

1 **Analysis of spatial and temporal risk of Peste des Petits Ruminants Virus (PPRV) outbreaks in**
2 **endemic settings: A scoping review**

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20 **Abstract**

21 Surveillance shows that Peste des petits ruminants (PPR) is endemic in both Africa and Asia due to its
22 continuous circulation. Several epidemiological factors work together to support PPR's geographical
23 spread. To investigate the risk of PPR transmission, analytical techniques based on spatial, spatiotemporal,
24 and transmission dynamics have been employed. The risk factors linked to the spatiotemporal distribution

20 and transmission dynamics of PPR at the regional level are extremely poorly understood. This study
26 assessed the risks of Peste des Petit ruminant virus (PPRV) epidemics during a comprehensive evaluation
27 of peer-reviewed literature, highlighting the differences between geographical and spatial-temporal
28 techniques used in endemic zones. Utilizing the PubMed and Google Scholar databases, a scoping
29 literature analysis of PPR research papers that evaluated PPR risks in endemic areas using spatial and
30 spatiotemporal techniques was conducted. Eight papers with a global perspective were chosen from 42,
31 20 of which were on Asia, and 14 on Africa. 35.7% employed spatial autocorrelation, while 61.9% used
32 clustering analysis. Of the research, the majority (71.2%) described temporal trends, whereas 13
33 publications (30%) used modelling methodologies. Geographic accessibility (n = 19), trade and commerce
34 (n = 17), environment and ecology (n = 12), socioeconomic variables (n = 9), and demography and
35 livestock–wildlife interactions (n = 20) are the five risk factors that were assessed. All the risk factors were
36 related, however only two papers discussed the transmission dynamics of PPR. Our understanding of PPR
37 outbreaks in endemic environments has improved because of the review, which also encourages
38 evidence-based decision-making to lessen the virus's effects on small ruminant populations. It has been
39 demonstrated that the association of additional risk variables with livestock trade, the primary force
40 behind livestock migration, significantly increases the probability of PPR outbreaks in endemic areas.
41 Since many studies are conducted in Asia rather than Africa, Africa should be considered in future
42 prediction model development to evaluate possible eradication strategies at the national and regional
43 levels.

44
45 **Keywords:** Peste des petit ruminant (PPR) epidemics, spatial methods, spatiotemporal methods, risk
46 factors

47

48 **1. Context**

49 A virus known as peste des petits ruminants (PPR) affects small ruminants, primarily sheep and goats, but
50 it can also infect other domestic animals (1). The PPR virus (PPRV) is a single-strand, non-segmented RNA
51 virus of the genus Morbillivirus in the family *Paramyxoviridae* (2). PPRV's genome spans 15,948
52 nucleotides (nt) and is structured into six open reading frames (ORFs). The six structural proteins that are
53 encoded by these ORFs are the polymerase (P) or large protein (L), fusion protein (F), phosphoprotein (P),
54 matrix protein (M), hemagglutinin protein (H), and nucleoprotein (N). Additionally, the non-structural
55 proteins C and V are encoded by the ORF transcription unit (3). Four lineages have been described from
56 two structural proteins N or F by phylogenetic studies using partial gene sequences (3). These lineages of
57 PPRV are distributed in several geographical areas including Africa, Asia and Europe (4). All four PPRV
58 lineages are present in Africa, where lineage I viruses have been confined in West African countries since
59 1940. Lineage II is predominantly present in West Africa, although it has recently been reported in the
60 Democratic Republic of the Congo (DRC) and Tanzania (5). Although Lineage III is prevalent throughout
61 northeastern, eastern, and central Africa as well as the Comoros islands, it has not been seen in the north
62 or west of the continent (5). To date, PPR has been identified in the northern, western, central and eastern
63 regions of Africa and is gradually moving southwards. Hundreds of millions of domestic small ruminants
64 and wildlife are at risk of infection as the PPRV continues to spread across previously uninfected regions
65 (6). However, the PPRV infection that has been found in previously uninfected areas and the admixture
66 of lineages in countries that have been infected jointly emphasize the geographically and temporally
67 dynamic character of PPR (7). With annual global economic losses estimated at approximately \$1.45 and
68 2.1 billion USD, half of these losses impact Africa, and a quarter affect Asia. Like other livestock respiratory
69 diseases losses are caused by mortality, which reaches up to 20% in naive population and morbidity, which
70 reaches up to 100% (8,9). A global program to eradicate PPR by 2030 has been formally launched by the
71 Food and Agriculture Organization (FAO) and the World Organization for Animal Health (WOAH), formerly

