situ measurements of particle sinking velocities.**

**(a) Sedimentation column**

A method that is cheap and simple to set up is to measure the time it takes for a particle to pass two marker points in a large measuring cylinder full of seawater (O’Brien et al., 2006; Riley et al., 2012; Cavan et al., 2015). To replicate in situ sinking behavior, the experimental setup should be set to the same temperature and salinity as the particle origin. While this method sounds straightforward, there are several caveats. (1) Subtle differences in temperature between the inside and outside of the experimental container introduce convection currents that strongly affect slow-sinking particles. (2) Salinity gradients can develop relatively quickly, sometimes leading to a complete halt of the settling particle. (3) Particles are often introduced into the settling column with peripheral water (e.g., when using a pipette), which may have a different density (temperature and/or salinity) to the water in the settling column. The measured sinking velocity will therefore reflect the behavior of the introduced water as much as that of the particle itself. (4) The downward motion of the particle and, potentially, water around it causes an upward motion of water elsewhere in the container. Depending on the size of the container, the resulting internal turbulence may affect particle sinking behavior. (5) When measurements are taken at sea, the motion of the ship may influence the measurement. Uncertainties in the final sinking velocity measurement are therefore potentially large.

To mitigate some of these problems, cold lights should be used as even short exposures of light instantly create turbulence flow in the sinking cylinder. The measuring cylinder should also be as large as possible in diameter to reduce wall effects. Microgradients of salinity can be used to shut down all convection in experimental cylinders (O’Brien et al., 2006). For all experiments, flow conditions in the cylinder are best tested with some dye or neutrally buoyant particles to ensure that the water in the cylinder is quiescent (Poupl et al., 2008a).

**(b) Ex-situ time-lapse**

A more sophisticated version of the sediment column is the combination of a sinking column or roller tank with cameras. There are several examples, and we here explain the FlowCAM method (Bach et al., 2012) and the Orbit method (Poupl et al., 2010). More elaborate setups may involve two video cameras, which can be aligned to give a 3D view of sinking (Poupl et al., 2008a).

The FlowCAM is a laboratory piece of equipment that is used to measure particle concentration, type and size. Conventionally particles are pushed through a cuvette past a camera that takes photographs, and particles are automatically counted. Depending on the magnification, particles down to 20 µm can be measured. The FlowCAM can be adapted to measure sinking velocities. For this, instead of being pushed past the camera, particles are left to settle in a tube leading to the cuvette and camera so that, by the time they reach the field of view (FOV), particles are settling at their natural velocity (Bach et al., 2012). Multiple images are taken (up to 12 per second) and, knowing the number of images taken per particle and the distance traveled through the FOV, sinking velocities can be calculated. As for the sedimentation column, the temperature and salinity should be the same as at the particle’s origin. Some of the benefits of this method are that it is very quick and semi-automatic, allows a relatively large particle size range to be measured, and gives individual particle characteristics (e.g., ESD, type, color) that can be associated with individual sinking velocities. Some drawbacks include wall effects as particles sink toward the cuvette, magnification limiting the maximum size of particle that can be measured, aggregation within the setting tube prior to entrance into the cuvette, and convection currents if the temperature is not constant.

The “Orbit method” measures sinking velocities of particles directly in roller tanks (Poupl et al., 2010). Particles are incubated in a rolling tank where solid body rotation has been established. Video capture of several orbits allows for “repeat” calculations of sinking velocities and thus present solid values. This method is non-destructive, and particles may be used for later analysis.

**(c) Flow Chamber**

The Flow Chamber allows measurements of all three particle dimensions (height, length, and width), sinking velocity and micro-sensor measurements while the particle is kept suspended by an upward-directed flow matching the settling velocity of the particle (Poupl and Jørgensen, 1999; Wekerle et al., 2018). The Flow Chamber system is filled with filtered seawater with in situ temperature and salinity. The upward flow is adjusted with a needle valve until the particle remains suspended at a distance of one particle diameter above the net, whereby the particle sinking velocity is equal the flow velocity. The sinking velocity of each particle is then calculated from the flow rate divided by the cross-sectional area of the flow chamber. Unfortunately, this method is time-intensive, allowing investigation of only a few (typically <10) particles per hour.
High temporal and vertical resolution sampling of particle concentration allows the calculation of “bulk” estimates of particle settling velocities using correlations between variations in concentration at different depths. This method works best when the sinking velocity of an identifiable group of particles is homogeneous, and when the concentration of the tracked particles is pulsed or highly variable in time, exceeding variability due to advection across spatial gradients. This method has been applied to optical backscattering, fluorescence, and beam attenuation data collected by ships and autonomous gliders (Briggs et al., 2011) and to LOPC data collected by an autonomous profiling float (Jackson et al., 2015).

The indented rotating sphere trap

The indented rotating sphere (IRS) trap allows particles to be separated in situ according to their sinking velocities. Particles land on a sphere and, after an amount of time (typically 12–24 h), the sphere rotates releasing the particles to be collected in a sample carousel that also rotates at specific time intervals. This setup results in sinking velocity bins between 0.88 and ~980 m d⁻¹. IRS traps can be mocred at any depth and usually deployed on a monthly time frame. This method is useful for determining the relative importance of sinking fluxes. For instance, the IRS was used to show that slow and fast sinking particle fluxes were larger than intermediate sinking fluxes in the Mediterranean (Alonso-González et al., 2010). In the northwest Pacific, it revealed how a power-law attenuation of flux with depth (Martin et al., 1987) can be explained by a proportionally higher loss of slow-sinking particles with depth (Trull et al., 2008).

