Corrigenda

WHO Classification of Tumours, 5th edition: Breast Tumours

July 2024 (after 3rd print run)

Updated corrigenda for this volume can be found at https://publications.iarc.who.int/Book-And-Report-Series/Who-larc-Classification-Of-Tumours/Breast-Tumours-2019.

Summary of corrections:

WHO classification tables (p. 10, 164, 188, 232, 250)

The following footnote has been added below the WHO classification (ICD-O coding) tables:

Subtype labels are indented.

Updated online: Yes

Updated in print: Yes (in 3rd print run), November 2020

Lobular carcinoma in situ (p. 73)

The text has been corrected as shown.

Original text	Corrected text
Staging According to the eighth editions of the Union for International Cancer Control (UICC) TNM classification {229} and the American Joint Committee on Cancer (AJCC) cancer staging manual {61}, LCIS is no longer staged as Tis. Pleomorphic LCIS is not included in the pTis classification.	Staging The eighth edition of the Union for International Cancer Control (UICC) TNM classification {229} recommends that LCIS be staged as pTis. However, the eighth edition of the American Joint Committee on Cancer (AJCC) cancer staging manual {61} considers LCIS to be a benign disease and therefore does not include it in staging.

References cited above:

61. Amin MB, Edge S, Greene F, et al., editors. AJCC cancer staging manual. 8th ed. New York (NY): Springer; 2017. **229.** Brierley JD, Gospodarowicz MK, Wittekind C, editors. TNM classification of malignant tumours. 8th ed. Oxford (UK): Wiley Blackwell; 2017.

Updated online: Yes

Invasive breast carcinoma: General overview (p. 92)

The word "internal" has been corrected to "international".

Original text	Corrected text
Histopathology Stromal response patterns and tumour microenvironment	Histopathology Stromal response patterns and tumour microenvironment
For quantifying TILs, it is recommended to follow the internal consensus scoring recommendations	For quantifying TILs, it is recommended to follow the international consensus scoring recommendations

Updated online: Yes

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Invasive breast carcinoma of no special type (p. 104)

A comma has been deleted to correct the meaning of the text, as shown.

Histopathology Special morphological patterns Histopathology Special morphological patterns	Original text	Corrected text
Oncocytic, lipid-rich, glycogen-rich, clear cell, and sebaceous carcinomas are rare tumours Oncocytic, lipid-rich, glycogen-rich clear cell, and sebaceous carcinomas are rare tumours	Special morphological patterns Oncocytic, lipid-rich, glycogen-rich, clear cell, and	Special morphological patterns Oncocytic, lipid-rich, glycogen-rich clear cell, and

Updated online: Yes

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Invasive lobular carcinoma (p. 117)

The term "subtype" has been corrected to "pattern" as shown.

Original text	Corrected text
Histopathology	Histopathology
Although the literature suggests ER was found to be expressed in the classic form and in subtypes, with the rate of positivity being highest (100%) in the alveolar subtype	Although the literature suggests ER was found to be expressed in the classic form and in patterns, with the rate of positivity being highest (100%) in the alveolar pattern

Updated online: Yes

Invasive lobular carcinoma (p. 117-8)

The term "subtype" has been corrected to "pattern" as shown.

Original text	Corrected text
Prognosis and prediction Despite the favourable prognostic features When the histological subtypes of ILC were analysed separately	Prognosis and prediction Despite the favourable prognostic features When the histological patterns of ILC were analysed separately
After neoadjuvant chemotherapy However, this relative resistance to cytotoxic therapy may be more related to the molecular characteristics (and especially the lower proliferation) of ILC rather than the histological subtype per se	After neoadjuvant chemotherapy However, this relative resistance to cytotoxic therapy may be more related to the molecular characteristics (and especially the lower proliferation) of ILC rather than the histological pattern per se

Updated online: Yes

Updated in print: Yes (in 3rd print run), November 2020

Invasive micropapillary carcinoma (p. 129)

The legend of Fig. 2.131C (Fig. #2496 online) has been amended as shown.

Original text	Corrected text
Fig. 2.131 Invasive micropapillary carcinoma C An example showing a moderate to intense but incomplete (U-shaped or basolateral) HER2 staining pattern (score: 2+). <i>ERBB2</i> (<i>HER2</i>) FISH showed <i>ERBB2</i> gene amplification (<i>ERBB2</i> /CEP17 ratio: 2.5).	Fig. 2.131 Invasive micropapillary carcinoma C An example showing a moderate to intense but incomplete (U-shaped or basolateral) HER2 staining pattern (score: 2+). ERBB2 (HER2) FISH showed ERBB2 gene amplification (ERBB2/CEP17 ratio: 2.5; mean number of ERBB2 signals/cell: > 6.0).

Updated online: Yes

Updated in print: Yes (in 3rd print run), November 2020

Adenoid cystic carcinoma (p. 145)

The word "local" has been corrected to "regional" as shown..

Original text	Corrected text
Prognosis and prediction	Prognosis and prediction
Classic AdCC: Despite the triple-negative phenotype, the classic subtype usually shows favourable behaviour. Local and distant metastases are rare	Classic AdCC: Despite the triple-negative phenotype, the classic subtype usually shows favourable behaviour. Regional and distant metastases are rare

Updated online: Yes

Phyllodes tumour (p. 174)

The high-power field area has been corrected from 0.5 mm² to 0.2 mm² in two places, and the field diameter has been added.

