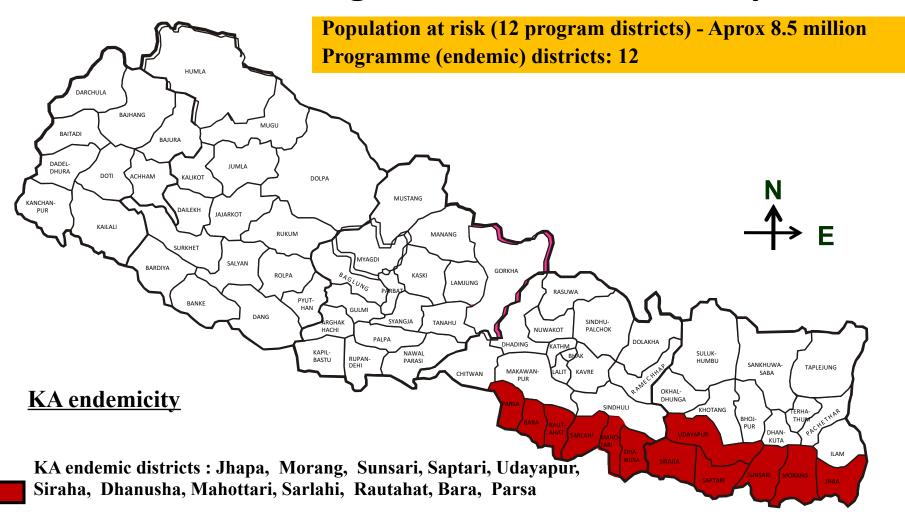
Elimination efforts in Nepal: lessons learned and challenges remaining

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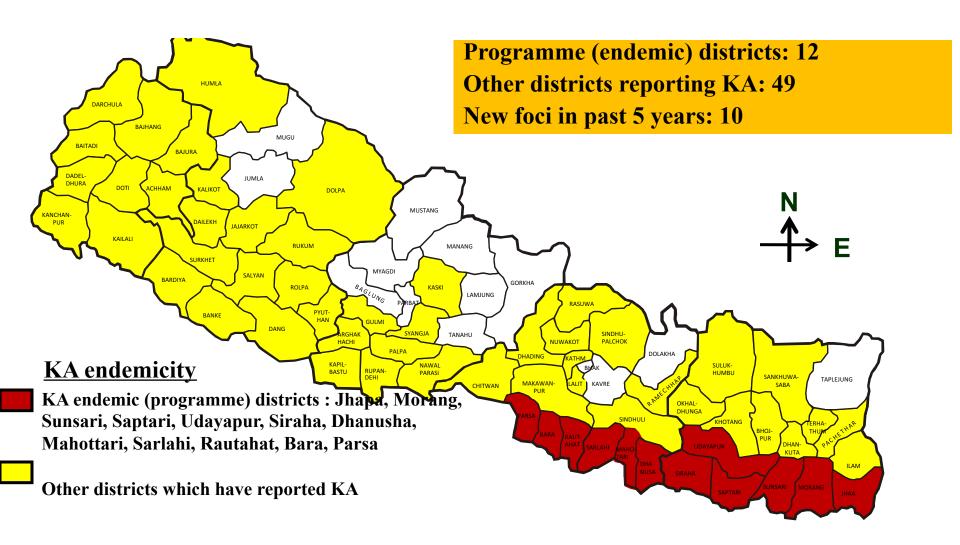
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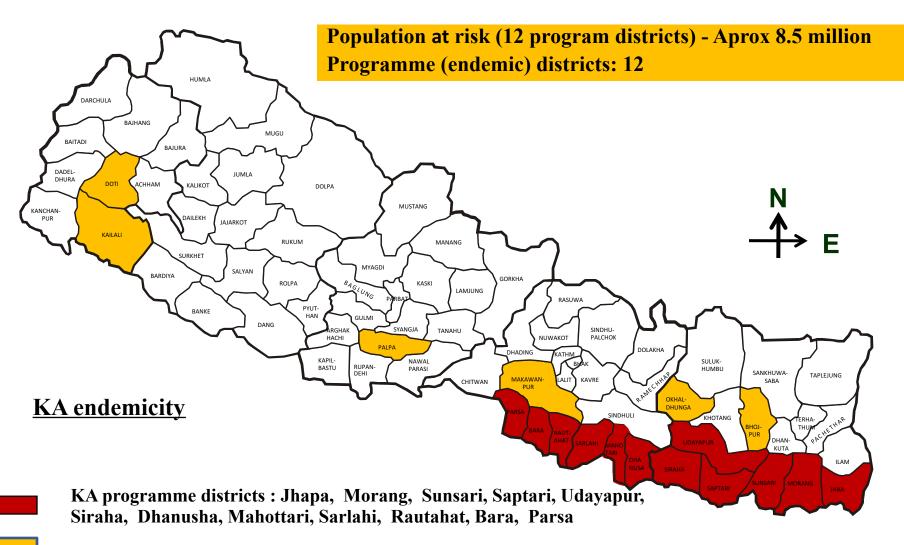
Place = KA Programme Districts in Nepal