An advantage of this method is that collecting particles in situ negates any imposed temperature or density changes when trying to replicate settings in the laboratory. Some limitations are that particles may partially collapse on the ball in the IRS, reducing porosity and potentially increasing sinking velocities (Peterson et al., 2005). The sphere rotation time could be reduced giving less time for particles to be altered whilst sitting on the sphere, but this would increase the lowest settling velocity, e.g., 3 h rotation = minimal sinking velocity of 5.4 m d⁻¹ (Peterson et al., 2009).

The radioactive pairs Thorium-234/Uranium-238 and Polonium-210/Luclid-219 allow the estimation of particle fluxes. A relatively novel method, a one-box inverse model, further uses ²¹⁰Po or ²³⁸Th to calculate in situ sinking velocities in the upper mesopelagic, allowing observations of changes in sinking rates with depth (Villa-Alfageme et al., 2014, 2016). Samples are collected using Niskin bottles or in situ pumps and average sinking velocities are calculated for each sample. The simple one-box model assumes a steady-state system, which is not always appropriate in changing conditions, such as blooms, as ²¹⁰Po or ²³⁴Th have relatively long half-lives (Villa-Alfageme et al., 2014). It is important to note that calculated sinking velocities tend to be <100 m d⁻¹ and represent an average sinking velocity for the entire particle population at each depth. Estimates also depend on how samples are collected, e.g., whether particles are filtered (e.g., using Stand Alone Pumping Systems), and whether they include the suspended non-sinking fraction or not.

The “SETCOL” method (Bienfang, 1981) is an ex situ bulk method that can be used to estimate settling rates of selected particle groups. This method is primarily used to measure phytoplankton cell sinking velocities, and the same principle is applied for the Marine Snow Catcher (Riley et al., 2012; Giering et al., 2016; Cavan et al., 2017). A homogenous water sample is placed or collected inside a settling column and an initial sample is taken. After a pre-determined time interval, the concentration of particles in different parts of the settling column (e.g., the bottom, middle, and top) is measured. The average sinking velocity can now be calculated by the difference in biomass in the different parts of the settling column between the start and end of the settling period. This method is relatively easy and fast, though the final estimate strongly depends on the settling time and dimensions of the settling column. It has useful applications in laboratory culture experiments, yielding precise sinking velocity estimates for culture species when comprehensively replicated (Waite et al., 1992). In addition, for field samples, SETCOL can be combined with size-fractionation or type-specific particle counts to provide sinking velocity estimates for selected groups (Waite and Nodder, 2001).

Optical settling tubes work on the same principle but monitor the settling process in the entire settling period using an optical device, such as a transmissometer, fitted near the bottom of the settling tube. This set-up allows the calculation of particle size distribution and settling velocity distribution (Zaneveld et al., 1982) and has been used in situ (e.g., Kineke et al., 1989; Spinrad et al., 1989; Murray et al., 1996), though these studies focussed on coastal or near-benthic particles.
Optical devices for particle measurements have the potential to provide high spatial and temporal resolution of particle fluxes, though we are still far away from accurately estimating fluxes based on optical devices alone. The biggest bottleneck is the uncertainty associated with translating an optical signal, such as an image, into an accurate flux estimate, which requires additional information such as the sinking velocity and density of the particle. Moreover, the parallel development of various devices and analysis routines currently hinders meaningful comparison between techniques. Such comparisons are urgently needed as well as the standardization across methods of data collection, analysis, and data archiving.

In situ optical methods provide us with the ability to identify particles in the water column with minimal disturbance, and the fast imaging rates offer high-resolution information on their spatial distributions and interactions. For zooplankton, images give much higher vertical and horizontal resolution than nets and provide much better taxonomic identification than acoustic methods. For aggregates, in situ optical methods inform on the structure of aggregates without fragmenting them or changing their shape as pump systems do or integrating them spatially and temporally as sediment traps do. While no single method captures the entire size range of particles in the ocean at once, a combination of devices can produce a particle spectrum from a few micrometers to several centimeters in size.

Qualitative observations can inform us about biological mechanisms driving particle structure and spatiotemporal variability. Potential insights include both qualitative information about which processes are important (e.g., direct observations of zooplankton–aggregate interactions) as well as quantitative model validation (e.g., using aggregate size distributions to constrain aggregation and disaggregation models). These insights are critical for improving our mechanistic, predictive understanding of processes controlling POC in the ocean.

Continuing developments in this field will move us closer toward the goal of estimating particle flux by constraining the carbon in each particle and its sinking velocity. Yet, this goal requires more ex situ and in situ experimentation, validation, and regional calibration to overcome the fundamental problem of translating optical information on particles (a standing stock) into useful estimates of carbon flux (a process). With a collective effort, optically based flux estimates may soon be feasible on large scales when we draw on the entire wealth of information contained in images from the interior of the ocean.

**AUTHOR CONTRIBUTIONS**

SG coordinated the writing of the manuscript. All authors provided valuable input on the full manuscript.

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