

# Breast cancer imaging: ductal carcinoma in-situ (DCIS) - Part I

**D**uctal carcinoma in-situ (DCIS) is a noninvasive malignancy and a potential precursor to invasive cancer. At pathologic analysis, DCIS shows proliferation of malignant epithelial cells that line the ducts (at the level of the terminal ductal-lobular unit) (Fig 1) without invasion through the basement membrane. The detection rate of DCIS has increased markedly over the past two decades with the advent of breast cancer screening. Early detection and assessment of extent of DCIS is important for planning successful conservative breast surgery. Half the cases of recurrent DCIS are associated with invasive ductal cancer. In addition, 20% of patients with DCIS develop metastases within 10 years of initial diagnosis.

In the following article, we will review the findings of DCIS on mammography and breast ultrasound (US) and also discuss the role of breast Magnetic Resonance Imaging (MRI) for improved detection of DCIS.

Over 90% of cases of DCIS are detected as microcalcifications on mammography. These calcifications are calcified cellular debris or secretions within the intraductal lumen. The uneven calcification of the cellular debris explains the fragmentation and irregular contours of the calcifications.

Figure 2 shows the distinction between linear and rounded microcalcifications that helps distinguish acinar (mostly benign) from ductal (suspicious for malignancy)-type calcifications. Calcifications are extremely variable in size, density and form; they may be amorphous (*morphus* means form in Greek, amorphus means no particular form), pleomorphic (*pleo* is Greek for more or many forms), heterogeneous (mixed density), rounded, coarse ( $\geq 5\text{mm}$ ) or fine ( $< 5\text{mm}$ ). Their distribution may be clustered, linear, or segmental

(Fig 3). The diagnostic approach to breast calcifications is to analyze the morphology, distribution and sometimes change over time.

Pleomorphic calcifications distributed in a linear fashion or in a cluster ( $>5$  calcifications in an area of 1cm diameter) should raise enough suspicion to advise biopsy (Fig 4). The presence of amorphous (Fig 5) or rounded calcifications in a linear or clustered distribution may also lead to biopsy, however the level of suspicion is lower in

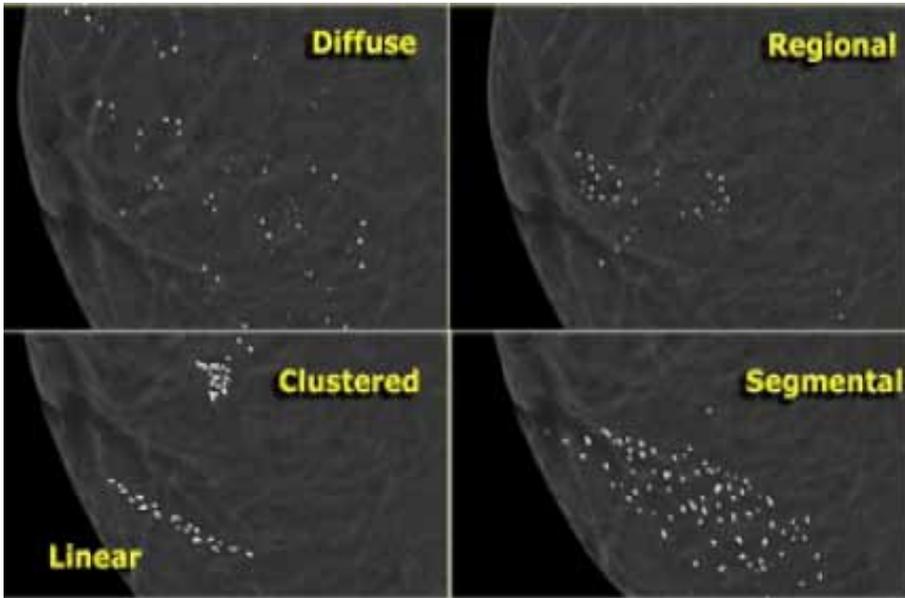
these instances. Dispersed or regional distribution of calcifications in one or both breasts or in multiple foci is usually indicative of benign disease (Fig 6). A segmental distribution of calcifications especially in the absence of pleomorphism is of indeterminate significance; in such situations or when calcifications are scanty in number, a close mammographic follow-up may be justified (Fig 7). Fine linear and branching calcifications, particularly when fragmented, require biopsy (Fig 8).  $\S$



**Figure 1.** Terminal lobular ductal unit (highlighted) includes the terminal duct with extralobular (\*\*) and intralobular (\*) portions and the acini.



**Figure 2.** Calcifications may be classified as acinar (round or crescentic) or ductal (elongated, linear distribution).



