



HAL
open science

A Proposal from the Montpellier World Health Organization Collaborating Centre for Better Management and Prevention of Anaphylaxis

Luciana Kase Tanno, Nidhal Touati, Salome Allichon, Bryan Martin, Motohiro Ebisawa, Ignacio Ansotegui, Mario Sanchez-Borges, Victoria Cardona, Paul Greenberger, Dermot Ryan, et al.

► To cite this version:

Luciana Kase Tanno, Nidhal Touati, Salome Allichon, Bryan Martin, Motohiro Ebisawa, et al.. A Proposal from the Montpellier World Health Organization Collaborating Centre for Better Management and Prevention of Anaphylaxis. *Journal of Allergy and Clinical Immunology: In Practice*, 2021, 9 (2), pp.676-683.e1. <10.1016/j.jaip.2020.09.062>. <hal-03641886>

HAL Id: hal-03641886

<https://hal.umontpellier.fr/hal-03641886v1>

Submitted on 13 Feb 2023

HAL is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers.

L'archive ouverte pluridisciplinaire HAL, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.



Distributed under a Creative Commons CC BY-NC 4.0 - Attribution - Non-commercial use - International License

A Proposal From The Montpellier World Health Organization Collaborating Centre For Better Management And Prevention Of Anaphylaxis

Luciana Kase Tanno^{1,2,3,4*} MD PhD, Nidhal Touati² MD, Salome Allichon² MD, Bryan Martin⁵ MD, Motohiro Ebisawa⁶ MD PhD, Ignacio Ansotegui⁷ MD PhD, Mario Sanchez-Borges⁸ MD PhD, Victoria Cardona^{9,10} MD PhD, Paul A. Greenberger¹¹ MD PhD, Dermot Ryan¹² MD, Guillaume Pouessel^{13,14} MD, Etienne Beaudouin¹⁵ MD, Jean-Marie Renaudin¹⁶ MD, Francis Thiens^{17,18} MD PhD, Yoon-Seok Chang¹⁹ MD PhD, Ruby Pawankar²⁰ MD PhD, Maximiliano Gomez²¹ MD PhD, Edgardo Jares²² MD, Patricia Latour Staffeld²³ MD, Ioana Agache²⁴ MD PhD, Antonella Muraro²⁵ MD PhD, Todd A. Mahr²⁶ MD, James Subett²⁷ MD, PhD, Thomas Casale²⁸ MD PhD, David Lang²⁹ MD PhD, Pascal Demoly^{2,3,4} MD PhD

1 Hospital Sírio-Libanês

2 University Hospital of Montpellier, Montpellier, France

3 Sorbonne Université, INSERM UMR-S 1136, IPLESP, Equipe EPAR, 75013, Paris, France

4 WHO Collaborating Centre on Scientific Classification Support, Montpellier, France

5 Medicine and Pediatrics, The Ohio State University in Columbus, Ohio, USA

6 Clinical Research Center for Allergy and Rheumatology, Sagami National Hospital, Japan

7 Department of Allergy and Immunology, Hospital Quirónsalud Bizkaia Erandio, Bilbao, Spain

8 Allergy and Clinical Immunology Department, Centro Medico Docente La Trinidad, Caracas, Venezuela

9 Allergy Section, Department of Internal medicine, Hospital Universitari Vall d'Hebron, Barcelona, Spain

10 ARADyAL research network, Madrid, Spain

11 Division of Allergy and Immunology, Department of Medicine, Northwestern University Feinberg School of Medicine, Chicago, IL, USA.

12 Usher Institute, University of Edinburgh, UK

13 Department of Pediatrics, Children's Hospital, F-59056 Roubaix, France

¹⁴ CHU Lille, Pediatric Pulmonology and Allergy Department, and Lille University, F-59000 Lille, France

15 Regional Institute for Allergic and Environmental Disease Clinical Immunology, Metz Regional Hospital, France

16 Réseau d'Allergo-Vigilance (Allergy Vigilance Network), Vandoeuvre les Nancy, France.

17 Faculty of Medicine, Nursing and Health Sciences, Monash University, Clayton, Vic Australia

18 Department of Respiratory Medicine, Eastern Health, Vic, Boxhill, Australia

19 Seoul National University Bundang Hospital, Seoul National University College of Medicine, Seongnam, Korea

20 Department of Pediatrics, Nippon Medical School, Tokyo, Japan.

21 Research & Education, Fundación Ayre. Allergy & Asthma Unit, Hospital San Bernardo. Salta, Catholic University of Salta, Argentina.

22 LIBRA Foundation and CMP SA Buenos Aires, Argentina

23 Centro Avanzado de Alergia y Asma, Santo Domingo, Dominican Republic

24 Transylvania University, Brasov, Romania

25 Food Allergy Referral Centre , Padua University Hospital , Padua , Italy

26 Pediatric allergy and clinical immunology, Gundersen Health System in La Crosse, USA

27 Managing Partner, Family Allergy & Asthma; Clinical Professor, Section of Allergy & Immunology, Department of Pediatrics, University of Louisville School of Medicine, Louisville, KY USA

28 Morsani College of Medicine, University of South Florida, Tampa, FL, USA

29 Department of Allergy and Clinical Immunology, Respiratory Institute, Cleveland Clinic, Ohio, USA

* Corresponding author: Luciana Kase Tanno MD, PhD, Division of Allergy, Department of Pulmonology, Hôpital Arnaud de Villeneuve, University Hospital of Montpellier, 371, av. du Doyen Gaston Giraud - 34295, Montpellier cedex 5, France. Tel.: +33 467336107 Fax: +33 467633645

E-mail: luciana.tanno@gmail.com

CONFLICT OF INTERESTS:

Pascal Demoly and Luciana Kase Tanno received an unrestricted Novartis and MEDA/Mylan Pharma grants through CHUM administration. LKT received a research AllerGOS grant. The other authors declare that they do not have conflict of interests related to the contents of this article.

CONTRIBUTIONS:

The first and last authors contributed to the construction of the document (designed the study, including the questionnaire, analysed and interpreted the data, and wrote the manuscript). All the authors critically revised and approved the final version of the manuscript and agree to be accountable for all the aspects of the work.

