Coprinopsis pannucioides (J.E. Lange) Örstadius & E. Larss. 2008

in Myc. Research 112 : 1180

Daniel Deschuyteneer

Synonymes :

Drosophila pannucioides (J.E. Lange) Kühner & Romagn. 1953, in Fl. anal. Champ. sup.: 362 (inval.) *Psathyrella pannucioides* (J.E. Lange) M.M. Moser 1967, in Gams, Kl. Krypt.fl. 2b/2, 2. éd. : 220

La description de cette espèce est basée sur plusieurs récoltes réalisées en Belgique en Brabant Flamand.

Trois récoltes ont été réalisées à Bertem, fin octobre et mi novembre 2017, ainsi que le 10/10/2018, dans le Bertembos un bois humide de feuillus divers essentiellement hygrophiles. L'une d'entre elle (DD2216 - GPS: 50.884158, 4.632601) a fait l'objet d'un séquençage.

Les basidiomes se présentaient en petits groupes de quelques exemplaires connés par leur base émergeant le long de rondins pourrisssants utilisés pour stabiliser le chemin boueux. Le long de celui-ci dans l'humus on pouvait observer de grandes touffes denses de spécimens réunis en faisceaux soudés par la base du stipe.

Une récolte complémentaire d'un trentaine d'exemplaires fasciculés a été effectuée le 24/11/2018 à la base d'un vieux saule vivant, à Zemst (Mechelen - GPS: 50. 998266, 4.479677).



Voucher DD2216 – Bertembos – GPS : 50.885479 - 4.634351 Bertem- Belgique





Chapeau mesurant 20(40) x 15 mm, non strié, conico-paraboloïde devenant sur le tard plan convexe, pourvu d'un large mamelon presque lisse, obtus et beige pâle, contrastant avec le reste de la surface du chapeau qui, étant largement recouverte par un voile aranéeux constitué de fibrilles à orientation radiaire, apparaît feutrée, soyeuse et brillante. Le voile déborde de la marge et reste appendiculé à celle-ci, lui donnant un aspect cotonneux.



Lames alternant avec lamelles et lamellules, serrées, largement adnées, brunes devenant brun grisâtre; arête fimbriée concolore ou blanche. La trame lamellaire est nettement pigmentée de brun.

Stipe mesurant 35-60 x 2-3 mm (Örstadius : 30-60 x 2-6 mm), cylindrique, creux, fragile, se brisant aisément, pruineux au sommet et conné par la base qui est parfois légèrement dilatée. Le voile abondant persiste sur certains exemplaires sous forme de fibrilles gris noirâtres teintées par la sporée.



Spores mesurant (8,5) 9,2 -**9,9**- 10,6 (11,1) × (0,6) 5,4 – **5,7**- 6,2 (6,6) μ m (Örstadius : 9-11,5 x 5-6,5 μ m, Qav 1,7 -1,9) ; non opaques, brunes dans l'ammoniaque et grise dans le KOH, oblongues à ellipsoïdes, asymétriques de profil et légèrement amygdaliformes ; dépression suprahilaire fréquente, pore germinatif large de 2 μ m, souvent tronqué.







Mesures réalisées avec Piximètre (8,5) 9,2 - 10,6 (11,1) × (0,6) 5,4 - 6,2 (6,6) μ m Q = (1,4) 1,6 - 1,8 (2,1) ; N = 100 Me = 9,9 × 5,7 μ m ; Qe = 1,7





Cheilocystides nombreuses ventrues, sublagéniformes, spatulées, mesurant 40-65 x 13-26 µm ;

(Örstadius : 30-80 x 11-25 µm) ; mêlées à de nombreuses basidioles et basides, les cellules marginales clavées et sphéropédonculées étant peu fréquentes.



cheilocystides







Pleurocystides peu nombreuses à nombreuses suivant les lames examinées, essentiellement utriformes, ventrues et spatulées, ainsi que parfois lagéniformes, stipitées ou non, semblables aux cheilocystides ; mesurant 40-83 x 16-30 μ m ; (Örstadius : 35-90 x 12-24 μ m).











Pileipellis constitué de cellules subglobuleuses, irrégulières et entremêlées, recouvertes d'une fine couche d'hyphes cylindriques légèrement pigmentées.
Basides tétrasporiques, mesurant 20-30 x 10-12 μm.
Boucles présentes.



Caulocystides très nombreuses, clavées ou analogues aux cystides.



Le séquençage ADN (ITS & LSU) été effectué par Pablo Alvarado Garcia ; laboratoire Alvalab référence: 2017-803-ALV13310 - DD2216.

