# ADVANCED OPTICAL METHODS IN TRANSIENT HEAT TRANSFER AND TWO-PHASE FLOW

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#### 1. INTRODUCTION

Optical methods have a long tradition in heat transfer, but their application to two-phase flow started much later. Their instant signals make them specially suited to transient situations provided that the registration method is quick enough and has a high sampling frequency.

A further benefit of optical methods is their non-invasive mode of operation, not interacting with the material to be investigated and thereby not affecting the physical process under study.

Optical methods use changes of light waves as sensoric signals, which are due to interaction between the light and the material. Such changes and interaction consequences can be

- attenuation,
- scattering,
- deflection or
- reflection.

Depending on the mode of registration we distinguish between

- image-forming and
- non-image-forming

methods. The latter ones allow only a spotwise record of the events and the first one is registers usually a two-dimensional picture of situations or processes on a surface or in a volume as well-known from photography. Since approximately twenty years a new image-forming method has been in use, that is, the holography, which found admission to measuring techniques in heat and mass transfer and two-phase flow in the last ten years. The holography uses two-dimensional registration tools, namely photographic plates, from which however three-dimensional information can be reconstructed. This has special advantages in studies of transient two-phase flow.

Holography combined with the well-known interferometry using the phase-shift of the light-wave allows the registration of temperature and concentration fields, and became a very valuable tool in heat and mass transfer.

LDA works spotwise, i.e. records the single phase velocity in a very small volume only. A modification of the LDA is the PDA, which is applicable for two-phase flow in dispersed mixtures. Mie-scattering is another method used in two-phase flow for example to detect spray characteristics.

It is not possible to present all optical methods being used in heat transfer and two-phase flow within a given time and a limited space. Even the concentration of such a survey on optical methods being applicable to transient conditions would cover a wide field of different techniques. Therefore, the author apologises for restricting this presentation on optical methods with which he has a longer experience and to which he contributed to their development to a certain extent.

## 2. HIGH-SPEED HOLOGRAPHY

In 1949 Gabor /1/ invented a new optical recording technique which he called "holography". In contrast to photography by which only the two-dimensional irradiance distribution of an object is recorded, holography allows the recording and reconstruction not only of the amplitude but also of the phase distribution of the wave-fronts. Making use of this unique property, completely new interference methods could be developed. As holography demands a highly coherent light source it can be only performed by using a laser.

The general theory of holography is very comprehensive and for a detailed description one must refer to the literature /2-4/. Here only the principals necessary for understanding the holographic measurement techniques can be mentioned. In fig. 1 the holographic two-step image-forming process of recording and reconstructing of an arbitrary wave-front is illustrated.

The object is illuminated by a monochromatic light source and the reflected, scattered light falls directly onto a photographic plate. This object wave usually has a very complicated wave-front. According to the principal of Huygens one can, however, regard it to be the superposition of many elementory spherical waves. In order to simplify

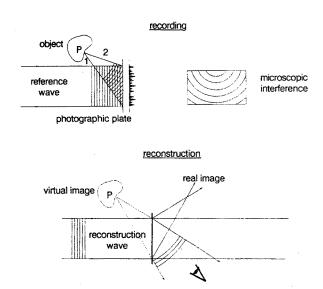


Fig. 1: Holographic two-step image forming process

the matter, only one wave is drawn in fig. 1. This wave is superimposed by a second one called "reference wave". If both waves are mutually coherent they will form a stable interference pattern when they meet on the photographic plate. This system of fringes can be recorded on the photographic emulsion. After the chemical processing of the plate –in a developing bath and in a fixer–, it is called "hologram". The amplitude is recorded in the form of different contrast of the fringes and the phase in the spatial variations of the pattern.

If the plate is subsequently illuminated by a light beam similar to the original reference wave the microscopic pattern acts like a diffraction grating with variable grating constant. The light transmitted consists of a zero-order wave, travelling in the direction of the reconstructing beam plus two first-order waves. One of these first-order waves travels in the same direction as the original object wave and has the same amplitude and phase-distribution. Thus a virtual image is obtained. The other wave goes in the opposite direction and creates a real image of the object. The virtual image can be looked at with the naked eye and the real image can be studied with reconstruction devices for example with a microscope.

This holographic technique can be used instead of the photography for example for recording a swarm of droplets produced in an injection nozzle. The holographic setup for such a study is shown in fig. 2.

It consists of a pulsed ruby laser emitting pulses of a period of 30 ns and a lens- and mirror-system for expanding, dividing and guiding the laser beam through the measuring object and onto the holographic plate. The laser beam is first expanded by means of the lens AL and then divided in the beam splitter ST to produce the object beam and the reference beam. The object beam travels via a collecting lens, two mirrors and a screen through the object - in this case the spray coming out of a nozzle - and

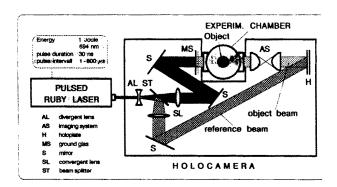


Fig. 2: Optical arrangement for the recording of pulse laser holograms

passes after the object an imaging lens before it falls ont the holographic plate. There it is superimposed by th reference wave which is splitted off by the beam split ter ST and falls via the collecting lens SL and a mirro onto the photographic plate by-passing the object. So a instantaneous picture of the situation in the spray ca be registered. If the electronic system of the ruby lase allows to emit more than one laser pulse within a ver short period of time sequences of the spray behaviour car be stored on the photographic plate from which the veloc ity of the droplets with respect to amount and direction as well as changes in the size and geometrical form c the droplets can be evaluated. This evaluation, however needs a very sophisticated and computerised procedure. For evaluating the hologram it first has to be reconstructed as demonstrated in fig. 3.

