

Designing a High-Resolution, LEGO-Based Microscope for an Educational Setting

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ABSTRACT Microscopy is an essential tool in many fields of science. However, because of costs and fragility, the usage of microscopes is limited in classroom settings and nearly absent at home. In this article we present the construction of a microscope made of LEGO® bricks and low-cost, easily available lenses. We demonstrate that the obtained magnification and resolution are sufficient to resolve micrometer-sized objects and propose a series of experiments that explore various biophysical principles. Finally, a study with students in the age range of 9 to 13 y shows that the understanding of microscopy increases significantly after working with the LEGO microscope.

KEY WORDS microscopy; LEGO; middle school

I. INTRODUCTION

The invention of microscopy in the 17th century by Antoni van Leeuwenhoek was the start of an era of research in the micro-world (1). Although we know by now what the “little animals” are that he observed, the micro- and nano-world is an inexhaustible topic for biophysical research, for which the light microscope is the instrument of choice.

Despite the simplicity of a basic light microscope, the fundamental working principles are beyond everyday intuition for pupils, but also for most adults. Although this lack of insight into simple optics might partially be because of the perception of microscopes as complex research instruments, hands-on experience promises to drastically improve the understanding of the working principles of microscopes by any interested person, including children.

Our aim is to introduce a microscope to individual students in a classroom setting, both as a scientific tool to access the micro-world and to facilitate the understanding of fundamental principles of the optical components of a microscope in a playful and motivating, yet precise approach. By basing the design on LEGO, we aim to make the microscope modular, cheap, and inspiring. The LEGO brick system provides a low entry level for children, because it is a common toy found in most homes. The modular design, flexibility, and high level of sophistication of the different building parts make it an ideal framework to demonstrate even complex instruments with simple

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means. Indeed, LEGO has been used before to demonstrate the working principle of scientific instruments, such as a conceptual atomic force microscope (2) and a watt balance (3), or in actual scientific setups (4). Furthermore, this article is in line with other initiatives to create high-performance, low-cost microscopes. Although a previous version of a LEGO microscope called the Legoscope was successful in generating high-quality images, it still required an objective and custom (3D-printed) parts, hindering its capacity to be used simply to demonstrate the principles of a microscope. Another initiative used very simple but smart designs to generate a \$1 foldable microscope (foldscope) (5), wherein the low adaptability of the microscope and the image quality were a trade-off for the extraordinarily low price. Finally, affordable, robust tools such as pocket microscopes and clip-ons for smartphones also allow students hands-on access to microscopy.

In this article we present the design of a fully functional microscope made entirely out of LEGO. The only non-LEGO components are the 2 optical parts, which can be purchased off the shelf at approximately €4 (\$4) each. We quantify the optical performance of the microscope and find submicrometer resolution. We show how, thanks to the flexible design, the magnification of a microscope can be tuned by combining different lenses and adjusting their distance, thus providing a simple tool for teaching fundamental optics and simple biophysical concepts. Furthermore, the simple and robust design allows long-term imaging without drift in the focus of the microscope.

We design a number of experiments that can be carried out with this LEGO microscope and easily accessible materials, while exploring various biophysical themes. Finally, we demonstrate that this LEGO microscope can be used in an educational setting by letting a group of 9- to 13-y-old students build the microscope and demonstrate their progress in understanding of microscopy with a questionnaire. To ease the usage of the LEGO microscope, we designed a step-by-step workflow for a school setting or for helping parents and children discover biophysics and microscopy in an autonomous

way. All materials, including parts lists, building plans, workflows and project suggestions are found in the Supplemental Material and on GitHub (6).

II. SCIENTIFIC AND PEDAGOGICAL BACKGROUND

Microscopy is a fundamental tool of modern science that is used on a daily basis to study and understand biological and biophysical questions. In sharp contrast to its importance, a layman's understanding of the fundamental principles remains superficial, and microscopes are often seen as complex instruments, exclusively usable in a science context. Although the working principles of a microscope are taught in high school settings, the usage of microscopes in the lecture setting often does not target an understanding of its principles, but simply of the use of ready-made microscopes.

Here we aim to teach in a playful way the magnifying properties of a single lens and a series of 2 lenses, which is effectively an understanding of the lens equation,

$$\frac{1}{f} = \frac{1}{d_1} + \frac{1}{d_2} \quad (1)$$

with f the focal length of a lens and d_1 and d_2 the distance between the lens and the object or the observer, respectively. Magnification occurs when the ratio $d_2/d_1 > 1$. We demonstrate that even children from 9 to 13 y of age can intuitively grasp this concept of optical systems. This knowledge is brought into practice with the aid of a widespread and brick-based modular toy system, LEGO. The flexibility and the already present experience of using LEGO to build complex assemblies lies at the core of the LEGO-based workflow. We combine 2 lenses with adequate sample illumination and a system to control precisely the position of the lenses, which are the essential components of an optical microscope. During the building process, the difficulties in object positioning, lens adjustment, and illumination become obvious and are solved in a simple way.

Besides an understanding of optics, we suggest a series of experiments that can be conducted with the home-built microscope and readily available samples. These experiments demonstrate some fundamental biophysical concepts, such as the drastic effects of osmotic pressure on plant cells. Furthermore, the motion of easily obtainable microswimmers can be observed, where the fundamental problem faced by swimming at low Reynolds number may be integrated in the experimental description.