known as OIE), due to the high impact PPR on sheep and goat farmers (10). Global PPR eradication campaigns adopt spatial and spatiotemporal risk analysis models for disease prevention, focusing on PPR risk factors and disease transmission dynamics in livestock contacts (11). Network analysis, prediction, and simulation models can be used to identify PPR clusters and their drivers, providing crucial information for disease investigation (12). Enormous availability of such information could support the development of locally adapted control and surveillance strategies (13). Combining such risk factor information with genetic and mobility network data may make it possible to identify disease hot spots, which are crucial for virus entry and dissemination to various regions (13). Identification of PPR hotspots will involve the use of spatial and Spatiotemporal tools to analyze PPR epidemic data. Such analysis will also explore the patterns and risk factors of the disease (14). Nevertheless, a variety of spatiotemporal tools are accessible for evaluating a range of spatiotemporal hypotheses to meet distinct goals. Either in testing a hypothesis, the outcomes of such a study can be directly compared if the analytical methods used are well understood (15). For instance, studies examining how local elements like topography, socioeconomics, demography, and environment can impact disease reporting, identification, and circulation changes over time and space (16). The important step in choosing a model's parameters is to consider the disease pathway that is thought to connect epidemiological factors with epidemics (15). In this aspect, disease transmission dynamics are contextualized to a specific location where the contact network is factored in disease diffusion to various areas. The need for this review remains critical because of the overall knowledge gap on the effectiveness of these tools in the analysis of PPR control in endemic situations (16). To our knowledge, no studies have reviewed the spatial and spatio-temporal methodologies used in PPR research. A narrative review by (17) summarizes the occurrence and distribution of PPR in Tanzania. In addition to estimating the prevalence of PPR in sheep and goats, systematic review publications assess probable contributing factors to the disease's heterogeneity in prevalence and distribution (4). The only Scoping review conducted by (18) was on PPR diagnostic platforms. Prior reviews did not discuss models,

96 variables, spatial, and spatiotemporal methodological frameworks for PPR risks, leading to a high lack of
97 information on risks-based spatial and spatiotemporal analysis (19). By filling this gap, PPR can be
98 controlled and eventually eradicated. This study evaluates spatial and spatial-temporal methodological
99 approaches in Peste des petit ruminant (PPR) epidemics, aiming to identify priority research areas, suggest
100 interventions, and standardize modelling inputs for improved comparability.

101

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Preprint

1.03 2. Data acquisition

1.04 2.1. Protocol

1.05 In compliance with the Preferred Reporting Items for Systematic Reviews and Meta-Analyses extension
1.06 for Scoping Reviews (PRISMA-ScR) standards, a scoping literature review was carried out. (Sup. Table 1),
1.07 in which a checklist for scoping reviews was adopted (20). The approach suggested by Arksey & O'Malley
1.08 was followed in this study, which included formulating the research question, finding pertinent sources,
1.09 choosing references, charting the data, compiling, summarizing, and reporting the findings (20) (Figure 1).

1.10

1.11 2.2. Research Questions Identification

1.12 The research questions listed below were the main focus of the scoping review.:

- 1.13 (a) What techniques have been applied to investigate PPRV's spatiotemporal and spatial distribution
1.14 in endemic environments?
- 1.15 (b) Based on the examination of those occurrences, what is known regarding PPR risk from the
1.16 literature (e.g., Finding clusters, hotspots, or seasonal or temporal tendencies)?
- 1.17 (c) Which risk variables have been examined and linked to PPR prevalence in endemic areas in Asia
1.18 and Africa?
- 1.19 (d) How do the risk factors under investigation affect the dynamics of PPRV transmission?

1.20 2.3. Search strategy and selection criteria

1.21 Spatial and spatial-temporal analysis served as the basis for a scoping evaluation of the PPRV threats. Risk
1.22 factors in endemic areas were discovered with the assistance of a library professional for electronic
1.23 bibliographic search, and the search strategy comprised a set of keywords on spatial and spatiotemporal
1.24 techniques. Two computerized bibliographic databases were systematically searched to find peer-
1.25 reviewed original publications published in English-language journals between January 1993 and June

126 2024. The peer-reviewed publications that were part of this review were found using the Google Scholar
127 and MEDLINE (PubMed) search engines. Boolean operators were used to select the papers based on
128 phrases like "pest des petit ruminant virus (PPRV) AND risks AND spatial and spatial-temporal analysis
129 AND epidemics AND endemic countries." In order to get results, essential terms were trimmed down
130 during the PubMed search. 766 documents were found overall through the searches of the two electronic
131 databases (Google Scholar: 648, and PubMed: 118, as shown in Figure 2).