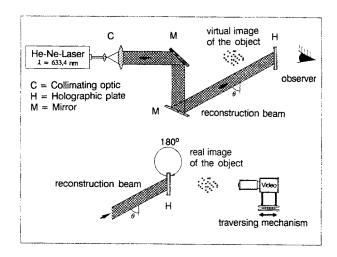


Fig. 3: Optical arrangement for the reconstruction of pulsed laser holograms

To do this for the special application of studying a nozzle spray, the holographic plate is replaced to the old position after chemical processing and is now illuminated by a continuously light emitting helium-neon-laser, sending its light via the path of the reference beam described in fig.

2. This new beam is now called reconstruction beam. If the holographic plate is replaced in the same orientation as it stood when the exposure occurred one can look at it with the naked eyes and can see a virtual image of the droplet spray exactly at the place where it was produced before by the injecting nozzle. For a quantitative evaluation one needs a closer examination by a camera, for example by a video camera. To do this the holographic plate has to be turned by 180°, when positioning to the old place and by illuminating with reconstruction beam a real image of the spray is produced, however, on the other side of the holographic plate. This real image has a three-dimensional extension and the video camera can be focussed to any plane within the image. For technical evaluation the camera, fixed at a certain position, is focussed to the mid-plane and then for getting information from other planes of the spray the camera is moved forward or backward with fixed focus of the lens. So plane by plane of the spray cloud can be evaluated. The virtual picture one would see with the naked eye is demonstrated in fig. 4.

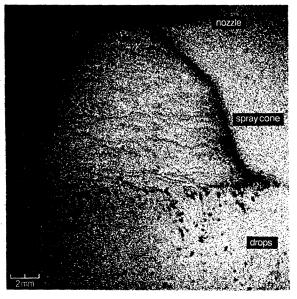


Fig. 4: Photography of a reconstruction of a pulsed laser hologram. Object: Spray of a hollowed cone spiral nozzle.

It looks like a photographic picture of a spray with a veil near the nozzle orifice and a swarm of droplets separating from this veil when it breaks up. The quantitative evaluation works via the above-mentioned video camera and a computer system as shown in fig. 5 and uses the real picture. Main components of this evaluating systems are a digitiser, a graphic monitor, a video camera and a PC. The video camera is scanning the real image of the reconstructed hologram O and sends its information to the digitiser D. It changes the electrical signal from a analogue character to a digital one and stores it in a frame memory. Now the computer C can use the digitalised

information for performing the image reprocessing. The digitiser simultaneously produces an analogue picture in a false-colour (red-green-blue) reproduction.

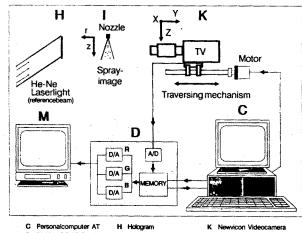


Fig. 5: Digital image processing system for the evaluation of pulsed laser holograms

After a first positioning of the camera, noise-signals are eliminated without suppressing the gradients of the greycolour. By this a first "clean" picture (with a minimum of noisy signals) is arising. Then the rest of pixels still originating from noisy signals is filtered out. In the next step gradients out of the less or more intensive grey-colours of the pixels surrounding the holographic reproduction of the droplets or of the contour of the veil are evaluated. So it can be distinguished between well focused parts and such ones which are out of focus. Then all pixels having a grey-colour below a certain pre-defined value are treated as zero. By this reproductions of droplets which are out of the interesting focus-plane can be eliminated and a "picture" is electronically produced containing only reproductions of droplets which were within a very narrow tolerance within the focussing plan of interest.

By this procedure it happens that the contours of some droplets do not have a closed and continuous outline, because pixels may have been extinguished spotwise during the gradient checking procedure. Therefore a next step follows in which the open contours are filled with colour to produce closed outlines of the droplet reproductions. To be sure not erroneously to create new spots by this process which could be interpreted as droplet reproductions, the situation after the contouring is compared with that which existed before reproductions of droplets out of focus were eliminated. A "droplet" in the new picture is only accepted if it existed already before in the old picture. Finally the remaining droplet-reproductions are filled with colour and now the evaluation with respect to droplet-size, -form and -concentration can go on.

After this plane in the holographic picture has been evaluated the video-camera is moved within a small step and the whole procedure starts again. So plane by plane of a spray-cloud can be evaluated and a three-dimensional picture of the two-phase flow situation is arising which was fixed on the holographic plate within a few nanoseconds. For more detailed information reference is made to the work by Chávez and Mayinger /5, 6/.

An example of an computerised reproduction of the veil and the droplets coming out from a nozzle with cone-like spray forming shows fig. 6. It is an evaluation of the same hologram which was reproduced in fig. 4 as photographic pick-up.



Fig. 6: Evaluation of single pulsed holograms by digital image processing

More downstream of the nozzle only droplet clouds can be observed. By evaluating a great number of such droplet clouds with the above-mentioned procedure information can be extracted with respect to the dependency of the droplet diameter from the mass-flow-rate through the nozzle and the pressure of the atmosphere into which the spray is injected. An example of such an evaluation gives fig. 7.

If two exposures of a droplet spray are illuminated onto the same holographic plate within a short period of time also the velocity of the droplets can be determined from such an hologram, however, with a much more complicated procedure which is described in detail in /6/. The data produced by this opto-electronic process are of high accuracy as fig. 8 demonstrates. Especially the suspicion that the computerised processing of the double exposure hologram produces a large scattering in the velocity-data is misproved by this figure. Even the influence of the pressure of the atmosphere in which the droplets are travelling is clearly brought out.

