

Presentation of the Digital Health Acceleration Strategy

French Ministry of Health and Prevention Ministerial Digital Health Delegation louisa.stuwe@sante.gouv.fr

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Agenda

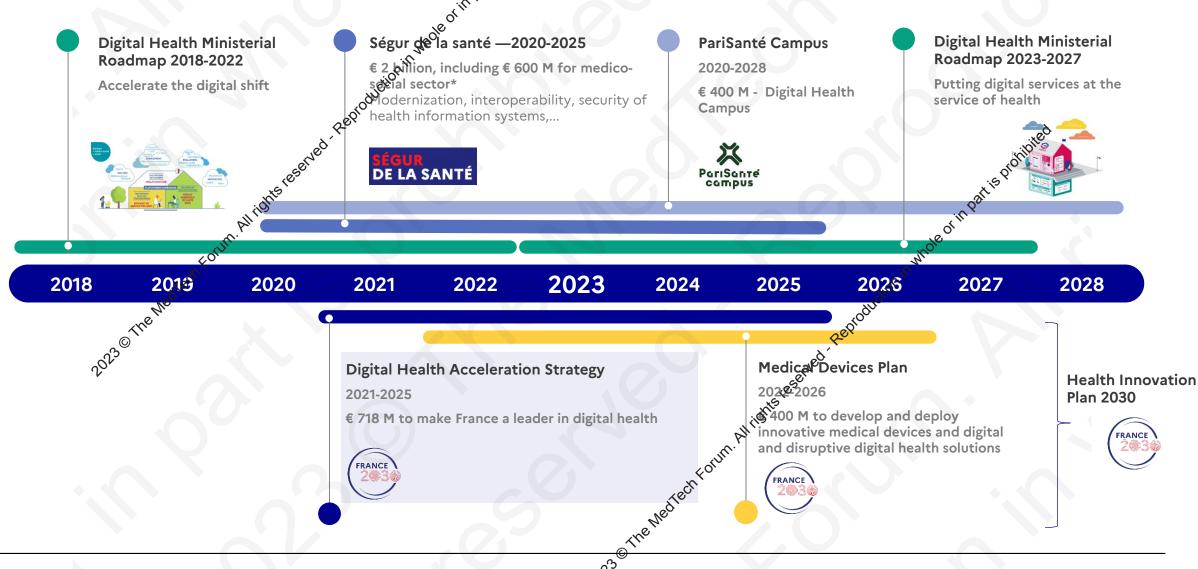
- 1. Digital health acceleration strategy
- 2. G_NJUS platform for digital health entrepreneurs

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3. Fast-track reimbursement in France for digital medical devices (« PECAN »)

French Digital Health Acceleration Strategy

Alignment with other national digital health funding strategies



Digital Health Acceleration Strategy (SASN)

*medico-social sector: institutes caring for children, medico-psychological teaching centres, medical education institutes, nursing homes for the dependent and elderly etc...

3

Digital Health Acceleration Strategy

A national funding program co-constructed by and for the health digital ecosystem



Digital Health Acceleration Strategy (SASN)

Digital Health Acceleration Strategy

SASN

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A strategy with € 718 M to support all levels of a digital health project cycle

Foster the conditions for the large-scale deployment of successful digital health projects

Sustain the implementation of experimentations in real-life conditions and first industrial steps

Support the development of structural projects to strengthen strategic territorial advantages

Prepare the future generation of key digital health technologies and facilitate rapid transfer mechanisms of research results

Develop stakeholder training, stakeholder trust and professional attractiveness of the sector

2023

Digital Health Acceleration Strategy

First assessment: an encouraging first year

First semester 2022: SASN "Tour of France"

- 9 stages to meet local digital health actors sil over France
- 27 roundtables and workshops with move than 600 actors involved

2nd semester 2022: One-year SASN celebration

- > 500 participants
- Featuring Minister of Education and Research, Minister of Health and Prevention, European Commission and General Secretary for Investment

€152 M committed in 2022

• at 31/12/2022 Ne. 1/3 of the France 2030 budget spent since the launch of the strategy

31 actions launched

• including 9 call for projects

61 winners of call of projects

More than 500 applicants for digital health call for projects

More ambition, more openness, more funding for digital health so far. A major component of France Santé 2030, to prepare the future of our healthcare system and the ecosystem of digital healthcare companies.