ABBREVIATIONS

AAI: adrenaline auto-injectors

BAT: basophil activation test

CAST: cellular allergen stimulation test

EAACI: European Academy of Allergology and Clinical Immunology

ICD: International Classification of Diseases

IgE: immunoglobulin E

WAO: World Allergy Organization

WHO: World Health Organization

1 **ABSTRACT**

2 Since the first description of anaphylaxis in 1902, its clinical importance as an emergency
3 condition has been recognized worldwide. Anaphylaxis is a severe, potentially life-threatening
4 systemic hypersensitivity reaction characterized by rapid onset and the potential to endanger life
5 through respiratory or circulatory compromise. It is usually, although not always, associated with
6 skin and mucosal changes. Although the academic/scientific communities have advocated to
7 promote greater awareness and protocols for management of anaphylaxis based on best evidence,
8 there are few efforts documenting feedback as to the success of these efforts. In this document, we
9 review the key unmet needs related to the diagnosis and management of anaphylaxis, propose a
10 public health initiative for prevention measures and a timetable action plan which intends to
11 strengthen the collaboration among health professionals and especially primary care physicians
12 dealing with anaphylaxis that can encourage enhanced quality of care of patients with anaphylaxis.

13 More than calling for harmonized action for best management of anaphylaxis to prevent
14 undue morbidity and mortality, the Montpellier World Health Organization Collaborating Centre
15 here proposes an action plan as a baseline for a global initiative against anaphylaxis. We strongly
16 believe these collaborative efforts are a strong public health and societal priority that is consistent
17 with the overarching goals of providing optimal care of allergic patients and best practices of
18 allergology.

19

20 **KEY WORDS:** anaphylaxis, classification, epidemiology, management, treatment, prevention;

21 adrenaline/epinephrine auto-injector

22

23 COPING WITH ANAPHYLAXIS

24 Anaphylaxis: a “118 years-old killing lady”

25 In the time capsule of research and clinical advances in the field of allergy, the years of 1902
26 and 1913 signify milestones in the history of our specialty; in 1902, Paul Portier and Charles Richet’s
27 described the experimental production of aberrant immunity, named “anaphylactique”, and the
28 Nobel Prize of Medicine and Physiology was subsequently awarded for this finding in 1913 (1). Our
29 understanding of the underlying mechanisms of anaphylaxis has evolved dramatically since the first
30 studies associating the physiological actions of histamine and the concept of histamine shock as the
31 basis of anaphylaxis in 1910 by Henry H. Dale (2). Half a century later, Kimishige Ishizaka, Teruko
32 Ishizaka, Gunnar Johansson and their teams demonstrated immunoglobulin E (IgE) as a distinct class
33 of immunoglobulin involved in allergic reactions (3). While Richet believed that anaphylaxis was a
34 “lack of protection”, it has become clear that an exaggerated immune reaction, involving IgE, was
35 the underlying pathomechanism in allergic anaphylaxis, besides immune complex reactions. Non-
36 immunologically mediated reactions leading to similar clinical symptomatology have been called
37 “anaphylactoid” or “pseudoanaphylaxis”, are now called “non-immune anaphylaxis”.

38 In the late 1990s, the allergy nomenclature was revised by Johansson et al., under the aegis of
39 the European Academy of Allergology and Clinical Immunology (EAACI). The published document (4)
40 has gained substantial international recognition and has been reviewed and endorsed by the World
41 Allergy Organization (WAO) (5). This nomenclature has been accepted widely and is currently used
42 worldwide. It was proposed that “anaphylaxis” would be the umbrella term for an acute reaction
43 defined as a severe, life-threatening generalized or systemic hypersensitivity reaction. The term
44 “allergic anaphylaxis” is used to define a reaction mediated by immunological, such as IgE, IgG and
45 immune-complex complement-related mechanisms. Anaphylactic reactions mediated by IgE
46 antibodies were referred to as “IgE-mediated allergic anaphylaxis.” Meanwhile, anaphylaxis from a
47 non-immunological cause should be referred to as “non-allergic anaphylaxis.” The terms
48 “anaphylactoid” and “pseudoanaphylaxis” are no longer used.

49 In the International Consensus on Anaphylaxis, published in 2014 (6), it was pointed out that
50 the correct term anaphylaxis would be preferred to anaphylactic shock since shock is not necessarily
51 present in patients with anaphylaxis. The term anaphylaxis should also be used in preference to
52 terms such as allergic reaction, acute allergic reaction, systemic allergic reaction, acute IgE-mediated
53 reaction, anaphylactoid reaction or pseudo-anaphylaxis (6,7).

54 Epinephrine (adrenaline) is the first medication of choice for the treatment of anaphylaxis (8).
55 It is a potentially life-saving non-selective adrenergic agonist that acts through vasoconstrictor
56 effects, preventing airway mucosal edema and hypotension, that also exerts bronchodilator activity
57 and has inotropic and chronotropic cardiac effects (9). Epinephrine was first discovered by Japanese
58 chemists Jokichi Takamine and Keizo Uenaka in 1900, two years before the first description of
59 anaphylaxis (10,11). The therapeutic potential of epinephrine was widely acknowledged, and it was
60 used before the molecule's mechanism of action was fully appreciated. Manufacturers then began
61 developing synthetic forms of epinephrine. It was first synthesized in 1904 (12). The discovery and
62 purification of epinephrine provided not only long overdue relief from anaphylactic reactions, but
63 also contributed to the beginning of our understanding of hormones and homeostasis. Adrenaline
64 auto-injectors (AAIs) are commercially available in many devices, in doses suitable for most, but not
65 all, adults and children. For instance, in France four commercial forms are available, but the
66 commercial availability varies in different countries (8) (Figure E1).

67 Since then, our progress in achieving further understanding of anaphylaxis has slowed such that
68 David B.K. Golden observed: "Portier and Richet would turn in their graves to know that we are little
69 more enlightened than a century ago on the real nature of anaphylaxis" (13).

70

71 [Anaphylaxis: what did we achieve so far?](#)

72 The awareness of anaphylaxis as a life-threatening medical condition and its incidence have
73 been increasing among different specialities. In recent years, evidence indicates its incidence has
74 been increasing (14). The reported increases probably reflect a true increase in the prevalence of
75 allergic disease, but are also confounded by cumulative incidence of anaphylaxis, better awareness
76 and recognition of anaphylaxis, and changes in anaphylaxis coding, in part due to modifications in
77 the international classification of diseases.

