

Time to revise classification of phyllodes tumors of breast? Results of a French multicentric study

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Abstract:

Objective. To assess prognostic factors of recurrence of phyllodes tumors (PT) of the breast.

Methods. We performed a retrospective, multicentric cohort study, including all patients who underwent breast surgery for grade 1 (benign), 2 (borderline) or 3 (malignant) PT between 2000 and 2016 in five tertiary University hospitals, diagnosed according to World Health Organization classification.

Results. 230 patients were included: 144 (63%), 60 (26%) and 26 (11%) with grade 1, 2 and 3 PT, respectively. Recurrence occurred in 10 (7%), 7 (12%) and 5 (19%) patients with grade 1, 2 and 3 PT, respectively. In univariate analysis, moderate to severe nuclear stromal pleomorphism (HR 8.00 [95% CI: 1.65 - 38.73], p<0.009) was correlated with recurrence in all groups including grade 1 (HR 14.3 [95% CI: 1.29 - 160], p=0.031). In multivariate analysis, surgical margin >5 mm, (HR 0.20 [95% CI: 0.06 - 0.63], p = 0.013) were significantly correlated with less recurrence in all PT grades. For grade 1 PT, there was also significantly less recurrence with surgical margin >5mm, (HR 0.09 [95% CI: 0.01 - 0.85], p=0.047) in multivariate analysis.

Conclusion. The surgical margin should be at least 5 mm whatever the grade of PT. Moderate to severe nuclear stromal pleomorphism identified a subgroup of grade 1 PT with a higher rate of recurrence. This suggests that the WHO classification could be revised with the introduction of nuclear stromal pleomorphism to tailor PT management.

Key words: phyllodes breast tumor; benign; borderline; malignant; prognostic factors; surgery.

Introduction

Phyllodes tumors (PT) of the breast are fibroepithelial tumors whose etiopathogenesis remains unclear. These are rare tumors accounting for around 0.3-1% of all primary breast tumors (1,2). They mostly affect women in their fourth decade of life. The classic clinical presentation is a supple, painless mass, well limited, with rapid growth without associated axillary adenopathy. In 1982, the World Health Organisation (WHO) classification distinguished three types of PT according to five factors: stromal cellularity, stromal atypia, stromal overgrowth, mitotic count, and character of the tumor borders (3) (4) (5) (6) (7). The stromal overgrowth was defined as definition: absence of epithelial elements in one low-power microscopic field containing only stroma (according WHO 2012 classification)

The three types of PT are: grade 1 or benign tumors, grade 2 or borderline tumors, and grade 3 or malignant tumors representing 64%, 18% and 18% of all PT, respectively (8).

The average risk for local recurrence and metastasis have been reported as being 15% and 0.1% for grade 1 PT, 17% and 0.2% for grade 2 PT, 28% and 22% for grade 3 PT (2). The recurrent tumor can be of a more aggressive histological form justifying surgical excision of all PT (9,10). Some prognostic factors have been identified (6,8,11,12) including: age, tumor size, grade, mitotic index, degree of cellular atypia, stromal necrosis and stromal cellularity and histologically negative margins for grade 1 PT, and margins greater than 10 mm for grades 2 and 3 PT for therapeutic factors. However, these prognostic factors are mainly based on old data. The largest study by Belkacemi et al (8) including 443 women with PT, dates from 1971 to 2003 prior to the WHO classification distinguishing 3 grades of PT. Moreover, PT management is based on low levels of evidence. Margin width was determined on a consensus from two studies by Spitaleri et al (2) and Kim et al (13) published in 2013. Thus, there is a need for a better determination of prognostic factors for optimal PT management.

Therefore, the main objective of the present study was to assess prognostic factors of PT recurrence based on clinical, pathological and surgical characteristics to tailor management of PT according to histological grade.

Material and methods

Patients

This was a retrospective, multicenter cohort study. Data of women with histologically proven PT operated on between January 2000 and December 2016 were abstracted from five institutions in France with prospective maintained breast databases (Tours, Tenon, Angers, Rennes and Pitié Salpêtrière University Hospitals). The research protocol was approved by the Institutional Review Board (N° 2088222).

The inclusion criteria were grade 1, 2 or 3 PT histologically confirmed from a surgical specimen. The exclusion criteria were: breast adenofibroma, breast cancer, other fibroepithelial tumors than PT.