132 **2.4. Eligibility criteria**

133 PPR research using spatial, temporal, or spatiotemporal techniques for data analysis and inference was
134 considered in this scoping review. Studies on various spatiotemporal analytical tools can be employed for
135 epidemiological research based on the classification put forward by (21). These can be categorized
136 according to the analysis's goals for (a) description and visualization, (b) pattern identification and
137 geographical or spatiotemporal dependence, (c) spatial smoothing and interpolation, and (d) modelling
138 and regression research. We were especially interested in research that included putative PPR risk factors
139 to forecast illness occurrence or that looked at epidemiological parameters associated with PPR outbreaks
140 at the population level within the modelling and regression category. All publications with original peer-
141 reviewed articles were chosen in order to meet the inclusion requirements. Based on data from the most
142 recent WOAHP list of members and areas designated as PPR-free (Resolution No. 17 (89th General Session,
143 May 2022), PPR endemicity was determined. However, research carried out in nations with recognized
144 free zones or PPR-free certifications, like Russia, was eligible for inclusion (22). Consequently, research
145 that (a) used data from population outbreaks in PPRV-endemic nations; (b) reported patterns or
146 distributions of PPR epidemics; or (c) used data that was geographically or temporally connected to model
147 PPR risk; (c) were published in an English-language peer-reviewed scientific publication between January
148 1990 and July 2024; and (d) confirmed cases of PPR in wildlife and livestock (such as cattle, small
149 ruminants, and camels) were taken into account for inclusion.

100 **Exclusion criteria**

101 Excluded were studies that used data from farm epidemic surveys that included risk factors and self-
102 reported PPR occurrences. Furthermore, this synthesis excluded studies that were meant to be risk
103 assessments, epidemic investigations, or narrative literature reviews that reported PPR occurrence and
104 trends (without further data analysis). Studies from nations that are known to be PPR-free, with or
105 without immunization, were not included in the review, as Figure 1 illustrates.

106

107 **2.5. Selection of relevant and reliable studies**

108 To balance sensitivity and specificity in research, a search method was created utilizing electronic
109 databases, Boolean, and proximity operators. Titles and abstracts were used to filter reports, removing
110 duplicates and irrelevant papers. Reasons for exclusion were noted, and full texts were obtained for
111 inclusion. Articles were examined for selection by two reviewers with expertise in molecular
112 epidemiology and PPR surveillance. Conflicts that arose during the screening process were settled.

113 Figure 1: PPR status depending on the acknowledgement of WOAHP PPR. The grey areas are either
114 unregistered PPR status, endemic, or reported occasional incidents.

115

116 **2.6. Data extraction from included studies**

117 Every study's data was extracted by a single reviewer using a pre-made, standardized form in a Microsoft
118 Excel Spreadsheet that contained the following:

119 (a) Study attributes include author, publication year, years examined, nation, geographic reach, and
120 species.

121 (b) PPR epidemic information and diagnosis, including the surveillance system, data source, and
122 diagnostic criteria; (c) The type of analytical tool used, the process for grouping and identifying patterns
123 in the data, the method for assessing the data based on time and season, and the list of epidemiological
124 factors that were investigated and their outcomes. JJM and JNH extracted and validated data from papers,

190 analyzing them for accuracy and quality appraisal. Spatial distribution, cattle commerce, weather, climate
191 change, and species/age/sex were among the topics found through thematic analysis.

192 **2.7. Collating, summarizing and reporting the findings**

193 The sizes and data formats suggested by (15) were used to categorise the methods for formally evaluating
194 the existence of clusters. As previously mentioned by (21) the framework was adapted to categorise
195 analytical tools for spatial or spatial-temporal analysis according to their function. Using five primary
196 categories, epidemiological models examine the variables that affect PPRV introduction, transmission,
197 survival, and the efficacy of control strategies. spatial accessibility; (b) the demography of livestock and
198 the interactions between livestock and wildlife; (c) the trade in livestock; (d) socioeconomic
199 considerations; and (e) ecology and environment. The term "other factors" was used to group
200 epidemiological covariates that did not fall into any of the categories (Sup. Table 2) and Figure 4). By
201 merging essential elements and presenting data narratively, the review investigates the relationship
202 between epidemiological determinants and PPR epidemics. R version 4.3.3 (*ggplot2*, *webr*, *tidyr*, and
203 *dplyr*) was used for the analyses (23).

204 **3. Results**

205 **3.1. General characteristics of the selected studies**

206 A total of 42 studies for the quantitative synthesis and 57 papers for the qualitative synthesis were
207 included using the electronic search approach. Out of the original pool of approximately 766
208 publications that were screened, 101 were examined in full text, and 57 of those were removed due to
209 their inability to meet the eligibility requirements. requirements (Figure 2).

210
211
212 Figure 2: Flowchart showing the study records' bibliographic search and eventual inclusion in the review

213

198 **3.2. Time intervals and geographic regions**

199 In Africa and Asia, the distribution of studies by geographic scale and scope differed (Figure 3). Included
200 research in Figure 3 spans 21 nations spread across two continents (Africa and Asia). Most research was
201 carried out on a national level using PPR pandemic data from official sources (95%), which covered the
202 entire country (45%). PPR epidemics were frequently identified based solely on clinical presentation in
203 most studies, where data were gathered as part of passive monitoring systems (37.5%) (Sup. Table 2).
204 Studies differed in the cattle species for which outbreaks were documented and the spatial unit employed
205 for analysis (Sup. Table 2). Epidemic reports from sheep and goats were included in certain research (50%,
206 n = 20). Nevertheless, PPR epidemics in other species categories were examined in 2.5–5% of the
207 investigations that were documented. Several studies looked at long-term epidemic data, the longest of
208 which was a 44-year PPR outbreak case series from India (24). Eleven studies looked at epidemic data
209 collected during a ten-year period, however, the length of each study varied (median = 3 years; range: 1–
210 44). Although a number of papers included a range of analytical tools and objectives, the majority of
211 studies concentrated on the description and visualization of epidemics.

