Indian J Med Res 158, September 2023, pp 217-254

DOI: 10.4103/ijmr.ijmr_3299_21



Verbal autopsy to assess causes of mortality among the economically productive age group in the tribal region of Melghat, central India

Ashish Rambhau Satav^{1,2}, Kavita Ashish Satav^{2,3}, Abhay Suresh Kelkar¹, Vrinda S. Sahasrabhojaney¹, Vibhawari Sharad Dani⁴, Dhananjay V. Raje⁴ & Eric A. F. Simoes^{5,6}

Departments of ¹Medicine, ²Community Health & ³Ophthalmology, Mahatma Gandhi Tribal Hospital, ⁴Department of Research, MAHAN Trust, Amravati, Maharashtra, India, ⁵Department of Paediatric Infectious Diseases, University of Colorado School of Medicine and Children's Hospital Colorado & ⁶Department of Epidemiology, Colorado School of Public Health, Aurora, Colorado, USA

Received November 11, 2021

Background & Objectives: Verbal autopsy (VA) is the systematic and retrospective inquiry (from relatives) about the symptoms of an illness prior to death. In tribal India, 67-75 per cent of deaths occur at home with an unknown cause of death (CoD). Hence, the aim of this study was to determine the CoD in the 16-60 yr age group utilizing VA.

Methods: A prospective, community based longitudinal study was conducted in 32 tribal villages in the Melghat region of Maharashtra, between 2004 and 2020. Number of deaths and VAs in 16-60 yr age group were collected by village health workers (VHWs) and supervisors, verified by five different persons (internal-external) and cross-checked by three VA interpretation trained physicians. A modified version of WHO VA was used. Cause-specific mortality fractions were calculated.

Results: Of the 1011 deaths recorded, mortality in males was significantly higher than females (P<0.001). A total of 763 VAs were conducted which revealed that tuberculosis was the leading CoD, followed by jaundice, heart diseases, diarrhoea, central nervous system infections and suicide. Suicides were significantly more common among males than in females (P=0.046). Significantly, more deaths occurred during the monsoon (P=0.002), especially diarrhoeal deaths (P=0.024).

Interpretation & conclusions: The findings of this study suggest that, in Indian tribal areas, infectious diseases are the leading causes of morbidity and one of the major causes of deaths in economically productive age group. Intensified VHW-mediated interventions are required to reduce the premature deaths.

Key words Economically productive age group - Melghat - tribal - verbal autopsy

Understanding the cause of death (CoD) is crucial for policy framing and planning programmes to improve health indicators. A reliable assessment of disease-specific mortality rates is nearly impossible in developing countries because the underlying cause (UC) is unknown or relevant information is not recorded due to poor reach and coverage of the death registration system¹. Verbal autopsy (VA) is

a retrospective inquiry systematically conducted involving the family members of the deceased to collect information pertaining to the symptoms before death through predetermined questions to describe the CoD for home deaths^{2,3}. VA will enable evidence-based decision-making and can guide public health priorities in communities without physician certified deaths^{4,5}.

In most resource constrained, hilly, forest, rural-tribal areas of world with high mortality, documentation of death information and medical certification of CoD is nearly impossible because of linited healthcare access, low health literacy in tribal populations and a high proportion (67 to 75%) of deaths which occur outside of the healthcare setting^{6,7}. The physician-certified verbal autopsy (PCVA) approach is reportedly more effective in determining CoD in community deaths⁸.

MAHAN trust, a non-governmental organization (NGO) is continually providing holistic medical services to poor tribals of Melghat for the last 25 years, conducted a 17 yr study which revealed that the age-specific mortality rate (ASMR) in Melghat was >400/100000 population², which is 2.25 times greater than the rest of India (178 in 2016). Deaths in the economically productive age group can have detrimental effects on the community, nation and family members, particularly children as it also reportedly increases malnutrition⁹ and deaths¹⁰.

This study was conducted to ascertain the CoD in the 16-60 yr age group in Melghat, which has previously been reported to have a disproportionately high age-specific mortality rate as compared to the Indian average. In order for intervention to prevent these premature deaths through the village based healthcare for the economically productive age group people (EPAGP), as a first step, VA in EPAGP was needed to know the CoDs to plan the programmes to reduce mortality. The objective of this study was to determine the underlying cause (UC) for cause-specific mortality and to determine contributory causes (CC) for all causes of mortality by VA in EPAGP in Melghat.

Material & Methods

A community based longitudinal prospective study was conducted from January 2004 to December 2020 in EPAGP in 34 villages of Melghat. The study was ethically approved by the Institutional Review Board of MAHAN trust. A written informed consent was obtained from all participants and the Gram Sabha of the villages.

Source population: Melghat is a hilly, forest, difficult to access, highly impoverished tribal area in Amravati district of Maharashtra. It is spread over 4000 km², across 320 villages. The population is 300,000 with 84 per cent poor tribal, 50 per cent semiliterate/illiterate, mostly small farmers/labourers. Melghat has poor transportation, grossly inadequate health services and low health seeking behaviour, leading to high mortality¹¹.

Sample size: In all 34 villages (10% of total 320 villages) from two blocks of Melghat were considered for this study (Fig. 1). A minimum of 50 per cent of total mortality was decided as the criteria for VA, considering the feasibility of such exercise in the tribal population. The World Health Organization (WHO) has recommended cluster sampling and minimum population size of 2000-20000. A two stage cluster sampling technique was employed. The first stage is a block and the second is the villages within each block. Eight clusters/villages from five zones were randomly picked up by the lottery method. Finally, 34 clusters were selected based on the willingness of the villagers and village health workers (VHWs) to participate and were block stratified. In the year 2016, two villages were dropped from the study due to administrative reasons and hence data was analysed from 32 villages. The sample unit comprised all deaths in the EPAGP in the selected villages. The total number of deaths during the period was 1011. Verbal autopsies were conducted in 763 (75.46%) deaths.

Inclusion and exclusion criteria: *De-facto* method (home and hospital deaths in the study area) was used¹². Only close relatives of all deceased individuals who gave consent were included in this study. Those who migrated (≥6 months ago) were excluded from the study.

Study tool and technique:

<u>Vital event capture methodology</u>: The VHWs, who were residents of the same villages, conducted a census (including population in EPAGP), that included a baseline door-to-door survey regarding deaths (16-60 yr) and demographic information in January 2004. They were supervised by data collection supervisors and project managers. It was reviewed by the principal investigator. Multiple methods were used for death data collection in the community to improve the accuracy of death statistics¹¹.

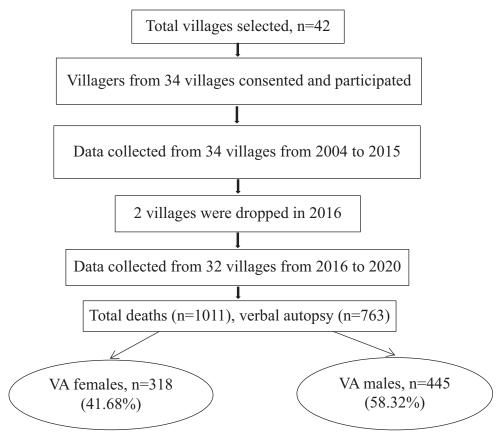


Fig. 1. Selection of study participants.

Standardized VA tool developed by MAHAN was used. It was the modified version of the WHO VA tool¹³ adapted to the local context. The MAHAN VA tool consisted of a standardized questionnaire (open and close ended) that gathered information on symptoms, medical history and circumstances preceding the death. The tool included a set of standard algorithms for determining major CoD among EPAGP. The CoD, or the sequence of causes that led to a death, was assigned based on the data. The tool was piloted for validation by a senior physician, among 20 deaths at a medical college in Sewagram (data not shown). The CoD by VA was compared with the hospital CoD and there was >75 per cent agreement between the two. The validity of this tool was confirmed with three different physicians/co-investigators and two field staff. From 2004 to 2010, the same VA data collection tool was used. Modifications were subsequently made based on our field experience and the WHO4 and IHME methods14.

The well trained team speaking local dialect included a local tribal semiliterate VHW from each village and data collection supervisor. Grief

counselling was done within 48 h of death and before conducting VA by the team, who conducted VA within seven to 15 days of the death of the person, after obtaining informed consent from the relatives. The process consisted of: (*i*) asking the family members open and close ended questionnaires; and (*ii*) its analysis¹⁵.