Clinical, surgical and pathological data were collected from the patients' medical charts. The histopathological data collected included: PT grade, tumor size (mm), stromal mitotic activity defined by the number of mitoses in high-magnification fields (less than 4, between 4 and 9 or higher or equal to 10), stromal cell atypia (absent; moderate i.e., difficile to spot; or severe i.e., of high nuclear grade), stromal cellularity (mild, moderate, severe), stromal overgrowth (mild, moderate, severe), stromal nuclear pleomorphism (classified as mild: small nuclei, regular chromatin, few nucleoli; moderate: larger nuclei, visible but small nucleoli; or severe: significant variation of nuclei in size and shape, prominent nucleoli), stromal necrosis, and surgical margins.

Histological evaluation

All women were classified according to the 2012 WHO classification on final pathology serving as the standard. A PT was defined as a fibroepithelial tumor with the following four criteria: high stromal cellularity (50% higher than that observed in adenofibromas), predominant stroma in the epithelial component, leaf-like projections into variably dilated elongated lumina and infiltrative margin (tumor border). The benign,

borderline and malignant grades (i.e 1,2 and 3 grades) were defined in accordance with the 2012 WHO recommendations (4) using these criteria: infiltrating periphery, stromal cellularity (categorized as mild / moderate / severe assessed in the most cellular area, mild is defined as increase in at least 50% of the stroma in PT compared with a typical adenofibroma, moderate is defined as the presence of stromal nuclear crowding or overlapping, severe is defined as marked stromal cellularity), mitotic activity (<5, 5-9, \geq 10 per 10 high power fields), and cellular atypia. The surgical specimens were analyzed by an expert breast pathologist.

Treatment, follow-up and endpoints

All women had undergone primary surgical treatment. Clinical follow-up consisted of physical examinations and the use of imaging techniques according to the findings. Recurrent disease was assessed by physical examination, histological findings, clinical follow-up and imaging. The diagnosis of recurrence was based on histological sampling. The date of the last recorded information corresponded to either the date of the last visit in the follow-up center, or the date of recurrence or death.

The primary endpoint was the ipsilateral recurrence of a PT during the follow-up period. The secondary endpoint was the PT grade.

Recurrence free survival (RFS) was defined as the time from the date of primary surgery to any PT recurrence and was censored at date of the last follow-up or date of death without recurrence. Overall survival (OS) was defined as time from primary surgery to death as a result of any cause.

Statistical analysis

For the descriptive analysis of population characteristics, a χ^2 or Fisher test was used for the qualitative parameters, and a Kruskal-Wallis test for the quantitative parameters. Variables were considered statistically significant when p < 0.05. A Receiver Operating Characteristic

(ROC) curve was used to define a surgical margin threshold. A univariate analysis of clinical, therapeutic and pathological recurrence risk factors on RFS by the log rank test was then performed. These results are presented by Hazard Ratio (HR) with 95% confidence interval (CI). Variables with a p < 0.20 were included in a multivariate model, followed by a step-by-step descending selection. Survival curves were performed using the Kaplan-Meier model.

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Results

Characteristics of the population and preoperative data

Two-hundred thirty patients were included: 144 (63%) had grade 1 PT, 60 (26%) grade 2 PT, and 26 (11%) grade 3 PT. The characteristics of the patients are shown in Table 1.

Surgery and post-operative data

The initial surgery for all PT grades consisted of enucleation (or local excision) for 38 patients (16.5%), wide excision for 172 patients (74.8%) and a total mastectomy for 20 (8.7%). The surgical and pathological data are shown in Table 2. Adjuvant treatment consisted of radiotherapy for 14 patients (6.1%), including one patient with grade 2 PT (1.7%), and 13 patients with grade 3 (50%). Chemotherapy was performed in three of the patients (11.5%) with grade 3 PT.

Follow-up

The mean follow-up was 42.5 months (\pm 46.8 months). During the follow-up period, 22 patients (9.6%) experienced PT recurrence with 17 of the recurrences (77%) occurring within the first 3 years of follow-up. Overall, the mean recurrence time was 28.7 months (\pm 21.6 months): 37 months (\pm 27.6 months) for grade 1 PT; 21.1 months (\pm 12.6 months) for grade 2 PT; and 22.6 months (\pm 13.7 months) for grade 3 PT (Not Significant (NS)). Patterns of recurrences and treatments performed are described in Figure 1.