212

213 Figure 3: Geographical distribution of the papers that were part of the scoping review.

214

215 Figure 4: Classification of risk variables for outbreaks of Peste des petits ruminants using conceptual
216 frameworks.

217

218 **3.3. Methods used to identify spatial and temporal variations of PPRV risk factors**

219 **Error! Reference source not found.** shows various spatial, temporal and Spatiotemporal methods that
220 were used to visualize patterns, explore spatial clusters, and model risks across space and time.
221 Although the results of some studies suggested the use of these techniques, they did not state clearly
222 how useful they were. Over half of the studies used spatial clustering analysis techniques to test the

223 non-randomness hypothesis of PPR epidemic distribution. 35.7% and 26.2% used spatial or spatial-
224 temporal tools. Four studies used various approaches for identifying clusters in parallel, including
225 Moran's I, Maximum entropy spatial statistic, Getis-Ord, and Clark Evans test. SaTScan was used for
226 simultaneous cluster detection (11). In three studies, the direction of PPR epidemic progression was
227 determined using spatiotemporal directionality tests.

228 Figure 5: Dimensions and data forms for classification of cluster analysis modified from
229

230 **3.4. Spatial autocorrelation or spatial clustering**

231 Of the investigations, 35.7% (n = 15) showed evidence of using PPR spatial autocorrelation approaches for
232 illness cluster identification. However, because different methods were employed to find these clusters
233 (unusual aggregation of epidemics) and hotspots (excess level of epidemics compared to a threshold
234 level), the data configuration on these findings tended to differ both within and between studies (16).
235 Heterogeneous results were reported by four research, which noted the discovery of random or clustered
236 patterns that differed depending on the time period or analytical technique. For instance, (25–27) found
237 that the identification of spatial association was impacted by annual change. The subsequent two
238 investigations (11,28) found random spatial patterns and clustering tendencies that changed over time,
239 as well as the different clustering techniques applied in the research. Sup. Table 5 summarizes the cluster
240 evaluation techniques and overall results of the assessed research. Possibly as a result of local disease
241 status, tool modifications, hypothesis testing, and assumptions made, cluster size, as indicated by
242 significant radius, varies greatly between research.

243 **3.5. Temporal and seasonal trends assessment**

244 Depending on the underlying data's temporal aggregation, the majority of research (71.2%, n = 30)
245 detailed the temporal patterns of PPR outbreaks aggregated each day, week, month, or year. The model
246 known as the Generalized Linear Negative Binomial Regression, (11), Generalized Linear Mixed Models
247 (GLMMs) (29), Negative Binomial (27), linear (13,24,30,31) or logistic regression models (32) Were used

248 to explore or test hypotheses related to temporal trends. Other studies resorted to Bayesian approaches
249 (2,3,33). The NAADSM (North American Animal Disease Spread Model (12,34), autoregressive integrated
250 moving average model (ARIMA) (35), Least Cost Path (LCP) (36), Random forest (26) Ensemble Algorithm
251 (37), Event-Driven Memoryless model of state transitions (38), Regression Tree Models (26) and Mantel
252 Correlograms (13), which were employed to investigate the relationship between genetic distances and
253 various measures of network and geographic distance. Moreover, 4 studies formally analyzed PPR
254 seasonality through the assessment of seasonal trends distributed geographically and socioeconomically
255 (1,22,24,25).

256

257 **3.6. Modelling approaches**

258 A variety of geographical and spatial-temporal tools were used to evaluate the relationship between a
259 number of epidemiological parameters and PPR epidemics, as well as to describe the patterns of PPR
260 epidemic distribution and identify disease hotspots using clustering approaches. Thirteen (30.9%) of the
261 research in this study used covariables to forecast PPR risk, create risk probability maps, or investigate
262 the relationship between population-level epidemiological factors and the outbreaks. Eleven research
263 modelled or projected PPR counts (19%, n = 8) using epidemiological parameters (11,14,27,33,35–37,39–
264 41). Six studies focused only on predicting PPR epidemics using epidemiological factors as covariates to
265 forecast the PPRV suitability area and estimate the spatial risk (number of outbreaks in PPR passive
266 surveillance data (25–27,37,42,43). Transmission dynamics were also evaluated using metapopulation
267 model (33). Before applying statistical models to evaluate the risks of the PPR epidemic, visualization and
268 descriptions of numerous PPR epidemics were conducted. These visualizations and descriptions provided
269 a perfect environment for understanding the distribution of the PPR epidemic over space and time. Data
270 visualization and descriptions were used by 34 studies (80.9%), and this was the most reliable method
271 used by all the studies to present the distribution of PPR epidemics. Various analytical approaches used

