Registre français
DE LA MUCOVISCIDOSE

REGISTRE FRANÇAIS DE LA MUCOVISCIDOSE

PUBLICATIONS

2021

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.	*Am J Respir Crit Care Med. 2021 Jul 1;204(1):64-73.*
Durieu I, Dalon F, Reynaud Q, Lemonnier L, Dehillotte C, Bérard M, Walther D, Viprey M, Van Ganse E, Belhassen M.	Temporal trends in healthcare resource use and associated costs of patients with cystic fibrosis.	J Cyst Fibros. 2021 Apr 14:S1569-1993(21)00107-7.
Coriati A, Sykes J, Lemonnier L, Ma X, Stanojevic S, Dehillotte C, Carlier N, Stephenson AL, Burgel PR.	The impact of the high emergency lung transplantation program in cystic fibrosis in France: insight from a comparison with Canada.	Eur Respir J. 2021 Jun 25:2100014.
Bain R, Cosgriff R, Zampoli M, Elbert A, Burgel PR, Carr SB, Castaños C, Colombo C, Corvol H, Faro A, Goss CH, Gutierrez H, Jung A, Kashirskaya N, Marshall BC, Melo J, Mondejar-Lopez P, de Monestrol I, Naehrlich L, Padoan R, Pastor-Vivero MD, Rizvi S, Salvatore M, Filho LVRFDS, Brownlee KG, Haq IJ, Brodlie M.	Clinical characteristics of SARS-CoV-2 infection in children with cystic fibrosis: An international observational study.	J Cyst Fibros. 2021 Jan;20(1):25-30.

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	Pregnancy outcome in women with cystic fibrosis and poor pulmonary	J Cyst Fibros. 2020 Jan;19(1):80-83.

S, Durieu I	function.	
Burgel PR, Munck A, Durieu I, Chiron R, Mely L, Prevotat A, Murris-Espin M, Porzio M, Abely M, Reix P, Marguet C, Macey J, Sermet-Gaudelus I, Corvol H, Bui S, Lemonnier L, Dehillotte C, Da Silva J, Paillasseur JL, Hubert D; French Cystic Fibrosis Reference Network Study Group.	Real-Life Safety and Effectiveness of Lumacaftor–Ivacaftor in Patients with Cystic Fibrosis	Am J Respir Crit Care Med. 2020 Jan 15;201(2):188-197.
McClenaghan E, Cosgriff R, Brownlee K, Ahern S, Burgel PR, Byrnes CA, Colombo C, Corvol H, Cheng SY, Daneau G, Elbert A, Faro A, Goss CH, Gulmans V, Gutierrez H, de Monestrol I, Jung A, Justus LN, Kashirskaya N, Marshall BC, McKone E, Middleton PG, Mondejar-Lopez P, Pastor-Vivero MD, Padoan R, Rizvi S, Ruseckaite R, Salvatore M, Stephenson AL, Filho LVRDS, Melo J, Zampoli M, Carr SB; Global Registry Harmonization Group.	The global impact of SARS-CoV-2 in 181 people with cystic fibrosis.	J Cyst Fibros. 2020 Nov;19(6):868-871.
Corvol H, de Miranda S, Lemonnier L, Kemgang A, Reynaud Gaubert M, Chiron R, Dalphin ML, Durieu I, Dubus JC, Houdouin V, Prevotat A, Ramel S, Revillion M, Weiss L, Guillot L, Boelle PY, Burgel PR.	First Wave of COVID-19 in French Patients with Cystic Fibrosis.	J Clin Med. 2020 Nov 10;9(11):3624.