Agnès Audier, **Ambassador France 2030**





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https://gnius esante.gouv.fr

Support the European digital health entrepreneurs to navigate through the digital health ecosystem and accelerate market access in the EU



G_NIUS is a French initiative, part of the 2021-2025 Digital Health Acceleration Strategy (670 B€) willing to connect to European colleagues. A single gateway to support digital health entrepreneurs navigating through the ecosystem, save time and accelerate access to the market of their solutions



Discover eHealth news



Decrypt pational digital health regulation



Identify actors of the digital health ecosystem



A podcast 100 days to succeed



G_NIUS services



Follow your healthcare pathway (MonEspaceSanté, Ségur numérique, National Health Identity (INS)...)



Decipher funding keys and trends



Commercialize a digital medical device

Identify ehealth events

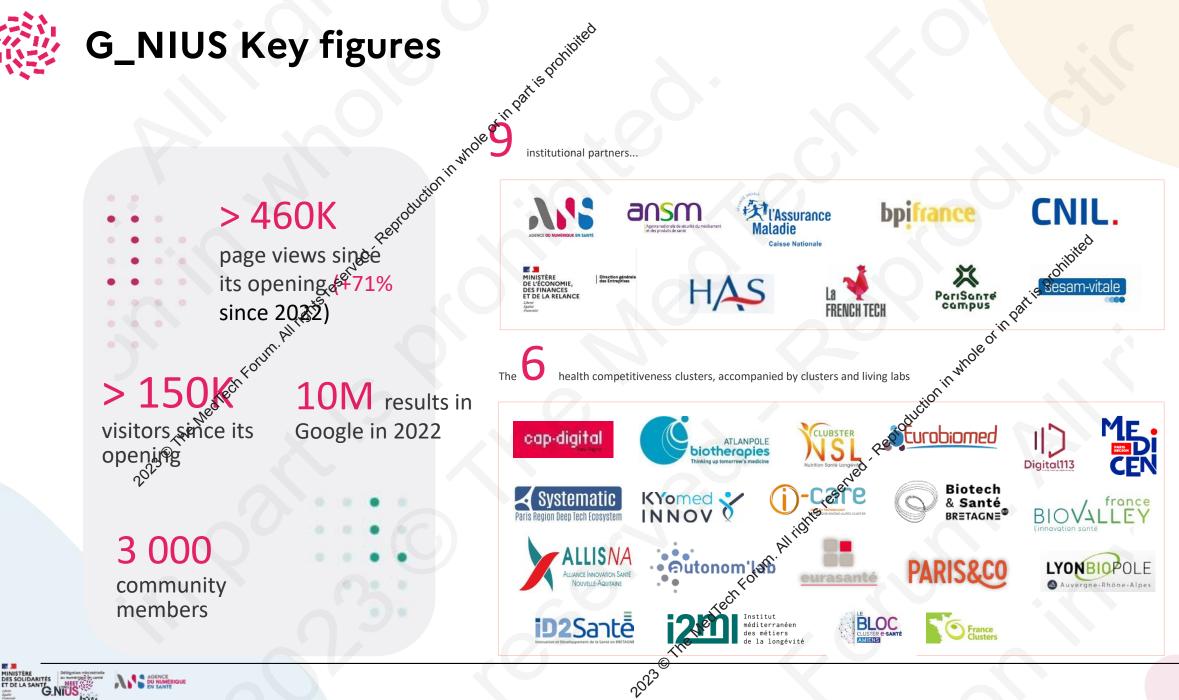


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New: funding search engine for eHealth projects Link



New: map of EU-level markets and hubs Link



Inite









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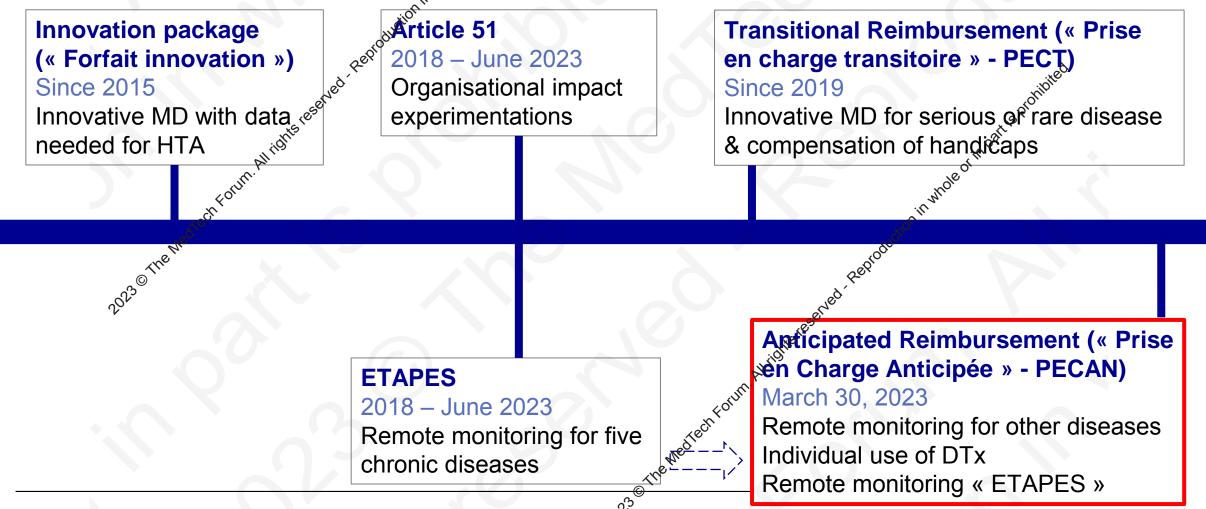