78 Anaphylaxis is recognized as a severe, life-threatening systemic hypersensitivity reaction,
79 characterized by rapid onset and the potential to endanger life through respiratory or circulatory
80 compromise. It is associated in most cases with skin and mucosal symptoms (15-19). It may present
81 with different combinations of symptoms, and early onset of mild cutaneous pruritus may rapidly
82 progress to entail a life-threatening reaction. This multi-faceted condition can manifest at any age
83 and any health professional may be faced with it. Difficulty recognizing anaphylaxis can lead to
84 delayed treatment with epinephrine and increase the risk of untoward outcomes including death
85 (20,21).

86 The incidence of anaphylaxis ranges from 1.5 to 7.9 per 100 000 person-years in European
87 countries (22) and 1.6 to 5.1 per 100 000 person-years in United States (23). Epidemiological data
88 are heterogeneous for a number of reasons. Most of epidemiological studies have addressed
89 subpopulations or specific triggers, which does not provide a global view from a public health
90 perspective that would lead to general recommendations for clinical practice.

91 Anaphylaxis is a recognized cause of death in all age groups, in both genders and regardless of
92 the ethnicity. The rate of anaphylaxis-related mortality is less than 1 per million per year in most
93 high-income countries (15-19). There are limited epidemiological data from middle and low-income
94 countries. One of the recognized difficulties in establishing accurate anaphylaxis data in population-
95 based studies is the misclassification of this condition under the World Health Organization (WHO)
96 International Classification of Diseases (ICD), but this may change with implementation of the ICD-
97 11, which includes a chapter focused on allergic and hypersensitivity conditions (24-27).

98 Recently, the definition of anaphylaxis by the WAO was reviewed, in order to capture not only
99 severe cases (26), but also to align this effort with the anaphylaxis definition proposed in the WHO
100 ICD-11: “Anaphylaxis is a severe, life-threatening systemic hypersensitivity reaction characterized
101 by being rapid in onset with potentially life-threatening airway, breathing, or circulatory problems
102 and is usually, although not always, associated with skin and mucosal changes” (26-28). From the
103 initial description of anaphylaxis as a clinical entity with acute onset of symptoms involving 2 or
104 more organs or with the association with hypotension or upper respiratory commitment, its
105 definition has evolved to a more mechanistic description based on precision medicine into
106 phenotypes with underlying endotypes supported by diagnostic biomarkers (29). Clinical and basic
107 research have been done since the initial description of anaphylaxis 118 years ago, which has
108 advanced the field and provided insight into its pathogenesis and management. For example, IL-33
109 has been described as key cytokine involved in anaphylaxis (30), neutrophils have been described as
110 potential cellular actors in defined types of anaphylaxis (31) and PAF has been described as a new
111 mediator of anaphylaxis among many other basic discoveries (32) and the fact that chemotherapy
112 and monoclonals have been described as the new most important drugs inducing anaphylaxis (33) .
113 Also, mastocytosis, hereditary alpha tryptasemia and non clonal mast cell activation syndrome have
114 been described as the essential clinical causes of anaphylaxis.

115 The WAO also reviewed the diagnostic criteria of anaphylaxis as follows (28) (Table E1).

116 Although the academic/scientific communities have made efforts to work on documents to advocate
117 in favour of awareness and management (15-19,34-37), there are few publications documenting the
118 populations’ feedback to the awareness efforts.

119

120 LESSONS FROM THE FIELD

121 Clinical vignette: “I can’t breathe well!”

122 “I can’t breathe well” was the chief complaint of a 14 year-old girl in 3 previous episodes of
123 anaphylactic reactions during the winter of 2018-2019 – each of which occurred with exercise. In
124 addition to dyspnea and cough, she presented with palpebral angioedema and rhinoconjunctivitis
125 symptoms. All episodes occurred with consumption of food including wheat or wheat-containing
126 items within the several hours prior to exercise. In all episodes, she required emergency department
127 management, receiving anti-histamines, inhaled Beta2-agonists and systemic corticosteroids. No
128 adrenaline was administered.

129 She developed asthma in childhood, and was treated with a Beta2 agonist inhaler to be used
130 on an as needed basis. Skin testing demonstrated wheal/flare reactions to multiple inhalant
131 allergens including house dust-mite, cat dander, *Alternaria alternata* and pollens (olive, plane and
132 birch trees, cypress, ambrosia, grass). Despite being multi-sensitized, sublingual allergen
133 immunotherapy with cypress pollen was started in October 2019 and was associated with
134 improvement in respiratory symptoms. The patient had no total serum tryptase measured. Her
135 medications included valproic acid for seizure disorder.

136 Food-dependent exercise-induced anaphylaxis was suspected based on the clinical history.
137 Serum specific IgE (4.5 KU/l) to wheat was detected with total IgE 3000 KU/l, and food provocation
138 test with wheat and subsequent exertion (Treadmill challenge) and 500mg of Aspirin did not
139 provoke generalized reaction. Despite this, the patient was advised to avoid wheat consumption
140 before exercise. She received a written action plan and prescription of AAI.

141 In January 2020, the patient had another episode of anaphylaxis at home, presenting with
142 ocular and axillary pruritus, abdominal pain, pharyngeal edema with cough, urticaria and
143 respiratory distress. Her mother administered EAI (Emerade® 300 µg) and emergency medical
144 services was called. She also received systemic corticosteroids. However, she was advised to avoid a
145 second administration of AAI. Her reaction progressed such that she developed vomiting and
146 hypotension (systolic AP 80mmHg) at the time of the arrival of emergency medical services. No
147 tryptase was measured during the acute phase of the anaphylactic reaction. Since our Allergy
148 Department was also contacted, we recommended obtaining serum tryptase in the emergency
149 department. Tryptase level was 7.2 µg/L (baseline: 3.7 µg/L) taken 4 hours after the acute phase,
150 supporting a diagnosis of anaphylaxis. No specific trigger or co-factor was identified.

151 We reviewed the patient's files in order to identify possible non-investigated food triggers.
152 Based on this evaluation, skin prick tests to pine nuts and tomato were performed and were
153 positive. Differential diagnoses such as vocal cord dysfunction and features of dysautonomia have
154 been ruled out. The patient is undergoing further investigation to revise management
155 recommendations and a provocation test is scheduled.

156 This case highlights the challenges we may face in identifying an etiology for anaphylaxis in
157 practice, and care in emergency departments that may not be consistent with best evidence and
158 recommendations in recent guidelines (35). An understanding of the emergency medical assistance
159 service for this patient has been set up but a standardized procedure for anaphylaxis with the
160 measure of total serum tryptase dosage would be desirable at the Emergency department. A key
161 point is the mistaken advice of not administering a second dose of epinephrine in the face of
162 progressively worsening symptoms. We take this case as an example to call for harmonized actions
163 for improved diagnosis and management of anaphylaxis.

