

REGISTRE FRANÇAIS DE LA MUCOVISCIDOSE

PUBLICATIONS

2022

Corvol H, de Miranda S, Dehillotte C, Lemonnier L, Chiron R, Danner-Boucher I, Hamidfar R, Houdouin V, Macey J, Marguet M, Murris-Espin M, Reynaud Q, Reix P, Reynaud Gaubert M, Burgel PR, French Cystic Fibrosis Reference Network study group

Cumulative Incidence and Risk Factors for Severe Coronavirus Disease 2019 in French People With Cystic Fibrosis

Clin Infect Dis. 2022 Apr 27

Martin C, Reynaud-Gaubert M, Hamidfar R, Durieu I, Murris-Espin M, Danner-Boucher I, Chiron R, Leroy S, Douvry B, Grenet D, Mely L, Ramel S, Montcouquiol S, Lemonnier L, Burnet E, Paillasseur JL, Da Silva J, Burgel PR.

Sustained effectiveness of elexacaftor-tezacaftor-ivacaftor in lung transplant candidates with cystic fibrosis.

J Cyst Fibros. 2022 Feb 2

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Burgel PR, Durieu I, Chiron R, Ramel S, Danner-Boucher I, Prevotat A, Grenet D, Marguet C, Reynaud-Gaubert M, Macey J, Mely L, Fanton A, Quetant S, Lemonnier L, Paillasseur JL, Da Silva J, Martin C; French Cystic Fibrosis Reference Network Study Group.

Rapid Improvement after Starting Elexacaftor-Tezacaftor-Ivacaftor in Patients with Cystic Fibrosis and Advanced Pulmonary Disease.

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2020

Denis A, Touzet S, Diabaté L, Durieu I, Lemonnier L, Poupon-Bourdy S, Iwaz J, Reynaud Q, Rabilloud M

Quantifying Long-term Changes in Lung Function and Exacerbations after Initiation of Azithromycin in Cystic Fibrosis

Ann Am Thorac Soc. 2020 Feb;17(2):195-201

Reynaud Q, Rousset Jablonski C, Poupon-Bourdy S, Denis A, Rabilloud M, Lemonnier L, Nove-Josserand R, Durupt S, Touzet S, Durieu I

Pregnancy outcome in women with cystic fibrosis and poor pulmonary function.

J Cyst Fibros. 2020 Jan;19(1):80-83.

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Carine L'Hostis, Clémence Dehillotte, Lydie Lemonnier, Gil Bellis, Gilles Rault, Sophie Ramel, Pierre-Régis Burgel, Claude Férec, Virginie Scotet	Estimation of survival of CF patients in France by two different methods	ECFS 2018 (e-poster)

2017

Anne Munck, Dominique Delmas, Marie-Pierre Audrezet, Lydie Lemonnier, David Cheillan and Michel Roussey	Optimization of the French cystic fibrosis newborn screening programme by a centralized tracking process	J Med Screen 0(0) 1–7
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Burgel PR, Bellis G, Elborn JS.	Modelling future trends in cystic fibrosis demography using the French Cystic Fibrosis Registry: update and sensitivity analysis	Eur Respir J. 2017 Aug 3;50(2).
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L'Hostis C, Dehillotte C, Lemonnier L, Bellis G, Rault G, Ramel S, Burgel PR, Ferec C, Scotet V	Survival of CF patients in France: application of two different methods on data of the French CF Registry	NACFC Abstract 2017
Denis A, Touzet S, Diabate L, Reynaud Quitterie, Lemonnier Lydie, Poupon-Bourdy S, Durieu I, Rabilloud M.	Long-term effects of azithromycin in cystic fibrosis	Abstract ECFS 2017
VONGTHILATH R, Dehillotte C, Lemonnier L, Guillien A, Richaud-Thiriez B, Degano B, Plésiat P, Dalphin JC	PYOnever study: Characteristics of cystic fibrosis adults free of Pseudomonas aeruginosa pulmonary colonization.	Abstract ECFS 2017
VONGTHILATH R, Dehillotte C, Lemonnier L, Guillien A, Richaud-Thiriez B, Degano B, Plésiat P, Dalphin JC	PYOnever : Caractéristiques des adultes atteints de mucoviscidose indemnes de colonisation pulmonaire par P. aeruginosa	Poster CPLF 2017
Reynaud Q, Poupon-Bourdy S, Rabilloud M, Al Mufti L, Rousset Jablonski C, Lemonnier L, Nove-Josserand R, Touzet S, Durieu I; participating centers of the French Cystic Fibrosis Registry.	Pregnancy outcome in women with cystic fibrosis-related diabetes.	Acta Obstet Gynecol Scand. 2017 Oct;96(10):1223-1227.
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2016

Kris De Boeck, Anne Munck, Isabelle de Monestrol, Vincent Gulmans, Lydie Lemonnier, Peter G. Middleton, Simeon Wanyama, Muriel Thomas.	Does newborn screening influence the young CF cohort included in national CF registries?	ERJ Research letters section 2016
Durieu Isabelle	Vivre avec la Mucoviscidose à l'âge adulte	JFM 2016