The recurrence rate was 7% (n = 10) for grade 1 PT, 11.7% (n = 7) for grade 2 PT, and 19.2% (n = 5) for grade 3 PT (NS). There were no deaths among the patients with grade 1 PT. The death rate was 1.7% (n = 1) for grade 2 PT and 15.4% (n = 4) for grade 3 PT (p < 0.001).

Overall, the 3-year RFS rate was 92.6%: 95.8%, 90.0% and 80.8% for grade 1, 2 and 3 PT, respectively (p = 0.018). The 10-year RFS rates were 93.1%, 88.3% and 80.8% for grade 1, 2 and 3 PT, respectively (NS).

The 3-year OS rate was 100%, 96.2% and 96.2% for grade 1, 2 and 3 PT, respectively (p = 0.072). The 10-year OS rate was 100%, 98.3% and 84.6% for grade 1, 2 and 3 PT, respectively (p < 0.001).

Prognostic factors

ROC curves were used to assess optimal surgical margins to prevent or decrease recurrence rate (Figure S1 – supplementary data). For grade 1 PT, a surgical margin value of 1 mm had a sensitivity, specificity, positive predictive value (PPV) and negative predictive value (NPV) of 41.0%, 90.4%, 22.2% and 91.8%, respectively; a surgical margin value of 5 mm had a sensitivity, specificity, PPV and NPV of 90.0%, 26.8%, 8.8% and 97.1%, respectively; and a surgical margin value of 10 mm had a sensitivity, specificity, PPV and NPV of 90.0%, 26.8%, 8.8% and 97.1%, respectively; and a surgical margin value of 10 mm had a sensitivity, specificity, PPV and NPV of 90%, 24.4%, 8.6% and 96.9%, respectively. For grade 2 PT, a surgical margin value of 1 mm had a sensitivity, specificity, PPV and NPV of 28.6%, 92.3%, 33.3% and 90.6%, respectively; a surgical margin value of 5 mm had a sensitivity, specificity, PPV and NPV of 42.9%, 53.8%, 11.1% and 87.5%, respectively; and a surgical margin value of 10 mm had a sensitivity, specificity, PPV and NPV of 42.8%, 47.2%, 9.7% and 13.8%, respectively.

For grade 3 PT, we were unable to demonstrate margins with a meaningful sensitivity, specificity, PPV or NPV.

The prognostic factors of recurrence for all grades of PT are described in Table 3. According to univariate survival analysis, age > 40 years (HR 0.38 [95% CI: 0.16-0.89], p = 0.026), a surgical margin of more than 1 mm (HR 0.29 [95% CI: 0.10 - 0.85], p = 0.044), and a margin of more than 5mm (HR 0.31 [95% CI: 0.11 - 0.90], p = 0.044) were significantly correlated with less recurrence. As a continuous variable, surgical margin size was also significantly correlated with less recurrence (HR 0.22, [95% CI: 0.05 - 0.91], p = 0.036). A margin of more than 10 mm (HR 0.85 [95% CI: 0.35 – 2.02], p = 0.706) was not significantly correlated with less recurrence.

The nuclear pleomorphism of the stroma was known for 97 patients and showed that moderate to severe stromal nuclear pleomorphisms (HR 8.00 [95% CI: 1.65 - 38.73], p = 0.009) were correlated with more recurrence.

According to multivariate analysis, a surgical margin of more than 1 mm (HR 0.26 [95% CI: 0.09 - 0.76], p = 0.013), and of more than 5 mm (HR 0.20 [95% CI: 0.06 - 0.63], p = 0.013) were significantly correlated with less recurrence for all grades of PT.

The prognostic factors for recurrence of grade 1 PT are described in Table 4. According to univariate survival analysis, age > 40 years (HR 0.24 [95% CI: 0.06 - 0.92], p = 0.038) and a surgical margin of more than 5 mm (HR 0.09 [95% CI: 0.01 - 0.85], p = 0.047) were significantly correlated with less recurrence. Moderate to severe stromal nuclear pleomorphisms (HR 14.3 [IC 95%: 1.29-160], p=0.031) were significantly correlated with recurrence. A margin of more than 10 mm (HR 0.25 [95% CI: 0.03 - 1.95], p = 0.184) was not significantly correlated with less recurrence. Stromal nuclear pleomorphism was known for 56 of the patients with grade 1 PT: recurrence rates were 2/49 (4%) for patients with low stromal nuclear pleomorphism and 2/7 (28%) for those with moderate to severe stromal nuclear pleomorphism.