If the droplets are moving in a vapour atmosphere of the same substance as the liquid and if the temperature of the liquid is below the saturation temperature condensation occurs at the phase-interface which makes the volume of the droplet growing. By using a simple energy balance the condensation heat transfer can be calculated from this growth of volume versus time. The accuracy and the reproducibility of the described opto-electronic measuring technique are good enough to determine these heat transfer coefficients at the veil and at the droplet-cloud as fig. 9 demonstrates. The heat transfer data are averaged values from all droplets, being reproduced in a hologram.

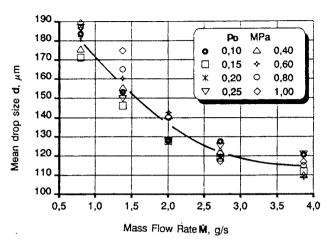


Fig. 7: Mean droplet diameter as a function of the mass flow rate at different ambient pressures

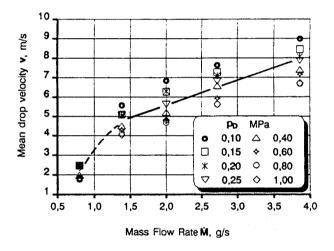


Fig. 8: Mean droplet velocity as a function of the mass flow rate at different ambient pressures

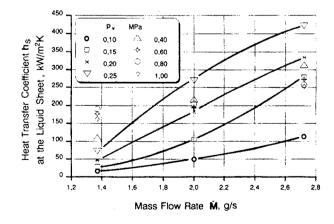


Fig. 9a: Heat transfer coefficient at the liquid sheet as a function of the mass flow rate at different vapor pressures

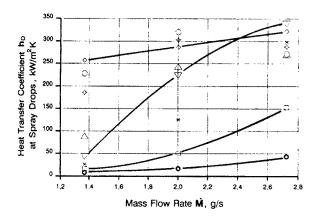


Fig. 9b: Heat transfer coefficient at the droplets as a function of the mass flow rate at different vapor pressures

#### 3. HOLOGRAPHIC INTERFEROMETRY

For convective heat transfer processes the temperature gradient in the boundary layer near the heat emitting or absorbing wall is of special interest because from the temperature gradient at the wall, supposing a laminar sublayer the heat transfer coefficient

$$h = \frac{-k \left(\frac{dT}{dy}\right)_w}{T_w - T_\infty} \tag{1}$$

can be directly derived if, in addition, the temperature of the wall or of the bulk and the thermal conductivity of the fluid are known. This temperature gradient can be measured by interferometric optical methods, like the Mach-Zehnder interferometry, as well-known from the literature.

A new interferometry method is the combination of holography and interferometry. A most commonly used arrangement of optical set-ups for this holographic interferometry is shown in fig. 10.

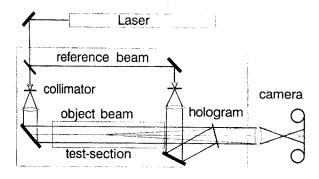


Fig. 10: Optical set-up for holographic interferometry

A helium-neon laser or an argon-laser serves as a light source emitting continuously monochromatic and coherent light. By means of a beam splitter the laser beam is divided into an object- and a reference-beam in a similar way as we learnt before. Both beams are then expanded to parallel waves by a telescope which consists of a microscope objective and a collimating lens. The object-wave passes through the test section in which the temperature is to be examined whereas the reference-wave directly falls onto the photographic plate. There are many possibilities for arranging the optical set-ups to form a holographic interferometer which cannot be discussed here in detail. Reference is, therefore, made to the literature for example 17, 8/.

Several procedures exist to produce interferograms; here only a sophisticated one will be explained, which can be used for high-speed cinematography and by which transient heat transfer phenomena can be examined. It is called "real-time-method" because it allows to observe a process to be investigated continuously in real time. The method is illustrated in fig. 11.

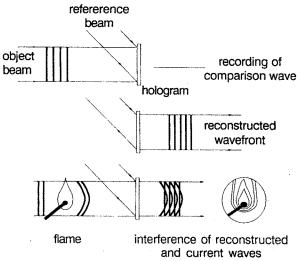


Fig. 11: Real time method for holographic interferometry

After the first exposure, by which the comparison-wave is recorded and during which there is no heat transfer in the test section –for two-phase flow experiments, even only single- phase flow may exist– the hologram is developed and fixed. Remaining at its place or repositioned accurately, the comparison-wave is reconstructed continuously by illuminating the hologram with the reference-wave. This reconstructed wave can now be superposed onto the momentary object-wave. If the object-wave is not changed and the hologram is precisely repositioned, no interference fringes will be seen at first (infinite-fringe-field adjustment).

Now the heat transfer process which is to be examined and for two-phase flow for example the boiling with bubble formation, or the condensation can be started. Due to the heat transport process a temperature field is formed in the fluid and the object-wave receives an additional phase shift passing through this temperature field. Behind the hologram both waves interfere with each other and the changes of the interference pattern can be continuously observed or photographed on still or moving film.

The real-time-method demands an accurate reconstruction of the comparison-wave; therefore, the hologram must be repositioned precisely at its original place. This can be done by using a well-adjustable plate-holder, which, nowadays, can be purchased from the market. It is recommended to use a plate-holder where the final adjustment can be done via a remote control for example with quartz crystals. The adjustment of the repositioned holographic plate gets its feed-back control signals on an optical basis because the adjustment has to be done in such a way that the interference fringes -at first visible due to non-precise position of the plate- disappear during this procedure. This, certainly, has to be done without the heat transfer process having started, however, the pressure and the temperature are existing under which the system is operated during the experiments.