2019

Burgel PR, Lemonnier L, Dehillotte C, Sykes J, Stanojevic S, Stephenson AL, Paillasseur JL	Cluster and CART analyses identify large subgroups of adults with cystic fibrosis at low risk of 10-year death.	Eur Respir J. 2019 Mar 14;53(3).
Mainbourg S, Durieu I, Dehillotte C, Reynaud Q.	Extra-respiratory comorbidities and transplantation in the French cystic fibrosis registry.	Expert Rev Respir Med. 2019 Aug;13(8):799-802.
Coriati A, Sykes J, Nkam L, Hocine MN, Burgel PR, Stephenson AL.	Validation of the French 3-year prognostic score using the Canadian Cystic Fibrosis registry.	J Cyst Fibros. 2019 May;18(3):396-398

Vongthilath R, Richaud Thiriez B, Dehillotte C, Lemonnier L, Guillien A, Degano B, Dalphin ML, Dalphin JC, Plésiat P.	Clinical and microbiological characteristics of cystic fibrosis adults never colonized by Pseudomonas aeruginosa: Analysis of the French CF registry.	PLoS One. 2019 Jan 8;14(1):e0210201

2018

Hubert D, Dehillotte C, Munck A, David V, Baek J, Mely L, Dominique S, Ramel S, Danner Boucher I, Lefeuvre S, Reynaud Q, Colomb-Jung V, Bakouboula P, Lemonnier L.	Retrospective observational study of French patients with cystic fibrosis and a Gly551Asp-CFTR mutation after 1 and 2 years of treatment with ivacaftor in a real-world setting	J Cyst Fibros. 2018 Jan;17(1):89-95.
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
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).
Nkam L, Lambert J, Latouche A, Bellis G, Burgel PR, Hocine MN.	A 3-year prognostic score for adults with cystic fibrosis	J Cyst Fibros. 2017 Nov;16(6):702-708.
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.
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?	Eur Respir J. 2017 Jan 11.

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
Reynaud, Quitterie 1, 2; Dehillote, Clemence 3; Lemonnier, Lydie 3; nove josserand, raphaele 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.
Burgel PR, Bellis G, Olesen HV, Viviani L, Zolin A, Blasi F, Elborn JS. ERS/ECFS Task Force on Provision of Care for Adults with Cystic Fibrosis in Europe.	Future trends in cystic fibrosis demography in 34 European countries.	Eur Respir J. 2015 Jul;46(1):133-41.
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, Raphaele; 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. Brussells 2015
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 paients 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.
Amstead J., Morris J., Denning D.W.	Multi-Country Estimate of Different Manifestations of Aspergillosis in Cystic Fibrosis.	PlosOne 2014, June 10.
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, Hatziagorou 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.

2013

Amoureux L.	Achromobacter xylosoxidans : épidémiologie au CRCM de Dijon et réservoir environnemental.	Thèse de Docteur de l'université de Bourgogne 2013. Discipline : Sciences de la Vie. École Doctorale : Environnements – Santé.
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.
Catherinot E, Roux A-L, Vibet M-A, Bellis G, Lemonnier L, Le Roux E, Bernède-Bauduin C, Le Bourgeois M, Herrmann J-L, Guillemot D, Gaillard J-L., OMA group.	Inhaled therapies, azithromycin and Mycobacterium abscessus in cystic fibrosis patients.	Eur Respir J. 2013 May;41(5):1101-6.
Catherinot E, Roux A-L, Vibet M-A, Bellis G, Ravilly S, Lemonnier L, Le Roux E, Bernède-Bauduin C, Le Bourgeois M, Herrmann J-L, Guillemot D, Gaillard J-L.	Mycobacterium avium and Mycobacterium abscessus complex target distinct cystic fibrosis patient subpopulations.	J Cyst Fibros. 2013 Jan;12(1):74-80.
Vandevanter DR, Pasta DJ.	Evidence of diminished FEV1 and FVC in 6-year-olds followed in the European cystic fibrosis patient registry, 2007-2009.	J Cyst Fibros. 2013 Dec;12(6):786-9.
M. Bournez*, G. Bellis, C. Pienkowski, M. Tauber, F. Huet	La croissance des fi lles suivies pour mucoviscidose de 8 à 20 ans : les données du registre français	Article Congrès de la Société Française de Pédiatrie