Place = geographical expansion of Kala-azar: 2009 - 2015

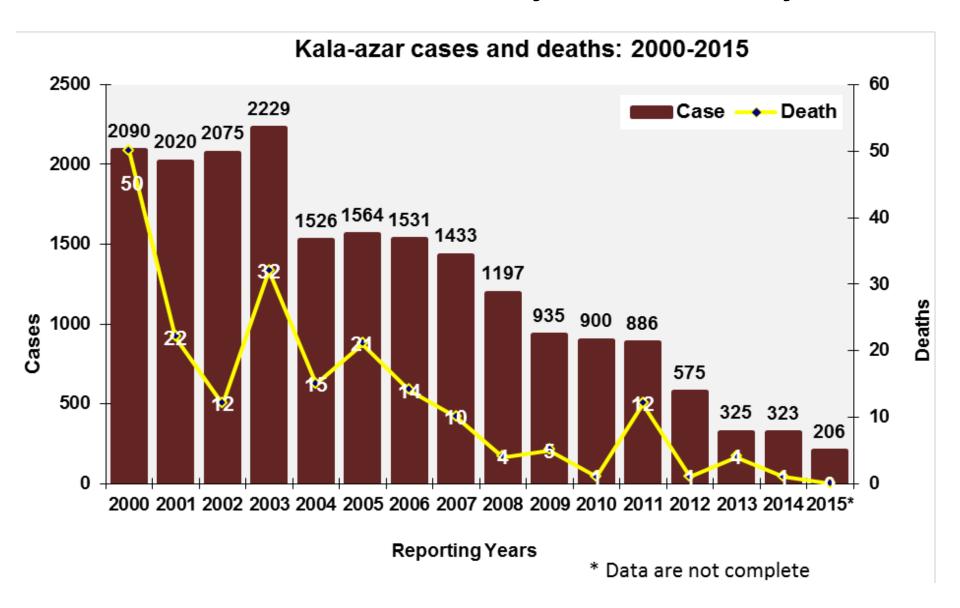


Place = KA Programme Districts in Nepal (2015)



New KA programme districts added in 2015

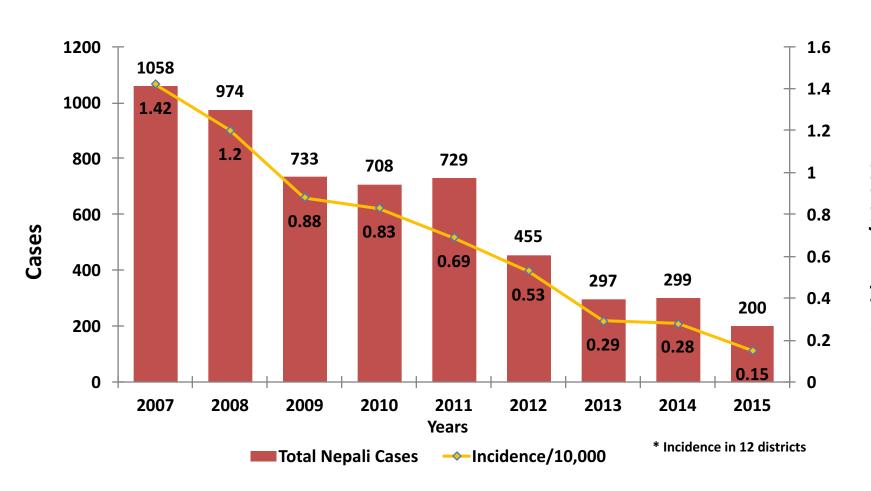
Trend in KA morbidity and mortality



Incidence/10,000 pop

National Annual Incidence: 2007-2015

Incidence against native cases



Implementation of the Kala-azar Programme in Nepal = phases

- Preparatory phase= 2005 2006 (2 years)
 - Development/review of national policy & strategic plans
 - Training to healthcare providers, IEC/BCC activities
- Attack phase = 2007 2011 (5 years)
 - Implementation and monitoring of activities = IRS,IVM, access to early diagnosis & Treatment.
 - ACD, PCD, Case-based surveillance, and vector surveillance
 - Access to early diagnosis, and complete treatment
 - IVM, vector surveillance etc

Implementation of the Kala-azar Programme in Nepal = phases

- Consolidation phase = 2012 2014 (3 years)
 - Total coverage by IRS
 - Active surveillance = no increase in the incidence rate at district
 - Limited IRS based on geographical location of cases, and in areas with high vector density
 - Intensified active-case detection, early diagnosis & complete
 Treatment.