A trained medical supervisor cross checked the VA forms. VA with its algorithms was checked by two other VA interpretation-trained doctors who independently assessed it for CoD. If the CoD determined by two doctors concurred, then that defined the CoD. The agreement level between two coders was 89.39 per cent. In 11 per cent of VA, there was discrepancy in CoDs by two doctors. Following underlying CoDs were different in the two coders: Tetanus, central nervous system (CNS) infection vs. intracranial space occupying lesions, intestinal obstruction vs. perforation peritonitis, diabetes mellitus, tuberculosis (TB) and acquired immunodeficiency syndrome. The authors sent discrepant VAs to a third coder, an experienced physician who acted as an arbitrator. The arbitrator personally checked the VA, verified the discrepancy

and finalized the CoD. There were 15 garbage codes in all deaths. However, only two of these were included in the final list and the rest were the part of a miscellaneous CoD.

As the CoD was determined by VA, it was not possible to conduct bacteriological confirmation for TB. TB was differentiated from chronic obstructive pulmonary disease (COPD) by the following history: presence of haemoptysis, significant weight loss, extreme fatigability and anorexia during the last one month, family/past history of TB, no BCG vaccination, any lump in cervicaland/axillary area, presence of any associated symptoms such as neck pain, headache, vomiting, change in voice, difficulty in deglutition, fever suggested that TB (pulmonary or extrapulmonary) was the CoD. The following history favoured COPD: excessive breathlessness leading to difficulty in dayto-day routine activities and patient had to sit for long time/days without sleeping in supine position, patient had cough with white mucoid expectoration for a minimum for two months every year which increased during winter for three consecutive years, cyanosis, recurrent breathlessness, history of smoking, etc. The table showing algorithms for TB and COPD and VA form is attached as Annexures 1 and 2.

Statistical analysis: Data entry was done using Microsoft Excel 2007. Cause-specific mortality fractions (CSMF) were calculated as the number of deaths from specific cause, divided by the total deaths, during a given time interval¹⁶. The homogeneity of event occurrence between gender types and seasons was tested using the Pearson's Chi-square test of homogeneity. The analysis was done independently for each study period. The computations were performed using the SPSS version 26.0 (IBM Corp., Armonk, NY, USA) software and the significance was evaluated at five per cent level.

Results

The completeness of death reporting was ~95 per cent. The population of the 32 villages was 29,356, with 15,002 (51.1%) males and 14,354 (48.9%) females. The minimum sample size needed was 500 VA. Seven hundred and sixty three VA were randomly undertaken to improve the confidence interval and accuracy. There were no significant differentials in the response rates over time or by age/sex of the deceased. Only response rates varied across surveyors.

Table I shows age sex distribution of the study population and occurrence of deaths during three study

		Table I. Age	-sex distribution	of study popule	tion and occurr	Table I. Age-sex distribution of study population and occurrence of deaths during the three study periods	periods	
Age	Gender	Total population#,	2004-2009	2010-2015	2016-2020	200	2004-2020	
(yr)		n (%)	Deaths	Deaths	Deaths	Deaths/100,000 population per yr	Total deaths	% (95% CI)
16-20	Male	1826 (21.3)	22 (10.2)**	25 (10.9)	8 (5.7)	177.18	55	3.01 (2.28-3.9)
	Female	1882 (23.2)	32 (16.8)	21 (15.4)	9 (8.9)	193.79	62	3.29 (2.53-4.2)
21-30	Male	2945 (34.3)	51 (23.7)	51 (22.4)***	$31 (21.9)^*$	265.65	133	4.52 (3.79-5.33)***
	Female	2893 (35.6)	46 (24.2)	21 (15.4)	16 (15.9)	168.76	83	2.87 (2.29-3.54)
31-40	Male	1999 (23.3)	53 (34.7)	36 (15.8)	15 (10.6)	306.04	104	5.2 (4.27-6.27)
	Female	1778 (21.9)	40 (21)	23 (16.9)	20 (19.8)	274.6	83	4.67 (3.74-5.75)
41-50	Male	1234 (14.4)	57 (26.5)	55 (24.2)*	32 (22.7)	686.43	144	11.67 (9.92-13.59)
	Female	1014 (12.5)	51 (26.8)	28 (20.6)	29 (28.7)	626.52	108	10.65 (8.82-12.71)
51-60	Male	575 (6.7)	32 (14.9)	61 (26.8)	55 (39)	1514.07	148	25.74 (22.21-29.52)***
	Female	550 (6.8)	21 (11)	43 (31.6)	27 (26.7)	973.26	91	16.55 (13.54-19.92)
Total	Male	8579 (100)	215 (53.1)	228 (62.6)	141 (58.5)	400.43	584	$6.81 (6.28-7.36)^{***}$
(16-60)	Female	8117 (100)	190 (46.9)	136 (37.4)	101 (41.7)	309.44	427	5.26 (4.79-5.77)
#As per co	ensus 2009; P *<	"As per census 2009; P *<0.05, "*<0.01, "**<0.001. CI, confidence interval	01. CI, confidence	ce interval				

		Table	e II. Age-	wise distri	bution of	cause-spec	cific morta	Table II. Age-wise distribution of cause-specific mortality during three study periods	ree study per	iods		
Age groups (yr)	Major causes of deaths	2004-2009	6003	2010-2015	2015	2016-2020	2020	2004-2020 17 (yr)	17 (yr)	Deaths/100,000 population per yr	100,000 on per yr	UC + CC, % (95% CI)
		UC	CC	UC	CC	UC	CC	UC	CC	UC	CC	
16-20	TB	10	5	12	10	0	-	22	16	34.9	25.38	1.02 (0.7-1.4)
	Diarrhoeal disease	9	9	11	9	2		19	13	30.14	20.62	0.8 (0.59-1.22)
	Suicide	4	4	3	3	0	0	7	7	11.1	11.1	0.38 (0.21-0.63)
	Jaundice	4	2	-	2	_		9	S	9.52	7.93	0.30 (0.15-0.53)
21-30	TB	17	6	21	19	2	3	40	31	40.3	31.24	1.22 (0.95-1.53)
	Suicide	7	7	8	8	_	-	16	16	16.12	16.12	0.55 (0.37-0.77)
	Jaundice	7	S	8	4	2	4	17	13	17.13	13.1	0.51 (0.35-0.73)
	Diarrhoeal disease	2	4	4	5	_	2	7	11	7.05	11.08	0.31 (0.18-0.49)
31-40	TB	10	7	12	13	4	2	26	22	40.49	34.26	1.01 (0.71-1.38)
	Diarrhoeal disease	5	4	9	9	_		12	11	18.69	17.13	0.61 (0.39-0.91)
	Jaundice	4	4	7	5	0	0	11	6	17.13	14.02	0.53 (0.32-0.82)
	Suicide	4	4	2	2	_	П	7	7	10.9	10.9	0.37 (0.2-0.62)
41-50	TB	16	9	20	19	4	7	40	32	104.67	83.73	3.2 (2.51-4.02)
	Heart diseases	13	11	10	6	4	9	27	26	70.65	68.03	2.36 (1.77-3.07)
	Diarrhoeal diseases	5	6	4	4	-	3	10	16	26.17	41.87	1.16 (0.76-1.69)
	Jaundice	2	2	10	9	_	_	13	6	34.02	23.55	0.98 (0.61-1.48)
	CNS infection	4	4					4	4	10.47	10.47	0.36 (0.15-0.7)
51-60	TB	0	0	20	18	∞	~	28	26	146.41	135.95	4.8 (3.63-6.22)
	Heart diseases	∞	15	20	13	4	4	32	32	167.32	167.32	5.69 (4.41-7.21)
	Diarrhoeal diseases	∞	7	4	9	-	-	13	14	67.97	73.2	2.4 (1.59-3.47)
	CNS infection	3	9	4	3	3	4	10	13	52.29	26.79	2.04 (1.3-3.05)
	Jaundice	Э	7	7	ж	_		11	9	57.52	31.37	1.51 (0.88-2.41)
UC, underlyi	UC, underlying cause; CC, contributory cause; TB,	y cause; T	B, tubero	tuberculosis; CNS, central nervous system	JS, central	nervous s	ystem					

Alcohol toxication

AIDS

RF

Tetanus

11

8

10

6

2

6

1

5

8

4

10

7

2

4

1

3

23

22

22

21

1.51 (0.96-2.25)

1.44 (0.91-2.17)

1.44 (0.91-2.17)

1.38 (0.85-2.1)

8.1

7.75

7.75

7.4

Contd...