164

165 [Anaphylaxis: unmet needs](#)

166 Since anaphylaxis entails the potential for rapidly developing life-threatening respiratory
167 and/or circulatory compromise, prompt management is imperative. Over the last decade, the
168 allergy/immunology community has intensified its efforts to encourage recognition and appropriate
169 management of patients with anaphylaxis. However, gaps in understanding and implementation of
170 management recommendations persist due to many factors including lack of a point-of-care
171 diagnostic test, limited understanding of which patient may progress to life-threatening
172 cardiopulmonary involvement, limited availability and appropriate use of first-line medications. Key
173 unmet needs are:

174 **Lack of adoption of standardized treatment protocols**

175 Although the number of publications on anaphylaxis has increased over the last decade,
176 anaphylaxis is still not consistently recognized by health care professionals and the optimal
177 management is still hampered by specific false medical beliefs. Difficulty in the recognition of
178 anaphylaxis is, in part, due to the variability of diagnostic criteria, and complicated by the
179 heterogeneity in recommendations made by different national and international guidelines. Rarity of
180 the event, multiple differential diagnoses and false medical beliefs also play a role. These factors
181 tend to perpetuate a delay in administration of appropriate treatments, increase the risk of

182 untoward outcomes from anaphylaxis, and encumber epidemiological studies of anaphylaxis since
183 medical records are the basis of national and international registries.

184 Many countries and regions have national anaphylaxis guidelines such as in France (34),
185 Europe (16), the USA (35), Australia (36) and Latin America (37). Recently, consistent efforts have
186 been made to reach a broader harmonization between these guidelines (6), but it is still necessary to
187 have a unified management system for the benefit of patients worldwide.

188 **Development of reliable markers for risk of severe/near-fatal/fatal anaphylaxis**

189 Limited comparable epidemiological studies or research to increase understanding and to
190 develop diagnostic and predictive tests remain key unmet needs. Data can differ widely depending
191 on the number of variables (38,39). The most widely discussed issues in the epidemiology of
192 anaphylaxis over the last 10 years are: (I) regional variations in concepts and definitions, (II) whether
193 prevalence or incidence is the best measure of the frequency of anaphylaxis in the general
194 population, (III) whether the frequency of anaphylaxis is higher than previously thought, and (IV)
195 whether the increasing incidence published is real or reflects different methodologies and
196 definitions used.

197 The etiology and risk factors/ co-factors for anaphylaxis described in epidemiologic studies are
198 not well characterized and may indeed be influenced by regional/national differences in allergen
199 exposures and genetic markers. In general, the most frequent triggers of anaphylaxis are drugs,
200 foods and insect venoms. The frequency varies with age. Currently, anaphylaxis phenotypes are
201 defined by clinical presentation into type-I-like reactions, cytokine storm-like reactions, and mixed
202 reactions. The endotypes underlying these phenotypes include IgE- and non-IgE-mediated
203 mechanisms, cytokine release, mixed reactions, and direct activation of immune cells (29). However,
204 further elucidation of specific underlying mechanisms of anaphylaxis is required in order to better
205 characterize anaphylaxis phenotypes and endotypes, and decrease the number of cases labelled as
206 idiopathic anaphylaxis.

207 **Frequent lack of appropriate follow-up from emergency departments**

208 Anaphylaxis can occur in every setting, and pre-hospital management plays a crucial role in
209 influencing outcomes of anaphylaxis (40-42). Therefore, patients and their caregivers have to be well
210 prepared for prompt treatment of anaphylaxis based on written emergency action plans (7).
211 Physicians, nurses and/or technicians working in ambulances should also be aware of first-line
212 anaphylaxis management protocols and align their actions accordingly.

213 Most cases of anaphylaxis are first seen by emergency department physicians or general
214 practitioners. However, only about 50% of patients are referred to allergists for further investigation
215 and/ or treatment. Recommendations for follow up and trigger avoidance at time of discharge from
216 the emergency room are provided infrequently (39). These data highlight the need for optimizing
217 and standardizing protocols for anaphylaxis management, and implementing effective education
218 and training programs. Also there should be specific programs in medical schools, residencies and
219 postgraduate training programs that include aspects of anaphylaxis and its management, as well as
220 funding for the postgraduate education of specialists.

221 **Development of a rapid point-of-care diagnostic test**

222 Although knowledge has evolved in specific areas, such as in food allergy, standardized
223 diagnostic procedures should be tailored to specific triggers, combination of manifestations and
224 specific age groups. Although standardized diagnostic procedures have been published, validation of
225 these clinical tests for all allergens does not exist and multi-center, multi-national studies are
226 needed. Generally speaking, diagnosis of allergen sensitization is made using skin tests (foods,
227 Hymenoptera venom, some drugs and aeroallergens), serum allergen-specific IgE (foods,
228 Hymenoptera venom, some drugs and aeroallergens), and provocation tests (foods, drugs) (43).
229 Other complementary tests, such as cellular allergen stimulation test (CAST) and basophil activation
230 test (BAT) and molecular diagnostic testing, are available in a number of countries, mainly for
231 research purposes (44-47).

232 Serum (or plasma) levels of total tryptase and mature tryptase measurements are
233 recommended in the diagnostic evaluation of anaphylaxis. However, the first measurement of
234 serum tryptase during the acute event is seldom performed, or in some areas of the world this
235 marker is not even available to be performed.

236 **Lack of availability, adherence and use of essential medicine for anaphylaxis**

237 Pharmacological treatment of anaphylaxis, including epinephrine, β 2 adrenergics,
238 antihistamines, corticosteroids, dopamine, glucagon, and oxygen are available in virtually all
239 countries. However, AAI are not always available in most of world countries. In countries in which
240 AAI are commercially available, national policies regarding the availability of AAI at public settings
241 are required (schools, public transport and etc).