272 for data visualization and description include GLMNB (11), GLMMs (29) and negative binomial to evaluate
273 the possible risk factors linked to each outbreak's PPR disease case count (1,27). Either linear (13,24,30,31)
274 or logistic regression models were used to explore or test hypotheses related to temporal trends. GeoDa
275 1.14.0 was used to perform Geographically Weighted Regression (GWR), with climate and geographical
276 variables acting as predictors and log-transformed PPR cumulative incidence serving as the dependent
277 variable (28). Bayesian techniques, such as Bayesian Time-Scaled Phylogenetic Analysis, were used in
278 other investigations (2,3,44) along with the empirical Bayesian kriging (EBK) technique for risk map
279 generation and geostatistical prediction. NAADSM (North American Animal Disease Spread Model (12),
280 autoregressive integrated moving average model (ARIMA) (35), least cost path (LCP) (36), The Naïve Bayes
281 (NB) and Random forest machine learning algorithms (26), ensemble algorithm(37), regression tree
282 models (26) and Mantel correlograms (13) which was employed to examine the relationship between
283 genetic distances and various measures of geographical and network distance. The seasonal population
284 matrix model was used to assess the ability of different vaccination schedules they can be incorporated
285 into the PPR control program (45). Using ArcGIS v10.4, hot spot and cluster analyses were performed to
286 determine the hot and cold spot regions (1). Maximum Entropy Ecological Niche (MaxEnt) modelling was
287 employed to detect suitable areas for PPR virus distribution (46). GIS-based multi-criterion decision
288 analysis was used to identify areas at risk of PPR occurrence and spread (43). To investigate how the virus
289 spreads during memoryless state transitions in Afghanistan, an event-driven model of PPR derived from
290 the susceptible-exposed-infectious-recovered (SEIR) model was employed (38).

291

292 **3.7. Epidemiological risk factors associated with PPR epidemics**

293 Nine studies found that socioeconomic factors significantly influence PPR risk, despite trade and
294 commerce, environment, ecology, animal demographics, spatial accessibility, and other epidemiological
295 features accounting for a larger proportion. Risk factors linked to PPR outbreaks in Asia and Africa are

296 identified in the review, as shown in Figure 6 and Sup. Table 4. Two studies were not examined, and
297 epidemiological parameters were incorporated across studies, indicating a consistent risk pathway linking
298 them to outbreaks. The study lists all examined covariables in detail in Figure 6 and Sup. Table 4.

299
300 Figure 6: Distribution of models, at regional and subregional, containing variables from every category in
301 the conceptual framework
302

303 **3.7.1. Livestock demographics and livestock–wildlife interactions**

304 According to Sup. Table 2 The primary objective of twenty studies (six in Africa, twelve in Asia, and one
305 worldwide) was to identify hotspots by examining the impact of susceptible livestock populations on PPR
306 risk. Sheep and goat populations were reported as a covariate more frequently than other susceptible
307 species, even though many other species were taken into consideration (e.g. camel, cattle and pigs) (Sup.
308 Table 2). In 3 out of 17 studies, there was a correlation between a small ruminant population size and
309 sheep and goat density, or a higher risk of PPR. (19,32,47). In general, 47.4% (7/19) of research done in
310 Asia and Africa linked this category of wildlife-livestock interaction to PPR risk.

311 312 **3.7.2. Livestock Trade**

313 Livestock trade-related factors were included in 19 studies, 9 in Africa and 10 in Asia, as in Sup. Table 2.
314 Eight studies (three in Africa and five in Asia) included the distance or adjacency to an international
315 boundary in their models, demonstrating the common goal of examining the impact of international
316 connections on PPR risk using proxy variables that represented the likelihood of an international trade
317 network or cross-border movements. Six investigations found that the incidence of PPR decreased with
318 increasing distance to an international border (11,13,27,36,48,49). Additionally, elements of the dynamics
319 of the market were taken into account, such as the movement of live animals or their products. Finding
320 the markets in China, Ethiopia, Uganda, and Comoros was crucial for determining the outbreaks in those

321 countries (1,11,47,50). Two investigations examined the relationship between PPR epidemics and human
322 demography (25,49).

323 **3.7.3. Accessibility and networks**

324 Using topographical and landscape factors such as water bodies and permanent transport networks, 14
325 research (five in Africa, seven in Asia, and two worldwide) investigated PPR risk (Sup. Table 2). Studies in
326 African countries show a link between spatial accessibility and PPR risk, while Asia has a different
327 association. The greater risk is found near major transportation routes or dense road networks (11,43).
328 According to one Ethiopian study, one of the primary indicators of PPR transmission was the quantity of
329 successful encounters per unit of time (at pastures or watering spots, as well as through live-animal
330 commerce) (33). Four studies assessed the influence of natural landscape elements on inland waterbodies
331 and rivers, with only one highlighting the buffering effect of adjacent water bodies against PPR (25).