Genbank Accession number/Version: MK400695.1

DD2216_Coprinopsis_pannucioides_ITS_final

DD2216_Coprinopsis_pannucioides_LSU

Sequencing and phylogenetic analysis

DNA Extraction, Amplification and Sequencing of the fungus was performed by Alvalab (Oviedo, Spain). The phylogenetic analysis was done by Dieter Wächter (Thiersheim, Germany). The genomic DNA was extracted from dried fruiting bodies. Amplification of the ITS region was performed with the ITS4 primer [1], amplification of the LSU region was performed with the LR0R primer [2]. The initial base calling was done with FinchTV [3]. The nucleotide sequences were checked manually for errors, as well as the base calling at unsafe regions (trails, low confidence scores, stutters and polymorphs) on the basis of existing sequences of the */canoceps-clade by divergence matrix and corrected if necessary. In the present case only a trimming of the trails and som minor* corrections were necessary. The following molecular phylogenetic markers were used for the phylogenetic analysis: ITS1 (Internal Transcribed Spacer 1), 5.8S (5.8S rRNA Gene), ITS2 (Internal Transcribed Spacer 2), LSU (Large Subunit 28S rRNA Gen), β-tub (exons of the β -tubulin gene), ef-1 α (exons of the ef-1 α gene). The nucleotide sequences for the tree inference were taken from NCBI [4] and Unite [5] (essential ones of the /cortinata, /Nivei, /canoceps and /Fragilissimae-clades see Table 1). Region boundaries for the ITS- and LSU-region were carried out with ITSx [6] and HMMER [7] including the databases. As outgroup, the sequence sets of the most closely related clades of the ingroup were used, i.e. from the Genera Lacrymaria, Homophron and Parasola. Due to the rapidly evolving, indel-rich areas of the ITS region, it can only be aligned veridical by using an iterative multigene-guide tree. The initial alignment of the ITS region was performed with Mafft [8] using the FFT-NS-2 method. The initial alignments of the LSU-, β-tub and ef-1α genes was carried out using E-INS-i method. The indel matrices for the ITS and LSU regions were each coded with SegState [9] using the SIC = "Simple Indel coding" [10] method. After each alignment step, an ML analysis with RAxML [11] (model: GTRCAT, refining under GTR+G for DNA, GTR2+G with acquisition bias correction according to Lewis [12] for indel partitions) was carried out and the resulting best tree was used as a guide tree for the refinement of the ITS1 and ITS2 MSA. The iterative alignments were done with Prank [13], whereby the switches -once and -uselogs were set. Tracing values were recorded, evaluated statistically and thus the end of the iteration loop of the alignment was determined. The partitioning of all alignments and the indel matrices as well as the model selection for the DNA alignments was done with Partitionfinder [14]. For the final partitioning, the guide tree of the last iteration step was used. As information criterion the Bayesian Information Criterion (BIC) [15] used was after comparison with the Corrected Akaike Information Criterion (AICc) [16] and evaluation with respect to over- or under-partitioning. The partitioning scheme of the final phylogeny was:

•DNA-partition 1: ITS1 + ITS2

•DNA-partition 2: 5.8S

•DNA-partition 3: LSU + β -tub Codon 1

•DNA-partition 4: β-tub Codon 1 + ef-1α Codon 1 + ef-1α Codon 2

•DNA-partition 5: β -tub Codon 3 + ef-1 α Codon 3

•Binary partition (gap matrices): ITS1 + ITS2 + LSU

The final maximum likelihood analysis was done with RAxML 8.2.10 [11]. For all DNA partitions, the GTR substitution matrix [17] under the CAT model [11] was used. The final optimization took place under gamma distribution [11]. For the binary partitions, the "Two State Time-Reversible Model" with acquisition bias correction [12] was used. 1000 ML bootstrap inferences were calculated. Of these, 1000 trees were sampled and the best tree was labeled with the ML bootstrap support values and collapsed to the ML bootstrap value of 50%. The phylogram in Fig 1 was edited with Treegraph [18]. The Outgroup and the upper Coprinopsis clades has been collapsed for a better view.

Arbre phylogénétique précisant la position de ma récolte de Coprinopsis pannucioides- DD2216



Fig 1 50% collapsed maximum likelihood consensus phylogram. The values on the branches are ML bootstrap values. Abbreviations: I: ITS region, L: LSU region, B: β -tubulin region, A: ef-1 α region.