242 Though there is no contraindication to epinephrine in the treatment of anaphylaxis and
243 intramuscular administration is recommended, subcutaneous and intravenous administration are in

244 use in 10 to 20% of cases (mainly during peri-operative anaphylaxis) and in many other cases
245 epinephrine is not even administered (34). There is also a lack of consensus regarding how long a
246 patient with anaphylaxis should be kept under observation at the healthcare setting after treatment
247 and resolution of the acute phase, especially in view of the possibility of biphasic reactions. Recently,
248 the Joint Task Force Practice parameter on anaphylaxis rated the recommendation of extended
249 clinical observation in a setting capable of managing anaphylaxis (to detect a biphasic reaction) for
250 patients with resolved severe anaphylaxis and/or those who need > 1 dose of epinephrine as having
251 very low evidence (35). This low rate score for the recommendation may reflect the lack of
252 comparable evidence-based data. The lack of research is directly due to the fleeting nature of
253 anaphylaxis and the difficulty in doing studies in humans that could induce anaphylaxis.

254

255 ACTION PLAN FROM THE WHO COLLABORATING CENTER

256 Public health's core mission is prevention of injury or disease. Taking the concept of
257 prevention levels and applying it to anaphylaxis (Table 1) facilitates understanding that the measures
258 proposed as primary and secondary preventions are addressed mostly to asymptomatic conditions,
259 in which the main concerns are identifying individuals or populations at risk (48).

260 Tertiary prevention strategies are the most familiar for physicians worldwide who are involved
261 in clinical medicine. When applied to anaphylaxis, it is intended to reduce the risk of another
262 reaction and/or manage it appropriately to avoid negative outcomes. Prevention of anaphylaxis
263 depends primarily on optimal management of patient-related risk factors, strict avoidance of
264 confirmed relevant allergens or triggers, and, where indicated, immunomodulation (e.g.,
265 Hymenoptera venom immunotherapy) (48).

266 In June 2018, the **WHO Collaborating Center (WHO CC) for the Scientific Classification of**
267 **Allergic and Hypersensitivity Diseases** was established at the University Hospital of Montpellier,
268 headed by Luciana Kase Tanno and Pascal Demoly (49). This designation is the result of recognition
269 by WHO of all the efforts of the ALLERGY in ICD-11 initiative (24,25,27, 50,51) and is intended to
270 provide academic, research and scientific support to WHO in the implementation, refinement and
271 maintenance of the WHO-FIC (Family of International Classifications) in our areas of expertise. WHO
272 CCs are institutions designated by the Director-General of the WHO and endorsed by the national
273 minister of health to carry out activities in support of the WHO programmes, such as communicable
274 diseases, nutrition, mental health, occupational health among others. Currently, there are 25 WHO
275 CCs responsible for the WHO-FIC and the Montpellier WHO CC is the only one with expertise in

276 allergy and clinical immunology. The WHO is a recognized specialized agency of the United Nations
277 concerned with international public health. Since the Montpellier WHO CC is aligned with WHO
278 actions to support the community, tailored actions for quality of care of patients, such as
279 management and prevention of anaphylaxis, are under this context.

280 As the only WHO Collaborating Center for classifications of allergic and hypersensitivity
281 conditions, we intend to establish close collaboration with national bodies in order to implement
282 actions for better patients' care, monitor and prevention, developments in research, and launch
283 measures in order to reduce avoidable deaths. Also, we intend to extend these actions
284 internationally with the support of the WHO-FIC, academic and scientific networks, the Joint Allergy
285 Academies, stakeholders and patients' organizations. Our WHO CC will provide the means through
286 which governmental and nongovernmental collaborating parties can combine their strengths to
287 achieve focused objectives, avoiding wasting of energy and resources.

288 The WHO CC can facilitate bilateral dialogue with these bodies and foster easier
289 communication with health organizations. Our aim is to use the action plan applied to anaphylaxis as
290 a model, but we may also extend this to other allergic and hypersensitivity conditions in the coming
291 period. Human and financial resources will be required and may be achieved via support from the
292 abovementioned bodies, robust research projects and structured collaborations. We intend to take
293 all the support to move on the proposed action plan. For that, structured collaborations are under
294 development.

295 Promoting increased awareness of anaphylaxis will be a key step forward. This will require
296 consistent and bilateral communication with general practitioners, emergency department
297 providers, primary care physicians, pediatricians and specialists, as well as other health
298 professionals.

299 In order to align international and national efforts for increasing awareness, collaborations
300 must be forged among professionals dealing with anaphylaxis. For this reason we propose a
301 timetable schedule in order to optimize the diagnosis and management of anaphylaxis (Figure 1). A
302 prioritized agenda should encapsulate all these steps in the frame of a global initiative against
303 anaphylaxis. Countries with different economic conditions have specific priorities and requirements.
304 The proposed action plan should support countries with different needs.

305 More than calling for harmonized action for best management of anaphylaxis to prevent
306 undue morbidity and mortality, we are proposing an action plan as a baseline for a global initiative
307 against anaphylaxis. We strongly believe these collaborative efforts are a strong public health and

308 societal priority, that is consistent with the overarching goals of providing optimal care of allergic
309 patients and best practices of allergology.