Cause of death		2	2004-20	010			2	011-20)15			2	2016-20	020	
	U	C (n)	C	C (n)	CSMF	U	C (n)	CO	C (n)	CSMF	U	C (n)	C	C (n)	CSMI
	Male	Female	Male	Female	(%)	Male	Female	Male	Female	(%)	Male	Female	Male	Female	(%)
ТВ	20	33	11	15	13.5	50	35	48	31	22.53	11	8	8	8	16.5
Jaundice	10	10	6	9	6	27	16	13	11	9.2	4	0	2	1	3.3
IHD	15	7	10	7	6.7	17	8	14	9	6.6	3	3	6	7	9
Diarrhoeal diseases	12	14	16	14	9.6	10	5	12	5	4.4	5	2	3	5	7.1
CNS infection	11	8	16	16	8.7	9	5	13	9	4.9	1	1	4	1	3.3
Suicide	13	4	13	4	5.8	17	6	17	6	6.3	1	1	1	1	1.9
Pneumonia	4	6	16	15	7	3	0	8	4	2.1	3	5	10	6	11.3
Cancer	5	7	4	4	3.4	13	11	10	10	6	4	3	2	2	5.2
CVE	5	5	5	7	3.8	3	8	4	8	3.2	3	1	1	0	2.4
COAD	7	4	7	6	4.1	3	3	5	3	1.9	4	3	1	2	4.7
RTA	1	0	1	0	0.3	8	0	8	0	2.2	2	0	4	0	2.8
Alcohol toxication	5	0	5	0	1.7	3	1	3	1	1.1	3	1	0	1	2.4
AIDS	4	2	1	1	1.4	3	2	3	1	1.2	1	2	0	2	2.4
Tetanus	3	1	3	1	1.4	5	0	5	0	1.4	2	0	2	0	1.9
RF	1	0	2	0	0.5	5	5	5	3	2.3	0	0	0	0	0
Homicide	5	2	5	2	2.4	2	1	2	1	0.8	0	0	0	0	0
Snake bite	2	0	2	0	0.7	5	1	5	1	1.7	0	0	0	0	0
Drowning	2	1	2	1	1	3	0	3	0	0.8	0	0	0	0	0
ALD	0	0	0	0	0	5	0	5	0	1.4	1	0	0	0	0.5
Miscellaneous	35	29	35	31	22.2	37	29	45	33	19.8	9	19	13	13	25.5
Total	160	133	160	133	100	228	136	228	136	100	57	49	57	49	100
Cause of death 2004-202		-2020													
_		UC ((n)		CC (1	n)	Tot	al (UC	+ CC)	CSM	IF, % (95% CI)	Γ	Deaths /10	00,000
	M	ale	Female	e Ma	ale l	Female	e M	ale + f	emale				po	opulation	per y
ТВ	8	31	76	6	7	54		278	}	18.22	2 (16.3	1-20.25)		97.9	5
Jaundice	4	11	26	2	1	21		109)	7.	14 (5.9	-8.55)		38.4	ŀ
IHD	3	35	18	3	0	23		106	ó	6.9	5 (5.72	2-8.34)		37.3	5
Diarrhoeal diseases	2	27	21	3	1	24		103	3	6.7	5 (5.54	1-8.13)		36.2	9
CNS infection	2	21	14	3	3	26		94		6.1	6 (5.01	1-7.49)		33.1	2
Suicide	3	31	11	3	1	11		84		5.3	5 (4.41	-6.77)		29.5	9
Pneumonia	1	10	11	3	4	25		80		5.2	4 (4.18	3-6.48)		28.1	9
Cancer	2	22	21	1	6	16		75		4.9	1 (3.88	3-6.12)		26.4	2
CVE	1	11	14	1	0	15		50		3.2	28 (2.4	4-4.3)		17.6	2
COAD	1	14	10	1	3	11		48		3.1	5 (2.33	3-4.15)		16.9	1
RTA	1	11	0	1	3	0		24		1.5	7 (1.01	1-2.33)		8.46	ó

Cause of death					2004-2020		
	U	C (n)	C	C (n)	Total (UC + CC)	CSMF, % (95% CI)	Deaths /100,000
	Male	Female	Male	Female	Male + female		population per yr
Homicide	7	3	7	3	20	1.31 (0.8-2.02)	7.05
Snake bite	7	1	7	1	16	1.05 (0.60-1.7)	5.64
Drowning	5	1	5	1	12	0.79 (0.4-1.37)	4.23
ALD	6	0	5	0	11	0.72 (0.36-1.29)	3.88
Miscellaneous	81	77	93	77	328	21.49 (19.46-23.64)	115.56
Total	445	318	445	318		100	

RTA, road traffic accident; CVE, cerebrovascular episode; AIDS, acquired immunodeficiency syndrome; RF, renal failure; ALD, alcoholic liver disease

periods. In the 21-30 yr category, during 2010-2015 and 2016-2020, the proportion of male deaths was significantly higher than that of females (P<0.001 and P=0.032, respectively). In the 41-50 yr category, the proportion of male deaths was significantly higher than females (P=0.034) for the period 2010-2015. In the 51-60 yr category, for the period 2016-2020, the male deaths were significantly higher than that of females (P=0.003). Overall, for the last two study periods, the male deaths were significantly higher than females. The male deaths (all age groups (16-60 yr), 21-30 and 51-60 yr groups) were significantly higher than that of females for total 17 yr (P<0.001, P<0.001 and P < 0.001, respectively). There was a significant increase in deaths per 10,0000 population per year in the age group of 41-50 yr and 51-60 yr as compared to previous age groups. Table II shows the age wise distribution of cause-specific mortality during three study periods.

CNS infections were more common among the older age groups as compared to the younger one. TB was the most common CoDs in all age groups except in the 51-60 yr age group.

Table III provides the gender-wise distribution of CoDs and CSMF during the three study periods. The subjects comprised of 41.68 per cent females and 58.32 per cent males.

TB was the leading UC as well as CC of deaths acros sall age groups, equally distributed across both genders. Jaundice was the second leading CoD followed by heart diseases and diarrhoeal diseases. Heart diseases, jaundice, homicide, alcohol intoxication, tetanus, drowning, snakebite and alcohol liver diseases were more common in males. Suicide and road traffic accidents (RTAs) were significantly more common in males than females (*P*=0.046 and

0.018, respectively). Deaths due to tetanus in Melghat were significantly higher (7.75 per 100000 population per year) as compared to the rest of India (1.12 per 100,000 UI) and equal to the highly impoverished countries such as south Sudan (7.62 per 100,000 UI)¹⁷. Out of total deaths, 50 per cent of deaths are due to preventable infections. Deaths due to TB were also higher in Melghat than rest of the India.

Table IV and Figure 2 provides the seasonal pattern of major UCs of death during the three periods. During the period 2011-2015, the proportion of deaths due to TB and jaundice was significantly higher in the winter and summer seasons as compared to the rainy season. Further, the deaths due to TB were significantly higher in winter as compared to other seasons (P<0.001) and the deaths due to diarrhoeal diseases were significantly higher in the rainy season as compared to other seasons (P=0.013). The proportion of CNS infection was also significantly higher in the rainy season as compared to other seasons (P=0.002) and deaths due to heart diseases were significantly higher during the summer season (P=0.044). Chronic obstructive airway disease (COAD) was higher during the monsoon.

Discussion

In this study, the deaths in EPAGP were found to be higher than the national average. Health outcomes in this population have been reported to be poor because of poverty, low heath literacy, inaccessible and grossly inadequate healthcare services, low health seeking behaviour¹⁸, poor hygiene¹⁹ and high prevalence of addiction²⁰.

A high rate of premature adult mortality is reported as a major issue at the population level as it creates a negative impact on families and communities²¹.

CoD across EPAGP in India and worldwide shows respiratory infections, diarrhoeal diseases and non-communicable diseases as the major CoD^{1,3,22}, a similar trend was also observed in Melghat.

In the present study, TB was found to be the most common underlying CoD across all age groups indicating shortfalls in outreach of the existing TB control programme and the need for extensive revision of such programmes suitable for the tribal areas. The prevalence of TB is high in the tribal area of Melghat (>0.4%), twice as compared to the national average due to low health literacy, crowded living conditions, delayed diagnosis, high prevalence of smoking and malnutrition, inaccessible health facilities and poverty^{23,24}. Jaundice was reportedly the second most common UC across all age groups. This could be due to abuse of single use unsterile injections to multiple patients²⁵ by unlicensed healthcare providers, multiple sexual partners²⁶ drinking impure water²² and blood transfusion without testing for hepatitis virus²⁷. Heart and diarrhoeal diseases were the third and fourth leading CoD. High diarrhoeal mortality is because of impure drinking water (90%) and impure food, inadequate hand hygiene and a high prevalence of malnutrition (body mass index<18.5)²⁸. Intensive vaccination can prevent many deaths due to infections.