A series of holographic interferograms taken with this method is illustrated in fig. 12, where the bubble formation at a heated surface in water with slow horizontal flow was studied. The water is slightly subcooled, i.e. the bulk temperature of the water was below the saturation temperature and, therefore, the bubble is condensing again after detaching from the heated surface and departing from the superheated boundary layer as one can observe by following the holographic interferograms versus a period of 7 ms in fig. 12. The horizontal black and white fringes in fig. 12 represent lines of constant temperature in a first approximation.

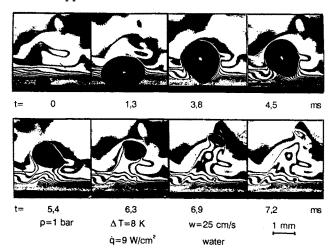


Fig. 12: Interferogram of subcooled boiling on a heated wall

From this figure, however, we can also learn something about the limitation with which this and all other interferometric methods are afflicted. The temperature field near the heated wall in the fluid is not only shifting the phase of the light-wave, it is also deflecting the light beam when travelling through it. This deflection has the conse quence that with very high gradients at the wall —as it i the case in subcooled boiling— the zone immediately ad jacent to the wall cannot be seen. With lower heat fluxe this deflection is not a problem in interferometry.

For learning the procedure of evaluating such an interfer ogram a more simple example shall be used as that in fig 12. Fig. 13 shows the interference fringes in a subchannel of a tube-shell heat exchanger formed by three heat emitting tubes. The interference fringes here represent exactly isotherms in the fluid around the tubes and it can be clearly seen that the spacing of these isotherms is different at positions of narrow and of wider gap-size in the subchannel. Wide-spreading of the isotherms means a low temperature gradient and as we know from experience low heat transfer coefficient.

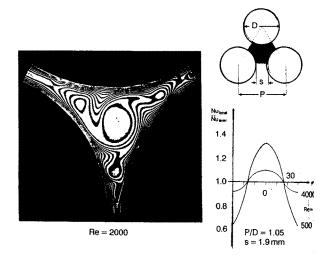


Fig. 13: Temperature field between three heated tube with coaxial flow

This temperature field in the subchannel is of two-dimens ional nature with good approximation. For the sake simplicity only the evaluation of the simple case of a two dimensional temperature field will be discussed here. It is also assumed that the holographic interferogram is produced with parallel object-waves. The evaluation of the interference patterns is than quite similar to that of the Mach-Zehnder-interferometer /9/. Therefore, here on the basic equations will be given. In the holographic in terferometry the object-waves passing through the tesection at different time are superposed and, therefore reveal the changes in optical path-length between the twexposures. Expressed in multiples S of a wave-length this change is calculated to

$$S(x,y) \lambda = l [n(x,y)_2 - n(x,y)_1]$$

where l is the length of the test section in which the fractive index n is varied because of temperature change. The refractive index distribution n(x, y) during the record

ing of the two waves is —as mentioned above— assumed to be two-dimensional (no variation in light direction). Equation (2) shows that initially only local variations can be determined. Only if the distribution of the refractive index  $n(x,y)_1$  during the recording of the comparison wave is known, absolute values can be obtained. Therefore, one usually establishes a constant refractive index field (constant temperature) while recording the comparison wave.

$$S(x,y) \lambda = l \left[ n(x,y)_2 - n_{\infty} \right]$$
 (3)

To obtain absolute values for the temperature field, the temperature at one point in the fluid has to be determined by thermocouple measurements. This is usually done in the undisturbed region or at the wall of the test chamber. Equ. (3) is the equation of ideal interferometry. It is assumed that the light beam propagates in a straight line. Passing through a boundary layer, the light beams, however, are deflected because of refractive index gradients. This deflection is used for the various Schlierenand shadowgraph-methods. The light deflection can be converted into an additional phase shift S, if a linear distribution of the refractive index is assumed to be within this small area.

$$\Delta S = \frac{n_0 \,\lambda \,l}{12 \,b^2} \tag{4}$$

In this equation b is the fringe width and  $n_0$  is the average refractive index.

In many applications an ideal, two-dimensional field cannot be found. Often the boundary layer extends over the ends of the heated wall, or there are entrance effects or temperature variations along the path of the light beam (axial flow in the test section). Therefore, only integrate values are obtained. Having corrected the interferogram, the obtained refractive index field can be converted into a density field. The relation is given by the Lorentz-Lorenz-formula where N is the molar refractivity and M the molecular mass.

$$\frac{n^2 - 1}{n^2 + 2} \frac{1}{\rho} = \frac{N}{M} \tag{5}$$

If there is only one component in the test section and the pressure is kept constant, the density variations can only be caused by temperature changes. If the fluid is a gas, the situation is very simple because its refractive index is very near to 1, which reduces Equ. (5) to the Gladstone-Dale-equation:

$$\frac{2}{3}(n-1)\,\frac{1}{\rho}\,=\frac{N}{M}\tag{6}$$

With the simple Boyle-Mariotte-law and R as the gas constant we then obtain the following formula, which relates the fringe shift to the temperature:

$$T(x,y) = \left[ \frac{S(x,y) \ 2 \ \lambda \ R}{3 \ N \ p \ l} + \frac{1}{T_{\infty}} \right]^{-1}$$
 (7)

For liquids the procedure is a little more complicated because we have to take in account the real behaviour of the thermodynamic properties as a function of temperature. Therefore, we have to use an equation of state for the refractive index n or we have to take the refractive index from tables interpolating the data with simple equations. Fortunately, there are good data banks available in the literature for most of the fluids. However, it is also not difficult to measure the refractive index in a simple optical set-up.