To make the work with the microscope an efficient experience, we provide a detailed work plan that can be followed at the individual or the class level. As we provide all information as freely accessible and extensible GitHub repositories (6), we hope that the instructions might be translated into other languages by the community. We have already provided versions in English, German, Dutch, Spanish, and French.

III. MATERIALS AND METHODS

A. Microscope assembly

We designed the microscope by taking advantage of the modular and flexible characteristics of the LEGO brick system. The design of the microscope is shown in Figure 1a,b. The building plans, including a list of components, can be found in Supplemental Files S1 and S2. The main components are the illumination part, the objective holder, including the objectives, and the ocular part.

1. Illumination

Illumination is one of the simplest parts of the microscope (Fig 1c). In case of a group work to construct the full microscope, this part can also be built by young children that are less used to complex building instructions. The illumination itself is a special LEGO brick integrating an orange LED. It is straightforward to replace the orange LED with a different color. A short video demonstrating the opening and replacement of the LED is given in Supplemental Movie S1. Because the light source is positioned close to the sample, no special condenser is included, which helps reduce the optical parts to a minimum. The

absence of a condenser means the illumination with the lower magnification objective can become slightly nonuniform. To avoid this, a diffuser can be made from a thin sheet of paper placed between the LED and the sample.

2. Objective holder

To allow precise focus adjustment even with a high-magnification objective, an accurate and smooth positioning with submillimeter precision is required (Fig 1d). Because LEGO is not designed to generate such precise motion, we had to push the limits of the brick system by combining a gear rack with a gear worm screw that results in approximately 3 mm of travel for every full rotation. This movement is sufficient for the final adjustment of the focus, even when looking at samples of only several micrometers in size. Although the objective holder is the smallest of all the individual parts of the microscope, it is the most demanding with respect to building skills. Hence, building this part should either be done by older and more experienced children, or the building should be supervised by an adult to prevent frustration in an early phase of constructing the microscope. Once built, the gear system will move the objective in the vertical direction when rotating the control wheel (black wheel in Fig 1d). The precision and repeatability of this threaded system suffices for both objectives presented below.

3. Ocular

The ocular is a key element of the microscope, which is first built in the proposed classroom or exploration setting and then used as a classical magnifying glass. Although the overall construction of the ocular remains quite simplistic, the positioning of the lenses is still demanding. Although the lenses used here are cheap yet powerful, mishandling plastic lenses will rapidly lead to scratches on the surface. Users need to be careful when handling the lenses, because scratches on the surface disturb the image quality.

4. Optics

The only non-LEGO components in the building plans are the lenses in the objectives and the ocular. The designs for 2 different