Heart diseases were the second leading CoD in the age group of 41-60 yr. The shift from infectious to non-communicable diseases in the adult age group shows the epidemiological transition in India²⁹. The risk factors for heart diseases in Melghat are high tobacco and alcohol use, high prevalence of hypertension (10% in EPAGP), low birth weight (43%)30,31 and severe malnutrition in the age group of 0-5 yr (20%)³⁰. Suicide is the sixth most common UC CoD mainly seen in 16-40 yr' group indicating significant mental health issues in younger population warranting attention³². Many of the suicides were under the influence of alcohol or due to family disputes. Depression and anxiety may go unnoticed in the tribal areas in the context of limited family support. The greatest burden of suicide is seen in young people in developing countries³³.

The probable causes of significantly high death rate in males in the present study were high risk behaviour of males *e.g.*, smoking, alcoholism, more outdoor activities, occupational hazards and mental health issues, *etc.* These findings are in coraboration with other studies⁷. In the present study, jaundice deaths were more frequent in males due to more high

			Table IV.		Seasonal pattern of major underlying causes of death during the three periods	r underlying	causes of	death duri	ng the thre	e peric	spo				
Causes of deaths		2004-2010			2011-2015			2016-2020	C		2004-2020	20	Deaths/1	Deaths/100,000 population per yr	pulation
	Rain (n=104), n (%)	Rain Winter Summer (n=104), (n=98), (n=91), n (%) n (%)	Summer (n=91), n (%)	Rain (n=151), n (%)	Winter (n=115), n (%)	Summer (n=98), n (%)	Rain (n=32), n (%)	Winter (n=35), n (%)	Summer (n=39), n (%)	Rain	Rain Winter	Summer	Rain	Winter	Summer
TB	16 (15.3)	16 (15.3) 22 (22.4) 15 (16.5)	15 (16.5)	9 (5.9)	29 (25.2)***	25 (25.5)***	6 (18.8)	7 (20)	6 (15.4)	31	58	46	32.77	61.30***	48.62
Diarrhoeal diseases 13 (12.5) 6 (6.1) 7 (7.7)	13 (12.5)	6 (6.1)	7 (7.7)	12 (7.9)	5 (4.4)	1(1)	4 (12.5)	1 (2.9)	2 (5.1)	29	12	10	30.65**	12.68	10.57
Heart disease	10 (9.6)	10 (9.6) 11 (11.2) 11 (12.1)	11 (12.1)	7 (4.6)	12 (10.4)	12 (12.2)	2 (6.3)	2 (5.7)	7 (17.9)	19	25	30	20.08	26.42	31.71*
CNS infection	(9.8) 6	6 (6.1)	4 (4.4)	21 (13.9)***	5 (4.4)	2 (2)	0	0	2 (5.1)	30	11	∞	31.71**	11.63	8.46
Jaundice	7 (6.7)	4 (4.1)	8 (8.8)	3 (1.9)	$14(12.2)^{**}$	8 (8.2)**	3 (9.4)	1 (2.9)	0	13	19	16	13.74	20.08	16.91
COAD	5 (4.8)	4 (4.1)	2 (2.1)	11 (7.3)**	1 (0.9)	2 (2)	0	4 (11.4)	3 (7.7)	16	6	7	16.91	9.51	7.4
Cancer	3 (2.9)	4 (3.1)	5 (5.5)	2 (1.3)	6 (5.2)	7 (7.1)	1 (3.1)	4 (11.4)	2 (5.1)	9	14	14	6.34	14.8	14.8
Tetanus	2 (1.9)	2 (2)	0	2 (1.3)	2 (1.7)	1 (1)	0	1 (2.9)	1 (2.6)	4	5	2	4.23	5.28	2.11
*Obtained using Pearson's Chi-square test. Rain; ra	arson's Ch	ni-square te.	st. Rain; rai	iny season. T	iny season. TB, tuberculosis; CNS, central nervous system; COAD, chronic obstructive airway disease	is; CNS, cen	tral nervo	us system;	; COAD, c	hronic	obstruct	ive airway	disease		

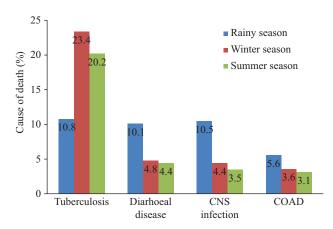


Fig. 2. Seasonal variation of causes of deaths during the study period (2004-2020). CNS, central nervous system; COAD, chronic obstructive airway disease

risk behaviour *e.g.*, multiple sex partners, alcoholism (50%), drinking impure water because of more frequent outdoor activities similar to previous reports^{4,7}. Diarrhoeal deaths were more common among males, because of more outdoor work particularly due to consumption of impure outdoor water and impure food, and lesser tendency for handwashing with soap. Similar findings have been found in other low-income settings, such as in Gaza, where the prevalence of diarrhoea is reportedly higher in males (5.4/100) as compared to females (1.3/100)³⁴. On the other hand in Melghat death due to heart diseases (coronary artery disease) are reportedly more common in males due to the higher prevalence of smoking, alcoholism and hypertension²⁰.

In the present study, suicide deaths were more common in males than females similar to previous reports, indicating significantly more personal/social reasons and mental disorderin males³⁵. Many suicides in Melghat were recorded as under the influence of alcohol. In present study, COAD (chronic obstructive airway disease) or COPD (chronic obstructive pulmonary disease) deaths were more common in males due to more smoking and outdoor pollution. RTA as CoD was significantly more in males than females. Motorbikes are the important mode of transport in Melghat due to inadequate public transport. Driving motorbikes, especially under the influence of alcohol is significantly higher amongst the tribal males. Similar findings were found in Iran³⁶.

In the present study, significantly higher deaths occurred during the rainy season. The difference in the proportion of deaths across season was significant (P=0.002). It is because of more deaths due to diarrhoeal diseases, CNS infections and COAD during the rainy season. This highlights that the onset of rainy season opens the door to various infections and diseases due to unhygienic conditions. Poor transport connectivity during this season further inhibits access to hospitals. In the present study, TB deaths were found to be maximum in the winter season in Melghat when the temperature reaches to 2°C and people live in enclosed huts without proper ventilation leading to a greater risk of TB exposure. It is in contrast to the majority of studies (n=49), where TB decreased during winter²⁴. In present study, diarrhoeal deaths were significantly more common during the rainy season, similar to previous studies³⁷. It is because of more impure drinking water and more flies infecting food during the rainy season. There is no system for water purification and control of flies in Melghat. Open defecation (>90%) exacerbates more contamination of water and contamination by flies (unpublished data). Deaths due to CNS infections were more common in Melghat during the rainy season because of unhygienic surroundings in villages, poor drainage systems and water logging leading to mosquito breeding and more cerebral malaria/viral encephalitis, similar to the findings from a rural hospital in Uganda³⁸.

In our study, COAD was more common during the rainy season with increased morbidity and mortality, due to the increased prevalence of respiratory viral infections⁶.

We used these CoD by VA to plan cluster randomized control trial (CRCT) to reduce ASMR. Our CRCT resulted in significant reduction in ASMR in intervention villages as compared to control villages (P<0.001), demonstrating the reliability of VA. Accurate information of CoD is essential for health policy, planning, monitoring, evaluation, comparisons, public attention and reducing premature deaths³⁹.

This study was not without limitations. The study team chose to use the physician-reviewer method for diagnosis. Diagnosis of TB was not made by bacteriological confirmation. The accuracy of VA to know the CoDs was 75 per cent. This was a long-term follow up study of 16 yr. During the study, the interviewers were changed and the quality of data obtained could have been affected due to the poor educational status of responders. Furthermore, some villages were dropped out of the study and new villages

were added in between the tenure. This may have limited the interpretation of trends in CoD over time.

Overall, our data showed that communicable diseases are the major CoDs (59.18%) in EPAGP. Providing oral rehydration solution (ORS) to all diarrhoea cases can save many lives immediately. Improving access to TB diagnosis and treatment services and ensuring treatment adherence can break the chain of transmission and prevent unnecessary deaths due to TB. It is essential to strengthen the routine death registration systems to cover all the locations and population groups across India.

As per our study finding from 2004 to 2021, the mortality rate in EPAGP among tribals in Melghat is more than 400 per 100,000 population of that age group per year. If we extrapolate our findings to all tribal areas of India, then, more than 208,846 people of EPAGP are dying every year⁴⁰.

Financial support & Sponsorship: The study was funded by Stichting Geron, Caring Friends, Mastek Foundation and the Tribal Development Department. The funders of the study had no role in study design, data collection, data analysis, data interpretation or writing of the report. The corresponding author had full access to all the data and had final responsibility to submit for the publication.