Fast-track reimbursement in France for digital medical devices (« PECAN »)

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Paving the way to a fast-track DMD access in France

PRIOR EXPERIMENTATIONS TO A NATIONAL TRANSITIONAL AND TEMPORARY ONE-YEAR REIMBURSEMENT ACCESS SCHEME FOR DIGITAL MEDICAL DEVICES AND TELEMONITORING IN FRANCE



Fast-track reimbursement in France for digital medical devices (« PECAN ») ϑ°

"PECAN", the new French DMD reimbursement process that accelerates the deployment of digital health solutions

IMPLEMENTED AS PART OF THE "DEPLOYNENT AXIS" OF THE DIGITAL HEALTH ACCELERATION STRATEGY

Launch of digital medical devices (DMD) reimbursement ("PECAN")

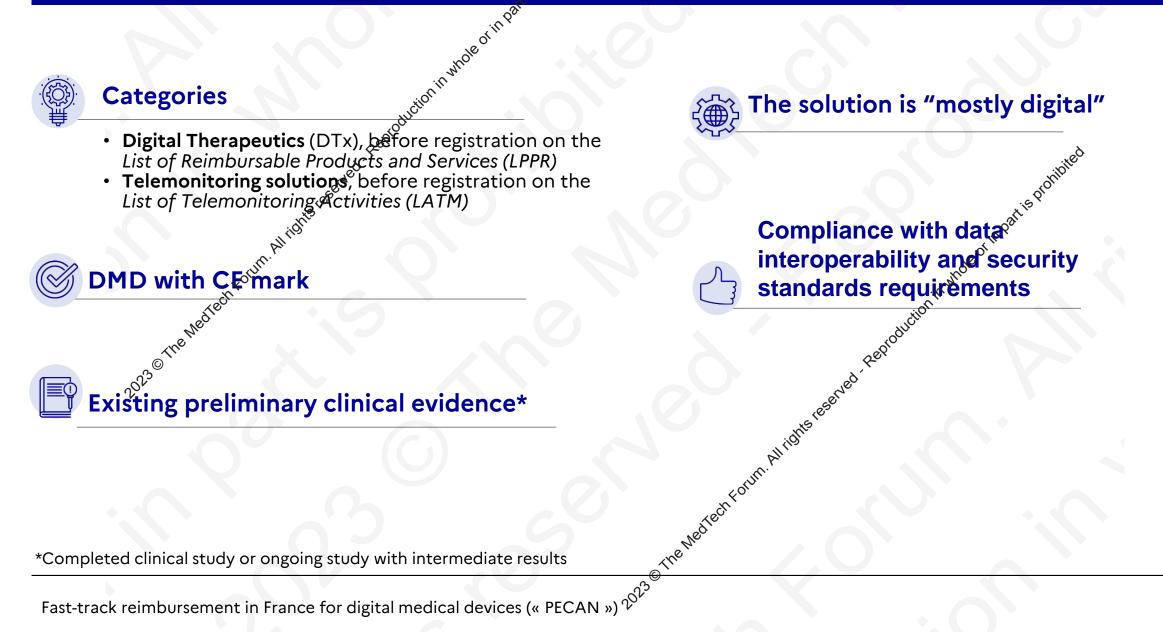
- Objective: to accelerate reimbursement by the French National Health Insurance for innovative DMDs
- Provision for "PECAN" (Prise en charge anticipée numérique) by French 2022 Social Security Funding Law (Årt. 58)
- Implementation throwgh <u>ministerial order of March 30, 2023</u> by means of service desks offered by the French Digital Health Agency (ANS) and HTA agency (HAS)
- This fast-track enables the manufacturer to **finalize the clinical trials while already being reimbursed during one-year** (non-renewable)

Scope Digital medical devices for therapeutic purposes or innovative medical remove monitoring solutions