332 **3.7.4. Association between environmental factors and PPR epidemics**

333 The review of 13 studies on PPR outbreak risks, including those in Africa, globally, and Asia, identified 10
334 environmental and ecological factors as key categories (Sup. Table 2). The most frequently analyzed
335 features were season, temperature and precipitation (1,22,24–26,30,37,49). Landscape was also assessed
336 by five studies (25,31,42,49). Seasonality, landscape, weather, and climate-related covariables are
337 strongly linked to PPR epidemics, with lower altitudes being more associated with PPR risk compared to
338 higher altitudes (25,27,49). Temperature, precipitation, humidity, sun radiation, wind speed, and other
339 variables that affect land cover status all have an impact on PPR epidemics. (25,26,37). Studies in Asia
340 (4/4) and Africa (3/15) found a correlation between PPR risk and ecological features, but the variables
341 investigated varied, indicating different analysis goals.

342 **3.7.5. Economic and social advancement**

343 Only 9 research (Sup. Table 2) (27) used socioeconomic data to evaluate the risks of PPR spread globally
344 and in the Republic of Kazakhstan, respectively, using socioeconomic and geographic factors such as

340 landscape characteristics. The literacy rate and the availability of veterinary services (such as the number
341 of veterinarians, technicians, and animal health professionals) were the subjects of two studies (47) (25),
342 one conducted in Asia and the other in Africa. Poor animal health and veterinary services assessed in
343 Mandi City, northwest of India, have increased the risk of PPR infection (25). The contribution of the urban
344 population to PPR epidemics was reported by two studies, one in Africa and the other in Asia (25,35).

300 **3.8. PPR transmission dynamics and respective control measures**

301 To illuminate the important PPR transmission dynamics, evaluation of PPR risks was done in almost all
302 studies except two (9,17), which have not explicitly evaluated this covariate. Four studies have explicitly
303 evaluated the transmission dynamics to the control (33,35,39,41). The first one was conducted in Ethiopia
304 and uses a metapopulation model that mimics PPRV spread to evaluate the degree of PPRV transmission
305 in endemic situations. (33). The second one uses a mathematical model to assess the impact of four
306 vaccination strategies implemented at different times in reducing the PPR burden (41). In India,
307 Susceptible-Exposed-Infectious-Removed (SEIR) were used to evaluate PPR transmission dynamics (35).
308 Another study conducted in Tanzania has provided evidence of the PPR transmission pattern (39).
309 However, the probability of PPR transmission is linked to some covariables evaluated in this review; only
310 13 studies are influenced by livestock contact. Livestock movement as the main contact parameter in
311 PPRV transmission has been evaluated using network analysis to identify PPRV hot spots (13). Those
312 covariables included temporal and spatial aspects, production systems and the use of PPR control
313 strategies such as vaccination and early reporting systems (9,25). Additionally, all six studies assessing the
314 effectiveness of immunization found that regions taking part in PPR vaccination programs had a lower
315 probability of PPR outbreaks (Sup. Table 2). Despite the indisputable significance of PPR transmission
316 dynamics, most studies have used temporal and spatial resolution during epidemiological data collection.
317 The diverse range of these linked covariables has included all epidemiological parameters in this review.

318

369 **4. Discussion**

370 PPR epidemics have significantly impacted African and Asian economies, requiring spatial and
371 spatiotemporal approaches for disease management. However, uneven load may be due to diverse data
372 sources and analytical approaches. An increasing trend is seen in studies using spatial and temporal
373 analysis for estimating PPR risks in endemic countries (21). This review highlights the improvement in
374 understanding disease risk tracking, control planning, and eventual elimination of PPRV through the
375 assessment of PPR transmission dynamics.

376 **4.1. Modelling approaches**

377 This review analyzes the spatiotemporal distribution of PPR using visualization and descriptive tools. It
378 improves understanding of PPR impact in endemic countries and aids in designing control strategies like
379 vaccines or surveillance buffer zones. As more data becomes available, model development and
380 refinement are crucial for understanding local PPR risks. Predictions need to account for livestock
381 mobility patterns.

382 **4.2. Spatial autocorrelation**

383 Spatial autocorrelation or spatial dependence, is a key component of PPR spatial epidemiology (25).
384 Quantification of spatial autocorrelation in some studies was done by using global spatial
385 autocorrelation indices (21) i.e., Moran's I (25), Mantel test (13), and Getis Ord (27). When evaluating
386 seasonal and temporal changes in disease, it is crucial to analyze spatial-temporal autocorrelation. The
387 identification of outbreaks, clusters, or hotspots might provide clues about the hidden reasons behind
388 the rise in disease incidence and related factors that contribute to endemicity (15). The usefulness of
389 these results is not limited to hotspot detection but it goes further to PPR targeted control measures
390 (proactive or reactive vaccination, culling or depopulating and/or quarantine).