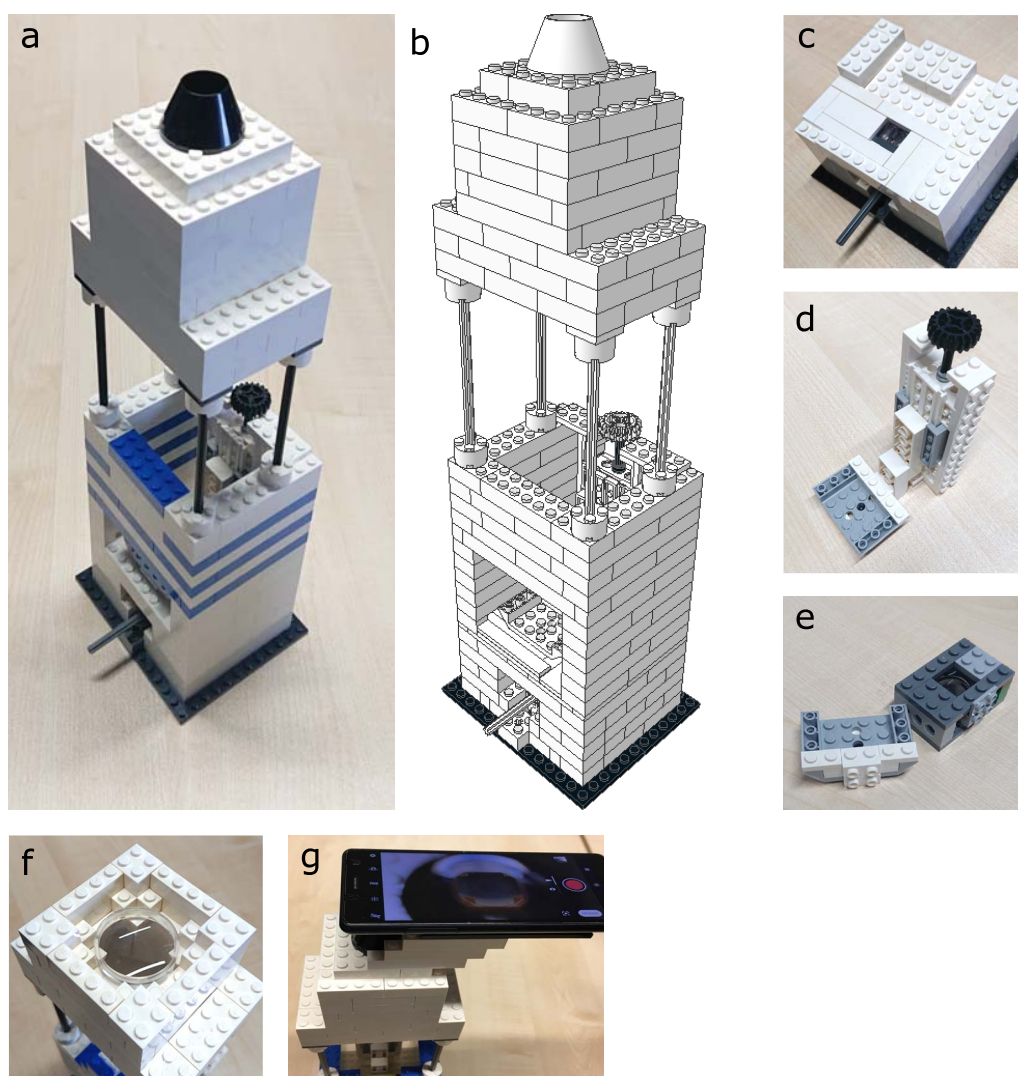


Fig 1. Design of the LEGO microscope. (a, b) A photograph and a schematic representation of the microscope, (c) the LED that illuminates the sample from below, (d) the threaded system that adjusts the focus of the microscope by moving the objective, (e) 2 objectives containing a replacement smartphone lens with a 3.85-mm focal distance (left) and a glass lens with a 26.5-mm focal distance (right), (f) the second lens consisting of 2 acrylic lenses in its holder just below the eyepiece, (g) a smartphone used as a camera by adapting the eyepiece.

objectives are included (Fig 1d), with a higher and a lower magnification. However, other lenses or even lens systems can be easily integrated. However, it is important that such adapted objectives position the optical elements in the same position, to keep the alignment of the microscope. For the high-magnification objective (Fig 1d, left), the lens of a replacement iPhone 5 camera module is used. Such highly specialized plastic lenses of non-spherical shape have outstanding optical performance compared with glass or polymer beads and are suitable for high-resolution

microscopy that approaches the diffraction limit (7), while also being available at a low price. The lens used here has a reported focal length of 3.85 mm, leading to a numerical aperture of 0.54 as discussed in the Results section. After carefully peeling the lens off the chip, it is attached to a LEGO brick with transparent tape and a glass cover slip. Detailed photographs of this procedure are shown in Supplemental File S1.

For the low-magnification objective (Fig 1e, right), a glass lens with a focal length of 26.5 mm and a numerical aperture of 0.29 was used,

which was built into a holder that shares the attachment brick and also has the same optical center as the high-magnification objective.

The ocular consists of 2 acrylic lenses with a focal length of 106 mm each, with 1 flat and 1 convex side. Placing the 2 lenses with their flat sides on top of each other, the focal length is reduced to 53 mm (see Fig 1f).

5. Sample holder

The sample holder is a surface of flat LEGO bricks that can be covered with rough tape to reduce the risk of accidentally moving the microscope slide. Furthermore, advanced students are encouraged to design a stage to position the microscope slide accurately. Either the gear rack and worm screws of the objective holder or a translation stage [e.g., as in Tsung-Han *et al.* (2)], can be used as a starting point.

Table 1 summarizes the list of non-LEGO components used in the assembly of this microscope, including their costs in euro at the time of writing. A smartphone is not included but recommended for image acquisition. In case no LEGO parts are already available, the typical cost of the full set of parts is also listed.

B. Sample preparation

1. Salt crystals

To create sodium chloride crystals, a 1 M sodium chloride solution is prepared by dissolving 5.8 g of kitchen salt in 100 mL of lukewarm water. A drop of the solution is placed on a microscope slide and left at room temperature for approximately an hour, during which the water evaporates and salt crystals are formed.

2. Onion cell monolayer

A monolayer of red onion cells is obtained by cutting a red onion with a sharp knife and carefully peeling off a thin slice with a pair of tweezers. This layer is then spread out on a microscope slide and covered with a drop of water and a coverslip. The coverslip is sealed off on 2 opposing sides with nail polish and left open on the 2 other opposing sides for the influx and efflux of liquid. Supplemental Movies S2, S3, and S4 show the preparation procedure.

Table 1. List of components used in the assembly of the microscope. The cost in euro is given at the time of writing and does not include shipping.

Part name	Price (€)
iPhone 5 replacement camera	4
Acrylic lens (2) (diameter 34.5 mm, $f = 106$ mm)	3
Glass lens (diameter 18.0 mm, $f = 26.5$ mm)	5
Set of microscope slides (50) and cover slips (100)	10
All LEGO parts	80

3. Microswimmers

A selection of water-based microorganisms such as amoeba, paramecia, and water fleas are obtained by collecting water from a local pond in a plastic bottle. Because water fleas feed on bacteria and algae, some grass was added as a source of nutrients for bacterial growth. The mixture was then left for 1 wk at room temperature, so that the water fleas could multiply. Afterward, a few drops were placed on a microscope slide, which was covered with a coverslip and sealed off with nail polish. Because harmful bacteria can also grow in the bottle, strict hygiene must be followed, and careful hand washing after contact with the water is important. Alternatively, paramecia and water fleas can be bought from pet shops.

4. *Artemia*

Eggs are obtained in powder form from a pet shop and put in water to revitalize them. After 2–3 d the organisms have hatched, and a drop of water with the *Artemia* is placed on a microscope slide, which is covered with a glass coverslip.

