

Clinical experience of the Magseed[®] magnetic marker to localize non-palpable breast lesions: a cohort study of 100 consecutive cases

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Background: The aim of this study was to report on a cohort of 100 patients where the Magseed[®] paramagnetic marker was used to localize non-palpable breast lesions.

Methods: Data were collected from a cohort of 100 patients with non-palpable breast lesions, who underwent localization using the Magseed[®] marker. This marker consists of a paramagnetic seed that can be seen on mammography or ultrasound and intraoperatively detected with the use of the Sentimag[®] probe. The data were collected over a period of 23 months (May 2019 to April 2021).

Results: All 111 seeds were successfully placed in the breasts of 100 patients under ultrasound or via stereotactic guidance. Eighty-nine seeds were inserted in single lesions or small microcalcification clusters in a single breast, 12 seeds were deployed to a bracket microcalcification clusters and 10 to help localize two tumors within the same breast. Most Magseed[®] markers (88.3%) were placed in the center of the lesion (≤ 1 mm). The re-excision rate was 5%. All Magseed[®] markers were successfully retrieved and no surgical complications were observed.

Conclusions: This study reports our experience in a Belgian breast unit using the Magseed[®] magnetic marker and it highlights the many advantages of the Magseed[®] marker system. With this system, we successfully identified subclinical breast lesions and extended microcalcification clusters, targeting multiple sites within the same breast.

Keywords: Surgery; breast cancer; localisation; Magseed

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Introduction

In recent years, the implementation of mammography screening programs as well as new medical imaging technologies such as tomosynthesis or artificial intelligence and women's increased awareness of breast cancer, have resulted in an increase in the detection rate of non-palpable lesions (1-5). These tumors are too small in size or located too deep in the breast parenchyma to be detected during clinical examination. Consequently, techniques for pre-

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operative identification of non-palpable cancerous lesions during breast conservation surgery have been developed. The success of breast conservation surgery requires effective communication between the radiologist and the surgeon, and precise guidance by the radiologist to allow the surgeon to perform complete removal of the lesion with healthy resection margins, while striving to preserve as much as possible the breast's aesthetic curve. However, this also requires operating schedule management and the practitioner availability. There are many guide techniques available. The standard localization method used is wireguided localization (WGL), in which wire hook is placed into the tumor with ultrasound or stereotactic guidance (6,7). However, this technique has several disadvantages. One of the most important being that the guide wire has to be placed within a few hours of surgery to minimize the risk of dislocation. This requires interdependence of radiology and surgery schedules, reducing operating room efficiency. Furthermore, wire guides cause discomfort for patients and migration or transection can occur (8). The WGL may also limit alternatives for placement and the distance between the entry site and the tip of wire may be greater, thus leading to extensive dissection and affecting potential aesthetic outcome. Consequently, alternative guide methods were developed such as radioactive seed localization (RSL), non-radioactive radar localization, radio frequency identification and magnetic seed localization (9). In this study, we evaluated the Magseed[®] magnetic marker

Highlight box

Key findings

 The Magseed[®] magnetic marker combined with the Sentimag[®] probe is an effective method to localize non-palpable breast lesions.

What is known and what is new?

- Several reports have demonstrated the feasibility and safety of this technique compared to the wire-guided localization (WGL) technique.
- We demonstrated our clinical experience on 100 consecutive patients with whom we used the Magseed[®] for identification of subclinical breast lesions and is effective when extended microcalcification clusters are present, or in targeting multiple tumors within the same breast.

What is the implication, and what should change now?

• Further studies are needed to validate this technique with all practitioners and also to prove its efficacy of other long-term applications for patients who require neoadjuvant treatment.

(Endomagnetics, Cambridge, UK). The Magseed[®] marker is a non-radioactive paramagnetic seed that can be inserted under ultrasound or with stereotactic guidance. During surgery, the seed is detected using a magnetic detection probe (Sentimag[®], Sysmex, GmBH, Hamburg, Germany). Several reports have demonstrated the feasibility and safety of this technique (10-16). The aim of this study is to report the experience of 100 consecutive cases where the Magseed[®] marker was used to localize non-palpable breast lesions. We present this article in accordance with the STROBE reporting checklist (available at https://gs.amegroups.com/ article/view/10.21037/gs-22-552/rc).