391 **4.3. Temporal trends**

392 Epidemics of PPR like other diseases tend to have seasonal variation due to continuous changes of
393 environmental and ecological factors. In our review, we have identified how these environmental and
394 ecological factors underlying the seasonal transmission of PPR are critical for predicting and
395 understanding the long-term environmental trends and effects on livestock health. PPR spread is
396 influenced by animal contacts, resource sharing, economic activities, and trade. Seasonal variations in
397 precipitation, temperature, and availability of pasture and water can affect livestock mobility and PPR
398 risk projection. Seasonal activities like festivals and dowries during harvest also increase the risk of PPR
399 spread. Control strategies include strategic vaccination and movement restrictions before disease
400 outbreaks. These risk factors vary geographically and demographically.

4.4. Risk factors associated with PPR epidemics

402 This review examines risk factors related to sheep and goat disease, focusing on demographics and
403 livestock interactions. It highlights the importance of livestock movement, geographic and
404 environmental factors, and the increased density of wildlife or livestock in interface areas. The review
405 also highlights the role of socioeconomic activities in causing interaction between people and animals.
406 Identifying PPR-prone areas is crucial for risk-based surveillance and control measures (43).

4.4.1. Livestock- wildlife Interactions

408 The impact of livestock-wildlife interactions on the risk of PPR epidemics is linked to the density of
409 susceptible species at livestock wildlife interface areas. In our review, several studies have shown
410 evidence of PPR outbreaks in a protected area to be due to its proximity to a PPR risk area. Few
411 evidence of PPR in wildlife species in Africa is surprising given the increasingly visible epidemics in
412 wildlife in Asia (48). Evidence from our models suggests that the spatiotemporal patterns of PPRV
413 outbreaks in wildlife were similar to those in livestock, suggesting evidence of PPR virus spillover from
414 livestock to wildlife. Global climate changes have resulted in shifts in species distributions and habitat
415 suitability consequently reducing resource availability and increasing wildlife-livestock interactions. In

our models, evidence of increased risk of PPR infection due to interfered relationship between species, habitat and climate leading to mortalities of wildlife species e.g saiga (*Saiga tartarica tartarica*) in Kazakhstan has been demonstrated (51).

4.4.2. Accessibility and network

Livestock movement is a key risk factor for PPR outbreaks, with most models indicating that geographical infrastructure links are crucial for contact between livestock populations (43). Large water bodies like lakes, rivers, and oceans pose PPR risk due to their accessibility, but poor infrastructure and proximity to water bodies may limit disease spread. Expanding vehicular transportation and limited access to resources may reduce disease contamination. Our review identifies critical sites for PPR transmission in livestock movement, improving local control and surveillance strategies by combining genetic and mobility network data (13).

4.4.3. Social Economic Factors

This review evaluates the impact of socioeconomic factors on the risks of PPR spread, focusing on political, economic, veterinary services, and stakeholders' knowledge. A herd-level event-driven PPR model was created to identify effective management strategies for different herd compositions and circumstances. For example, in Afghanistan, the Lack of scientific data due to due to the political situation has impacted large-scale disease mitigation strategies. A cost-benefit analysis was conducted to assess the economic significance and impact of PPR. Stakeholders' knowledge assessment is crucial for involving livestock keepers in control and eradication programs. The study reveals that social-economic vulnerability, climate change, and risk mitigation strategies, such as livestock mobility and herd diversification, are key drivers of the spread of PPR, influenced by factors such as cultural events, husbandry methods, and economics. Other social practices such as livestock marriage dowries can also promote the spread of PPR to other areas through livestock mobility. The social and political status of PPR endemic countries significantly influences future regional transmission dynamics, necessitating

careful consideration of control benefits in cost-benefit analysis for resource mobilization and political will.

4.4.4. Livestock trade

The review examines market dynamics of sheep and goat production linked to PPR risks, examining the likelihood of livestock trade facilitating PPR spread in specific areas. It examines infrastructure like road and railway networks and slaughter facilities. For instance, in Bangladesh, the livestock movement within herds and the central market of the local small and large markets has been connected to the PPR outbreak (28). Another model which evaluates urbanization and habitat characteristics has shown naked evidence of PPR risk to the livestock trade (29). The review highlights regional variations in livestock trade infrastructure, highlighting the need for different control strategies in local contexts. It emphasizes the importance of empowering sub-Saharan countries with livestock trade infrastructure for PPR surveillance and control to meet the 2030 PPR eradication target.