In multivariate analysis, only a margin over 5 mm was significantly correlated with less recurrence (HR 0.09 [CI 95%: 0.01 - 0.85], p = 0.047) for patients with grade 1 PT. In univariate survival analysis for grade 2 PT, only age > 40 years was significantly correlated with less recurrence (HR 0.12 [95% CI: 0.03 - 0.53], p = 0.005). No margin width threshold was significantly correlated with recurrence for grade 2 PT, probably due to lack of power. No factors were found to be significantly correlated with recurrence for grade 3 PT in univariate analysis, probably due to lack of power.

Discussion

The overall PT recurrence rate was 9.6%. Surgical margin size was found to be an independent prognostic factor of recurrence for all grades of PT, including grade 1. Furthermore, stromal nuclear pleomorphism was significantly correlated with recurrence for grade 1 PT in univariate analysis, identifying a subgroup of grade 1 PT with a similar or higher recurrence rate to grade 2 PT (28% *vs* 12%, respectively).

Although we did not find a significant correlation of stromal cellularity with recurrence (as others have shown (8,13,14)), in accordance with Sawalhi (2013) we found a significant correlation between stromal nuclear pleomorphism and recurrence (15). In theory, stromal nuclear pleomorphism is defined as mild (small nuclei, regular chromatin, few nucleoli); moderate (larger nuclei, visible but small nucleoli), or severe (significant variation of nuclei in size and shape, prominent nucleoli). However, in practice, these definitions appear to vary, and no inter- or intraobserver variability data are available, which could weaken reproducibility of diagnosis. In accordance with others (2,8,13,14) we found that margin status is an independent prognostic factor: the wider the margin the lower the recurrence rate (in a significant manner). Kim et al (13) and others (9,10) advocate only *in sano* surgery for grade 1 PT, in contrast to the present study which showed that a margin size of more than 5 mm is required for grade 1 PT (with an NPV of 97.1% for recurrence). This would suggest that optimal surgical treatment for all grades of PT requires local surgical excision with a more severe grade, as shown in the present study and others (15,19).

Some limitations of the present study deserve to be mentioned. First of all, there were some missing data (such as tumor border) and a risk of underestimating recurrence rates because of the retrospective nature of the study and the duration of the inclusion period. This, coupled with a low rate of recurrence, hampered the possibility of building a scoring system to predict

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recurrences. Another limitation is the relatively short mean follow-up of 42.5 months (\pm 46.8 months), *versus* 85 months in the Spitaleri study (2) and 106 months in the Belkacemi study (8). Thus, the use of ROC curves to find margin size threshold according recurrence could be hampered. However, local recurrences are mainly observed during the first two years of follow-up (29, 30) and even earlier for grade 3 PT (31–32).

Conclusion

This large retrospective study analyzing data from prospectively managed databases from five major university hospitals in France, suggests that the recurrence rate of PT is around 10% and that nearly one third of these recurrences occur in patients with a higher grade PT conveying a poorer prognosis. Surgical margin size remains the main prognostic factor, including in patients with grade 1 PT, with a required surgical margin of over 5 mm. Moderate to severe stromal nuclear pleomorphism identified a group of grade 1 PT patients with a higher risk of recurrence (28%).

The authors declare that they have no conflict of interest

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Table 1. Preoperative characteristics of patients