310

311 **References**

- 312 1. Portier P, Richet C. De l'action anaphylactique de certains venins. C R Séances Soc Biol 1902;54:170.
- 313 2. Dale HH, Laidlaw PP: The physiological action of beta-imidazoylethylamine. J Physiol 1910;41:318-344.
- 314 3. Ishizaka K, Ishizaka T, Hornbrook MM. Physicochemical properties of reaginic antibody. Presence of a
315 unique immunoglobulin as a carrier of reaginic activity. J Immunol 1966;97:75-85.
- 316 4. Johansson SG, Hourihane JO, Bousquet J, Brujnzeel-Koomen C, Dreborg S, Haahtela T, et al.; EAACI (the
317 European Academy of Allergology and Clinical Immunology) nomenclature task force. A revised
318 **nomenclature** for allergy. An **EAACI** position statement from the **EAACI nomenclature** task force. **Allergy**
319 2001 Sep;56(9):813-24.
- 320 5. Johansson SG, Bieber T, Dahl R, Friedmann PS, Lanier BQ, Lockey RF et al. Revised nomenclature for
321 allergy for global use: Report of the Nomenclature Review Committee of the World Allergy Organization,
322 October 2003. **J Allergy Clin Immunol**. 2004 May;113(5):832-6.
- 323 6. Simons FER, Arduzzo LR, Bilò MB, Cardona V, Ebisawa M, El-Gamal YM, et al. International consensus on
324 (ICON) Anaphylaxis. *World Allergy Organ J* 2014; 30; 7: 9.
- 325 7. Kase Tanno L, Demoly P. Anaphylaxis and pseudoanaphylaxis. *Revue de Médecine Int* 39 (2018) A2-A4.
- 326 8. Tanno LK, Demoly P; Joint Allergy Academies. Action Plan to Reach the Global Availability of Adrenaline
327 Auto-Injectors. *J Investig Allergol Clin Immunol*. 2018 Nov 13:0.
- 328 9. Tanno LK, FER Simons, Sanchez-Borges M, Cardona V, Moon HB, Calderon M, Sisul JC, Casale T, Demoly P.
329 Applying Prevention Concepts to Anaphylaxis: A Call For Worldwide Availability Of Adrenaline Auto-
330 Injectors. *Clin Exp Allergy*, 2017, in press.
- 331 10. Yamashima T (May 2003). "Jokichi Takamine (1854–1922), the samurai chemist, and his work on
332 adrenalin". *Journal of Medical Biography*. **11** (2): 95–102.
- 333 11. Bennett MR (June 1999). "One hundred years of adrenaline: the discovery of autoreceptors". *Clinical*
334 *Autonomic Research*. **9** (3): 145–59.
- 335 12. Arthur G. Epinephrine: a short story. *Lancet Respir Med*. 2015 May;3(5):350-1.
- 336 13. Golden DB. What is anaphylaxis? *Curr Opin Allergy Clin Immunol*. 2007 Aug;7(4):331-6.
- 337 14. Lieberman P. Recognition and first line treatment of anaphylaxis. *Am J Med* 2014; 127: S6-S11
- 338 15. Sampson HA, Munoz-Furlong A, Campbell RL, Adkinson NF Jr, Bock SA, Branum A et al. Second symposium
339 on the definition and management of anaphylaxis: summary report – Second National Institute of Allergy
340 and Infectious Disease/Food Allergy and Anaphylaxis Network Symposium. *J Allergy Clin Immunol*
341 2006;117:391– 397.
- 342 16. Muraro A, Roberts G, Worm M, Bilò MB, Brockow K, Fernández Rivas M et al. Anaphylaxis: guidelines from
343 the European Academy of Allergy and Clinical Immunology. *Allergy* 2014;69:1026–1045.
- 344 17. Lieberman P, Nicklas RA, Oppenheimer J, et al. The diagnosis and management of anaphylaxis practice
345 parameter: 2010 update. *J Allergy Clin Immunol* 2010; 126:477–480.
- 346 18. Simons FER, Arduzzo LRF, Bilò MB, et al. World Allergy Organization guidelines for the assessment and
347 management of anaphylaxis. *World Allergy Organ J* 2011; 4:13–37.
- 348 19. Brown SG, Mullins RJ, Gold MS. Anaphylaxis: diagnosis and management. *Med J Aust* 2006; 185:283–289.
- 349 20. Tanno LK, Alvarez-Perea A, Pouessel G. Therapeutic approach in anaphylaxis. *Curr Opin Allergy Clin*
350 *Immunol*. 2019 Aug;19(4):393-401.
- 351 21. Alvarez-Perea A, Tanno LK, Baexa ML. How to manage anaphylaxis in primary care. *Clin Transl Allergy*.
352 2017 Dec 11;7:45.
- 353 22. Panesar SS, Javad S, de Silva D, Nwaru BI, Hickstein L, Muraro A, Roberts G, Worm M, Bilò MB, Cardona V,
354 Dubois AEJ, Dunn Galvin A, Eigenmann P, Fernandez-Rivas M, Halken S, Lack G, Niggemann B, Santos AF,
355 Vlieg-Boerstra BJ, Zolkipli ZQ & Sheikh A on behalf of the EAACI Food Allergy and Anaphylaxis Group. The
356 epidemiology of anaphylaxis in Europe: a systematic review. *Allergy* 2013; 68: 1353–1361.
- 357 23. Neugut AI, Ghatak AT, Miller RL. Anaphylaxis in United States: an investigation into its epidemiology. *Arch*
358 *Intern Med*. 2001 Jan 8;161(1):15-21.
- 359 24. Tanno LK, Calderon MA, Demoly P; on behalf the Joint Allergy Academies. New allergic and
360 hypersensitivity conditions section in the International Classification of Diseases-11. *Allergy Asthma*
361 *Immunol Res* 2016;8:383-8.
- 362 25. Tanno LK, Chalmers R, Bierrenbach AL, Simons FER, Molinari N, Annesi-Maesano et al. on behalf Joint
363 Allergy Academies. Changing the history of anaphylaxis mortality statistics through the World Health
364 Organization's International Classification of Diseases (ICD)-11. *J Allergy Clin Immunol*. 2019
365 Sep;144(3):627-633.