Methods

Patient selection

This retrospective study includes 100 patients who were managed at the Center of Senology Drs Crèvecoeur (a private radiology center) and the CHC MontLegia hospital between May 2019 and April 2021. Patients with nonpalpable lesions undergoing breast-conservation surgery were deemed suitable for seed insertion following discussion among the multidisciplinary team. We also included patients with complete or partial response of neoadjuvant chemotherapy (NAC). For these patients, an intramammary clip was placed before their scheduled NAC. The followups during their chemotherapy assessing response were performed by mammography and ultrasound. At the end of treatment, magnetic resonance imaging (MRI) was performed to obtain the precise dimension of the residual tumor, or the marker clip released before NAC in case of complete tumor response, was then marked by the Magseed[®] marker. Exclusion criteria were pregnancy/ lactation or any mental condition with might render the patient incapable of giving written consent and patients who denied to sign the consent form. The study was approved by the CHC MontLegia Ethics Committee (reference No. 20/40/1052) and was conducted in accordance with the Declaration of Helsinki (as revised in 2013). Written informed consent was obtained from all participants included.

Description of guide method

We used the Magseed[®] marker (Endomagnetics, Cambridge, UK), a magnetic seed localization technique.

It consists of a 5 mm \times 1 mm cylindrical non-radioactive paramagnetic surgical steel marker that can be visualized via mammography or ultrasound. It is supplied in sterile packaging preloaded into an 18-gauge 7-cm steel needle. The marker can be inserted at any time before surgery. In the operating room, the Magseed[®] marker is detected using the Sentimag[®] probe which generates an alternating magnetic field, transiently magnetizing the seed and subsequently measuring the magnetic field emitted by the seed.

Technique

Seeds were inserted using an aseptic technique by ultrasound guidance. All seeds were inserted by the same breast radiologist. Xylocaine 2% was used to anesthetize the skin and a small incision was made with a blade to facilitate insertion of the introducer needle. Two mammograms (craniocaudal and mediolateral views) were carried out to confirm the position of the seed at the target site and successful localization. X-ray dimensions of the tumor, as well as the distance between the seed and the tumor were documented. This procedure was performed for solid tumors. In the cases of lesions with microcalcifications, seeds were inserted by ultrasound guidance using the biopsy clip as a target. For some cases, no biopsy clip was present, so the radiologist used the post-biopsy haematoma to insert the Magseed[®] marker on the ultrasound guidance. If we did not have a biopsy clip or ultrasound image (like hematoma) to visualise the lesion, we took an X-ray with the introducer needle in the breast before dropping the magseed. In case of incorrect positioning, we had the possibility to correct the trajectory. The stereotactic guidance to target residual calcification was not used because the displacement or migration of the clip during decompression was frequently observed. Finally, if the patient had two non-palpable lesions or extensive microcalfication clusters, two Magseed® markers were inserted.

Surgical excision

Once in the operating room, the surgeon used the Sentimag[®] system's probe to locate the Magseed[®] marker with precision. The surgeon was guided by the two mammograms, which allowed placement with the Sentimag[®] probe in the correct place on the breast's skin, and with the Sentimag[®] units that give the numerical counter along with its audio tone. This enabled the surgeon to gauge the distance between the seed and the probe. As soon as the lesion was localized (maximum magnetic signal), the surgeon made an incision and inserted the probe to confirm the seed's position. Once excised, the surgeon then used the probe to confirm that the seed was included in the specimen. For this study, surgical excision was carried out by two breast surgeons trained in the technique.

Specimen radiography was carried out for each patient with microcalcifications to provide reassurance that all the lesions and seeds had been excised. The examination of the specimen radiography was performed by the breast radiologist who inserted the Magseed[®] marker.

Pathological examination

The pathologist analyzed the specimen, both the dimensions of the specimen and the tumor were documented. Macroscopic extemporaneous examination was performed for all solid tumors for detection of the seed and margins evaluation. Then the tissue was formalin-fixed, paraffin-embedded and entirely sectioned at 4-µm intervals for the definitive histopathological examination.