Table 1 List of relevant sequences used in this publication

Art	Beleg	ITS1	LSU	β-Tub	ef-1α
Coprinopsis aesontiensis	LZ P-7614	KY554753.1	KY554752.1		
Coprinopsis afronivea	SFSU BAP 619	NR_148105.1			
Coprinopsis candidata	HB19840914A	follows			
Coprinopsis canoceps	LO148-95	KC992964.1	KC992964.1		
Coprinopsis cerkezii	CNF 1/7253	KX869912.1	KX869913.1		
Coprinopsis marcescibilis	SZMC-NL-0629	FM878021.1	FM876278.1	FN396267.1	FM897256.1
Coprinopsis marcescibilis	LO31-03	DQ389728.1	DQ389728.1	KJ664919.1	KJ732829.1
Coprinopsis marcescibilis	SZMC-NL-2140	FM878020.1	FM876277.1	FN396271.1	FM897257.1
Coprinopsis musae	JV06-179	KC992965.1	KC992965.1	KJ664920.1	
Coprinopsis musae	JV06-180	KC992966.1	KC992966.1	KJ664921.1	KJ732830.1
Coprinopsis nivea	TU118721	UDB019531	UDB019531		
Coprinopsis nivea	4585	JF907848.1			
Coprinopsis nivea	SZMC-NL-0847	HQ847032.1	HQ847117.1	HQ847182.1	
Coprinopsis pannucioides	SZMC-NL-3528	FN396143.1	FN396202.1	FN396341.1	FN396238.1
Coprinopsis pannucioides	10143-03	DO389727.1	D0389727.1	KI664917.1	
Continonsis nseudocortinata	HB20161119A	follows			
Continonsis pseudomarcescibilis	AH-33775	KY698006 1			
Coprinces as a section of the sectio	AU-32710	KY608000.1			
	All.23711	KYC08008.1	ME02224E 1		
	AII:33/11	K1696008.1	WIF035345.1		
Coprinopsis pseudomarcescipilis	AH:33712	KY698007.1	FN 44 CO 700 4	512052004	51420500.4
coprinopsis pseudonivea	SZMC-NL-2340	FM163181.1	FW160728.1	FN396288.1	FN430698.1
Coprinopsis sp.	CBM-FB41367	LC259499.1			
Coprinopsis sp.	CBM-FB42007	LC259498.1			
Coprinopsis sp.	CBM-FB39186	AB854626.1			
Coprinopsis sp.	CBM-FB38829	AB854625.1			
Coprinopsis sp.	Mushroom Observer 260358	MF163178.1			
Coprinopsis sp.	TU124456	UDB028407	UDB028407		
Coprinopsis sp.	HB19881002A HB20101002A	follows			
Coprinopsis submicrospora	AH27055	KC992959.1	KC992959.1	KJ664918.1	
Coprinopsis udicola	AH:33715	KY698002.1	KY698003.1		
Coprinopsis udicola	AM1240	KC992967.1	KC992967.1	KJ664922.1	KJ732831.1
Coprinopsis udicola	AH:33714	KY698004.1	KY698005.1		
Coprinopsis utrifer	SZMC-NL-0591	FN396140.1	FN396209.1	FN396356.1	
Coprinus bellulus	SZMC-NI -2341 A	FM163176.1	FM160680.1	FN396274.1	
Coprinus bellulus	SZMC-NI -2341 B	FN430682.1			
	SZMC-NI -3414	FN396122.1	FN396207 1	EN396354 1	
Coprinus cortinatus	16428	15907847 1	11055020712	1105055412	
Coprinus cortinatus	57MC NI 1601	EN206121.1	EN206171 1	EN1206246 1	ENI206224 1
Reathyrolla off, canoconc	TI1119496	1000017029	UDB017029	110390340.1	FN350224.1
Psathyrella all. calloceps	10110400 BBNN4705625	008017928	008017928		
Psati yreia an. nuronensis		AWI712291.1	AW1712291.1		
Psathyrella submicrospora	SZMC-NL-0635	HQ847053.1	HQ84/133.1		
Uncultured Agaricaceae	0194	AMU/6653.1			
Uncultured Agaricaceae	C191	AMU76650.1			
Uncultured Agaricaceae	C193	AM076652.1			
Uncultured Agaricaceae	C192	AM076651.1			
Uncultured Agaricales	HPm2c251	JN802317.1			
Uncultured fungus	15962456	15962456.1			

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Elewijt $- \frac{24}{10} / \frac{2020}{-A}$ la base d'un vieux saule - At the basis of an old willow tree.





Spores bossues - Hunchbacked spores





Spores mesures : N = 50 (9,5) 9,54 - 10,5 (10,9) × (5,1) 5,3 - 5,9 (6) μ m ; Me = 10,1 × 5,6 μ m ; Q = (1,7) 1,71 - 1,87 (1,9) ; Qe = 1,8 Cheilocystides denses, clavées et spatulées - Cheilocystidia densely packed, clavate and spatulate.



Cheilocystides mesures (N = 29) : (28,3) 34,7 - 46,4 (51) × (11,4) 12,1 - 18,5 (19,3) μm ; Me = 40,3 × 14,9 μm. Basides tétrasporiques. Pleurocystides clavées analogues aux cheilocystides.



Pleurocystides mesures : N = 10 (42,3) 46,5 - 49,1 (52,8) × (16,6) 16,9 - 18,5 (20,7) μ m ; Me = 47,8 × 17,8 μ m.

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Remerciements :

à Pablo Alvarado Garcia pour le séquençage ADN de cette récolte, à Dieter Wächter pour l'étude des séquences ADN et le création de l'arbre phylogénétique ainsi qu'à Marcel Lecomte pour la relecture de cet article sur le plan de la forme.

Note:

Le présent article ainsi que d'autres, relatifs aux *Psathyrellaceae* sont disponibles au format pdf sur le site de l'Association des Mycologues francophones de Belgique (AMFB) : http://www.amfb.eu/Myco/Psathyrelles/psathyrella.html

Je suis intéressé par l'examen de toute récolte de *Psathyrelle* que vous voudrez bien me confier.

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