Conflicts of Interest: None.

References

- Mishra N, Barde PV, Awasthi S, Kumawat A, Gaur R, Singh P, et al. Profile of diseases prevalent in Datia district of Madhya Pradesh, India. Front Public Health 2022; 10: 926000.
- Thomas LM, D'Ambruoso L, Balabanova D. Verbal autopsy in health policy and systems: A literature review. BMJ Glob Health 2018; 3: e000639.
- Palanivel C, Yadav K, Gupta V, Rai SK, Misra P, Krishnan A. Causes of death in rural adult population of North India (2002-2007), using verbal autopsy tool. *Indian J Public Health* 2013; 57: 78-83.
- Nichols EK, Byass P, Chandramohan D, Clark SJ, Flaxman AD, Jakob R, et al. The WHO 2016 verbal autopsy instrument: An international standard suitable for automated analysis by InterVA, InSilicoVA, and Tariff 2.0. PLoS Med 2018; 15: e1002486.
- Indian Council of Medical Research. Study on causes of death by verbal autopsy in Tamilnadu State. New Delhi: ICMR; 2005. p. 37.
- Simões EAF, Dani V, Potdar V, Crow R, Satav S, Chadha MS, et al. Mortality from respiratory syncytial virus in children under 2 years of age: A prospective community cohort study in rural Maharashtra, India. Clin Infect Dis 2021; 73 (Suppl 3): S193-202.
- 7. Jha P, Gajalakshmi V, Gupta PC, Kumar R, Mony P,

- Dhingra N, et al. Prospective study of one million deaths in India: Rationale, design, and validation results. PLoS Med 2006; 3: e18.
- 8. Benara SK, Sharma S, Juneja A, Nair S, Gulati BK, Singh KJ, *et al.* Evaluation of methods for assigning causes of death from verbal autopsies in India. *Front Big Data* 2023; 6: 1197471.
- 9 Shah PS, Zao J, Ali S. Maternal marital status and birth outcomes: a systematic review and meta-analyses. *Matern Child Health J* 2011; 15: 1097-109.
- Finlay JE, Moucheraud C, Goshev S, Levira F, Mrema S, Canning D, et al. The effects of maternal mortality on infant and child survival in rural Tanzania: A cohort study. Matern Child Health J 2015; 19: 2393-402.
- Satav AR, Satav KA, Bharadwaj A, Pendharkar J, Dani V, Ughade S, et al. Effect of home-based childcare on childhood mortality in rural Maharashtra, India: A cluster randomised controlled trial. BMJ Glob Health 2022; 7: e008909.
- Van Rossem T, Deboosere P, Devos I. [De jure and de facto deaths]. The impact of unregistered attendees and absentees on urban death rates in early twentieth-century Belgium. *Population* 2018; 73: 89-114.
- 13. World Health Organization. *WHO technical consultation on verbal autopsy tools*. Geneva: WHO; 2007.
- 14. Murray CJL, Lopez AD, Black R, Ahuja R, Ali SM, Baqui A, et al. Population Health Metrics Research Consortium gold standard verbal autopsy validation study: design, implementation, and development of analysis datasets. Population Health Metrics 2011; 9:27.
- Murray CJL, James SL, Birnbaum JK, Freeman MK, Lozano R, Lopez AD. Simplified symptom pattern method for verbal autopsy analysis: multisite validation study using clinical diagnostic gold standards. *Popul Health Metr* 2011; 9:30.
- Murray CJ, Lopez AD, Barofsky JT, Bryson-Cahn C, Lozano R. Estimating population cause-specific mortality fractions from in-hospital mortality: Validation of a new method. *PLoS Med* 2007; 4: e326.
- Kyu HH, Mumford JE, Stanaway JD, Barber RM, Hancock JR, Vos T, et al. Mortality from tetanus between 1990 and 2015: Findings from the global burden of disease study 2015. BMC Public Health 2017; 17: 179.
- Busch SLP, Houweling TAJ, Pradhan H, Gope R, Rath S, Kumar A, et al. Socioeconomic inequalities in stillbirth and neonatal mortality rates: Evidence on Particularly Vulnerable Tribal Groups in eastern India. Int J Equity Health 2022; 21:61.
- 19. McGuinness SL, O'Toole JE, Boving TB, Forbes AB, Sinclair M, Gautam SK, *et al.* Protocol for a cluster randomised stepped wedge trial assessing the impact of a community-level hygiene intervention and a water intervention using riverbank filtration technology on diarrhoeal prevalence in India. *BMJ Open* 2017; 7: e015036.
- Madavanakadu Devassy S, Webber M, Scaria L, Amuthavalli Thiyagarajan J, Fendt-Newlin M, Joubert J, et al. Social and behavioural risk factors in the prevention and management of cardiovascular disease in Kerala, India: A catchment area population survey. BMC Cardiovasc Disord 2020; 20: 327.
- 21. Cao B, Bray F, Ilbawi A, Soerjomataram I. Effect on longevity of one-third reduction in premature mortality

- from non-communicable diseases by 2030: A global analysis of the Sustainable Development Goal health target. *Lancet Glob Health* 2018; 6: e1288-96.
- 22. Rathour RK, Sakhuja D, Bhatt AK, Bhatia RK. Municipal wastewater connection for water crisis and jaundice outbreaks in Shimla city: Present findings and future solutions. *Int J Environ Res Public Health* 2022; *19*: 1266.
- 23. Kashyap RS, Nayak AR, Husain AA, Shekhawat SD, Satav AR, Jain RK, *et al.* Impact of socioeconomic status and living condition on latent tuberculosis diagnosis among the tribal population of Melghat: A cohort study. *Lung India* 2016; *33*: 372-80.
- 24. Tedijanto C, Hermans S, Cobelens F, Wood R, Andrews JR. Drivers of seasonal variation in tuberculosis incidence: Insights from a systematic review and mathematical model. *Epidemiol* 2018; 29:857-66.
- Panda S, Kamei G, Pamei M, Sarkar S, Sarkar K, Singh ND, et al. Clinical features of HIV infection in drug users of Manipur. Natl Med J India 1994; 7: 267-9.
- Singh S, Prasad R, Mohanty A. High prevalence of sexually transmitted and blood-borne infections amongst the inmates of a district jail in Northern India. *Int J STD AIDS* 1999; 10: 475-8.
- 27. John TJ, Carman RH, Hill PG. Hepatitis B antigen and viral hepatitis type B in India. *Bull World Health Organ* 1974; 51:617-20.
- Guerrant RL, Kirchhoff LV, Shields DS, Nations MK, Leslie J, de Sousa MA, et al. Prospective study of diarrheal illnesses in Northeastern Brazil: Patterns of disease, nutritional impact, etiologies, and risk factors. J Infect Dis 1983; 148: 986-97.
- Nair M, Bherwani H, Mirza S, Anjum S, Kumar R. Valuing burden of premature mortality attributable to air pollution in major million-plus non-attainment cities of India. Sci Rep 2021; 11: 22771.
- 30. Dani V, Satav K, Pendharkar J, Satav A, Ughade S, Adhav A, et al. Community-based management of severe malnutrition:

- SAM and SUW in the tribal area of Melghat, Maharashtra, India. *Clin Epidemiol Glob Health* 2017; 5: 62-9.
- Mukherjee I, Kuiri S, Mahato KP, Dutta S, Bisai S. Low birth weight scenario in India: A systematic review and metaanalysis. *IJFANS Int J Food Nutr Sci* 2022; 11: 15.
- India State-Level Disease Burden Initiative Suicide Collaborators. Gender differentials and state variations in suicide deaths in India: The Global Burden of Disease Study 1990-2016. Lancet Public Health 2018; 3: e478-89.
- 33. Hawton K, van Heeringen K. Suicide. Lancet 2009; 373: 1372-81.
- 34. Abuzerr S, Nasseri S, Yunesian M, Hadi M, Mahvi AH, Nabizadeh R, et al. Prevalence of diarrheal illness and healthcare-seeking behavior by age-group and sex among the population of Gaza strip: A community-based cross-sectional study. BMC Public Health 2019; 19: 704.
- 35. Dandona R, Bertozzi-Villa A, Kumar GA, Dandona L. Lessons from a decade of suicide surveillance in India: Who, why and how? *Int J Epidemiol* 2017; *46*: 983-93.
- 36. Delavary Foroutaghe M, Mohammadzadeh Moghaddam A, Fakoor V. Time trends in gender-specific incidence rates of road traffic injuries in Iran. *PLoS One* 2019; *14*: e0216462.
- Nengminza Sangma RV, Rasania SK, Prasuna JG, Das R, Jais M. Seasonal variation of diarrheal diseases and drinking water quality in resettlement colony in Delhi, India: A community-based crosssectional study. *Indian J Public Health* 2021; 65: 387-90.
- 38. Williams EH, Hayes RJ, Smith PG. Admissions to a rural hospital in the West Nile District of Uganda over a 27 year period. *J Trop Med Hyg* 1986; 89: 193-211.
- Bang AT, Bang RA. Diagnosis of causes of childhood deaths in developing countries by verbal autopsy: Suggested criteria. The SEARCH team. *Bull World Health Organ* 1992; 70: 499-507.
- 40. Dubey M, Mohanty SK. Age and sex patterns of premature mortality in India. *BMJ Open* 2014; 4: e005386.