4.4.5. Ecology and environment

Studies on landscape and climatic features linked to PPRV environmental circulation, stability, and survival have been assessed in this review. Results have shown geographical variation in risk magnitude. For example, an area with good precipitation will have good pasture and water supply (42). Shrinkage of Grazing land due to factors like the expansion of conserved areas, agricultural activities and climate change has been linked to PPR risk. In this review, we have been able to identify prediction models which can be used in designing control strategies according to the environment and ecology of the respective area. For example, in China, seasonality pattern identification is critical for vaccination schedules because during summer high temperatures lower animal immunity (36). The ecological niche model discussed in this study has demonstrated how ecological and environmental characteristics are related to PPR outbreaks. The annual maximum temperature was inversely correlated with PPR outbreaks because PPRV is more susceptible to hot temperatures. Other elements, such as seasonal

464 variations in precipitation (warm vs. dry season), exhibited a substantial positive connection with PPR
465 outbreaks, whereas wind speed had a negative association (37). Prediction models allow us to identify
466 the right time for control measures deployment, it becomes ineffective if done outside the time range
467 hence 2030 PPR eradication target has been identified using prediction models (37).

468 **4.4.6. PPR transmission Dynamics**

469 The dynamics of PPR transmission depend on the rate of transmission from PPRV-infected animals to
470 susceptible hosts. In disease models, this rate of transmission is captured in one parameter called the
471 probability of transmission. In our review, several studies have captured spatial transmission dynamic
472 modelling approaches to investigate PPR transmission dynamics and control. These models have been
473 useful in generating scenario analyses of the potential course and severity of PPR epidemics by
474 characterizing and forecasting the spatiotemporal transmission patterns of PPR epidemics, or assessing
475 the effectiveness of interventions and the feasibility of achieving elimination targets (24). In addition to
476 attempting to capture pertinent mechanisms of PPR transmission, such as the possible influence of
477 environmental factors, our review has been able to include important epidemiological characteristics of
478 PPR infection (48). A specific kind of spatial dynamic model known as a metapopulation model divides
479 the population into a number of interacting population groupings based on demographic or spatial data
480 (33). For instance, in Ethiopia, PPRV incursions from lowlands into highlands occur as a result of unequal
481 pasture, water, and animal market distribution, necessitating immunization that targets interfaces
482 between various population sites. Our understanding of PPR transmission dynamics has the potential to
483 achieve a high level of communication speed between the response team in the outbreak event. The
484 outbreak will prompt coordinated events from the point of origin and contributors, immediate
485 epidemiologic characterization on the ground, evaluation to establish spread pathways, and specimen
486 collection to molecular characterization of the virus, all while executing spatial prediction models, using
487 the understanding of PPR transmission dynamics (33).

488 **4.5. Strength**

489 This scoping review provides detailed information on risk-based spatiotemporal techniques for PPR
490 spread in endemic situations. It highlights a promising trend in using spatial epidemiology tools to
491 understand PPR's transmission mechanism. The review identifies areas for future research and
492 highlights methodological limitations in existing studies. It also provides a fair depiction of PPR risk
493 mapping initiatives using a thorough search method in compliance with PRISMA Scoping criteria.

494

490 **4.6. Limitations.**

496 The study acknowledges the challenge of modelling PPR risks in endemic situations due to the lack of real
497 data and the complexity of approaches. Some models are hypothetical, focusing on mathematical tools
498 for new ideas. The study acknowledges the potential overlooking of relevant publications and studies
499 published in other languages due to the inclusion of only English-published studies.

500 **4.7. Conclusion**

501 Spatial and Spatiotemporal approaches have played a critical role in shifting and improving our
502 understanding of the available disease management options for PPR. However, the uneven burden of PPR
503 may be attributed to the diverse data sources, covariates, and analytical approaches employed. Future
504 development of prediction models to assess potential eradication efforts at national and regional levels
505 should also take Africa into consideration, as many studies have been conducted in Asia as opposed to
506 Africa.

507 **Ethics**

508 Not relevant

509 **Authorization for publishing**

510 Inapplicable

511 **Materials and data accessibility**

012 The corresponding author can provide the datasets used and/or analysed in this study upon reasonable
013 request.

014 **Conflict of interest**

015 There are no conflicting interests, according to the authors.

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021 **Author Contributions**

022 Concept and design of the study: JJM, SK, GM, AC

023 Data collection: JJM, JNH

024 Data analysis and interpretation: GM, AC, JJM, JNH

025 The manuscript was drafted by JJM

026 The manuscript has been critically revised for significant intellectual content by JNH, GM, AC, SK, DM, EM,

027 EO, GO, and GPO.

028 Analysis of statistics: JJM

029 SK, GM, and AC provide administrative, technical, and material support.

030 Study oversight: SK, GM, and AC

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037

- 038 **Supplementary tables (Sup. Tables)**
039
040 Sup. Table 1: Preferred Reporting Items for Systematic reviews and Meta-Analyses extension for Scoping
041 Reviews (PRISMA-ScR) Checklist
042 Sup. Table 2: A classification system that includes a comprehensive list of the epidemiological factors
043 that have been reported in the included studies
044 Sup. Table 3: Features and synopsis of the studies that are included
045
046 Sup. Table 4: Index of research papers mentioning at least one covariate associated with the
047 likelihood of PPR outbreaks, broken down by region and subregion.
048
049 Sup. Table 5: An overview of techniques for utilizing PPR epidemic data to study spatial, temporal, or
050 spatiotemporal clustering.

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