	All PT	РТ	РТ	РТ		
Data	n = 230	Grade 1 n = 144	Grade 2 n = 60	Grade 3 n = 26	p value	
Age (mean +/- SD)	47.2 ± 14.1	44.3 ± 12.5	53.2 ± 16.0	49.6 ± 13.7	< 0.001	
(range)	(16-95)	(16, 83)	(17, 95)	(26, 80)		
Parity (mean +/- SD)	1.7 ± 1.5	1.6 ± 1.4	1.7 ± 1.5	2.3 ± 2.0	0.240	
≤ 1	102 (44.3%)	67 (46.5%)	27 (45.0%)	8 (30.8%)	0.328	
≥ 2	128 (55.7%)	77 (53.5%)	33 (55.0%)	18 (69.2%)		
Contraception	65 (45.1%)	51 (48.6%)	11 (44.0%)	3 (21.4%)	0.158	
Menopause	85 (37.0%)	39 (27.1%)	34 (56.7%)	12 (46.2%)	< 0.001	
Hormonal Replacement Therapy	21 (9.1%)	12 (8.3%)	8 (13.3%)	1 (3.8%)	0.331	
Smoking	27 (13.2%)	21 (16.0%)	3 (5.8%)	3 (13.6%)	0.198	
Breast adenofibroma history	70 (30.4%)	49 (34.0%)	13 (21.7%)	8 (30.8%)	0.217	
PT history	2 (0.9%)	1 (0.7%)	0 (0.0%)	1 (3.8%)	0.295	
Breast cancer history	6 (2.6%)	1 (0.7%)	4 (6.7%)	1 (3.8%)	0.044	
PT family history	1 (0.4%)	1 (0.7%)	0 (0.0%)	0 (0.0%)	1	
Pregnancy (at diagnosis)	2 (0.9%)	1 (0.7%)	1 (1.7%)	0 (0.0%)	0.609	
Single localisation unifocal PT	198 (86.1%)	118 (81.9%)	56 (93.3%)	24 (92.3%)	0.077	
Duration of symptoms					0.017	
< 2 months	55 (33.1%)	30 (28.3%)	16 (34.8%)	9 (64.3%)		
2 - 12 months	56 (33.7%)	38 (35.8%)	13 (28.3%)	5 (35.7%)		
> 12 months	55 (33.1%)	38 (35.8%)	17 (37.0%)	0 (0.0%)		
Pain	42 (18.3%)	30 (20.8%)	7 (11.7%)	5 (19.2%)		
Skin changes	10 (4.3%)	3 (2.1%)	3 (5.0%)	4 (15.4%)	0.015	
Nipple retraction	1 (0.4%)	0 (0.0%)	0 (0.0%)	1 (3.8%)	0.113	
Clinical tumor size (mm, mean +/- SD)	43.9 ± 60.7	32.6 ± 20.8	55.5 ± 95.2	80.7 ± 88.4	< 0.001	
(range)	(6, 700)	(10, 150)	(6, 700)	(20, 400)		
Clinical tumor size					< 0.001	
< 30 mm	101 (45.9%)	82 (59.4%)	17 (28.8%)	2 (8.7%)		
\geq 30 mm	119 (54.1%)	56 (40.6%)	42 (71.2%)	21 (91.3%)		
Ultrasound tumor size (mean +/- SD)	38.8 +/- 43.1	32.8 +/- 21.4	35.6 +/- 29.8	86.1 +/- 107.8	0.007	
(range)	(6, 400)	(8, 130)	(6, 170)	(11, 400)		
Ultrasound tumor size					0.003	
< 20 mm	50 (25.8%)	31 (26.1%)	18 (32.1%)	1 (5.3%)		
20 - 50 mm	105 (54.1%)	71 (59.7%)	26 (46.4%)	8 (42.1%)		
\geq 50 mm	39 (20.1%)	17 (14.3%)	12 (21.4%)	10 (52.6%)		