C. Workflow for classroom and individual exploration

To support the usage of the LEGO microscope and to optimize the learning experience, we designed a workflow plan. Such workflows can be included either in a classroom environment by separating a class into several groups, where 2 or 3 pupils work on a single LEGO microscope set and the project is split into different sets. Alternatively, the workflow can also be followed in an individual setting. Because the help of an experienced adult is not guaranteed, the workflow includes help sections to ensure that certain steps are

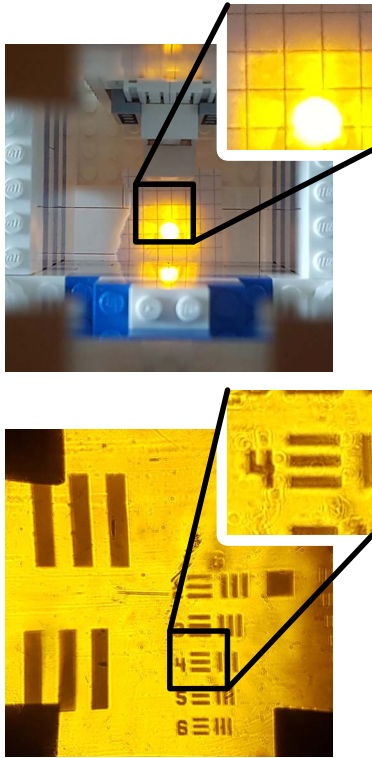


Fig 2. Experimentally determining the magnification of the microscope. (Top) Imaging of squared paper without lenses in place. The width of each square is 5 mm. (Bottom) Imaging of the USAF Resolution target with the high-magnification objective and an $f = 53$ mm lens near the eyepiece. The length of the bar next to the number 4 is $27.6 \mu\text{m}$. Note that for a correct determination of the magnification, the (digital) zoom of the camera has to be the same for both images.

successful before the next steps are initialized. This step is important to avoid frustration, in case some questions or tasks turn out to be difficult. The full workflow, translated in several languages, is found in Supplemental File S3 and on GitHub (6).

IV. RESULTS AND DISCUSSION

The microscope can be used both for direct observation by eye, or images can be recorded with a smartphone camera. For the images in this article, a Sony Xperia XZ2 Compact smartphone was used. Its rear camera has a chip of 5057×3796 pixels with a pixel size of approximately $1.2 \mu\text{m}$. To record images or movies, the Android camera app was used. In all experiments, the autofocus setting was used.

The following subsections quantify the specifications of the microscope. Although this is

important to assess the capabilities of the microscope, it might be beyond the interest of middle school students.

A. Measuring magnification

To start the characterization of the LEGO microscope, its magnification is experimentally determined. To achieve that, images are taken of objects with known length, with and without lenses in the microscope (Fig 2). The magnification is then given by the ratio of the respective millimeter-to-pixel ratios:

$$M = \frac{L_{\text{no lens}}(\text{mm})/L_{\text{no lens}}(\text{px})}{L_{\text{lens}}(\text{mm})/L_{\text{lens}}(\text{px})} \quad (2)$$

where $L_{\text{no lens}}$ and L_{lens} are the lengths of the structure in an image without and with lenses, respectively, measured in units of millimeters (mm) and pixels (px). The millimeter-to-pixel ratio was obtained from squared paper with 5-mm squares for the microscope without lenses, whereas a 1951 USAF Resolution Test Target (Thorlabs, Dachau, Germany) was used to obtain the millimeter-to-pixel ratio when the lenses were in place. For the high-magnification objective, we found $M = 254\times$. For the low-magnification objective we found $M = 27\times$.

B. Calculating the magnification

The magnification of the microscope can also be calculated from the distances between the object, lenses, and observer. In Figure 3, the light path of the microscope is drawn schematically. The magnification M of the first lens is simply given by the ratio of the lens to the intermediate image distance $d_{\text{intermediate}}$ and the lens to object distance d_{object} :

$$M = \frac{d_{\text{intermediate}}}{d_{\text{object}}} \quad (3)$$

The distance between the object and the second lens was measured with a ruler and found to be 240 mm. Because the intermediate image is located in the focus of the second lens, the object-to-intermediate image distance $d_{\text{object}} + d_{\text{intermediate}} = 240 - 53 = 187$ mm. To determine d_{object} and $d_{\text{intermediate}}$ individually, we have to use the lens formula (Eq. 1), which can also be expressed as:

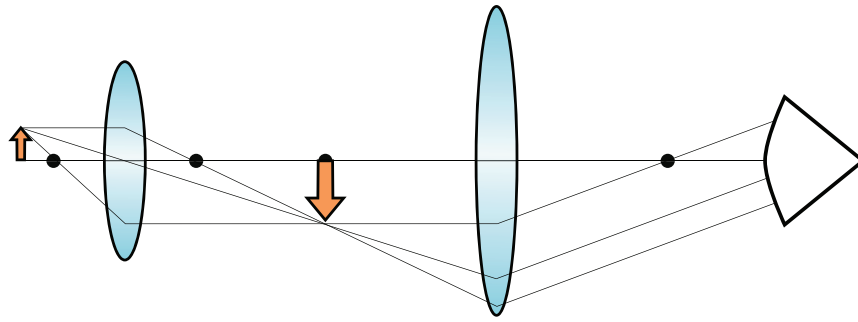


Fig 3. Schematic overview of the light path in the microscope. The object (here depicted as an arrow) forms an inverted intermediate image in the focus of the second lens. The second lens then sends collimated light to the observer.