For correspondence: Dr Ashish Rambhau Satav, Mahatma Gandhi Tribal Hospital, Karmgram, Utavali, Dharni, Amaravati 444 702, Maharashtra, India

e-mail: drashish@mahantrust.org

Annexures

Annexure 1: Algorithm for respiratory diseases (pneumonia, TB and COPD) in verbal autopsy format.

II) Respiratory system disorder:		
1. Did the patient has cough	?	
Yes ()	No ()	Don't know ()
2. Was the patient breathless? W	Vas there any difficulty in breathing?	
Yes ()	No ()	Don't know ()
If both answers are no, then sk	ip subquestions 3 to 7 of question II a	and move to question III.
But if one or both answers are	yes, then continue the subquestions 3	3 to 7 of question II.
3. Since how long patient h	nad the cough?	
E (more than 24 hours)		
4. Was red expectoration so	een in cough?	
C Yes()	No ()	Don't know ()
5. Since when was the shortne	ess of breath?,	
C (more than 6 hours)		
6. Was there a fever?		
S Yes ()	No ()	Don't know ()
7. Was there any chest wal	l indrawing?	
C Yes ()	No ()	Don't know (
8. Was there chest pain?		
S Yes ()	No ()	Don't know ()
Did chest pain increased w	with deep inspiration or breathing?	
C Yes ()	No ()	Don't know ()

Pneumonia Possible= 1E + 1S Most probable = 1E + 1C + 2S or 1E + 2C

If the symptoms (cough or fever) of sub questions 3 to 6 from question II, lasted for more than 15 days, then ask sub questions 9 to 25 of question II. If not, then go to question III)

1	, 2 1	,
9. Since when the patient had cough	?	
E Yes (if more than 15 days, t	hen 1E) () No ()	Don't know ()
10. Was there blood in the cough?		
C Yes ()	No ()	Don't know ()
11. Was there history of weight lo	ss? (significant during last 1	5 days or more)
E Yes ()	No ()	Don't know ()
12. How was the patient's health in the	e last 15 days?	
E Sick()	was good ()	Don't know ()
13. Was patient severely fatigued du	ring last one month?	
S Yes()	No ()	Don't know ()
14. How was the appetite of the patient	for the last 30 days before death	?
S Less ()	good()	Don't know ()
15. Did the patient had TB (tuberc	ulosis)?	
S Yes ()	No ()	Don't know ()
16. Did the patient come in co	ontact with someone wh	no had TB? (tuberculosis)
S Yes ()	No ()	Don't know ()
17. Was the patient administered T	TB vaccine (BCG)?	
Yes ()	S No ()	Don't know ()
18. Was there any lump in the neck/s	armpit of the patient for more	than 15 days?
C Yes ()	No ()	don't know ()
19. Did the patient had neck pain o	or headache and vomiting fo	or more than 15 days?
S Yes ()	No (),	don't know ()

20. Was there any change in the voice	ce of patient, l	noarser	ness and difficulty in swallowing?
S Yes ()	No (),	don't know (),
21. Was there any breathlessness / sl	hortness of bro	eath/ in	tercostal indrawing? And since when?
C Yes ()	No (),	don't know (),
TB Possible = E1+ 1S +1C / 2E + 3S Most probable = 3E + 2C or 2E + 1C +5S or 2E+2C+2S		•	If cough persists for more than 15 days, then there is a robability of TB)
22. Was there any history of breathless for a long time? (because of which			
E Yes ()	No (),	don't know () ,
23. Was there any history of cough for	a long time?		
E Yes ()	No (),	don't know () ,
24. Did the patient sit day and nig position?	ht because of	f breatl	hlessness? Was he not able to sleep in the supine
C Yes ()	No (),	don't know () ,
25. Did the patient had cough with white e	expectoration for	at least	two months every year, for three consecutive years?
C Yes () No ()	, don't	know	()
Did the cough increase during cold/win	nter?		
C yes	No (),	don't know ()
S yes	No (),	don't know ()
26. Did the patient's lips, palms and	soles turn blu	e/cyan	osed ?
C Yes ()	No (),	don't know ()
27. Did the patient has frequent brea	thlessness?		
S Yes ()	No (),	don't know ()

28. Was the patient addict to smoking chillum/Bidis/cigarettes?

C Yes (-----) No (), don't know ()

COAD

Possible = E1+ 1S / 1E + 1C

Most probable = 2E + 2C / 1E +1C +2S / 1E +1C+3S

(Questions 22 to 28)

Annexure 2

महान ट्रस्ट, कममग्राम उत**ावली, म**ेलघाट k ctkt vilks fyfeVsM vikj vkfnoklh fodk1 folkkx] e- 'kk- & 1kStU; 1k (Verbal Autopsy Form) १६ – ६० वर्षे पयंत मृत्यु सर्वेक्षण (कारणमममाांसा)

ददनाांक: पावती क्र.

गाांव घर पररवार व्यक्त ी पहचान क्रमाांक ()()() - ()()() - ()()()()() - ()() १) मृतक का पूरा नाम : मपता का नाम जाटो
२) जन्मतारीखं (पता है तो): () ()/ () ()/ ()()() (ददन /ममहना/साल) ३) तलगः पुरुष ()स्नी
४) ग ा ंव क ा न ाम (हमेश ा रहन ेक ा) :
५) मरने दक तारीख: () ()/ () ()()()()()(ददन /ममहना/साल) ६)मृत्यू के समय उम्रः ()(
)()
७) जहााँ मृत्यू हुआ वह गााँव का नाम :
८) मुलाखत दक तारीख: () ()/ () ()/ ()()()() (ददन /ममहना/साल)
९) मरने का कारण :-

४)- कः सः। क्या मरं गया? क्यः। तकलाफ हुई था यह सम्बस्तर बताइयः। बतायः। हुआ वर्णमा मलमखए। बाहरकः।
मबम ार ी ह यह कारण न बताते हुए क्या तकलीफ थी, यह समजाने में मदत दकजीये _।
141011016 16 41111 1 41111 84 411111 11, 16 (11011 11 1141 41 9111

<i>१०) क््या ममह</i>	ला अर <i>ोग</i> ्यदत उपमस्थत थ ी ?	<i>ह</i> ा ()	
११)	वाले क ा न ाम:	मत क के साथ ररश्ताः	_
जानक ार ी दन			
सचनाः-			

१) कोई भी सवाल के सामने (- संे कम समय रहन े पर) दक			मरने के दकतन	ने ददन पहले से थी,(त	कलीफ एकददन
२) बड़े करिन सवालो क ा	भ ाग करक े	दो –मतन सवालों द्वार	ा जानका	र ी प <i>ु</i> यीये	
I) क्या मररज नीचे मलखे दकसी	कारण से मर गर	या			
१) अपघात	()	२) पानी में डूबकर	()	३) चोट	()
४) मगरने से	()	५) साांप काटने से	()	६) जहर खाने से	()
५) जगली ज ानवरों न े मार		ॉॅं. (क ीय ी
अपघात वरे रे बह ुत े =1 C					
II) सास दक मबमारी					
1. क्या मररज को खा ा ँसी र्थ	गे?				
ह⊺োঁ. (•)	नह ी ()	पत	ा नह ी ()	
2. क्या मररज को दम लगता थ	ग? सााँस फु	लती थी? क्या मररज को	सााँस लेने में	में कोई ददक्कत	
थी?हााँ. (यदी दोनों नहीं तो III पर		नह ी ()	पत	ा नह∂ी ()	
एक या दोनों हााँ होंगे पुमयये 3. खााँसी कब से थ					
E (२४ घांटों से ज्यादा)					
4. क्या खााँसी में लाल	। (ग ा ंजन े जै	ेसा) फे फड़ा जात	ा था?		
C हााँ.()	नही ()	ч	ता नही ()	
5. दम/सााँस फु लना कब से	ं था? ()			
C (६ घांटो से ज्यादा)					
6. क्या मररज बुखार था?					
s हााँ.()	नही ()	पर	ता नहीं ()	

