An Easily Made, Low-Cost, Tissue-Like Ultrasound Phantom Material

Ronald O. Bude, MD, and Ronald S. Adler, MD, PhD

Ultrasound phantoms are generally of two types. One mimics the acoustic properties of tissue (with regard to the speed of sound, average attenuation, etc.). The main purpose of the other is to approximate the sonographic appearance of tissue. The latter is often used as a biopsy training aid. Those which mimic the acoustic properties of tissue have been constructed of agar with suspended graphite, polyurethane foam, and magnesium silicate gels, and are used chiefly as test phantoms for assessing diagnostic ultrasound imaging equipment or for studying the interaction of sound with tissue. They are generally not used for biopsy phantoms as they are either expensive or time-consuming to produce. Biopsy phantoms are simpler in construction, contain simulated cysts or masses, and are either echogenic or sonolucent. Echogenic media have consisted of a flour or cornstarch in gelatin suspension, a silicium carbide powder in agar suspension, or agar. When properly made, their echogenicity simulates parenchymal tissue (except agar, which is only weakly echogenic and does not simulate parenchymal echo texture well); however, they can be laborious to produce, requiring stirring or rotation during cooling to insure that the scatterers or biopsy targets remain suspended and may require materials that are not widely available (agar, silicium carbide). Sonolucent media, on the other hand, consist of gelatin with suspended scatterers and are very easy and inexpensive to prepare. Unfortunately, they are also transparent (unless deeply colored), which allows the biopsy needle and biopsy target to be seen from the exterior. This does not mimic the in vivo situation, and makes the procedure artificially easy to perform. Additionally, the biopsy needle is artificially easy to demonstrate sonographically in a sonolucent or hypoechogenic medium compared to in vivo, as it stands out as a much more brightly echogenic structure in those media than in most tissues, again making the procedure artificially easy to perform. These shortcomings limit their usefulness in training for in vivo biopsy procedures.

We describe an easily made, low-cost preparation that possesses the advantages of both types of previously reported biopsy phantoms, with none of the disadvantages. It is visually opaque with an echogenicity which simulates parenchymal echo texture, yet is as easy and inexpensive to produce as clear gelatin phantoms.

MATERIALS AND METHODS

Ingredients in the following proportions, in the total volume desired, are stirred until completely dissolved, which takes only 1 or 2 minutes: 250 mL (approximately 1 cup) boiling water, 20 g of unflavored gelatin, and 10 g (approximately 1 tablespoon) of sugar-free psyllium hydrophilic mucilloid fiber (brand name: sugar-free Metamucil). If the sugar-containing variety is used, approximately three times the volume (3 tablespoons) will be required to obtain the same amount of psyllium fiber. To form a phantom without internal inclusions such as simulated “cysts” or “masses,” this mixture is poured into the container selected (plastic bag, milk carton, etc.) and cooled until firm. One “unit volume” as described above, roughly cubic in shape, takes 1 hour to 2 hours to congeal in a refrigerator at 6°C.

Forming the phantom in stages is the most satisfactory method to produce internal “cysts” or masses. A layer of mixture is poured into the container of choice and cooled until the surface is firm enough to support the intended inclusion, which is placed on it (the remaining mixture has
been partially submersed in a container of warm water to keep it from congealing). If the inclusion does not float, the remainder of the mixture is poured into the container and allowed to congeal. If the intended inclusion floats, a thin layer of mixture is poured into the container and the inclusion placed within this layer. The layer is allowed to congeal, trapping the inclusion. The remaining mixture is then poured over the trapped inclusion and cooled until firm.

“Cysts” are simulated with water-filled balloons, tips of examining gloves, grapes, or glycerine suppositories. “Masses” are simulated with carrot pieces, macaroni, olives, or hot dog pieces. If refrigerated, the phantom will not begin to undergo significant microbial degeneration for several weeks. We have found this mixture to be very “forgiving,” and the gelatin concentration can be reduced by approximately 50% without a substantial change in firmness.

DISCUSSION
We have produced a tissue-like ultrasound phantom using a previously unreported material, psyllium hydrophilic muciloid fiber, as the scattering medium (Figure 1). The echo texture of this material simulates thyroid or testicular parenchyma. The chief advantage that the use of this material provides, compared with previously reported tissue-like phantoms, is that after the mixture is initially prepared, further mixing is not required to maintain an even suspension of the scattering medium. This is in contrast to phantom materials using flour, cornstarch, or silicon carbide as the scattering medium, which require intermittent stirring during cooling until congealing of the mixture occurs, a process which can take more than 1 hour. Phantoms prepared using psyllium hydrophilic muciloid fiber can be prepared much more easily, more quickly, and more reproducibly than phantoms made with previously reported materials. This material is very inexpensive, costing less than $1 per unit volume (approximately 250 mL), uses widely available materials, and is opaque (so that biopsy needles and targets can only be seen sonographically and not with the naked eye). A strip of gelatin-impregnated gauze, which simulates skin and
TISSUE-LIKE ULTRASOUND PHANTOM

helps prevent surface laceration, can be coated onto the scanning surface with a small amount of gelatin to prolong the life of the phantom.

REFERENCES