$$d_1 = \left(\frac{1}{f_1} - \frac{1}{d_2} \right)^{-1} \quad (4)$$

Because the sum $d_{\text{intermediate}} + d_{\text{object}}$ is known, d_2 can be substituted,

$$\begin{aligned} d_1 &= \left(\frac{1}{3.85} - \frac{1}{187\text{mm} - d_1} \right)^{-1} \\ &= \frac{1}{2} \left(187 \pm \sqrt{187^2 - 4 \times 187 \times 3.85} \right) \text{mm} \end{aligned} \quad (5)$$

which shows that $d_{\text{object}} = 3.9 \text{ mm}$ and $d_{\text{intermediate}} = 183 \text{ mm}$. When substituting these numbers into Eq. 3, the magnification of the smartphone lens is $47\times$. This magnification is then further enlarged by the second lens, which magnifies it by a factor of the near distance to the eye [around 250 mm for adults (8)], divided by the focal length of the second lens, here 53 mm . That makes the total magnification $M = 220\times$, in good agreement with the value that was determined experimentally.

Total magnification with the low-magnification objective and the glass lens is determined in a similar way, substituting $f = 26.5 \text{ mm}$ in Eq. 5 for the focal length. In this case, $d_{\text{object}} = 32 \text{ mm}$ and $d_{\text{intermediate}} = 155 \text{ mm}$, leading to a total magnification of $M = 23\times$, again in good agreement with the value that was determined experimentally.

C. Quantifying the resolution

To quantify the optical performance of the microscope, the width of a sharp black-white transition is measured. To this end, a 1951 USAF

Resolution Test Target (Thorlabs) was used as shown in Figure 4a,b. The intensity profile along the red lines in Figure 4a,b is shown in Figure 4c,d, where the 3 black bars can be clearly recognized as dips in the intensity profile. In Figure 4e,f, a close-up of the intensity profile of a single bar is shown. This is a convolution of a step function with the point spread function (PSF) of the microscope. The PSF describes how a point-like object is imaged by a microscope. It depicts a fundamental limit of a microscope: when the PSFs of 2 objects overlap, they can no longer be optically resolved as different objects, but are imaged as 1 object. We approximate the shape of the PSF with a normal distribution, so that the intensity profile can be fitted to a cumulative distribution function of a normal distribution,

$$\Phi(x) = a \left(1 + \operatorname{erf} \left(\frac{x - \mu}{\sigma\sqrt{2}} \right) \right) + b \quad (6)$$

where σ is the deviation; μ the center of the normal distribution; a and b a prefactor and offset, respectively; and erf the error function. The full width at half-maximum (FWHM) of the normal distribution is then given by $\text{FWHM} = 2\sqrt{2 \ln 2} \sigma \approx 2.36\sigma$. Finally, because the width of the lines in the resolution test target is known, the FWHM can be converted from pixels to micrometers (μm). From the high-magnification objective (Fig 4a,c,e), $\text{FWHM} = 0.78 \pm 0.12 \mu\text{m}$ ($n = 6$). For the low-magnification objective (Fig 4 b,d,f), $\text{FWHM} = 4.3 \pm 0.9 \mu\text{m}$ ($n = 6$).

To compare this result to the maximal obtainable resolution, the Abbe diffraction limit is estimated for both objectives. The Abbe diffraction limit (9) states that 2 spots can be