7. क््या स ीन ा/पासलीया अांदर गखचरी	जाती थी?	•	
C हर्गां ()	नह ी ()	पत ा नहर्) ()
8. क्या स ी ने में दद थ ा ?			
C हांंं. ()	नह ी ()	पत ा नह ं) ()
क्या सााँस लेने से बढ़ता था?			
C हर्ााँ. (ह ोन े पर 1S)	ੀ ()	पत ा नह ी () (द ो नः	ो ं में स े दकतन े भ ी s
न् य ु ोमनय ा सांभव = 1E + 1S बहुदा =1E + 1C + 2 S अथवा 1E + 1S + 2	С		
सवाल 3 से 6 में तकलीफ (खााँसी या बुखा नहीं होंगी तो III पर जाइये	र) 15 ददन से ज्य	गदा समय तक होंगी तो सवा	ल ९ से २५ तक पुयीये यदी
9. क्या खा ाँ सी कब से थी? ()		
E हााँ (यदी 15 ददनसे ज्यादा होगा, त	तो 1E)	नहीं () पत	ग नहीं ()
10. क्या खााँसी में खून जाता था?			
C हााँ ()	नही ()	पता नहीं	()
11. क्या मररज ददन ब ददन सुखता ज ज्यादा समय से सतत) E हााँ(ा थ ा? (गत 15 ददन स े) पता नही (
12. आमखर 15 ददन में मररज की तबीयत कै	से थी?		
E मबमारी थी? ()	अ न	च्यी थी ()	पता नहीं ()
13. क्या मररज को एक माह से बहोत कमजो	री आयी थी?		
S हााँ ()	नही ()	पता नहीं (()
14. मरने के पहले 30 ददन के अांदर तक म	ररज का खाना कै	से था?	
s कम()	अच्या()	पता नही	()
15. क्या मररज क ो T.B. (क्षयर ो ग/स	खी) क ी मबम ा	र ी थ ी?	

s हााँ ()	नहीं ()	पता नही ()
16. क्या T.B (क्षयरोग) के मररज के स	ाथ इस मर ी ज क ा स	ना ंपक (स ाथ) आया थ ा ?
s हााँ ()	नहीं ()	पता नही ()
17. क्या मरीज को T.B. की लस (मब.सी.जी.)	लगाई थी?	
हााँ ()	s नहीं ()	पता नही ()
18. क्या मरीज के गले/ काख में 15 ददन से ज्य	ादा समय तक गािन थी?	
C हााँ ()	नहीं()	पता नही ()
19. क्या मरीज को 15 ददनसे ज्यादा	समय तक गदन। म्सर में	दद और उलरटया थी?
s हााँ ()	नहीं ()	पता नही ()
20. क्या मर ी ज के आवाज में बदल, घोग	रेपन तथा मनगलने मे	ं तकल ीफ ह ोत ी थ ी?
s हााँ ()	नहीं ()	पता नही ()
21. क्या दम/सााँस फ्ुलना/ पसमलया	अा ंदर मखच ी ज ान ः	ा थ ा? और कबसे ?
C हााँ ()	नहीं ()	पता नही ()
T.B. स वता = E1+1S+1C/2E+3S बहुदा = 3E+2C अथवा 2E+1C+5S अथवा 2E+2C+2S	(सुचना:1 <i>5</i> दर	इनसे १ माहके ऊपर खा ा ँसी
	फु लना रहनेप	र T. B. की सभावना ज्यादा ह
22. क्या मरीज को बहोत ददन से दम लगता थी?(मजसके वजह से वह काम नहीं कर	· ·	Ì
E हााँ ()	नहीं ()	पता नही ()
23. क्या मरीज को बहोत ददन से खााँसी थी?		
E हााँ ()	नहीं ()	पता नही ()
24. क्या मरीज दम के वजह से ददन रात बैि	ा रहता था? सो नहीं स	कता था?
C हााँ ()	नहीं ()	पता नही ()

	25. क्या मरीज को लगातार तीन साल से प्रत्येक था?	वषम हररोज	दो माह तक खााँ	सी में सफ़े द मचकट बलगम फे फड़ा आता
	C हााँ ()	नहीं ()	पता नही ()
	क्या वह खााँसी थांडी के ददन बढती थी?	नों		
	s हााँ ()	नहीं ()	पता नही ()
2	26. क्या मरीज के िओ, पैर, नीलें हुयें थे?			
	C हााँ ()	नहीं ()	पता नही ()
2	27. क्या मरीज को बार-बार दमा होता था?			
	s हााँ ()	नहीं ()	पता नहीं ()
2	28. क््या मर ीज क ो ददन स े म्बड ी /मचर	लम/मसग ा	र ेट मपन े क ी आव	रत थी?
	C हााँ ()	नहीं ()	पता नही ()
	COAD स वत ा = E1+ 1S / 1E + 1 C बहुदा = 2E + 2C / 1E +1C +2S / 2E +1C+2S	S		
	(सवाल 22 से 28 तक)			
Ш) हृदयरोग (IDH)			
1	ı. क्या मर ीज क ो स ीन े में (ब ाय ह ुआ थ ा (वजन रख ा ज ैस े लगत ा		ाबचमें) दद	
	E हााँ ()	नहीं ()	पता नही ()
	2. क्या मर ीज क े स ीन ेमे ं घुटन जं स े बढत ी थ ी ? और आर ाम करने से क	_	_	ाहट ह ो रह ी थ ी? ज ो कम करन े
	C हााँ ()	नहीं ()	पता नही ()
3	3. क्या स ीन े क ा दद बाये ह ाथ में	ं गया थ ा	?	
	C हााँ ()	नहीं ()	पता नहीं ()

4. क्या मरीज को बहोत पसीना आता	था?		
s हााँ ()	नहीं()	पता नही ()	
5. क्या मरीज को चलने से कम करने र	से या सीडी चढ़ने से दम लगता १	भा?	
s हााँ ()	नहीं()	पता नही ()	
 क्या मरीज चक्कर आके गगर ग 	या थ ा ?		
s हााँ ()	नहीं ()	पता नही ()	
7. क्या मरीज की ददल की धड़कन बढ़	ती थी?		
s हााँ ()	नहीं()	पता नहीं ()	
8. क्या मरीज को उलटी हुई थी?			
S हााँ ()	नहीं ()	पता नही ()	
9. क्या मर ीज क ो ता ंबाख ू, ¹	बड ी,मचलम, मसग ार ेट य ः	ा ग ा ंज ा मपन े क ी आदत थी ?	
s हा ँ ()	नहीं ()	पता नहीं ()	
10. क्या मरीज को पैर से उपर तक सर	जन (उलटी सजन) आई थी?		
s हााँ ()	नहीं ()	पता नही ()	
हृदयरोग (IDH) स वत ा = 1E+ 1S/1E + 1C बहूदा = 1E + 1C + 2S/या 2C अथवा	1E +2C		
IV) पतली दस्त (दस्तकी बीमारी)			
1. क्या मरीज को पतली तही व	पत ा नह ी () यद ी ह ा ाँ, त ो एक ददन मे	
रात में ज्यादा से ज्यादा दकतने में और रात तीन से ज्यादा बार पतले		कम ह ो न $ ceil$ पर $ m V$ पर जाइय $ ceil_{ m E}$ एक पें $ m $	ददन
2. क्या तृी में खुन या बहोत	ऑव गगरत ी थ ी?		
C हााँ ()	नहीं (पता नहीं ()	

मडसट्रूं ी बह ्त ं = 1E + 1C यद ी ह ा त ो VI पर जाइय े यदी पता नहीं तो सवाल 3 पमू यय े			
3. दकतने ददन से पतली दस्त शुरू थी?	()		
लबी हगवन पतली दस्त बहुुते = 1E + 1C मसफ सवाल 1 और 3 क मलय। C १५ ददन से ज्यादा ददन तक			
१५ ददन से कम समय होगा तो सवाल	४ से ११ पूमयये		
4. क्या दस्त पाणी जैसे पतली (ढाळ) हो	ती थी?		
C हााँ ()	नहीं()	पता नही ()	
5. क्या मरीज को उलटी होती थी?			
s हााँ ()	नहीं()	पता नही ()	
6. प्यास कै सी थी?			
C बढ़ गई थी ()	नहीं()	पता नही ()	
7. क्या ऑखे गहरी अांदर गई थी?			
C हााँ ()	नहीं()	पता नहीं ()	
 आखरी दो ददनमें दकतनी पेशः 	ाब ह ोत ी थ ी ?		
C कम∣कुय भी नह ी ं () हमेशा जैसी () पत ा नह ी ()	
9. पेशाब का रंग कैसा था?			
C गद य ा बह ोत मपली () हमेश ा ज ैस ी () पत ा नह ी ()
10. क्या मरीज को दस्त के समय	प पान ी / शरबत/ जीवनरक्षक घोत	न मपलान ा शुर ु रख ा था ?	
बहोत () ९ कम	नहीं ()	पता नहीं ()	