More information on the application process on the G_NIUS website G_NIUS

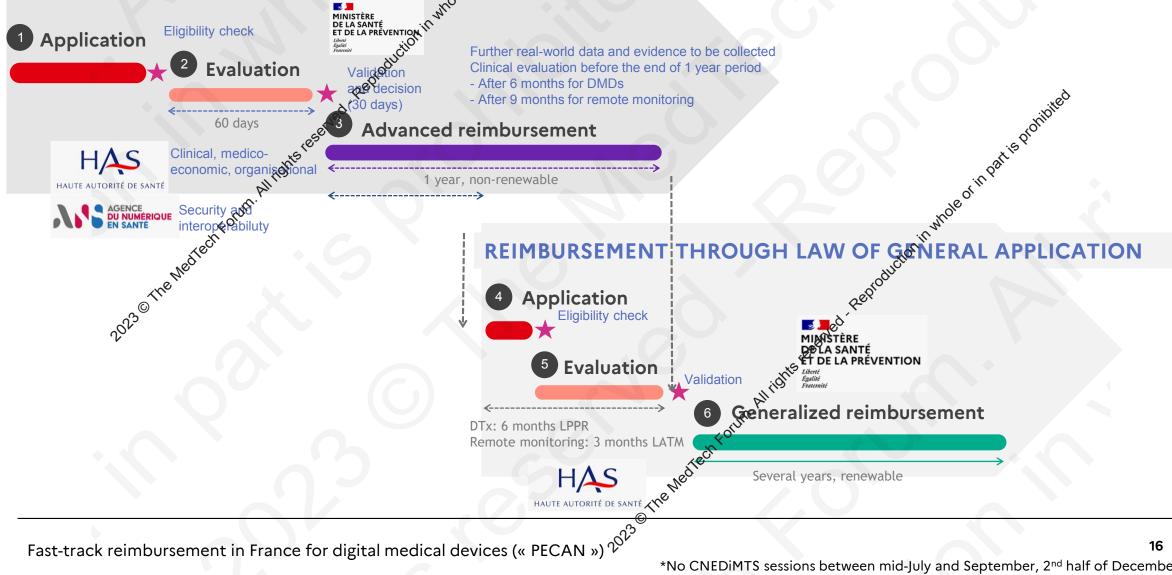


Which types of DMDs are in the scope of PECAN?



What are the steps to access DMD reimbursement?

FAST-TRACK DIGITAL MEDICAL DEVICES REIMBURSEMEN



^{*}No CNEDiMTS sessions between mid-July and September, 2nd half of December

Different approaches for fast track access for DMDs in the EU

BOTH GERMAN AND FRENCH NATIONAL REGULATIONS SUPPORT BROADER AND FASTER ACCESS FOR PATIENTS ACCESS TO INNOVATIVE DMDS, USING DUFFERENT SCOPES AND CATEGORIES OF DIGITAL HEALTH APPLICATIONS

German Digital Healthcare Act

DiGA Fast Track :

- DMD for therapeutic uses (DTx)
- DMD class I and la
- BfArM evaluates DiGA applicants on their potential positive health effects
- One year transitional rebate at manufacturer's rate
- Possibility of additional clinical studies during provisional listing

Social Security Funding Law 2022 + March 2023 order

PECAN:

- DMD for therapeutic uses (DTx) of telemonitoring
- DMD class I, IIa, IIb and III
- DMD presumed to be innovative, especially in terms of clinical benefit or improvement for the organization of care, as assessed by HAS (CNEDIMTS)
- Need to pursue data collection throughout deployment

One year transitional reimbursement period, predefined price **package**

Other European countries are actively working on the implementation of similar procedures

Fast-track reimbursement in France for digital medical devices (« PECAN ») $^{
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17

European Taskforce for Harmonised Evaluation of Digital Medical Devices

A European digital medical device (DMD) taskforce has been launched in April 2022, chaired by the Ministerial Digital Bealth delegation of the French Ministry of Health and Prevention and co-chaired by the European Network for Health Technology Assessment (EUnetHTA), coordinated by EIT Health and supported by contributors of other European Ministries of Health and/or national responsible authorities and agencies.

Mission: To classify innovative DMDs and align their EU-level health technology assessment procedures in the view of harmonizing national assessment in the view of reimbursement by national health insurance organisations for distinct subclasses of DMDs.