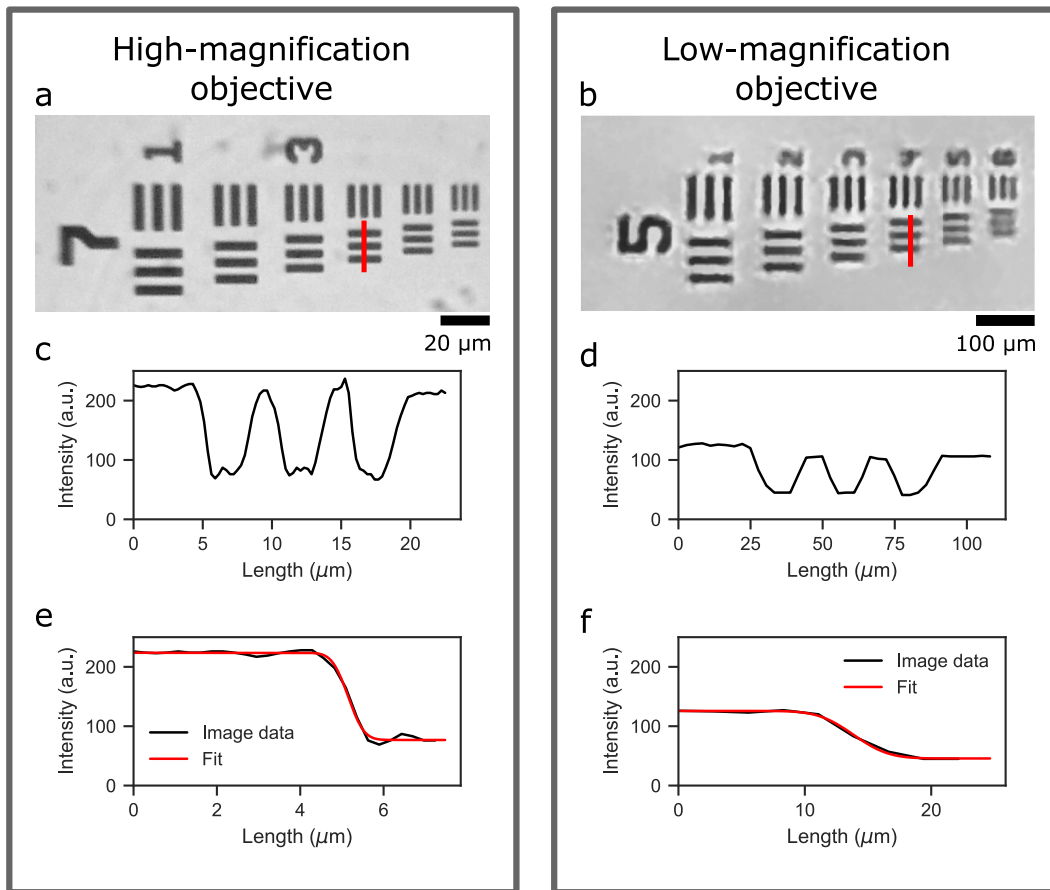


Fig 4. Quantifying the resolution of the microscope with a high-magnification and a low-magnification objective. (a, b) Using the known width of the lines in the USAF Resolution Test Target, the conversion factor from pixels to micrometers can be calculated. In these panels, the width of bars crossed by the red lines are 2.76 and 11.05 μm , respectively, and the corresponding intensity profiles along the lines are shown directly below. (c, d) Intensity profiles along the red lines using the pixel-to-micrometer conversion factor obtained from the calibration slide, (e, f) intensity profile along a single bar (black) that has been fitted (red) with a cumulative distribution function of a normal distribution.

optically separated if they are separated by a distance larger than d , given by:

$$d = \frac{\lambda}{2n \sin(\theta)} = \frac{\lambda}{2\text{NA}} \quad (7)$$

where λ is the wavelength of light, n is the refractive index of the medium through which the light travels, and θ is the maximum angle at which that light can still be collected by the lens. The numerical aperture NA is defined as $\text{NA} = n \sin(\theta)$. For the LEGO microscope, $n = 1$ because the sample and the objective are separated by air.

The angle θ needed to calculate the numerical aperture is obtained by taking the arctangent of the lens radius r_{lens} divided by the focal length f :

$$\theta = \arctan\left(\frac{r_{\text{lens}}}{f}\right) \quad (8)$$

For the high-magnification objective, we found a numerical aperture of 0.54, which, combined with an average wavelength of light of approximately $\lambda = 550 \text{ nm}$, leads to a resolution of $d = 0.51 \mu\text{m}$, which is only slightly smaller than the measured $0.78 \mu\text{m}$ that was estimated from the intensity profile in Figure 4e. For the low-magnification objective, the radius of the aperture of the LEGO brick that holds the lens needs to be used, rather than the radius of the actual lens. We found a numerical aperture of 0.29, which led to a resolution limit of $d = 0.95 \mu\text{m}$, a factor of 4.5 smaller than what was experimentally estimated from the intensity profile in Figure 4f. The poorer performance

of the low-magnification objective compared with the high-magnification objective is likely because a single lens is used in the low-magnification objective rather than the composite lens system of the smartphone camera lens used in the high-magnification objective. Indeed, a smartphone camera lens is designed to provide good image quality for low and high angles of incidence; here, we show that this lens functions close to the theoretically possible limit.

As an alternative to calculating the numerical aperture from the lens radius and its focal length, the numerical aperture is experimentally determined in Supplemental Figure S1. The measured value of the NA was comparable with the calculated value.

V. USE IN EDUCATIONAL SETTING

The goal of this project is to provide an accessible, hands-on microscope that can be used in an educational setting. In a preliminary attempt to quantify the effectiveness of the LEGO microscope as a learning tool, 8 students in the age range of 9–13 y were tested before and after following the provided workflow instructions. The tests were identical and contained 5 questions on the subject of microscopy as well as 5 general questions related to natural sciences that served as a benchmark (see Supplemental File S4). Although the subject of the microscopy questions was discussed in the workflow instructions, the answers were not included and had to be obtained by working through the different steps.

A. Workflow

The workflow instructions were organized in a 13-step procedure during which the students built the microscope step by step. The manual explores various aspects of the microscope. The workflow consists of the following basic steps:

1. Building of the 3 main parts of the microscope

The relevant parts should be built individually. At this point, the students are not

necessarily aware that the final outcome will be a microscope. Objectives are not yet added in this stage. However, the lens of the ocular is already mounted.

2. Exploration of the different parts

The students are asked to play with the different parts and to guess what they can be used for.

3. Focus on the ocular

The students are asked to inspect the ocular closely. They should find out that it can be used as a magnifying glass. They should realize that because of the closed construction, typically not enough light is delivered to the sample.

4. Combination of light source and ocular

To overcome the problem of low light, the students should realize that using a light source, built in a previous step, produces a better view of the magnified sample.

5. Exploration of individual samples

The students are asked to inspect arbitrary samples with the magnifying property of the ocular.