2012

Boëlle PY, Viviani L, Busson PF, Olesen HV, Ravilly S, Stern M, Assael BM, Barreto C, Drevinek P, Thomas M, Krivec U, Mei-Zahav M, Vibert JF, Clement A, Mehta A, Corvol H; French CF Modifier Gene Study Investigators	European CF Registry Working Group. Reference percentiles for FEV(1) and BMI in European children and adults with cystic fibrosis.	Orphanet J Rare Dis. 2012 Sep 7;7:64.
Bellis Gil, Le Roux Evelyne, Chalem Ylana, Parant Alain, Ravilly Sophie.	Projection of lung transplantation for cystic fibrosis needs in France from 2001 to 2010.	Saint-Louis Abstract
Thomas M, Castellani C, Cuppens H, Gulmans V, Lemonnier L, Norek A, Vermeulen F, De Boeck K.	Who is reported in the Belgian, Dutch and French CF registries?	ECFS Abstract 2012
Bournez M, Bellis G, Huet F.	Growth during puberty in cystic fibrosis: a retrospective evaluation of a French cohort.	Arch Dis Child. 2012 Aug;97(8):714-20.

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
2011		
KD Lu, C Engmann, F Moya and M Muhlbach.	Cystic fibrosis in premature infants.	J Perinatol. 2011 Jul;31(7):504-6.
Bellis G.	Registre de la mucoviscidose : des données encourageantes.	Vaincre, le Magazine de l'Association. N°128 (février 2011)
2010		
Burgel PR, Fajac I, Hubert D, Grenet D, Stremler N, Roussey M, Siret D, Languepin J, Mely L, Fanton A, Labbé A, Domblides P, Vic P, Dagorne M, Reynaud-Gaubert M, Counil F, Varaigne F, Bienvenu T, Bellis G, Dusser D.	Non-classic cystic fibrosis associated with D1152H CFTR mutation.	Clin Genet. 2010 Apr;77(4):355-64.
Bellis G., Cazes M. Hélène, Lemonnier L., Ravilly S.	Épidémiologie de la mucoviscidose en France : le rôle positif des CRCM.	Le Concours Médical (132), p.634.
Bellis G., Lemonnier L., Ravilly S.	Registre français de la mucoviscidose : les tendances 2007.	Vaincre, le Magazine de l'Association. N°124 (février 2010)
2009		
Lemonnier Lydie, Ravilly Sophie, Munck Anne, Roussey Michel and the participating centres of the French CF Registry.	Criteria for diagnosis of CF in the French Registry.	Poster and oral presentation at the 32nd European CF Conference (2009).
Huet F., Bensignor 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)

Roux A-L., Catherinot E., Gaillard J-L., Herrmann J-L. and al for the OMA Group.	Multicenter Study of Prevalence of Nontuberculous Mycobacteria in Patients with Cystic Fibrosis in France.	J Clin Microbiology, Dec. 2009, p. 4124–4128
2008		
Durieu I. and Josserand RN.	Cystic fibrosis in 2008.	Rev Med Interne. 2008 Nov;29(11):901-7.
Fauroux B, Le Roux E, Ravilly S, Bellis G, Clément A.	Long-Term Noninvasive Ventilation in Patients with Cystic Fibrosis.	Respiration. 2008;76(2):168-74
2007		
Bellis Gil, Cazes Marie Hélène, Parant Alain, Gaimard Maryse, Travers Cécile, Le Roux Evelyne, Ravilly Sophie, Rault Gilles.	Cystic fibrosis mortality trends in France.	Journal of Cystic Fibrosis, 6, p. 179-186.
Bellis Gil, Parant Alain. 2007.	Mucoviscidose, mise en couple et procréation.	14ème Colloque national de démographie, Cudep, Bordeaux, France.
Bellis Gil.	Intérêt des bases de données pour la recherche épidémiologique.	Atelier de l'Association française contre les myopathies : les gènes modificateurs dans les maladies rares, Paris, France.
Gaimard Maryse, Bellis Gil, Parant Alain. 2007.	L'observatoire national de la mucoviscidose : quelle couverture de la population ?	14ème Colloque national de démographie, Cudep, Bordeaux, France.
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