Maintenance phase (duration to be decided)

- Strong surveillance against re-introduction of Kala-azar cases
- Incidence of KA should be less than 1/10, 000 population at district
- Sustain the elimination target at district level
- Certification of elimination status

Elimination strategies adopted in Nepal

- Early case detection and completion of treatment
- Vector control activities
- Capacity building of health workers
- Maintenance of the supply chain
- Surveillance and research
- Partnership
- Pharmacovigilance
- Cross-border collaboration
- Non-health interventions (a financial incentive equivalent to US\$ 10 for cases having completed treatment).

Surveillance and Information system

- Kala-azar patient and laboratory register at district level,
- Monthly Kala-azar reports by district public health office,
- Health management and information system (HMIS)
- Line listing of cases
- Epidemiological and entomological verification in districts/new foci.

Diagnosis of VL at different healthcare levels

Level I: FCHV, HP

Screening and referral

Monitoring of compliance/ side effects by FCHV

Active case finding

Level II: PHC

Diagnosis by RDT

Treatment if doctor available

Referral: microscopy diag. & side effects

Level III: DH, ZH, Tertiary

Diagnosis: RDT and microscopy

Treatment,

Manage
complications

Pharmacovigilance

FCHV: Female community health volunteer, HP: health post, DH: district hospital, ZH: zonal hospital

Active case finding strategies

- Active case detection activities in VL endemic districts and pockets;
 - Camp based approach
 - Mobilization of female community health volunteers

Collaboration with leprosy programme:

- Orientation and training to leprosy service providers on KA and PKDL;
- Collaborative active case search in endemic districts (Combined search)
- Rapid diagnostic test kits provided to selected leprosy service centres/hospitals
- IEC and awareness activities

Treatment regimen

- 1st line treatment
 - Liposomal amphotericin B (L-AmB)
 - Combination regimens
 - ➤ Miltefosine + Paromomycin
 - L-AmB + Paromomycin
- 2nd line treatment
 - Amphotericin B
 - Miltefosine monotherapy

Diagnostic tools available

(i) rK39 dipstick





(ii) Bone Marrow Aspiration



Introduction of Liposomal Amphotericin-B

- AmBisome donation (2000 vials) was received in September 2015,
- L-AmB introduced in December 2015 after training to doctors and nurses.
 - Aprpx 60 doctors and nurses from endemic districts
- Rolled out to major hospitals in endemic districts.

Vector control

- Strategy: Focal spraying in priorityselected areas
- Insecticide: Lamda cyhalothrin and Deltamethrin
- Two rounds per year
- Coverage of spraying: >80%
- Monitoring: Onsite spraying monitoring, insecticide resistance, bioassay-research purpose
- Sporadic distribution of Long lasting insecticidal nets (LLINs)





Factors contributed to reduction in the number of cases and deaths

- Availability of diagnosis and treatment
- Capacity building through orientation/training physicians and health workers
- Indoor residual spraying and distribution of longlasting insecticidal nets
- Provision of transport incentives to patients after completion of treatment
- Social mobilization and awareness
- Socioeconomic interventions from non-health sectors.

Lessons learned

- For improved surveillance, Kala-azar should be made a <u>notifiable disease</u> in the affected areas.
- Passive case detection should be supplemented with <u>active case detection</u> with <u>laboratory diagnosis</u>
- Active case detection should be done at least once a year or ++ if possible
- ACD case detection becomes more important as the number of KA cases reported by passive case detection declines
- Disease surveillance for Kala-azar should be comprise monthly reporting with line listing of cases and feedback mechanisms at district level
 - HMIS
 - EWARS
 - Line listing of KA cases
 - Report of active case detection/case-based surveillance

Collaboration with other programs

- Combined active search for PKDL, KA and leprosy
- Training/orientation conducted to leprosy service providers/hospitals of KA endemic districts.
- Diagnostic kits (rK39) provided to leprosy hospitals.

Challenges

- Sustaining the achievement- Frequent outbreaks in endemic and non endemic areas
- Changing pattern in distribution: geographical expansion of cases- rural to urban and plain areas to up hills
- Detection of PKDL cases- even confused with other skin infections
- Diagnosis and reporting of relapse
- Drug resistance- Regular monitoring is required
- Cross border information sharing

Housing conditions in KA-endemic region of Nepal