6. Interruption to recapitulate experiences

The students are asked to collect the main experiences gathered so far. This ensures that the important lessons to work on the following steps have been learned. In a classroom setting this should be done together with all groups.

7. Identifying possible ways to increase magnification

Besides using stronger lenses, the students should figure out that 2 magnifying glasses might improve the total magnification. If this step was not suggested by the students, it will be introduced either by the teacher or by the help in the manual.

8. Introduction of the objectives

To begin, the students are exposed to the low-magnification objective. They should try to use both the ocular and the second lens to have an improved magnification. Because this step requires very precise alignment, students will mostly fail, thus introducing the requirement of stable mounting. Several helping strategies are included for this step.

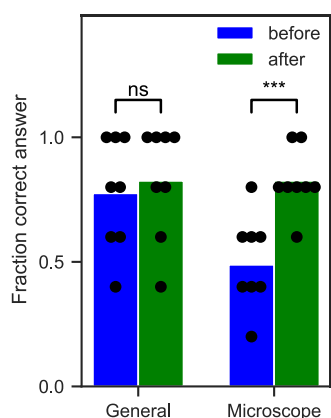


Fig 5. Fraction of correctly answered questions on subjects of natural sciences and microscopy, before and after using the LEGO microscope. The bars show the average results of 5 questions on either of the subjects, filled in by 8 students in the age range between 9 and 13 y. The black dots correspond to the results of individual students. There was no significant (ns) change in the results of the general science questions: from $78 \pm 21\%$ to $83 \pm 21\%$ correctly answered questions, but a statistically significant improvement (***, $P < 0.001$) was observed for the questions on microscopy, from $50 \pm 17\%$ to $83 \pm 12\%$ correct answers.

9. Interruption to collect experiences

The idea of fixing the lens position is discussed or introduced in the help section of the manual.

10. Combination of all parts

In this step, the students combine all parts to get a working microscope. Now they are also told that they have successfully constructed a LEGO microscope.

11. Exploration of samples with the microscope

Samples are explored, as suggested in the sample part of this paper or provided by the teacher.

12. High-magnification objective

To improve the magnification, the second objective is introduced at this stage. Because of the appearance of the lens, the similarity with a magnifying glass is lost. However, after integration in the microscope, micrometer-sized objects can be visualized.

13. Inspection of micrometer-sized samples

In the final step, the students are asked to inspect samples from the micro-world. Biophysical effects, such as osmotic pressure and the motion of microswimmers, are introduced here.

After having finished this workflow, the second test was taken, asking the same questions as the initial test. The general questions serve as a control benchmark, and, as shown in Figure 5, the fraction of correct answers of this control changes only marginally after the students have worked with the microscope. The fraction of correct answers on the microscopy questions however, almost doubles. Although the group size was relatively small, the effect is significant.

B. Example experiments

One aim of the LEGO microscope is to provide children and teenagers an easy and playful access to optics and biophysics. After building the microscope, simple objects of everyday use can be imaged, such as hairs from humans and animals or natural and artificial fibers from clothing. To improve the learning effect, we furthermore propose a series of simple experiments that can be performed at home that allow discovering some of the main biophysical principles. Specifically, we suggest discovering the process of crystallization, to observe the effect of osmosis on plant cells and to discover the motion of microswimmers.

Among the simplest, yet impressive samples is a salt crystal (Figure 6a). The shape of a sodium chloride (NaCl) crystal reflects the face-centered cubic lattice formed by the larger chloride ions that are surrounded by the smaller sodium ions. As described in the material section, these crystals can be prepared and observed.

Alternatively, when a thin film of the salt solution is placed on a microscope slide, the concentration of salt will slowly increase from evaporation of water so that the crystallization process itself can be recorded.

Osmosis is an important biophysical effect that regulates shape and pressure in many biological systems, such as the turgor pressure in plant cells that presses the plasma membrane against the cell wall and gives plant tissue its rigidity. In Figure 6b, a time-lapse is shown of red onion epidermal cells exposed to an osmotic shock. The full movie is shown in Supplemental