Postoperative histology was discussed at a multidisciplinary breast meeting and re-excision or mastectomy was proposed to patients if positive margins has been observed.

Analysis

Data were extracted from the electronic medical record after written consent of patient. Data included patient age, radiological features, placement/localization of seed, identification/retrieval of seed, surgical features and pathologic results. Percentages for categorical variables, mean and median were calculated as appropriate.

Results

Patient characteristics and Magseed® marker insertion

In the study, one hundred patients were included; their baseline characteristics are shown in *Table 1*. A total of 111 Magseed[®] markers were placed into the breasts of 100 patients using ultrasound guidance. No infection, hematoma or migration problems were observed. All Magseed[®] markers were retrieved, as confirmed by specimen radiography or by the pathologist during extemporaneous analysis. The average age of women was 63 years. Magseed[®]

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Table 1 Patient characteristics and basic information regarding $\mathsf{Magseed}^{\circledast}$ insertion

Patients characteristics	Number
No. of women	100
Age (years), mean [range], median	63 [32–78], 66
No. of seeds	111
Number of days between Magseed [®] marker placement and surgery, mean [range], median	5.3 [0–42], 4
Placement success, n (%)	
≤1 mm	98 (88.3)
2–5 mm	7 (6.3)
6–10 mm	6 (5.4)

markers were placed a median of 4 days before surgery. Majority of the Magseed[®] markers (88.3%) were inserted successfully into the target lesion.

Figure 1 shows an example of a Magseed[®] marker placed very precisely next to the biopsy clip. This clip was placed during macrobiopsy via stereotactic biopsy of the microcalcification clusters. Radiography of the surgical specimen (*Figure 1C*) confirmed the resection of the microcalcifications, the Magseed[®] marker and the biopsy clip.

Histological examination of the surgical specimen demonstrated the presence of microcalcification and neoplastic cells corresponding to ductal carcinoma *in situ* (*Figure 2*). Actin combined with P63 showed the persistence

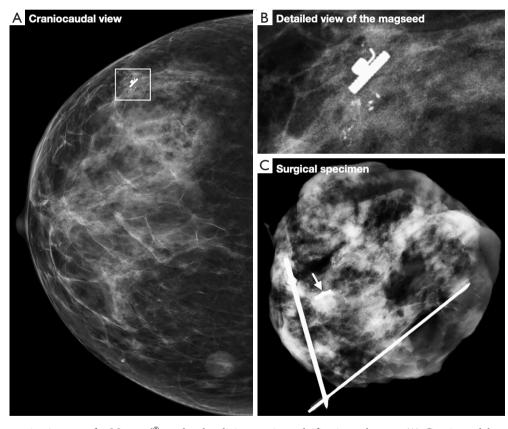


Figure 1 Representative images of a Magseed[®] marker localizing a microcalcifications clusters. (A) Craniocaudal mammogram image confirms placement of the Magseed[®] marker adjacent to the biopsy clip. (B) Detailed view of the Magseed[®] marker placed next the microcalcification clusters and the biopsy clip. (C) X-ray of the surgical specimen. The Magseed[®] marker and biopsy clip are indicated by the white arrow.

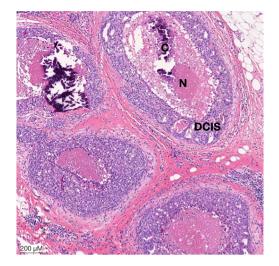


Figure 2 Histology study of the case illustrate in *Figure 1*. Tissue section stained with hematoxylin and eosin showing C, N and DCIS (original magnification ×50). C, microcalcifications zone; N, necrosis zone; DCIS, ductal carcinoma in situ.

of myoepithelial layer cells around all glandular structures and the absence of invasive tumor processes.

