



Standing Group of Experts (SGE) on African swine fever (ASF)
Global Framework for the progressive control of Transboundary Animal Diseases (GF-TADs)
Africa chapter.

The Declaration of Lomé on the use of vaccines against African swine fever (ASF) in Africa.

As a result of the fifth meeting of the *Standing Group of Experts* (SGE) on *African swine fever* (ASF) of the *Global Framework for the progressive control of Transboundary Animal Disease*'s Africa chapter (GF-TADs for Africa), dedicated to vaccines and vaccination, and held in Lomé, Togo on 14 October 2025, the Members of the ASF SGE have deliberated and agreed the following:

Recognising that :

- a) There are currently no ASF vaccines that fully meet the WOAH international standards for safety, efficacy, purity, and prior licensing.
- b) DIVA (*Differentiating Infected from Vaccinated Animals*) compatible and next-generation vaccines are essential for traceability, safety and long-term protection. DIVA capability is not available in any of the currently licensed vaccines, complicating disease surveillance and control post-vaccination.
- c) Any future use of the vaccine candidate should be based on a thorough risk benefit assessment considering all safety and efficacy features as well as the vaccination scenarios envisaged.
- d) Vaccination effectiveness should be evaluated based on performance in real-field settings and preferably with involvement of independent technical institutions.
- e) *Live attenuated vaccines* (LAVs) are currently the most promising ASF vaccine candidates, offering protection against homologous strains, but exhibiting very limited cross-protection against genetically distinct variants.
- f) All currently licensed vaccines are based on genotype II, whereas the Africa region hosts 24 distinct genotypes.
- g) Reversion to virulence and recombination risks are real and documented and may have a continental and global impact. Field monitoring and genomic surveillance in line with international standards are critical to detect and mitigate these risks.

The Members of the ASF SGE strongly encourage :

a) Member(s) Countries/States in Africa

▪ as of now (October 2025):

- to ensure appropriate evaluation of any new vaccine through AU-PANVAC, based on WOAH standards before granting any (marketing) authorization for its use or distribution.
- to be vigilant and not to import ASF vaccines, not conduct vaccinations as currently available commercial vaccines are not safe and not effective against the genotypes circulating in the region.

▪ as and when ASF vaccines are made available:

- Member(s) Countries/States should monitor for, and ensure that only vaccines registered for use, and that meet WOAH standards are used within their territories.
- Member(s) Countries/States to ensure mechanisms in place for field evaluation and vaccination monitoring based on guidance provided by AU-IBAR, AU-PANVAC, FAO and WOAH and should refrain from such evaluations until such regional and international guidance is available.
- Member(s) Countries/States note that vaccination can be counterproductive in settings with poor biosecurity (which increases the risk of virus circulation and hampers data reliability) and in lack of untraceable pig populations
- Member(s) Countries/States should recognise that vaccination strategies complement, and must not replace, strong biosecurity and other complementary ASF control measures. Implementation requires controlled vaccination areas and strict Veterinary Authority supervision.
- Veterinary authorities of Member(s) Countries/States should oversee vaccinations, based on knowledge of ASF disease spread and ASFV sequencing.
- Member(s) Countries/States implement well defined official control programmes rather than voluntary vaccinations without monitoring by the authorities.

b) AU-PANVAC

- to work with Member(s) Countries/States to confirm there is not fraudulently or unauthorized ASF vaccines circulating within countries.

c) AU-IBAR, AU-PANVAC, FAO, and WOAH

- to continue supporting research on ASF vaccine development and or validation in Africa.
- to continue sensitizing African Member(s) Countries/States on risks with importation of untested vaccines of unknown quality into Africa.

d) GARA Africa Chapter (GAC)

- to catalog ASF genotypes across Africa to guide vaccine development and selection.

e) ILRI

- to continue research on their genotype IX ASF vaccine candidate and conduct cross-protection studies against other genotypes.

Adopted and endorsed on 14 October 2025 in Lomé, Togo

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