With an equation for the refractive index as function of the temperature we then can use Equ. (2) and we get the connection between the pattern of the interference fringes and the temperature field, as shown in Equ. (8):

$$S(x,y) \lambda = \frac{dn}{dT} [T(x,y) - T_0]$$
 (8)

Often local heat transfer coefficients are of special interest. In this case the temperature gradient at the wall is determined and assuming a laminar boundary layer next to the wall or the phase-interface the heat transfer coefficient is obtained by using Equ. (1).

The assumption that the temperature field is two-dimensional and constant along the path of the beam travelling through the fluid is not valid in case of temperature fields around curved surfaces, like bubbles. The refractive index n is then a function of the radius r, and we have to use Equ. (2) in its differential form

$$S \lambda = \int_0^1 (n - n_0) dz$$
 (9)

and we write it in spherical or cylindrical co-ordinates:

$$S(y) \lambda = \int_0^z \left[ n(r) - n_0 \right] dz \qquad (10)$$

For spherical and cylindrical symmetry Equ. (10) can be solved and integrated as described, for example, in /8/ after transforming it in the form

$$S(y)\lambda = 2 \sum_{k=i}^{N-1} \Delta n_k \left[ \left( r_{k+1}^2 - r_i^2 \right)^{1/2} - \left( r_k^2 - r_i^2 \right)^{1/2} \right]$$
(11)

In temperature fields with very high gradients the deflection of the laser beam, which is demonstrated in fig. 14, too, cannot be neglected, as it is done in the evaluating procedure described in /10/. High temperature gradients are especially found in the liquid boundary layer around vapour bubbles, in particular if condensation occurs. In such a case a complicated correction procedure for this deflection has to be used which is described by Nordmann

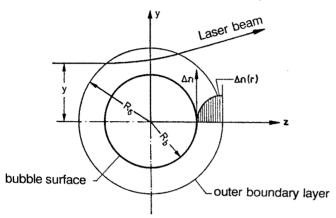


Fig. 14: Beam deflection and optical conditions in a temperature field around a spherical bubble

and Mayinger /11/ and by Chen /12/. With the equations and corrections described there, even temperature fields around very tiny bubbles can be evaluated.

With high heat transfer coefficients the boundary layers at the phase-interface are usually very thin –in the order of a few hundredth of a millimeter– and it is difficult to investigate them with the interferometric procedure described up to now, because only a few interference fringes would be observed within this narrow area. Therefore, here another interference method has to be used, the so-called "finite fringe method". In this method, after the reference hologram was produced, a pattern of parallel interference fringes is created by tilting the mirror in the reference-wave of fig. 15, or by moving the hologram there within a few wave-lengths. The direction of the pattern can be selected as one likes and it is only depending on the direction of the movement of the mirror or of the holographic plate.

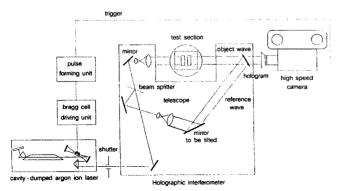


Fig. 15: Arrangement for holographic interferometry with finite fringe method

This pattern of parallel interference fringes is then distorted by the temperature field due to the heat transfer process. The distortion or deflection of each fringe from its original –parallel– direction is, in a first rough approximation, the temperature gradient and gives by using Equ.(1) the heat transfer coefficient. A short description how these interference patterns are evaluated is given in fig. 16. This figure also demonstrates for the example of a temperature field around a burning flame how these distorted interference fringes look like, depending on the original orientation of the parallel pattern. A detailed description of the mathematical procedure how to evaluate these pattern of finite fringes and how to determine the temperature gradients from it, is given in /11/ and /12/.

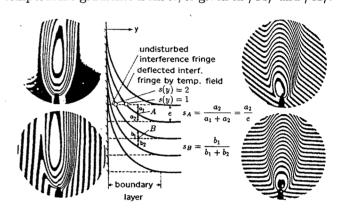


Fig. 16: Finite fringe interferograms (temperature field around a flame) and their evaluation

Fig. 17 demonstrates the possibilities of using these techniques in a flow with a bubble condensing in a liquid. By combining this method with the high-speed cinematography it allows an inertialess and precise evaluation of the heat transfer coefficient at the phase-interface of a condensing bubble. Holographic interferometry certainly can only be used if the flow situation is not too complicated and if the bubble population is not too numerous, so that individual bubbles can be identified. It is not possible to look inside the bubbles, because the light is totally reflected at the phase-interface.

As explained in the Equs. 2 - 5 the phase shift is a function of the change in the density of the fluid. In pure substances the density is a function of temperature and pressure. In multi-component systems also the concentration influences the density. In our evaluation procedures up to now we worked on the assumption that the alteration of the density is affected by a temperature change only and, therefore, the variations in the refractive index within the test section could be treated as temperature distributions. When the variations in the refractive index are caused not only by a temperature-, but also by a concentration- or pressure-change these simple evaluation of the interference pattern, described up to now is not possible any more. Therefore, coupled heat and mass transfer processes can be examined by interferomet-

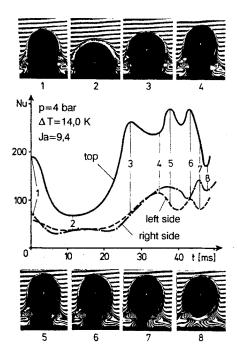


Fig. 17: High speed cinematography of interferograms around a condensing bubble and the evaluated heat transfer at the phase interface

ric methods only if one of the two fields is obtained by an additional measuring method. Only by assuming identical profiles of the temperature and of the concentration the interferograms can be evaluated without additional measurements as El-Wakil /13/ did.