Although the majority of cases involved identification of the lesion with one Magseed[®] marker some cases required the use of two markers. For 89 women, one Magseed[®] marker was inserted into their lesion which corresponded to either a mass or microcalcifications. When two lesions were diagnosed in the same breast, they were located with two Magseed[®] markers. In cases with extended microcalcification clusters, two seeds were deployed to bracket the lesion (*Figure 3*). Histological analysis of the tumorectomy specimen shown in *Figure 3* demonstrate the presence of invasive process (*Figure 4*).

The Magseed[®] technique also allows the identification of non-palpable lesions such as a radial scar (*Figure 5*). The case illustrated in *Figure 5* corresponds to a patient with breast implants. The radiologist successfully placed the seed despite the presence of breast implants.

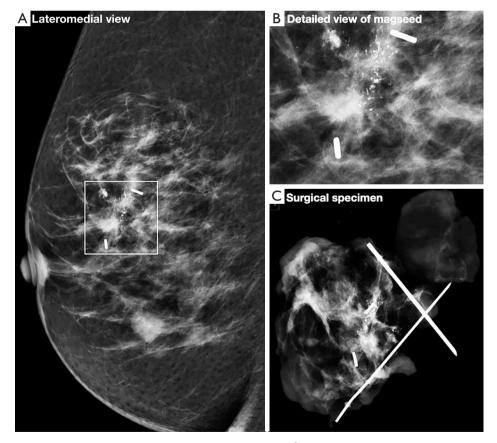


Figure 3 Localization of extended microcalcification clusters by two Magseed[®] markers. (A) Lateromedial view of a right breast depicting microcalcification clusters. (B) Detailed view of two Magseed[®] markers surrounding microcalcifications. (C) Radiography of the surgical specimen confirming successful resection.

Surgical excision

All Magseed[®] markers were located with a Sentimag[®] probe by the surgeon. Intraoperative identification and excision of the localized lesion was successful in 100% of patients. In patients where Magseed[®] markers was used for bracketing microcalcifications, the median size of the lesion was 27.5 mm and the mean size was 36.3 mm. Regarding patients with two solid tumors, the median size between Magseed[®] markers was 25 mm and the mean size 24.6 mm. For two patients, two Magseed[®] markers were placed at a distance of less than 10 mm and this small distance complicated their detection. The diagnosis for these patients was made by a radiologist from outside our

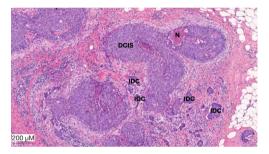


Figure 4 Histology study of the case illustrate in *Figure 3*. Tissue section stained with hematoxylin and eosin showing DCIS, IDC, N (original magnification ×50). DCIS, ductal carcinoma in situ; IDC, invasive ductal carcinoma, N, necrosis.

team and described a bi-focal lesion. As a precaution, the surgeon preferred to use two Magseed[®] markers. In one case, we also performed a skin surface marking.

Pathological examination

During examination of the specimens by the pathologist, identification of the Magseed® marker was not always straightforward. Whenever necessary, an X-ray of the surgical specimen was performed. From the time of their placement to the time of surgery, no migration of the Magseed[®] marker in the breast was observed. Majority of resected tumors were solid tumors. From a histological point of view, 57.6% of the patients had invasive ductal carcinoma, 16.2% ductal carcinoma in situ, and 9.0% invasive lobular carcinoma (Table 2). The radiological dimensions for the masses were on average of 7.0 mm and for microcalcifications 24.6 mm. The majority of the microcalcification clusters were small. One patient presented with an extensive microcalcification clusters corresponding of 100 mm. A good correlation between radiological-pathological dimensions were observed for solid tumors. Regarding dimensions of microcalcifications, sizes of lesions described by the pathologist and the radiologist were different. Radiological dimensions were performed before the diagnostic biopsy leading to reduction in lesion extent.

Complete excisions of the different lesions were

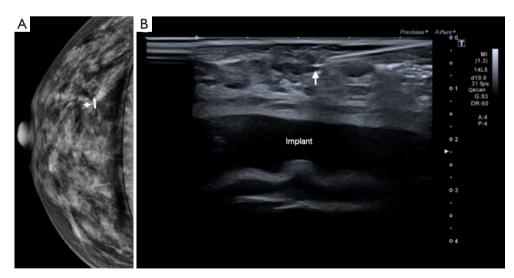


Figure 5 Representative craniocaudal view shows the Magseed[®] marker inserted in a non-palpable lesion corresponding to a radial scar (A). Insertion of the Magseed[®] marker under ultrasound guidance (B). Radial scar is indicated by the white arrow.