Abbreviations: SD standard deviation; PT: phyllode tumor

Table 2. Surgical and pathological characteristics of patients

Data	All PT n = 230	PT Grade 1 n = 144	PT Grade 2 n = 60	PT Grade 3 n = 26	р
Initial Surgery					
Enucleation (or local excision)	38 (16.5%)	30 (20.8%)	7 (11.7%)	1 (3.8%)	< 0.001
Wide excision	172 (74.8%)	111 (77.1%)	48 (80.0%)	13 (50.0%)	
Total mastectomy	20 (8.7%)	3 (2.1%)	5 (8.3%)	12 (46.2%)	
Histological size (mm, mean +/- SD)	38.9 ± 29.0	30.9 ± 21.5	47.2 ± 29.2	65.7 ± 43.7	< 0.001
(range)	(5, 210)	(5, 170)	(6, 130)	(15, 210)	
No. of mitoses					
< 4	143 (68.8%)	120 (92%)	20 (35.7%)	3 (13.6%)	< 0.001
4 à 9	31 (14.9%)	10 (8%)	19 (33.9%)	2 (9.1%)	
≥10	34 (16.3%)	0 (0.0%)	17 (30.4%)	17 (77.3%)	
Stromal cellular atypia					
Mild	141 (75.4%)	112 (95.7%)	27 (55.1%)	2 (9.5%)	< 0.001
Moderate	27 (14.4%)	4 (3.4%)	18 (36.7%)	5 (23.8%)	
Severe	19 (10.2%)	1 (0.9%)	4 (8.2%)	14 (66.7%)	
Stromal cellularity					
Mild	45 (22.7%)	38 (30.2%)	6 (12.0%)	1 (4.5%)	< 0.001
Moderate	76 (38.4%)	55 (43.7%)	19 (38.0%)	2 (9.1%)	
Severe	77 (38.9%)	33 (26.2%)	25 (50.0%)	19 (86.4%)	
Stromal overgrowth					
Mild	24 (29.3%)	19 (32.8%)	5 (27.8%)	0 (0.0%)	0.002
Moderate	44 (53.7%)	34 (58.6%)	9 (50.0%)	1 (16.7%)	
Severe	14 (17.1%)	5 (8.6%)	4 (22.2%)	5 (83.3%)	
Stromal nuclear pleomorphism					
Mild	68 (70.1%)	49 (87.5%)	16 (57.1%)	3 (23.1%)	< 0.001
Moderate	16 (16.5%)	6 (10.7%)	9 (32.1%)	1 (7.7%)	
Severe	13 (13.4%)	1 (1.8%)	3 (10.7%)	9 (69.2%)	
Tumor necrosis					
Yes	11 (5.1%)	1 (0.7%)	2 (3.5%)	8 (38.1%)	< 0.001
No	206 (94.9%)	138 (99.3%)	55 (96.5%)	13 (61.9%)	
Margins after first surgery (mm, mean+/- SD)	3.7 ± 5.4	3.7 ± 4.9	3.7 ± 5.3	3.7 ± 8.3	0.198
(range)	(0, 40)	(0, 30)	(0, 20)	(0, 40)	
Second surgery	58 (25.2%)	19 (13.2%)	25 (41.7%)	14 (53.8%)	< 0.001
Mastectomy	18 (31.0%)	1 (5.3%)	6 (24.0%)	11 (78.6%)	< 0.001
Second lumpectomy for marge status	40 (69.0%)	18 (94.7%)	19 (76.0%)	3 (21.4%)	
Residual disease after second surgery	7 (12.1%)	1 (5.3%)	4 (16.0%)	2 (14.3%)	0.589
Margins after second surgery (mm, mean +/- SD)	6.0 ± 7.4	4.4 ± 5.3	9.1 ± 9.6	7.5 ± 9.0	0.002
(range)	(0, 50)	(0, 30)	(0, 50)	(0, 40)	

Margins after second surgery					
< 1 mm	25 (11.4%)	18 (13.1%)	6 (10.2%)	1 (4.2%)	0.002
[1-5] mm	116 (52.7%)	84 (61.3%)	21 (35.6%)	11 (45.8%)	
> 5 mm	79 (35.9%)	35 (25.5%)	32 (54.2%)	12 (50.0%)	
Final performed surgery (including second or third surgery)					
Breast conservative surgery	195 (84.8%)	140 (97.2%)	51 (85.0%)	4 (15.4%)	< 0.001
Total mastectomy	35 (15.2%)	4 (2.8%)	9 (15.0%)	22 (84.6%)	

Abbreviations: SD standard deviation: PT: Phyllode Tumor

Table 3. Univariate analysis of prognostic factors of recurrence-free survival for all phyllodes tumors