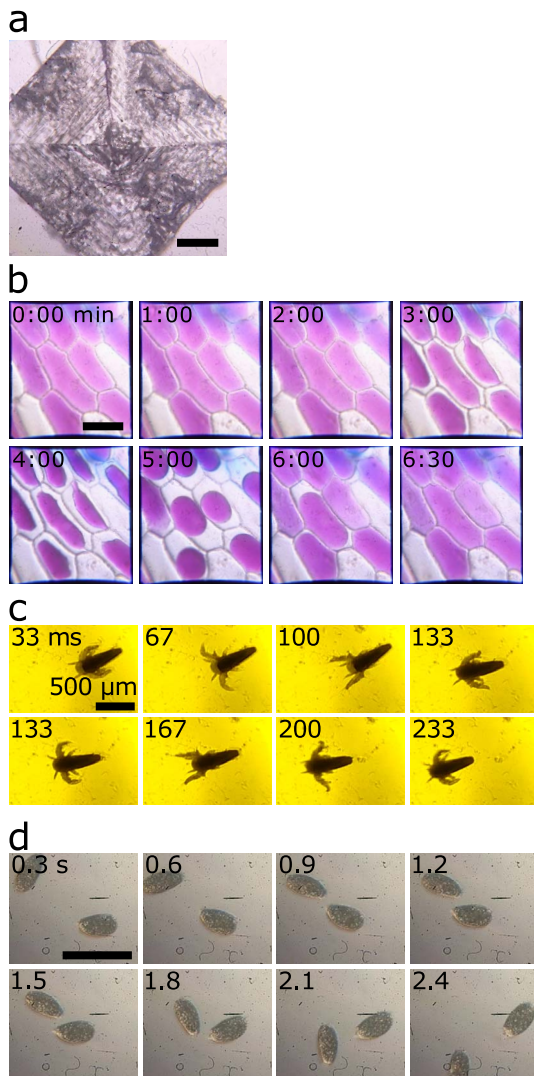


Fig 6. Examples of experiments conducted with the LEGO microscope. (a) Image of a sodium chloride crystal. (b) Time lapse of an osmotic shock in red onion cells. After approximately 30 s, a 1 M NaCl solution is flowed in. Subsequently, water leaves the cells, causing the cell membranes to detach from the cell walls. After approximately 5 min, distilled water is flowed in, washing away the 1 M NaCl solution, and the cells return to their original volume. (c) Time lapse of the movement of an *Artemia* shrimp in water. (d) Time lapse of the movement of 2 water fleas in water. The scale bars in panels a, b, and d are 100 μm .

Movie S5. At approximately $t = 30$ s, a drop of 1 M NaCl solution is placed on one open side of the coverslip, which, because of capillary forces and a piece of absorbing paper on the other side of the coverslip, ensured that the cells were rapidly surrounded by the NaCl solution, creating a large difference in osmotic pressure between the inside and outside of the cells and resulting in a loss of water. As a conse-

quence of the volume reduction, the plasma membrane detaches from the cell walls starting around $t = 3$ min, and the pigment becomes more concentrated. At approximately $t = 5$ min, the NaCl solution is replaced with distilled water, which enters the cells and returns them to their original volume. During the recording, care was taken not to touch the microscope to avoid movement of the microscope slide. No drift in focus was observed during our measurements, avoiding the need to adjust the height of the objective, which in turn would lead to motion of the microscope.

To discover and image systems that are currently the research focus of many biophysics groups, a series of different microswimmers can be imaged with the LEGO microscope. The propulsion of microswimmers is fundamentally different from macroscopic swimmers because it occurs in a low Reynolds number regime, where friction is the dominating force (10). Relatively large microswimmers are *Artemia* shrimps. Their movement is imaged with the low-magnification objective and is shown in Figure 6c and Supplemental Movie S6. *Artemia* shrimps can swim relatively fast by using their antennae for regular strokes. In contrast, unicellular organisms apply different strategies for propulsion. These organisms can be grouped in two categories: *flagellates* that use a small number of long flagella for movement and *ciliates* that have a large number of small, hair-like flagella (called cilia) covering the entire organism (11). Flagellates are typically smaller than ciliates, and they are hard to resolve with the LEGO microscope. On the other hand, the movement of ciliates can be imaged well. An example is the propulsion of paramecia, unicellular organisms that are covered with cilia and are commonly found in ponds. Their movement is imaged with the high-magnification objective and is shown in Figure 6d and Supplemental Movie S7.

VI. CONCLUSION

With LEGO and low-cost, easily available lenses, it is possible to construct a microscope that can resolve micrometer-sized objects with a resolution that is close to the diffraction limit

of light. A series of experiments is suggested covering different fields of natural sciences that can be conducted with household ingredients. The modular design of the microscope itself also lets it be incorporated easily into a curriculum on optics. A preliminary study with 8 students in the age range of 9–13 y shows that the understanding of microscopy increases after working with the LEGO microscope.

Because the design of the microscope presented here is only one of many possible configurations, customization is highly encouraged.

SUPPLEMENTAL MATERIAL

Supplemental materials are available at: <https://doi.org/10.35459/tbp.2021.000191/>. Supplemental File S1 has step-by-step building instructions for the LEGO microscope. File S2 is a list of parts for the LEGO microscope. File S3 is a manual to explore the LEGO microscope step by step. File S4 has a series of questions that were used to test knowledge on microscopy before and after using the LEGO microscope.

Supplemental Movie S1 has instructions for replacing the LED in the LEGO brick to change the color of illumination. Movie S2 has instructions for the preparation of red onion epidermal cells for microscopy. Movie S3 has instructions for closing the sides of a sample slide for flow experiments. Movie S4 has instructions for performing an osmotic shock experiment. Movie S5 shows red onion epidermal cells exposed to an osmotic shock by the addition of a 1 M sodium chloride solution. Movie S6 shows the movement of *Artemia* shrimps observed with the low-magnification objective. Movie S7 shows the movement of paramecia observed with the high-magnification objective.

Supplemental Fig S1. Experimental determination of the numerical aperture of the high-magnification objective. A green laser is refracted by the smartphone camera lens and illuminates the back side of a paper box. By measuring the distance from the lens to the box and the diameter of the refracted circle, the maximum angle θ under which light left the lens is estimated. The average of multiple measurements results in a numerical aperture of 0.53, close to what was calculated from the lens diameter and its focal length.

AUTHOR CONTRIBUTIONS

BEV, EBB, and TB designed the microscope. BEV and TB carried out experiments, analyzed data, and wrote the manuscript.

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