Target lesion characteristics	Number (%) –	Radiological dimensions (mm)			Pathological dimensions (mm)		
		Mean	Median	Range	Mean	Median	Range
Type of lesion			·				
Solid tumor	89 (80.2)	7.0	7.5	0–40	9.5	9.0	0–50
Microcalcifications	22 (19.8)	24.6	20.0	3–100	19.4	15.0	3–100
Specimen volume (cm ³)							
Solid tumor	_	-	-	-	27.6	21.0	4.7–141.3
Microcalcifications	-	-	-	-	59.6	26.0	5.5-403.0
Surgical pathology findings							
Invasive ductal carcinoma	64 (57.6)	-	-	-	-	-	-
Ductal carcinoma in situ	18 (16.2)	-	-	-	-	-	-
Invasive lobular carcinoma	10 (9.0)	-	-	-	-	-	-
Response after neoadjuvant chemotherapy	5 (4.5)	-	-	-	-	-	-
Other invasive tumor	5 (4.5)	-	-	-	-	-	-
Radial scar	2 (1.8)	-	-	-	-	-	-
Fibroadenoma	2 (1.8)	-	-	-	-	-	-
Benign	3 (2.7)	-	-	-	-	-	-
Lobular carcinoma in situ	1 (0.9)	-	-	-	-	-	-
No atypical epithelial hyperplasia	1 (0.9)	-	-	-	-	-	-

Table 2 Surgical excision data and pathological characteristics

 Table 3 Pathological findings for patients with positive margins

Pathological findings for patients with positive margins	Number [%]
DCIS	2 [40]
IDC and DCIS	2 [40]
ILC and LCI	1 [20]

DCIS, ductal carcinoma in situ; IDC, invasive ductal carcinoma; ILC, invasive lobular carcinoma; LCI, lobular carcinoma in situ.

performed with disease-free margins of at least 1 mm in 95% of patients. Five patients underwent further surgery for positive margins and pathological finding are described in *Table 3*. Of these, two patients had a further re-excision margins and three had radical mastectomy. For these five patients, each Magseed[®] marker was placed in the center of the tumor lesion (≤ 1 mm). None of the cases were attributed to the localization technique.

Discussion

The present study illustrates the use of the Magseed® marker, a promising technique allowing localization of nonpalpable breast lesion. Previous reports have demonstrated safety, feasibility and clinical use of magnetic seeds (10-18). We report our experience using the Magseed® magnetic marker and the Sentimag[®] probe in a Belgian breast unit. Our team had been using the standard WGL technique for intraoperative tumor localization for 30 years. Several disadvantages and limitations linked to the use of WGL, including the risk of migration or the need to place the wire on the same day as surgery, have led to the development of new guide techniques. This study demonstrates that the Magseed[®] marker offers a convincing clinical alternative to the wire hook. Moreover, this technique allows us greater flexibility in our scheduling, while allocating personnel more effectively and optimizing theater utilization, reducing costs and delays.

Our study describes our experience with the Magseed[®] marker in a cohort of 100 patients. Implementation of the Magseed[®] System was straightforward, with a short period of adaptation for the surgeon and the radiologist. Consequently, in a short period, this method has effectively replaced guide wire use. One of the main advantages of the Magseed[®] marker is that the seed can be placed several days before surgery allowing dissociation of the radiology and surgery schedules. In addition, the Magseed® marker has now been licensed for use over 30 days before the surgery. Timing is undefined provided there is the intention to excise. There were no apparent problems associated with the length time of the Magseed[®] marker was inserted prior to surgery. In a few cases, the Magseed[®] marker was placed on the day of surgery. The longest time between placement and removal was 42 days. All the lesions were successfully removed along with the Magseed[®] marker suggesting no migration of the markers. No other problems were reported. The Magseed[®] marker is principally used for non-palpable breast lesions but since there is no defined time limit for the removal of the seed, it could be inserted before NAC. In this case, it is of real benefit for the patient who would only need one type of guide system to localize their tumor instead of two. The follow-up of the patient during their chemotherapy to assess response could be carried out solely by mammography and ultrasound. MRI use is not recommended to perform because the Magseed® marker can create a bloom artefact which measures up to 4 cm (19,20). Further studies must be carried out in order to validate the Magseed® marker for tumor localization in patients undergoing NAC.