- 366 26. World Health Organization, ICD-11 Beta Draft website. (cited, available:
367 <http://apps.who.int/classifications/icd11/browse/l-m/en> January 2020).
- 368 27. Tanno LK, Chalmers R, Jacob R, Kostanjsek N, Bierrenbach AL, et al. Global Implementation of the World
369 Health Organization's International Classification of Diseases (ICD)-11: The Allergic and Hypersensitivity
370 Conditions Model. *J Allergy Clin Immunol*. 2020, in press.
- 371 28. Turner PJ, Worm M, Ansotegui IJ, El-Gamal Y, Rivas MF, Fineman S, et al.; WAO Anaphylaxis Committee.
372 Time to revisit the definition and clinical criteria for anaphylaxis? *World Allergy Organ J*. 2019 Oct
373 31;12(10):100066.
- 374 29. Castells M. Diagnosis and management of anaphylaxis in precision medicine. *J Allergy Clin Immunol*. 2017
375 Aug;140(2):321-333.
- 376 30. Claire Galand, Juan Manuel Leyva-Castillo, Juhan Yoon, Alex Han, Margaret S. Lee, Andrew McKenzie, et al.
377 IL-33 promotes food anaphylaxis in epicutaneously-sensitized mice by targeting mast cells. *J Allergy Clin
378 Immunol*. 2016 Nov; 138(5): 1356–1366.
- 379 31. Francis A, Bosio E, Stone SF, Fatovich DM, Arendts G, Nagree Y, et al. Neutrophil activation during acute
380 human anaphylaxis: analysis of MPO and sCD62L. *Clin Exp Allergy*. 2017 Mar;47(3):361-370.
- 381 32. Kasperska-Zajac A, Rogala B. Platelet function in anaphylaxis. *J Investig Allergol Clin Immunol*. 2006;16:1–
382 4.
- 383 33. *Reber LL, Hernandez JD, Galli SJ. The pathophysiology of anaphylaxis. J Allergy Clin Immunol*. 2017
384 Aug;140(2):335-348.
- 385 34. Gloaguen A, Cesareo E, Vaux J, Valdenaire G, Ganansia O, Renolleau S., et al. Management of Anaphylaxis
386 in Emergency Medicine. French Society of Emergency Medicine (SFMU) Guidelines with the Contribution
387 of French Allergology Society (SFA) and the French Speaking Group in Pediatric Intensive Care and
388 Emergency (GFRUP), and the support of the French pediatric pneumology and allergology society (SP2A).
389 *Ann. Fr. Med. Urgence* (2016) 6:342-364.
- 390 35. Shaker MS, Wallace DV, Golden DBK, Oppenheimer J, Bernstein JA, Campbell RL et al. Anaphylaxis-a 2020
391 practice parameter update, systematic review, and Grading of Recommendations, Assessment,
392 Development and Evaluation (GRADE) analysis. *J Allergy Clin Immunol*. 2020 Apr;145(4):1082-1123.
- 393 36. Vale S, Netting MJ, Ford LS, Tyquin B, McWilliam V, Campbell DE. Anaphylaxis management in Australian
394 schools: Review of guidelines and adrenaline autoinjector use. *J Paediatr Child Health*. 2019
- 395 37. Cardona V, Álvarez-Perea A, Ansotegui-Zubeldia IJ, Arias-Cruz A, Ivancevich JC, González-Díaz SN, et al.
396 Clinical Practice Guide for Anaphylaxis in Latin America (Galaxia-Latam). *Rev Alerg Mex*. 2019;66 Suppl
397 2:1-39.
- 398 38. Tejedor Alonso MA, Moro Moro M, Múgica García M V. Epidemiology of anaphylaxis. *Clin Exp Allergy*
399 2015;45:1027–1039.
- 400 39. Tanno LK, Bierrenbach AL, Simons FER, Cardona V, Thong BY, Molinari N, et al.; on behalf the Joint Allergy
401 Academies. Critical view of anaphylaxis epidemiology: open questions and new perspectives. *Allergy
402 Asthma Clin Immunol*. 2018 Apr 4;14:12.
- 403 40. Pouessel G, Dumond P, Liabeuf V, Kase Tanno L, Deschildre A, Beaumont P, et al. Gaps in the management
404 of food-induced anaphylaxis reactions at school. *Pediatr Allergy Immunol*. 2019 Nov;30(7):767-770.
- 405 41. Lejeune S, Deschildre A, Beaudouin E, Labreuche J, Meininger C, Lefort H, et al. Pre-hospital management
406 of paediatric anaphylaxis by French Emergency Medicine physicians: Still to be improved. *Clin Exp Allergy*.
407 2019 Jul;49(7):1047-1050.
- 408 42. Harper NJ, Cook TM, Garcez T, Farmer L, Floss K, Marinho S, Torevell H, Warner A, Ferguson K, Hitchman J,
409 Egner W. Anaesthesia, surgery, and life-threatening allergic reactions: epidemiology and clinical features
410 of perioperative anaphylaxis in the 6th National Audit Project (NAP6). *British journal of anaesthesia*. 2018
411 Jul 1;121(1):159-71.
- 412 43. Tanno LK et al. Management of Anaphylaxis: the World Allergy Organization Survey Identifies
413 Achievements and Remaining Challenges. *World Allergy Organ J*. 2020, in press
- 414 44. Sala-Cunill A, Björkqvist J, Senter R, Guilarte M, Cardona V, Labrador M et al. Plasma contact system
415 activation drives anaphylaxis in severe mast cell-mediated allergic reactions. *J Allergy Clin Immunol*
416 2015;135:1031-1043.e6.
- 417 45. Khodoun M V, Strait R, Armstrong L, Yanase N, Finkelman FD. Identification of markers that distinguish
418 IgE- from IgG-mediated anaphylaxis. *Proc Natl Acad Sci U S A* 2011;108:12413–12418.
- 419 46. Ansotegui IJ, Melioli G, Canonica GW, Caraballo L, Villa E, Ebisawa M, et al. IgE allergy diagnostics and
420 other relevant tests in allergy, a World Allergy Organization position paper. *World Allergy Organ J*. 2020
421 Feb 25;13(2):100080.
- 422 47. Ansotegui IJ, Melioli G, Canonica GW, Caraballo L, Gomez RM, Ebisawa M, et al. A WAO-ARIA-GA2LEN
423 consensus document on molecular-based allergy diagnosis (PAMD@): Update 2020 *World Allergy Organ J*.
424 2020 Mar 7;13(2):100091.
- 425

- 426 48. Tanno LK, Simons FER, Sanchez-Borges M, Cardona V, Moon HB, Calderon MA, Sisul JC, Muraro A, Casale
427 T, Demoly P; Joint Allergy Academies. Applying prevention concepts to anaphylaxis: A call for worldwide
428 availability of adrenaline auto-injectors. Clin Exp Allergy. 2017 Sep;47(9):1108-1114.
- 429 49. World Health Organization, Collaborating Centres list website. (cited, available:
430 [http://apps.who.int/whocc/Detail.aspx?cc_ref=FRA-](http://apps.who.int/whocc/Detail.aspx?cc_ref=FRA-133&designation_date1=1/6/2018&designation_date2=18/7/2018&)
431 [133&designation_date1=1/6/2018&designation_date2=18/7/2018&](http://apps.who.int/whocc/Detail.aspx?cc_ref=FRA-133&designation_date1=1/6/2018&designation_date2=18/7/2018&) December 2019)
- 432 50. Tanno LK, Ganem F, Demoly P, Toscano CM, Bierrenbach AL. Undernotification of anaphylaxis deaths in
433 Brazil due to difficult coding under the ICD-10. Allergy 2012; 67: 783–789.
- 434 51. Tanno LK, Calderon MA, Goldberg BJ, Gayraud J, Bircher AJ, Casale T et al. Constructing a classification of
435 hypersensitivity/allergic diseases for ICD-11 by crowdsourcing the allergist community. Allergy 2015; 70:
436 609-15.