Bellis Gil, Rault Gilles.	Espérance de vie : mesure et interprétation.	La Lettre de la Mucoviscidose, 62, p. 1.
Cazes Marie-Hélène, Bellis Gil, Nourry Annie, Le Roux Evelyne, Ravilly Sophie, Munck Anne, Huet Frédéric, Marguet Christophe.	Is there a clinical benefit from neonatal screening for cystic fibrosis.	Cystic Fibrosis Worldwide Newsletters, 7 (1), p. 32-33.
Cazes Marie-Hélène, Bellis Gil, Nourry Annie, Parant Alain, Popa Ruxandra.	Observation de la population française atteinte d'une maladie rare : la mucoviscidose.	Chaire Quételet 2006. Les systèmes d'information en démographie et en sciences sociales. Nouvelles questions, nouveaux outils, Louvain-la-Neuve, Belgique.
Duguépéroux I, Hubert D, Dominique S, Bellis G, De Braekeleer M, Durieu I.	Paternity in men with cystic fibrosis: a retrospective survey in France.	J Cyst Fibros. 2006 Dec;5(4):215-21.
Dyard François, Ravilly Sophie, Bellis Gil. 2006.	Pancreatic enzymes consumption by CF patients in France.	29th European cystic fibrosis conference, Copenhague, Danemark.
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, Copenhague, Danemark.
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, Copenhague, Danemark.
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,

		Copenhague, Danemark.
2005		
Bellis Gil, Parant Alain.	Projections démographiques. Contribution de l'INED au Livre Blanc sur la mucoviscidose.	
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.
Duguépéroux I, De Braekeleer M.	The CFTR 3849+10kbC->T and 2789+5G->A alleles are associated with a mild CF phenotype.	Eur Respir J. 2005 Mar;25(3):468-73.
Munck Anne, Sahler Cécile, Desgeorges Marie, Briard Marie-Louise, Farriaux Jean-Pierre , Bellis Gil, Ravilly Sophie	CF newborn screening: diagnosis saving time and presenting symptoms according to the genotype.	19th North american cystic fibrosis conference, Baltimore, États-Unis.
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.
McCormick J, Sims EJ, Green MW, Mehta G, Culross F, Mehta A.	Comparative analysis of Cystic Fibrosis Registry data from the UK with USA, France and Australasia.	J Cyst Fibros. 2005 May;4(2):115-22.
2004		
Badet F, Bellis G, De Braekeleer M, Nove Josserand R, Vital Durand D; The Cystic Fibrosis Centre of ONM, Durieu I.	Phenotype and genotype of French cystic fibrosis patients with long survival and follow-up.	Eur J Intern Med. 2004 Jul;15(4):238-241.

Bellis Gil.	Observatoire National de la Mucoviscidose : quels enseignements, quel avenir ?	5èmes Assises des réseaux de soins de la mucoviscidose, Versailles, France.
Cazes Marie-Hélène, Bellis Gil, Parant Alain.	Essai d'estimation de la contribution globale du gène de la mucoviscidose à la population française actuelle.	13ème Colloque national de démographie, Cudep, Dijon, France.
de Braekeleer Marc, Rault Gilles, Bellis Gil. 2004.	Reproductive attitudes of couples having a child with cystic fibrosis in Brittany (France).	Journal of Human Genetics, 49, p. 285-289.
Duguépéroux I, Bellis G, Lesure JF, Renouil M, Flodrops H, De Braekeleer M.	Cystic fibrosis at the Reunion Island (France): spectrum of mutations and genotype-phenotype for the Y122X mutation.	J Cyst Fibros. 2004 Aug;3(3):185-8.
Duguépéroux I, De Braekeleer M; The Participating Centres to the French National Cystic Fibrosis Registry.	Genotype-phenotype relationship for five CFTR mutations frequently identified in western France.	J Cyst Fibros. 2004 Dec;3(4):259-63.
Duguépéroux Ingrid, Bellis Gil, de Braekeleer Marc 2004.	Are the Y1092X and R1162X stop mutations, both located in the transmembrane segment 11, associated with various phenotypes?	27th European cystic fibrosis conference, Birmingham, Royaume-uni.
Duguépéroux Ingrid, Bellis Gil, de Braekeleer Marc	The CFTR 2789+5G>A allele: a mutation associated with a milder cystic fibrosis phenotype.	27th European cystic fibrosis conference, Birmingham, Royaume-uni.
Duguépéroux Ingrid, Bellis Gil, de Braekeleer Marc.	Étude de la mortalité (1992-2001) : bilan d'après l'Observatoire National de la Mucoviscidose.	5ème Colloque des jeunes chercheurs de la mucoviscidose. Association Vaincre la mucoviscidose, Paris, France.
Duguépéroux Ingrid, Bellis Gil, de Braekeleer Marc.	The CFTR 3849+10kbC>T/DeltaF508 genot ype is usually associated with a milder phenotype.	27th European cystic fibrosis conference, Birmingham, Royaume-Uni.
Duguépéroux Ingrid, Hubert Dominique, Dominique Stéphane, Bellis Gil, de Braekeleer Marc, Durieu Isabelle, Centres the French CF. 2004.	Paternity in men with cystic fibrosis: a retrospective study in France.	18th North american cystic fibrosis conference, Saint-Louis, États-Unis.
Guéganton Laëtitia, Bellis Gil, Chaventré André, Rault Gilles.	A database for knowledge of pedigrees and genotypes of CF patients and their parents in France.	27th European Cystic Fibrosis Conference, Birmingham, Royaume-Uni.