There is, however, one method to determine the temperature and the concentration field by optical means alone, which is called the "two-wave lengths interferometry". This is done by applying their dependence of the refractive index on the wave-length of the light to determine the temperature- and concentration-fields by means of separate interferograms, taken at different wave-lengths. Pross and El-Wakil /14/ used this two wave-lengths interferometry in a modified Mach-Zehnder-interferometer for the study of the evaporation and combustion of fuels. Panknin and Mayinger /7, 8/ used this basic idea and developed a two-wave lengths method for the holographic interferometry. The problem with the two-wave-lengths interferometry is that the two interferograms have to be superimposed very accurately. Here, the peculiarity of the holography allowing the recording of different interference pattern on one and the same plate is a great help to overcome these difficulties. A simple set-up for the holographic two-wave lengths interferometry is shown in fig. 18. It resembles very much the arrangement of fig. 10 and actually the only difference is that two lasers are used as light sources in fig. 18. The beams of the He-Ne-laser  $(\lambda_i = 6.328 \dot{A})$  and of the Argon laser  $(\lambda_k = 4.579 \dot{A})$  intersect and, therefore, only one shutter is needed and equal exposure times at both wave lengths are guaranteed. The beams are then superimposed by means of a beam splitter. By this one gets two object - and two reference-waves at the different wave-lengths.

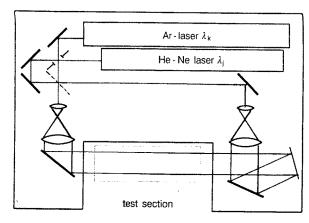


Fig. 18: Optical set-up for holographic two-wavelenght interferometry

It has to be mentioned that not only the object wave  $\lambda_j$  is reconstructed by the reference-wave  $\lambda_j$ , but also a false objective-wave  $\lambda_k$  is obtained and vice versa. These unwanted waves, however, emerge at different angle from the hologram and can, therefore, easily be separated from the original waves. For the evaluation of the interferograms here only some simple equations shall be presented. For a more detailed study reference is made to the work by Panknin /8/. In gas we can use the Gladstone-Dale-equation, Equ. (6), and the ideal gas law which relates the fringe shift to the temperature- and concentration-distribution in a heat and mass transfer boundary layer.

$$S(x,y) \lambda = \frac{3 p l}{2 R} N_m \left[ \frac{1}{T(x,y)} - \frac{1}{T_{\infty}} \right]$$
 (12)

The molar refractivity  $N_m$  for a mixture of two gaseous components is given by

$$N_m = N_a C_a + N_b C_b$$
; with  $C_a + C_b = 1$  (13)

where  $N_a$  and  $N_b$  are the molar refractivities of the components in their pure form and C is the concentration of the component in the mixture. During the recording of the comparison wave the temperature distribution T in the test section is constant and there are only two components of the mixture.

Combining Equ. (12) and (13) we obtain for each wavelength

$$S(x,y) \lambda = \frac{3 p l}{2 R} \left[ \frac{1}{T(x,y)} (N_a + C_b(x,y)) (N_b - N_a) - \frac{N_a}{T_\infty} \right]$$
(14)

eliminating  $C_b(x,y)$ . The temperature T(x,y) can be calculated:

$$S_{j}(x,y) \frac{\lambda_{j}}{N_{bj} - N_{aj}} - S_{k}(x,y) \frac{\lambda_{k}}{N_{bk} - N_{ak}} =$$

$$\frac{3 \ p \ l}{2 \ R} \left[ \frac{1}{T(x,y)} - \frac{1}{T_{\infty}} \right] \left[ \frac{N_{aj}}{N_{bj} - N_{aj}} - \frac{N_{ak}}{N_{bk} - N_{ak}} \right]$$
(15)

After determining the temperature distribution only one interferogram is used to calculate the concentration profile.

Equ. (15) shows that there is a difference between the phase shifts for the two wave-lengths which is used for the measurement of the temperature. This difference is usually very small. Therefore, the two wave-lengths used should be as far apart as possible. The dependence of  $N_{(a,b)}$  is also very small and gets larger proportions only in the vicinity of an absorption line which, however, is usually not in the visible range. This limits the choice of substances to those lengths used. Some test fluids suitable for this technique are naphtalene, carbondisulphide, bencene and hexane. The position of the fringes has to be determined very accurately at the same plate in the two interferograms.

A simple application example of the two-wave-lengths technique is given in fig. 19. In order to demonstrate the differences in the phase shift only the upper and lower halfes of each interferogram are shown and the evaluation is made at the intersection of the pictures. The interferograms show the heat and mass transfer boundary layer at a heated vertical wall with free convection. The mass transferred was naphtalene into air.

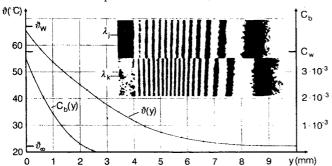


Fig. 19: Temperature and concentration profiles in a laminar boundary layer

Heat and mass transfer experiments were also performed in a burning flame with hexane flowing out of a horizontal porous cylinder and oxidising after evaporation. Fig. 20 shows the two interferograms obtained with the two wave-lengths as emitted by the He-Ne-laser and the Argon-laser. In this interferogram the finite fringe method was used as described before. The temperature field in this flame evaluated from the interferograms in fig. 20 is given in fig. 21, which clearly demonstrates the benefit of

this optical method by instantaneously presenting a complete information about the temperature distribution i the flame. However, for flame research there are toda newer methods available which will be described in th next chapter.