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Reynaud, Quitterie 1, 2; Dehillote, Clemence 3; Lemonnier, Lydie 3; nove josserand, raphaela 1; Colomb, Virginie 3; DURIEU, Isabelle	Extra pulmonary comorbidities in transplanted and non-transplanted adult patients in the French Cystic Fibrosis Registry	NACFC 2016
Réchana VONGTHILATH	PYOnever : Caractéristiques des adultes atteints de mucoviscidose indemnes de colonisation pulmonaire par Pseudomonas aeruginosa	Thèse de médecine 2016
Margot SIRVEN	La primo-dénutrition chez les enfants atteints de mucoviscidose – une approche démographique.	Mémoire de master 2 2016

2015

Martin C, Hamard C, Kanaan R, Boussaud V, Grenet D, Abély M, Hubert D, Munck A, Lemonnier L, Burgel PR.	Causes of death in French cystic fibrosis patients: The need for improvement in transplantation referral strategies!	J Cyst Fibros. 2015 Sep 18.
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Férec C.	Cystic fibrosis: from a monogenic disease to a multifactorial disorder.	12th Russian Cystic Fibrosis Congress, 22-24 avril 2015, Moscou.
Roux A-L, Catherinot E, Soismier N, Heym B, Bellis B, Lemonnier L, Chiron L, Fauroux B, Le Bourgeois M, Munck A, Pin I, Sermet I, Gutierrez C, Veziris N, Jarlier V, Cambau E, Herrmann J-L, Guillemot D, Gaillard J-L.	Comparing Mycobacterium massiliense and Mycobacterium abscessus lung infections in cystic fibrosis patients.	J Cyst Fibros. 2015 Jan;14(1):63-9.
Al mufti, Lina; Reynaud, Quitterie; Lemonnier, Lydie; Touzet, Sandrine ; Durupt, Stéphane ; Nove-Josserand, Raphaela; Rabilloud, Muriel; Durieu, Isabelle.	Predicting factors of FEV and BMI deterioration in CF pregnant women.	Abstract NACFC.
V. Scotet, I. Duguépérout, C. L'Hostis, MP. Audrézet, G. Rault, L. Lemonnier, G. Bellis, C. Férec.	Towards a better understanding of survival data in CF.	Abstract ECFS.
Durieu Isabelle	Advanced lung disease How to personalize treatment ?	38th European Cystic Fibrosis Conference. Brussels 2015

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Ségolène Bouet, Gil Bellis, Lydie Lemonnier, Marie Sponga, Virginie Colomb-Jung	Mortalité par mucoviscidose : analyse des données du registre français, 1992-2012	BEH
P.R. Burgel , G. Bellis	Modélisation des tendances démographiques de la mucoviscidose en France entre 2010 et 2025	CPLF

2014

Burgel PR, Hamard C, Carlier N, Kanaan R, Abely M, Hubert D, Munck A, Boussaud V, Grenet D, Lemonnier L, Sponga M, Martin C	Specific causes of mortality in CF patients in France 2007-2010	ECFS Abstract 2014
Thomas M, Lemonnier L, Gulmans V, Naehrlich L, Vermeulen F, Cuppens H, Castellani C, Norek A, De Boeck K.	Is there evidence for correct diagnosis in cystic fibrosis registries?	J Cyst Fibros. 2014 May;13(3):275-80.
Jaouen F., Duny Y., Pageaux G.-P., Sponga M., Lemonnier L., Schiesser C., Lemaitre F.	Impact of enteral feeding with nasogastric tube or gastrostomy on pulmonary function for malnourished adult patients with cystic fibrosis.	Communication orale JFHOD 2014 Poster DDW2014.
Jaouen F.	Intérêt de la nutrition entérale sur la fonction respiratoire des patients adultes dénutris atteints de mucoviscidose,	Thèse de médecine 2014. Université de Montpellier.
Cordier J.	Facteurs associés à la colonisation par mycobactéries non tuberculeuses dans la mucoviscidose.	Thèse de médecine 2014. Université de Lille.
Jacob E	Analyse du registre de la mucoviscidose, Étude de l'impact de l'asthme sur le développement à l'âge de 3 ans.	Master Santé Publique 2014. Faculté de médecine Laennec Lyon.
Férec C.	Compound CFTR and other mutations in children.	8th International Symposium on Inherited Diseases of the Pancreas, 2014, Jérusalem.
De Boeck K, Zolin A, Cuppens H, Olesen HV, Viviani L.	The relative frequency of CFTR mutation classes in European patients with cystic fibrosis.	Journal of Cystic Fibrosis 2014, Jul;13(4):403-9.

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Rollet-Cohen Virginie.	Infections respiratoires a mycobacterium bolletii dans la mucoviscidose - caractérisation clinique et microbiologique	DIU pathologie infectieuse pédiatrique. Année 2013-2014.
Kerem E, Viviani L, Zolin A, Macneill S, Hatzigorou E, Ellemunter H, Drevinek P, Gulmans V, Krivec U, Olesen H; on behalf of the ECFS Patient Registry Steering Group.	Factors associated with FEV1 decline in cystic fibrosis: analysis of the data of the ECFS Patient Registry.	Eur Respir J. 2014 Jan;43(1):125-33.