→ By developing a **joint approach** duplication of assessments can be avoided, patient access to innovative and proven digital health solutions can be accelerated & health systems improved.

examples



The recommendations for harmonizing clinical criteria and methodologies for evaluating DMDs will result from several work

	nackagos	
Work package 1	Work package 2	Work package 3 & 4
Harmonize the taxonomy of DMDs based on their application scope and evaluation categories	Consensus on determining quantity, quality and the type of evidence that is needed for assessing DMDs	Propose a social health evaluation framework based on pre-requirements (technical, technological, ethical)
External Advisory Soard		
Contribute to the final suggestions by sharing perspectives of different stakeholders and experiences from real-world		

PECAN ministerial order of March 30, 2023 Link Partie Provided PECAN information sheet on G_NIUS Link ECAN Webinar Replay Link Links to go further

More information on PECAN

- HAS guidance Link (in French only)

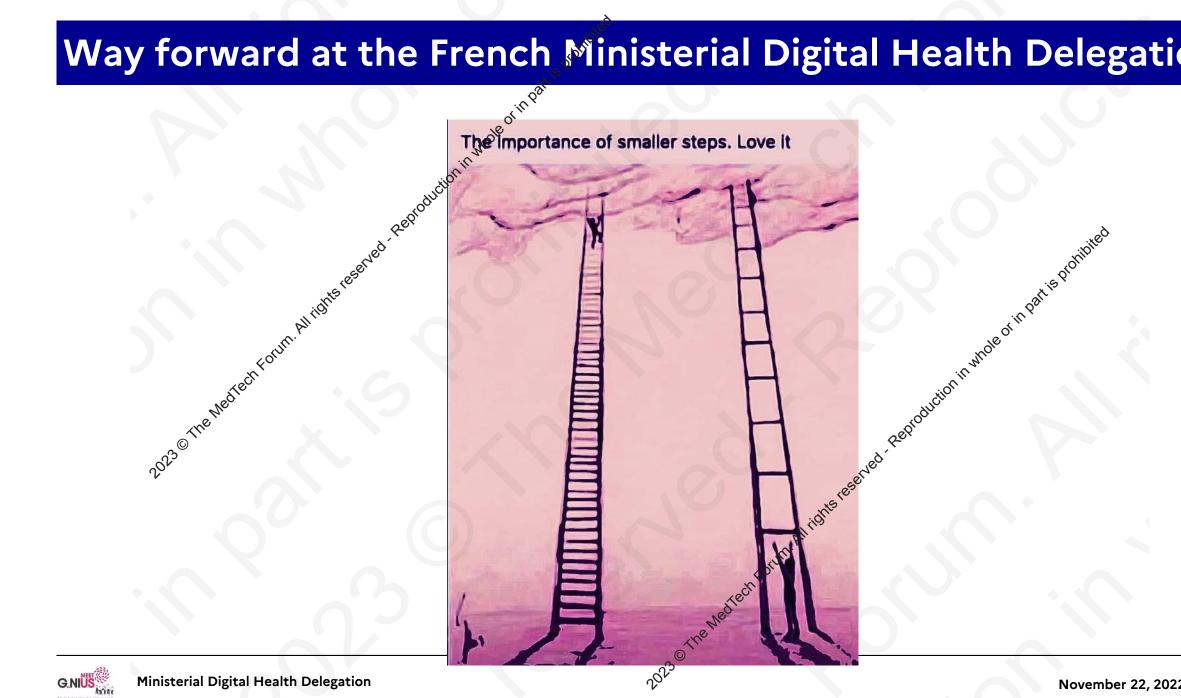
Guidelines by the National Health Technology Assessment Authority (HAS):

- Submission guidelines for medical devices (as of June 1, two different platforms, for clinical evaluation: Evatecho Link
- Evaluation principles by the CNEDIMTS of individual use medical device for their access to reimbursement Link in English / Link in French (May 2019)
- Methodology for the clinical development of medical devices (including digital) Link in English / Link in Evench
- Real-world studies for the assessment of medicinal products and medical devices Link
- Organisational impact for map for health technology assessment Link
- Analysis grid for the AI component in medical devices Link
- Remote monition uses Link in French
- Medical device evaluation by the CNEDIMTS (Medical Device and Health Technology Evaluation Committee): Guide to the specific features of clinical evaluation of a connected medical device (CMD) in view of its application for reimbursement Link
- Guidelines for telemonitoring on 5 chronic diseases (as in ETAPES) Link (In French)
- Functional classification, according to their intended use, of digital solutions used in the context of medical and paramedical care Link

Guidelines by the National eHealth Agency (ANS):

- Repository of Interoperability and Security of Digital Medical Devices Line
- Interoperability and Security standards for Digital Medical Devices (DMDs) Link and requirements Link
- Compliance certification platform Link

Way forward at the French Ministerial Digital Health Delegation











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