2003

Duguépéroux Ingrid, Bellis Gil, de Braekeleer Marc.	The DeltaI507 mutation: a rare allele associated with a severe phenotype.	26th European cystic fibrosis conference, Belfast, Irlande du Nord.
Duguépéroux Ingrid, Bellis Gil, Lesure Jean- François, Renouil Michel, de Braekeleer Marc.	Genotype/phenotype analysis of Y122X, a mutation particularly frequent in the Reunion Island (France).	26th European cystic fibrosis conference, Belfast, Irlande du Nord.
Huet Frédéric, Bellis Gil, de Monleon Jean-Vital, de Braekeleer Marc, Ravilly Sophie.	Auxological evolution of the french cystic fibrosis patients: a national survey.	26th European cystic fibrosis conference, Belfast, Irlande du Nord.
Le Roux Evelyne, Ravilly Sophie, Bellis Gil, Parant Alain.	Une estimation des besoins de transplantation pulmonaire pour la mucoviscidose.	3ème Congrès de la Société francophone de transplantation, Paris, France.

2002

Duguépéroux I, Bellis G, Férec C, Gillet D, Scotet V, De Braekeleer M; Participating centres of the French CF registry.	Relationship between genotype and phenotype for the CFTR gene W846X mutation.	J Med Genet. 2002 Jun;39(6):E32.
D Gillet, M De Braekeleer, G Bellis, I Durieu	Cystic fibrosis and pregnancy. Report from French data (1980-1999).	British Journal of Obstetrics and Gynecology 2002; 109: 912-918
Duguépéroux Ingrid, Bellis Gil, Lesure Jean- François, Renouil Michel, de Braekeleer Marc.	Genotype/phenotype analysis of Y122X, a mutation particularly frequent in the Reunion Island (France).	2002 Abstract

2001

Ravilly S, Rault G, Bellis G, de Braekeleer M	The French CF Registry general data.	ECFS Abstract
de Braekeleer M, Bellis G,	Clinical aspects of CF in adults. Report from The French CF Registry	ECFS Abstract

D Gillet, I Durieu, M De Braekeleer, G Bellis	Pregnancy in patients with CF. Report from French data (1980-1999).	ECFS Abstract
Gaimard Maryse, Gillet D, Duguépéroux Ingrid, Parant Alain, Bellis Gil, Ravilly S, Rault Gilles, de Braekeleer Marc.	Mortality of patients suffering from CF in France since 1994	Poster
2000		
Guilloud-Bataille M, De Crozes D, Rault G, Degioanni A, Feingold J.	Cystic fibrosis mutations: report from the French Registry.	The Clinical Centers of the CF. Hum Hered. 2000 Mar-Apr;50(2):142-5.
1999		
Feingold J, Guilloud-Bataille M. Clinical Centers of the French CF Registry.	Genetic comparisons of patients with cystic fibrosis with or without meconium ileus.	Ann Genet. 1999;42(3):147-50.
1998		
Delaisi B, Grosskopf C, Reignault E, Goehrs JM, Navarro	International registry on mucoviscidosis: comparison of the French data with the European data for 1995.	J. Arch Pediatr. 1998 Apr;5(4):384-8. French.