Our findings demonstrate a relatively low re-excision rate of 5%. This rate is significantly lower than that reported by other studies comparing WGL to the Magseed[®] marker (19). Zacharioudakis et al. observed no significant difference between the Magseed[®] cohort (16%) and the WGL cohort (14%) (13). This low re-excision rate observed in our study can be explained in several ways. Firstly, the same radiologist placed all 111 Magseed[®] markers. Secondly, both surgeons involved in this study were specialized in oncological breast surgery. In addition, for 87% of the patients included in our study, the radiologist carried out the diagnosis, the placement of the Magseed[®] marker and the radiography of the surgical specimen when necessary. Good preoperative communication between the radiologist and the surgeon is essential. For each patient, the radiologist explained to the surgeon how the Magseed[®] was inserted (distance between tumor and seed,

insertion point, etc.) using mammogram images. Among the patients with positive resection margins, MRIs were carried out for three patients. MRIs did not highlight tumor extension and radiological dimensions were similar to those on mammography and ultrasound. In one patient, the diameter of the tumor was 8 mm, therefore not meeting MRI criteria, and the pathological analysis of the specimen showed a non-calcified ductal carcinoma in situ. With regard to the last patient with positive resection margins, diagnosis was not made by the radiologist who inserted the Magseed[®] marker. This confirms the influence on patient care if several radiologists are involved. Another difference

between the WGL and the Magseed[®] marker lies in the initial incision. Indeed, surgical approach and incision placement is independent of localization access. This allows a more aesthetic result on the breast's curve.

We also demonstrated the feasibility of targeting multiple lesions in the same breast. Five patients had two Magseed[®] markers placed in the same breast to localize two tumors. Two markers were deployed to bracket extended microcalcifications clusters for six patients. There was no difficulty in discriminating between the different sites. For patients with microcalcifications, the distance between Magseed[®] markers was superior to 20 mm. However, two patients had two lesions less than 1 cm apart. For safety reasons, two Magseed[®] markers were placed. Despite the complexity in differentiating between the markers, the surgeon proceeded with successful removal of both. Some reports have demonstrated that the Magseed[®] marker was not detectable if the tumor lesion was deeper than 6 cm (10-22). In our study, one Magseed[®] was placed at a depth of 3.6 cm. The diameter of the tumor was 11 mm (Figure S1). Although the surgeon experienced some difficulty in localizing the Magseed[®], he successfully removed the Magseed[®] and the tumor with negative margins.

One disadvantage of this method is that, if deployed in the wrong place, the incorrectly placed seed would need to be surgically excised in addition to the correctly sited device.

For each patient, identification of the Magseed[®] markers in specimens was performed by the pathologist and by radiography whenever necessary. The pathologist reported some difficulties in finding the seed. A change in size or color would be desirable to optimize detection.