Data	N*	HR [95% CI]	р
Age ≥ 40 years	230	0.38 [0.16 - 0.89]	0.000
Mananausa	220	0.46 [0.19 1.19]	0.026
A dan of iknome history	230	1.00 [0.14 - 1.18]	0.100
	230	1.09 [0.44 - 2.67]	0.855
Single localisation	230	0.95 [0.28 - 3.21]	0.933
Duration of symptoms ≥ 2 months	166	1.29 [0.40 - 4.20]	0.671
Pain	230	0.99 [0.29 - 3.35]	0.984
kin changes	230	0.69 [0.09 - 5.12]	0.716
Jrade	230	1	0.421
1		1	
2		1.22 [0.46 - 3.20] 2 07 [0 70 - 6 15]	
Clinical tumor size (mm)	220	1.09 [0.97 - 1.21]	0.143
Ultrasound tumor size (mm)	194	1 90 [0 55 - 6 62]	0,600
No. of mitoses	208	1.50 [0.55 - 0.02]	0.427
< 4		1	
4 à 9		1.28 [0.35 - 4.68]	
≥ 10		2.07 [0.69 - 6.16]	
Stromal cellular atypia	187		0.591
Mild	107	1	0.071
Moderate		0.88 [0.10 3.05]	
Source		0.88 [0.19 - 3.93]	
	109	1.70 [0.55 - 5.57]	0.590
tromai cellular	198		0.589
Mild		1	
Moderate		0.76 [0.18 - 3.19]	
Severe		1.33 [0.36 - 4.84]	
tromal overgrowth	82		0.804
Mild		1	
Moderate		1.51 [0.16 - 14.60]	
Severe		2.58 [0.15 - 42.95]	
tromal nuclear pleomorphism	97		0.022
Mild		1	
Moderate		5.55 [0.92 - 33.35]	
Severe		12.89 [2.09 - 79.61]	
Stromal nuclear pleomorphism mild vs.	07	8 00 [1 65 29 72]	0.000
noderate to severe	71	0.00 [1.03 - 30.73]	0.009
Fumor necrosis	217	2.22 [0.49 - 9.94]	0.298
Surgery: Breast conserving surgery vs.	230	0.73 [0.27 - 2.01]	0.544
otal mastectomy	220	0.22 [0.05 0.01]	0 034
Surgical margin \$10mm after second	220	0.22 [0.03 - 0.91]	0.030
urgery (mm)	220	0.85 [0.35 - 2.02]	0.706
urgical margin size after second	220		
urgery (mm)	220		0.044
< 1mm [1_5] mm		l 0.29[0.10_0.85]	
ر ۲-۵٫ mm		0.29 [0.10 - 0.83] 0 31 [0 11 - 0 90]	
× 5 mm		0.31 [0.11 - 0.90]	

HR: Hazard Ratio; * could be less than 230 patients because of missing

Table 4. Univariate analysis of prognostic factors of recurrence-free-survival for grade 1phyllodes tumors

N*	HR [95% CI]	р
144	0.24 [0.06 - 0.92]	0.038
144	0.22 [0.03 - 1.71]	0.147
144	2.42 [0.70 - 8.42]	0.165
144	0.95 [0.20 - 4.47]	0.945
106	2.76 [0.32 - 23.68]	0.355
144	0.77 [0.10 - 6.09]	0.803
144	0.00 [0.00]	0.994
144	1.67 [0.42 - 6.70]	0.468
119	1.26 [0.15 - 10.30]	0.977
130		0.456
	1	
	0.00 [0.00]	
	. []	
117		0.995
	1	
	0.00 [0.00]	
	. []	
126		0.636
	1	
	0.49 [0.10 - 2.43]	
	0.51 [0.08 - 3.11]	
56		0.075
	1	
	9.95 [0.62 - 159.6]	
	29.63 [1.33 - 662.2]	
56	14.37 [1.29 - 160.1]	0.031
144	129 [0.0]	0.994
138	0.14 [0.01 - 1.47]	0.102
137	0.25 [0.03 - 1.95]	0.184
137		0.048
	1	
	0.28 [0.07 - 1.03]	
	0.09 [0.01 - 0.85]	
	N* 144 144 144 144 144 144 144 144 144 144 144 144 144 145 146 147 130 117 126 56 56 56 144 138 137	N* HR [95% CI] 144 $0.24 [0.06 - 0.92]$ 144 $0.22 [0.03 - 1.71]$ 144 $0.22 [0.03 - 1.71]$ 144 $0.42 [0.70 - 8.42]$ 144 $0.95 [0.20 - 4.47]$ 106 $2.76 [0.32 - 23.68]$ 144 $0.07 [0.10 - 6.09]$ 144 $0.00 [0.00]$ 144 $0.00 [0.00]$ 144 $1.67 [0.42 - 6.70]$ 119 $1.26 [0.15 - 10.30]$ 130 1 $0.00 [0.00]$ $.[]$ 117 1 $0.00 [0.00]$ $.[]$ 117 1 $0.00 [0.00]$ $.[]$ 117 1 $0.00 [0.00]$ $.[]$ 126 1 $0.49 [0.10 - 2.43]$ $0.51 [0.08 - 3.11]$ 56 1 $9.95 [0.62 - 159.6]$ $29.63 [1.33 - 662.2]$ 56 $14.37 [1.29 - 160.1]$ 144 $129 [0.0]$ 138 $0.14 [0.01 - 1.47]$ <

HR: Hazard Ratio; * could be less than 144 patients because of missing data