437

438

439

440 Figure 1: Timetable action plan proposal for the optimization of diagnosis and management of anaphylaxis

Table 1: Prevention concepts from a public health perspective, applied to anaphylaxis and intervention actions (adapted from 35)

LEVELS OF PREVENTION	APPLYING PREVENTION CONCEPTS TO ANAPHYLAXIS	INTERVENTIONS AND ACTIONS APPLIED TO ANAPHYLAXIS
Primary Prevention	<p>Primary prevention addressed to anaphylaxis would imply the identification of individuals or populations at risk in order to avoid sensitization. For this, specific risk factors should be identified in unsensitized individuals with no history of an anaphylactic reaction.</p>	<ul style="list-style-type: none"> – Increase health professionals’ awareness through education and continuing education programs (e.g., breastfeeding, latex avoidance, early food diversification for infants). – Support dissemination of accurate information to the public (e.g., EAACI Anaphylaxis campaign, WAO Allergy week). – Specific interventions with early introduction of specific foods in the infant diets (e.g., peanut). – Remove strong sensitizers from public places and workplace environments (e.g., remove powdered latex gloves to prevent occupational latex allergy/ anaphylaxis, remove OTC use of pholcodine to prevent neuromuscular blocking agent anaphylaxis). – Labelling industrial products – Availability of adrenaline auto-injectors in countries in which it is not available in the health system – Availability of adrenaline auto-injectors in public settings (schools, public transport and etc..) – Strengthen standard international definitions, notification, classification and coding (e.g., International Classification of Diseases) to support monitoring morbidity and mortality

<p>Secondary Prevention</p>	<p>Secondary prevention in the context of anaphylaxis includes sensitized individuals with no history of an anaphylactic reaction. The aim is to prevent the development of an allergic disorder in patients previously sensitized. Screening of the general population for sensitization is not recommended. Sensitization is common and does not imply the diagnosis of an allergic disease. Screening should be applied to individuals with known risk factors.</p>	<ul style="list-style-type: none"> – Individualized screening in order to identify sensitized individuals and support specific measures (e.g., those with occupational latex sensitization). – Individualized indication of adrenaline auto-injectors – Increase health professionals’ awareness through education and regular information programs. – Strengthen standard international definitions, notification, classification and coding (e.g., International Classification of Diseases) to support monitoring morbidity and mortality.
<p>Tertiary Prevention</p>	<p>Tertiary prevention: This concept should be focused on patients who have experienced an anaphylactic reaction. After initial clinical presentation, it is intended to reduce the risk of another reaction and/or manage it appropriately and avoid negative outcomes.</p>	<ul style="list-style-type: none"> – Complete allergological work-up to confirm triggers (inducers) and support specific immunomodulation (e.g., allergen immunotherapy for Hymenoptera venom anaphylaxis or full drug allergy work up as indicated) and provide a written documentation of the diagnosis and the confirmed triggers/agents. – Individualize patient’s education and provide specific information: environmental or behavior modifications to reduce patient’s exposure to allergens, provide a written anaphylaxis emergency action plan. Adrenaline auto-injectors (individualized indication) – Support accurate food allergen labelling to protect consumers. – Support the emergency training of health professionals to rapidly identify and manage anaphylaxis. – Correct notification of new cases (e.g., as new allergens arise,

		<p>support large cohort analysis).</p> <ul style="list-style-type: none">– Increase health professionals' awareness through education and continuous information programs.– Strengthen standard international definitions, notification, classification and coding (e.g., International Classification of Diseases) to support monitoring morbidity and mortality.
--	--	---

OTC, over-the-counter

Figure E1: Adrenaline auto-injectors commercially available in France (2018) and their differences (8).

HOME/ PUBLIC SETTING

EMERGENCY TRANSPORT

EMERGENCY DEPARTMENT/ HOSPITAL/ PRIMARY CARE

SPECIALIST (ALLERGIST)

Strengthen standard international definitions, notification, classification and coding (e.g., International Classification of Diseases) to support monitoring morbidity and mortality.

Education of medical students, residents, nurses and other key health care personal should be key to any improvement in the recognition of anaphylaxis and its management

- Identification of symptoms
- Prompt treatment and call the emergency following the written anaphylaxis action plan
- Availability of adrenaline auto-injectors in public settings (schools, public transport and etc..)
- Support dissemination of accurate information to the public (e.g., EAACI Anaphylaxis campaign, WAO Allergy week).
- Specific interventions with early introduction of specific foods in the infant diets (e.g., peanut).
- Remove strong sensitizers from public places and workplace environments (e.g., remove powdered latex gloves to prevent occupational latex allergy/ anaphylaxis, remove OTC use of pholcodine to prevent neuromuscular blocking agent anaphylaxis).
- Labelling industrial products
- Individualize patient's education and provide specific information: environmental or behavior modifications to reduce patient's exposure to allergens, provide a written anaphylaxis emergency action plan. Adrenaline auto-injectors (individualized indication)

- Identification of symptoms
- Prompt treatment and call the emergency following the written anaphylaxis action plan
- Availability of adrenaline and/or adrenaline auto-injectors
- First measure of total serum tryptase measure
- Support the emergency training of health professionals to rapidly identify and manage anaphylaxis.
- Increase health professionals' awareness through education and continuous information programs.

- Identification of symptoms
- Prompt treatment and call the emergency following the written anaphylaxis action plan
- Availability of adrenaline and/or adrenaline auto-injectors
- First measure of total serum tryptase (if the blood sample has not been collected previously and if among first 2 hours of the symptoms)
- Individualized indication of adrenaline auto-injectors and avoidance of potential elicitors
- Refer to the allergist for further investigation
- Support the emergency training of health professionals to rapidly identify and manage anaphylaxis.
- Increase health professionals' awareness through education and continuous information programs.
- Correct notification of new cases.

- Individualized screening in order to identify sensitized individuals and support specific measures (e.g., those with occupational latex sensitization).
- Individualized indication of adrenaline auto-injectors
- Second measure of total serum tryptase
- Complete allergological work-up to confirm triggers (inducers) and support specific immunomodulation (e.g., allergen immunotherapy for Hymenoptera venom anaphylaxis or full drug allergy work up as indicated) and provide a written documentation of the diagnosis and the confirmed triggers/agents.
- Individualize patient's education and provide specific information: environmental or behavior modifications to reduce patient's exposure to allergens, provide a written anaphylaxis emergency action plan. Adrenaline auto-injectors (individualized indication)
- Increase health professionals' awareness through education and continuous information programs.
- Correct notification of new cases (e.g., as new allergens arise, support large cohort analysis).