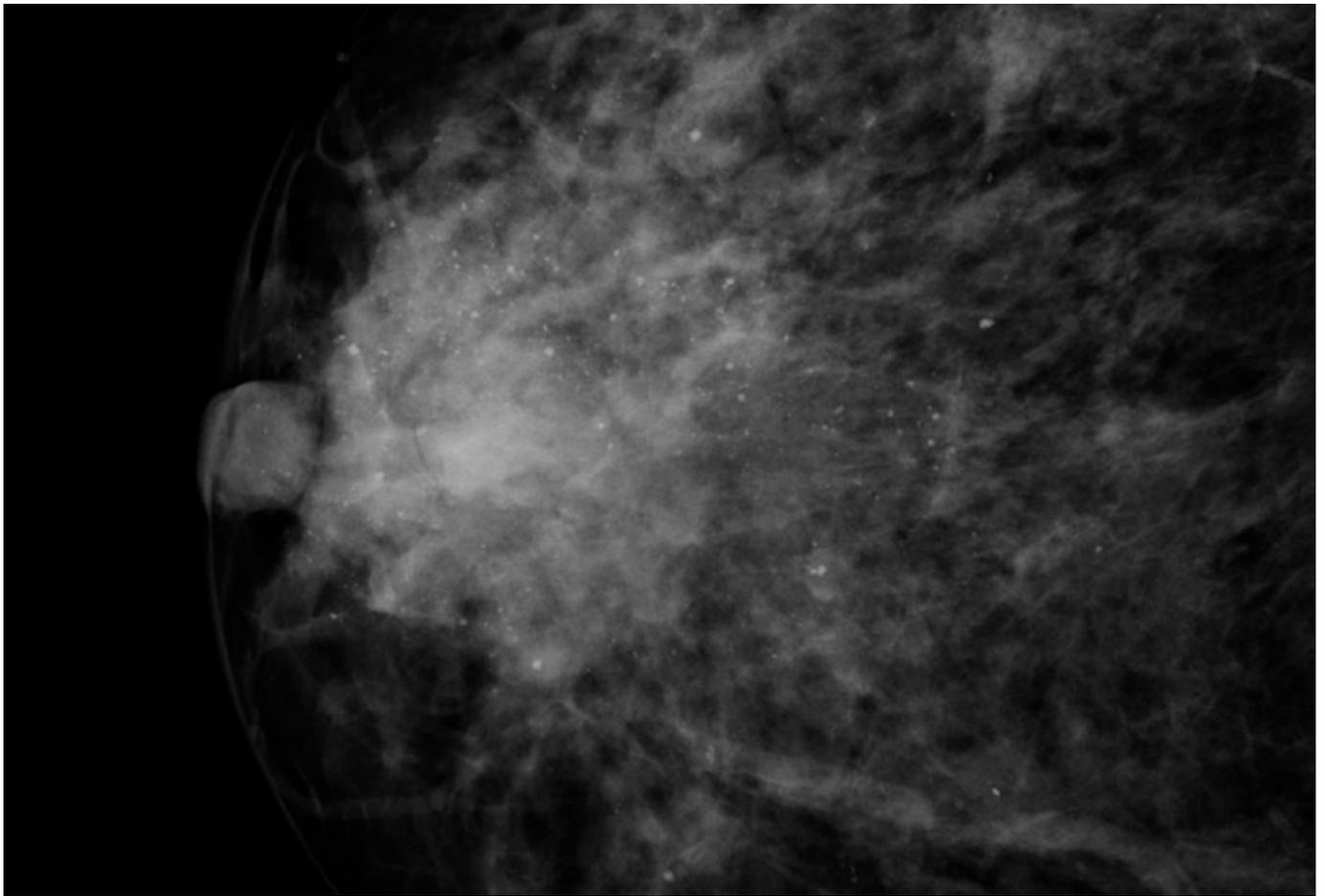
**Figure 3.** Distribution of calcifications may be diffuse, regional, segmental or clustered/linear.



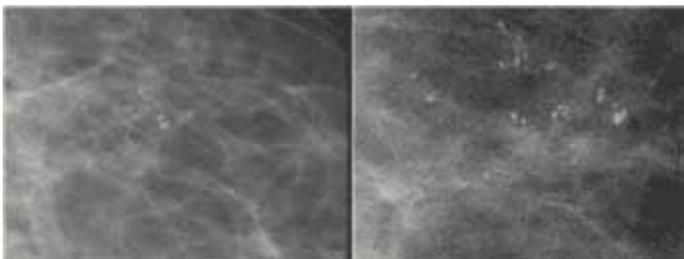
**Figure 4.** Mammogram showing pleomorphic calcifications in a linear distribution.



**Figure 5.** Mammogram showing amorphous calcifications in a clustered distribution.



**Figure 6.** Mammogram showing dispersed calcifications.



**Figure 7.** Mammographic follow-up of scanty calcifications; an increase in number of microcalcifications noted after 6 months should herald a biopsy.



**Figure 8.** Mammogram showing fine linear, branching and fragmented calcifications that are strongly indicative of malignant disease and required biopsy.