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Roussey-Bihouee T., Auteur, David V.	Impact du dépistage néonatal de la mucoviscidose sur l'état de santé des patients atteints par la maladie : comparaison d'une cohorte de patients mucoviscidosiques bretons dépistés avec une cohorte de patients non dépistés de Loire-Atlantique-Vendée.	Thèse de médecine 2013, qualification en pédiatrie. Université de Nantes.
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M. Bournez*, G. Bellis, C. Pienkowski, M. Tauber, F. Huet	La croissance des fil les suivies pour mucoviscidose de 8 à 20 ans : les données du registre français	Article Congrès de la Société Française de Pédiatrie

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VanDevanter.	Epidemiology of Cystic Fibrosis Lung Disease progression in adolescents.	ECFS Book on Healthcare Issues and challenges in Adolescents in CF, December 2012.
Bellis G, Cazes MH, Parant A	Improved life expectancy in cystic fibrosis patients: real progress or reduced asurement bias?	ECFS Abstract + Poster

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Huet F., Besignor C., Ravilly S. and Bellis G.	Growth during puberty in cystic fibrosis patients. What can we learn from registries?	Hormone Research, 2009 , vol 72 , 263-263 Endocrinology & Metabolism
	Les données du registre français de la mucoviscidose : évolutions et tendances depuis 10 ans.	Vaincre, le Magazine de l'Association. N°120 (février 2009)
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Huet Frédéric, Bellis Gil.	Growth during puberty in cystic fibrosis.	30th European cystic fibrosis conference, Belek, Turquie.
Munck Anne, Bellis Gil, Ravilly Sophie, Huet Frédéric.	Do overweight and obesity improve clinical outcome in the French cystic fibrosis population?	30th European cystic fibrosis conference, Belek, Turquie.
Munck Anne, Bellis Gil, Ravilly Sophie, Huet Frédéric. 2007.	Impact of overweight and obesity on the clinical outcome in the French cystic fibrosis population.	40th Annual meeting of the european society for paediatric gastroenterology, hepatology and nutrition, Barcelone, Espagne.
Munck Anne, Bellis Gil, Ravilly Sophie, Huet Frédéric. 2007.	Impact of overweight and obesity on the clinical outcome in the French cystic fibrosis population.	21st North american cystic fibrosis conference, Anaheim, États-Unis.
Ravilly Sophie, Le Roux Evelyne, Bellis Gil, Dufour Franck.	Epidémiologie et physiopathologie de la mucoviscidose.	Revue Francophone des Laboratoires, (397), p. 25-36.

2006

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Fauroux Brigitte, Le Roux Evelyne, Ravilly Sophie, Bellis Gil, Clément Annick.	Long term non-invasive positive pressure ventilation is associated with a stabilisation in the decline of lung function.	29th European cystic fibrosis conference, Copenhagen, Denmark.
Munck Anne, Sahler Cécile, Desgeorges Marie, Briard Marie-Louise, Farriaux Jean-Pierre, Bellis Gil, Ravilly Sophie.	Severe and mild CFTR genotypes: age and presenting symptoms in patients diagnosed through newborn screening versus clinical symptoms.	29th European cystic fibrosis conference, Copenhagen, Denmark.
Popa Ruxandra, Nourry Annie, Ravilly Sophie, Bellis Gil.	School and CF in France: a cohort study between 1999 and 2003.	29th European cystic fibrosis conference, Copenhagen, Denmark.

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Bellis Gil.	Apports de l'ONM dans le diagnostic et le suivi de la mucoviscidose.	4ème Atelier sur le diagnostic moléculaire de la mucoviscidose, Montpellier, France.
Bourdy Stéphanie, Touzet Sandrine, François Sabrina, Bellon Gabriel, Cracowski Claire, Pin Isabelle, Bellis Gil, Colin Cyrille, Durieu Isabelle.	Évolution de la prise en charge thérapeutique de patients atteints de mucoviscidose entre 1996 et 2003.	6ème Colloque des jeunes chercheurs de la mucoviscidose. Association Vaincre la mucoviscidose, Paris, France.
Cazes Marie-Hélène, Bellis Gil, Nourry Annie, Le Roux Evelyne, Huet Frédéric, Marguet Christophe, Munck Anne, Ravilly Sophie.	Evaluation of clinical advantage from neonatal screening for cystic fibrosis.	28th European cystic fibrosis conference, Hersonissos, Grèce.
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Ravilly Sophie, Olesen Hanne Vebert, Quinton Hebe, Viviani Laura, Wiedemann Bärbel, Le Roux Evelyne, Parant Alain, Bellis Gil.	Study on cystic fibrosis mortality in 5 countries.	28th European cystic fibrosis conference, Hersonissos, Grèce.
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