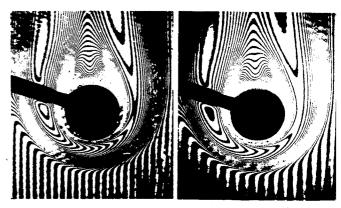


Fig. 20: Finite fringe interferogram of a flame taker with two different laser beams of the wavelength  $\lambda_j$  (right and  $\lambda_k$  simultaneously

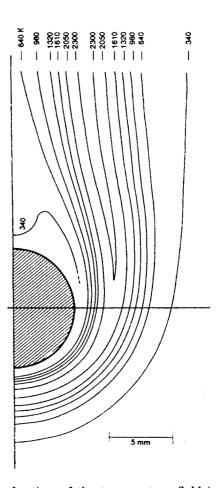


Fig. 21: Evaluation of the temperature field in a flat from the interferogram of fig. 20

## 4. HOLOGRAPHIC TOMOGRAPHY

The holographic-interferometric methods discussed up to now do not allow to evaluate a 3-dimensional temperature field of any local structure, because each beam travelling through the field under investigation is integrating all local situations on its way. With improving electronic data-acquisition-systems an expansion of this optical measuring technique became of interest which overcomes the disadvantage of the lack of local resolution. Following the methods well known in medicine also the "Optical Tomography" is based on a simultaneous irradiation of the volume under investigation, from different directions.

An optical set-up used for such a holographic-interferometric-tomographic method is shown in figure 22. In the centre of this holographic arrangement, the vessel is placed, in which the processes of interest - for example temperature field or concentration field in a mixing device - take place. The vessel with its volume to be investigated is irradiated via 4 different object waves spanning over an angle of 135°. The experimental set-up and the method to evaluate the interferograms are described in detail in /15 - 17/. Here only the guidance of the beams is briefly discussed.

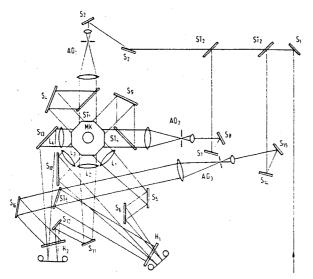


Fig. 22: Beam paths of an tomographic interferometer

To avoid deflection of the light at curved surfaces of the glass-vessel, the outer surrounding of the vessel is arranged in an octagonal form. So each of the beams hits a plane surface in orthogonal direction. Only the inner vessel is of cylindrical shape. To avoid a deflection of the light at its surface, a liquid is filled in the annular space between the outer and the inner vessel, which has exactly the same diffractive index as the glass of these vessels.

An Argon-Ion-Laser is used as coherent light source and its beam at first hits the mirror  $S_1$  and from there the beam is travelling to 2 adjustable beam splitters,  $ST_2$  and  $ST_3$ . The first beam splitter is fading out the reference wave and the second beam splitter is producing 4 different

object waves which are expanded via micro lenses. A prefocusing of the object waves is produced by the 4 lenses  $L_1$  -  $L_4$ . This makes it possible to store two interferograms of high quality on one holographic plate.

The experimental arrangement shown in fig. 22 has the benefit that the lenght of the optical paths of the reference waves and of the 4 object waves are equal. So the demand on the lenght of coherence of the laser can be moderate. By focusing the object waves in front of the hologram the density of the light-flux is enhanced. This allows short exposure-time and therefore this method can be applied also to transient processes. This procedure can be combined with real time holography and also cinematographic recording is possible.

# 5. LIGHT SHEET METHOD AND LASER-IN-DUCED-FLUORECENCE

There is not only the possibility of using phase shift to get information about temperature- and concentrationdistribution in a fluid, also scattering effects - elastic and non-elastic ones - can be used for detecting material properties as well-known since many years. A rough idea which scattering effects can serve for material investigations mediates fig. 23. From all these possibilities only a special measuring technique, based on the fluorescence will be discussed here. Compared to Rayleighor Raman-scattering the laser-induced-fluorescence (LIF) has the benefit that the signals from this method are by 5 to 15 orders of magnitude stronger than the Rayleighor Raman-scattering. The draw-back is that it is difficult to evaluate the fluorescence signals quantitatively. Usually calibration-tests are needed. It gives only information about the concentration and not about the temperature whilst the Raman-spectroscopy can be used for detecting concentration and temperature fields if the laser pulse is strong enough. Finally for each substance to be detected a very special frequency of the incoming monochromatic light is needed to produce fluorescence.

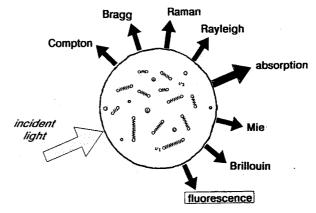


Fig 23: Scattering in a volume of particles

The laser-induced fluorescence is physically based on the fact that the molecule under investigation is absorbing

a photon of the incoming light by which the molecule is transferred to a state of higher energy for a very short period  $-10^{-5}$  to  $10^{-10}s$  and afterwards falls back approximately to the former energy level by emitting light of a frequency which is different from that of the incoming light. The radiation emitted is characteristic for the substance and its density is a measure for the concentration of this special substance in a mixture. Usually it is not possible to detect various substances unless several lasers with different light emitting frequency are available. The radiation emitted is characteristic for the concentration and the temperature of the substance. Pre-condition for the use of this method, however, is that the structure and the distribution of the energy level of the molecules and possible electron transfer processes with its probabilities have to be known.

The optical arrangements for measurements with laser-induced fluorescence is quite simple. The laser-beam is extended to a light sheet by means of cylindrical lenses and this light sheet travels through the substance in which a mass transfer process is under investigation. Such a mass transfer process of interest can be the burning zone of a flame for example. The fluorescence induced by the light sheet of the laser is detected by a CCD-camera. The camera transforms the optical signals into electronical ones and transmits them to a computer system. Fig. 24 presents an example of such an arrangement.