Original text	Corrected text
Histopathology	Histopathology
In benign phyllodes tumours The spindle cell stromal nuclei are monomorphic and mitoses are rare, usually < 2.5 mitoses/mm ² {2038} (corresponding to < 5 mitoses per 10 high-power fields of 0.5 mm ²).	In benign phyllodes tumours The spindle cell stromal nuclei are monomorphic and mitoses are rare, usually < 2.5 mitoses/mm² {2038} (corresponding to < 5 mitoses per 10 high-power fields of 0.5 mm in diameter and 0.2 mm² in area).
Malignant phyllodes tumours increased mitoses (≥ 5 mitoses/mm²; ≥ 10 mitoses per 10 high-power fields of 0.5 mm²)	Malignant phyllodes tumours increased mitoses (≥ 5 mitoses/mm²; ≥ 10 mitoses per 10 high-power fields of 0.5 mm in diameter and 0.2 mm² in area)

Updated online: Yes

Updated in print: Yes (in 2nd print run), January 2020

Tumours of the nipple: Introduction (p. 179)

In the second paragraph of the introduction, the unit "cm" has been corrected to "mm" as shown.

Original text	Corrected text
Tumours of the nipple: Introduction	Tumours of the nipple: Introduction
The skin lining of the nipple rests on a thin layer The squamous epithelium of the nipple lining extends into the major ducts for 1–2 cm before being abruptly replaced by the	The skin lining of the nipple rests on a thin layer The squamous epithelium of the nipple lining extends into the major ducts for 1–2 mm before being abruptly replaced by the
luminal/myoepithelial cell lining of the ducts	luminal/myoepithelial cell lining of the ducts

Updated online: Update pending

Updated in print: No (pending next print run)

Nodular fasciitis (p. 202)

The text has been amended as shown, in order to clarify the precise meaning.

Original text	Corrected text
Histopathology Nodular fasciitis Extravasated red blood cells and lymphocytes are commonly seen	Histopathology Nodular fasciitis Lymphocytes and extravasated red blood cells are commonly seen

Updated online: Yes

WHO classification of tumours of the male breast (p. 250)

The ICD-O code for Paget disease of the nipple has been added.

Original text	Corrected text
Epithelial tumours	Epithelial tumours
Gynaecomastia	Gynaecomastia
Florid gynaecomastia	Florid gynaecomastia
Fibrous gynaecomastia	Fibrous gynaecomastia
8500/2 Intraductal carcinoma, non-infiltrating, NOS	8500/2 Intraductal carcinoma, non-infiltrating, NOS
Ductal carcinoma in situ	Ductal carcinoma in situ
Lobular carcinoma in situ	Lobular carcinoma in situ
Paget disease of the nipple	8540/3 Paget disease of the nipple
8500/3 Infiltrating duct carcinoma NOS	8500/3 Infiltrating duct carcinoma NOS

Updated online: Yes

Updated in print: Yes (in 3rd print run), November 2020

Gynaecomastia (p. 252)

In Table 7.01 (Table #5225 online), in the row for **Spinobulbar muscular atrophy**, under the **Mechanism of production of gynaecomastia** column heading, the nucleic acid sequence "GAG" has been corrected to "CAG" as shown.

Original text Corr	rrected text
Similar to in Klinefelter syndrome – an increased Simi	echanism of production of gynaecomastia milar to in Klinefelter syndrome – an increased mber of CAG repeats in the AR gene

Updated online: Yes

Updated in print: Yes (in 3rd print run), November 2020

BRCA1/2-associated hereditary breast and ovarian cancer syndrome (p. 272)

A typo in the text has been corrected as shown.

Original text	Corrected text
Histopathology	Histopathology
Tumours arising in BRCA1 and BRCA2 mutation	Tumours arising in BRCA1 and BRCA2 mutation
carries are more likely than sporadic breast cancers	carriers are more likely than sporadic breast
to have certain histological and molecular	cancers to have certain histological and molecular
characteristics	characteristics

Updated online: Update pending

Updated in print: No (pending next print run)

BRCA1/2-associated hereditary breast and ovarian cancer syndrome (p. 273)

In Table 9.03 (Table #2466 online), under the **Typical findings in mutation carriers: BRCA2** column heading, in the row for **Histological types**, the term "ductal" has been corrected to "IBC-NST", and "NST" has been added to the abbreviations footnote, as shown.

Original text	Corrected text
Histological types Ductal, tubular, cribriform, mucinous, classic/pleomorphic lobular	Histological types IBC-NST, tubular, cribriform, mucinous, classic/pleomorphic lobular
DCIS, ductal carcinoma in situ; IBC, invasive breast carcinoma; LCIS, lobular carcinoma in situ.	DCIS, ductal carcinoma in situ; IBC, invasive breast carcinoma; LCIS, lobular carcinoma in situ; NST, of no special type.

Updated online: Yes

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Li-Fraumeni syndrome, CHEK2-associated (p. 282)

Please note the following:

Recent studies have demonstrated that *CHEK2* mutation does not in fact predispose individuals to Li– Fraumeni syndrome, as mentioned in the *Genetic tumour syndromes* volume of this series, in the section titled "*CHEK2*-related hereditary (breast) cancer predisposition syndrome (*CHEK2*)" [[WHO Classification of Tumours Editorial Board. Genetic tumour syndromes [Internet; beta version ahead of print]. Lyon (France): International Agency for Research on Cancer; 2023. (WHO classification of tumours series, 5th ed.; vol. 14). https://tumourclassification.iarc.who.int/chapters/67.]].

Updated online: No (pending 6th edition)
Updated in print: No (pending 6th edition)