Other alternatives to WGL and Magseed[®] have become available. These include radio-guided occult lesion localization (ROLL) (23), RSL (24), the infrared radar technique SCOUT (20) and LOCalizer using radiofrequency identification (25). Each system has its advantages and disadvantages. For example, the main disadvantage of the ROLL and RSL techniques is the use of radioactivity. One particularity of the Magseed[®] marker, unlike the localization systems, is that it requires the use of plastic or titanium surgical instruments. The iron contained in metal instruments disrupts the Sentimag[®] probe. However, it is possible to use metal instruments just be sure to take away the instruments when the probe is in use so as not to interfere the signal. The large size of the SCOUT (12 mm) and the LOCalizer (11 mm) tags represents a disadvantage in comparison to the size of the Magseed[®] marker (5 mm) and RSL (4.5 mm) (26). In addition, these techniques are significantly more expensive than WGL leading to obstacles in broadening their use in practices. Davis et al. compare the cost device for wire localization and non-wire localization like I-125 radioactive seed, Radar reflector, Magnetic seed, radio frequency identification tag (26). A comparison of the price of each device clearly demonstrates the low cost of WGL. The average device cost for WGL is around 18 € which makes it an economical choice compared to other techniques except the carbon marking which has a similar cost. Regarding the ROLL technique a randomized study was performed an economic evaluation comparing ROLL with WGL (27). It appeared that ROLL is not more cost-effective than WGL (28). This study took into account all the medical costs associated with the technique as well as the changes in quality of life. Regarding the RSL technique, Zhang et al. demonstrated that RSL had lower costs than WGL for breast-conserving surgery. But the nuclear regulatory issues and management can increase the costs (29). Devices prices for the Magseed[®] system and the SCOUT technique are quite similar (around 300 €). The LOCalizer is a little more expensive. Moreover, it is important to take into account the price of the detector which also vary (15,000 €–25,000 €). Like all new techniques, these are always more expensive at the beginning. But the increase in the number of users will have a significant impact on the decrease in the price of devices. It would also seem that the price of devices is different from one country to another. It is also necessary to take into account the reimbursement of these devices by the insurance system of the country. Further studies should be performed to determine if these techniques would lead to more cost-effective care. The implementation of nonwire localization techniques allowed more efficient use of radiology resources. To evaluate these new techniques

allowing localization of impalpable breast lesions, a National UK group collects all data from units in the United Kingdom through the iBRA-NET study (30-32). This will allow large scale evaluation of clinical outcomes, ensuring device safety and efficacy, to define new guidelines for pre-operative detection of subclinical breast lesions.

Conclusions

In conclusion, this study reported our clinical experience on 100 consecutive patients with whom we used the Magseed[®] magnetic marker to localize non-palpable breast lesions. Compared to the wire-guided technique we have been using for many years, the Magseed[®] combined with the Sentimag[®] probe is a safe and effective method for identification of subclinical breast lesions and is effective when extended microcalcification clusters are present, or in targeting multiple tumors within the same breast. One of the major advantages of the Magseed[®] marker is that it can be placed in advance allowing disassociation of radiology and surgery schedules. Further research is required to validate efficacy of other long-term applications for patients who require neoadjuvant treatment.

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Footnote

Reporting Checklist: The authors have completed the STROBE reporting checklist. Available at https://gs.amegroups.com/article/view/10.21037/gs-22-552/rc

Data Sharing Statement: Available at https://gs.amegroups. com/article/view/10.21037/gs-22-552/dss

Peer Review File: Available at https://gs.amegroups.com/ article/view/10.21037/gs-22-552/prf

Conflicts of Interest: All authors have completed the ICMJE uniform disclosure form (available at https://gs.amegroups.com/article/view/10.21037/gs-22-552/coif). The authors have no conflicts of interest to declare.

Ethical Statement: The authors are accountable for all

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aspects of the work in ensuring that questions related to the accuracy or integrity of any part of the work are appropriately investigated and resolved. The study was approved by the CHC MontLegia Ethics Committee (reference No. 20/40/1052) and was conducted in accordance with the Declaration of Helsinki (as revised in 2013). Written informed consent was obtained from all participants included.

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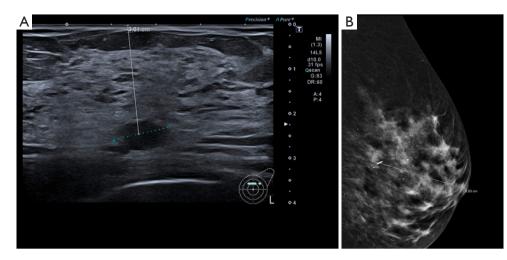


Figure S1 Ultrasound picture demonstrates the distance between the skin and the center of the tumor lesion was 36 mm (A). Lateral view shows a distance of 8 cm from the nipple (B). This measure was performed on mammograms images which is an error because mammograms give false information about the depth as this is performed with compression. L, left.