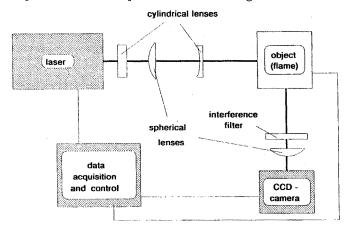


Fig 24: Set-up for laser-induced fluorescence

How such a laser-induced fluorescence information looks like is show in fig. 25.



Fig 25: Reaction front of a nonpremixed  $H_2$ -air-flame detected by laser-induced fluorescence

The process under investigation there was a simple Bunsen flame with hydrogen as fuel. The laser light sheet activating the fluorescence consisted of a monochromatic radiation with a frequency of 308 nm. The substance activated to fluorescence is the OH-radical one of the main intermediate products of hydrogen combustion and its maximum of concentration gives the position of the flame. Fig. 25 conveys the information that combustion takes place in a narrow zone at the outer surface of the  $H_2$ -jet where air and hydrogen are mixed.

One can also use the self-induced fluorescence of molecules due to the high temperature which, however, in gases only occurs if the temperature is high enough. A evaluation of the fluorescence signal also goes via CCD-camera and a computer. The intensity of the fluorescence can be documented in the reproduction by using a discoloured method which means that different values of concentration are assigned to various colours. By this for example the development of a flame in a high-speed flow can be studied Fig. 26 presents the evaluation of a hydrogen flame in highly turbulent flow where the at first a channelised air stream expands into a free jet with a wall at one side With increasing Mach-number that flame becomes more and more compressed and the beginning of the ignition

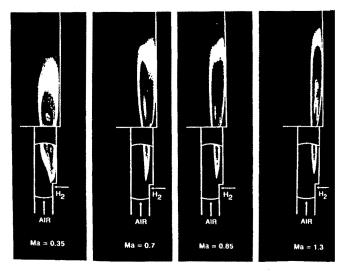


Fig. 26: H<sub>2</sub>-air-flame stabilization in the vicinity of rearward facing step at different air Mach-numbers

is blown downstream. The combustion in the channel stabilised by a recirculation zone behind a step. Finally one can combine various optical methods for example the holographic interferometry and the self-induced fluorescence. Fig. 27 and 28 present such an example. Again the combustion zone of a hydrogen flame under high air velocity was studied. At first the mixing of the air and the gaseous fuel was of interest. Therefore, in modelling experiment helium was injected into the air flow and its concentration in the mixing zone was measured by using the holographic interferometry.

Fig. 27 presents the concentration distribution in the mix ing zone. For a high-speed and high turbulence flame the minimum concentration of the fuel in the air must not be too low in order to avoid that the turbulent mixing is extinguishing the flame instead of agitating it. So on

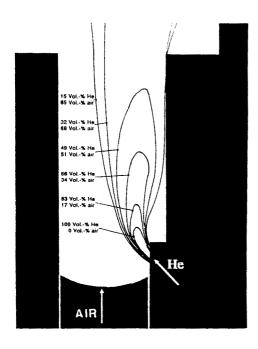


Fig. 27: Concentration distribution in the mixing zone

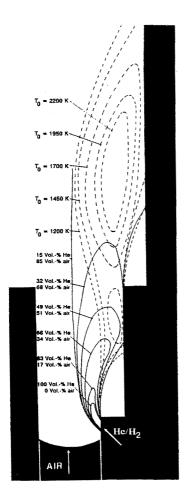


Fig. 28: Readings of the helium-mixing experiments superimposed by the isotherms of the burnig flame

can assume that for a hydrogen flame a fuel concentration of 10 - 15 volumetric percent may be a minimum for flame acceleration. Therefore, in fig. 27 the line of 15 volumetric percent helium can be roughly regarded as the outer border of the high-speed flame. In fig. 28 the readings of the helium-mixing experiments are superimposed by isotherms of the burning flame -with hydrogen as fuel- which result from a combined measuring technique of holographic interferometry and self-induced fluorescence method. The figure clearly conveys the information that the ignition starts in the recirculation zone behind the step, where the air concentration is rather low, but good enough for a first reaction to form radicals like OH. The main combustion takes place far downstream of the hydrogen injection place when the hydrogen concentration in the air is between 10 and 15 volumetric percent. An impression of the interference fringes in a holographic picture of a mixing jet shows fig. 29 where the finite fringe method was used. In this figure also the shock front can clearly be seen.

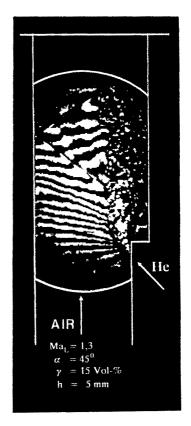


Fig. 29: Interferogram of a mixing jet with a shock front

## 5. Concluding Remarks

Optical Methods are expected to experience a powerfull revival due to three reasons:

Sophisticated theoretical treatment of heat and mass transfer processes with large computer codes needs very detailed information about temperature- and concentration-fields in the areas of interest - like in boundary layers - for assessing and improving the physical models used in the code and for verifying the code itself. An optically determined pattern of isotherms is a very stringent touchstone for the reliability and accuracy of the code. Modern developments in power and process engineering make transient situations more and more interesting, especially for controlling procedures and safety deliberations. Optical measuring techniques work inertialess.

A former draw back of the image-forming optical measuring techniques, the laborious and time-consuming evaluation does not exist anymore. A personal computer can evaluate a hologram or an interferogram within a few seconds which took manpowers of several hours before. The costs of such an evaluating equipment are relatively moderate.

So theoretical analysis and optical measuring techniques could form a new phalanx for investigating heat and mass transfer processes.

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