नतली दस्त

संभव = 1E बह_ु ते = 1E + 2C य**ा** 1E + 1C + 2S

V) रॅबीज

1. क्या मरीज को कु त्ते, जांगली जानवर ने, क	न्टा था?		
E हााँ ()	नहीं ()	पता नही ()
2. क्या मर ीज मरन े स े क ु य ददन वजह स े प ान ी नह ीं मपत ा थ			
)		
3. क्या मरीज मरने से पहले हवा से डरता थ	Π?		
C हााँ ()	नहीं ()	पता नही ()
* यदी 1 भी हााँ होगा तो 4 से 11 तक	सवाल पुयीये		
st यदी सभी नहीं होंगे तो $ m VII$ पर जाइये $ m I$			
4. क्या मरीज मरने से पहले पागल जैसे कर	ता था? काटता था?		
C हाा ँ ()	नहीं ()	पता नही ()
5. क्य ा मर ीज मरने के क ु य घांटे ब	वहोश हुआ थ ा य ा म्बल	कु ल शाांत हुॐ	॥ था?
s हााँ ()	नहीं ()	पता नही ()
6. क्या मरीज को मजस जगह कु ते न	े कट ा थ ा वहााँ इज आ	ग/जलन/स्पशमवेदन	ा ह ोन ा य ा
स्राय ु दक हलचल थ ी? C हााँ () नहीं ())	पता नही (
7. क्या मरीज के मुाँह से फेस मनकलता था?			
C हााँ ()	नहीं ()	पता नही ()
8. एक ाश, स्पश य ा आवाज स े क्या मरी	ज डरत ा थ ा?		
s हााँ ()	नहीं ()	पता नही ()
9. क्या मरीज म्बच-मबचमें मानमसक र	रुपस े म्वकलाांग ह ो त ा	थ ा?	
s हााँ ()	नहीं ()	पता नही ()

क्या मरीज को दो-दो चीजे ददखत	न ी थ ी'	? क्या मरीज क	चेहरा गतरय	ग्रा था?
S हा ा ँ ()	नहीं ()	पता नही ()
क्या मरीज को मनगलने में ददक	त ह ोतर्	ો થ ી ?		
S हाा ँ ()	नहीं ()	पता नही ()
ਦੱਕੀज संभव = 1E + 1C / 1C +2S ਕੁਫ਼ _ੂ ਜ ੇ = 1E + 3C ਧ ਾ 1E + 2C + 2S	/3C+3S			
ग्ड्स				
या एक माह से ज्यादा ददन तक बुखार था?				
E हा ँ ()	नहीं ()	पता नही ()
				म ह _ु आ? (सुखी पता नही (
त्या मरीज को मरने से पहले एक माह से ज्या	दा समय र	से लगातार पतली दर	स्त शुरु थी?	
E हा ाँ ()	नहीं ()	पता नही ()
या मरीज को 1 माह से ज्यादा समय तक खा	ा ँ सी थी [,]	?		
हााँ ()	नहीं ()	पता नही ()
ज्य ा मर ीज क ो नामगन क ी मबम ार ्	ी (चमड़	ी पर अांग ार ,ल	ालसर च्ः औ	र प ान ीवाल े फोड़) थी?
हााँ ()	नहीं ()	पता नही ()
त्या मरीज के जबान (जीभ) पर सफ़े द दाग ((बुरशी) अ	ायी थी?		
हााँ ()	नहीं ()	पता नही ()
्य ा मर ीज क ो टी.बी. क ी म्बम ार ्	ी थी?			
हााँ ()	नहीं ()	पता नही ()
_् य ा मर ीज क ो पेश ाब क ी जगह ह ग	⊣ेशा फ	ोड़ा/फस ी ह ोत	ી ય ી ?	
हााँ ()	नहीं ()	पता नही ()
	हांँ (हााँ () नहीं (क्या मरीज को मनगलने में द्यक्त होतर् हााँ () नहीं (रँबीज संभव = 1E + 1C / 1C +2S बहुते = 1E + 3C या 1E + 2C + 2S/3C+3S ह्स या एक माह से ज्यादा ददन तक बुखार था? हााँ () नहीं (या मरीज का वजन आखरी य माह में १९०० हो मबमारी थीं? हहााँ () या मरीज को मरने से पहले एक माह से ज्यादा समय तक खााँ सी थीं हााँ () नहीं (या मरीज को 1 माह से ज्यादा समय तक खााँ सी थीं हााँ () नहीं (या मरीज को नामगन की मबमारी (चमड़क हााँ ()) नहीं (या मरीज के जबान (जीभ) पर सफ़ेद दाग (बुरशी) अहााँ () नहीं (या मरीज को चीं तीं. की मबमारी थीं? हााँ (क्या मरीज को मनगलने में दरकत होती थी? रंबीज संभव = 1E + 1C / 1C +2S बहुते = 1E + 3C या 1E + 2C + 2S/3C+3S ड्स या एक माह से ज्यादा ददन तक बुखार था? हााँ () नहीं () या मरीज को मरने से पहले एक माह से ज्यादा समय से लगातार पतली दर्स हााँ () नहीं () या मरीज को मरने से पहले एक माह से ज्यादा समय से लगातार पतली दर्स हााँ () नहीं () या मरीज को 1 माह से ज्यादा समय तक खााँ सी थी? हााँ () नहीं () या मरीज को वजन आखर विवास समय तक खााँ सी थी? हााँ () नहीं () या मरीज को वजन जामगन की मबमारी (चमड़ी पर आंगार, लहां () नहीं () या मरीज के जबान (जीभ) पर सफ़ेद दाग (बुरशी) आयी थी? हााँ (क्या मरीज को मनगलने में द्वकत होता थी? हाँ (

9. क्या मरीज के पेशाब से सफ़ेद मपप जाता था?				
s हााँ ()	नहीं ()	पता नही ()
10. क्या मरीज के शारीर पर बहोत गािने आयी	થી?			
C हााँ ()	नहीं ()	पता नही ()
11. क्या मरीज वेश्या के साथ या म	ववाहबाह्य	सांभोग करता	थ ा ?	
s हााँ ()	नहीं ()	पता नही ()
12. क्या मरीज को मरने के एक साल पहले कभी	खुन लग	ाया/चढ़ाया था?		
s हााँ ()	नहीं ()	पता नही ()
13. क्या मरीज के पती/पत्नी को या	बच्चे क ो	एड्स क ी म्बमान	री है	
s हााँ ()	नहीं ()	पता नही ()
14. क्या मरीज को पीमलया था?				
s हााँ ()	नहीं ()	पता नही ()
15. क्या मरीज को भूख नहीं लगती थी?				
s हााँ ()	नहीं ()	पता नही ()
एड् स संभव = 2E + 2S या 2E + 1C बहुते= 3E + 1C या 3E + 2S या 2E+2C+3S अथवा 2E+1C	2 +5S			
1. क्या मरीज को झटके /फीट (दाटी) आये थे? कब				
C हााँ ()	नहीं (पता नही ()
2. क्या मरीज का शरीर कड़क हुआ थ			ियेे झुका?	
C हााँ ()	नहीं ()	पता नही ()

3. क्या मरीज गुँह हमेशा जसे पूरी तरह से खोल नहीं सकता था?

हााँ ()	E नहीं ()	पता नही ()
क्या मरीज की दातखील लगातार	बैिी?			
E हााँ ()	नहीं ()	पता नही ()
एक भ ी मबम ार ी क ा लक्षण न त ो सव ाल स े 4 से 14 तक पूमय		/III पर [ः]	जाईये∣ और एक भी म्ब	बम ार ी क ा ल क्षण ह ोग ा
4. क्या मरीज के झटके आवाज या प्रक	गश बढ़ते थे?			
S हांाँ ()	नहीं ()	पता नही ()
5. क्या मरीज आखरी क्षण तक होश में	था?			
C हा ा ँ ()	नहीं ()	पता नही ()
6. क्या मर ीज क ो लोहे क			_	
जखम गांद ी ह ुई थ ी? क ्या गभग उस जगह पर गोबर लगाया था?	मवता महाला का जच	का क स	तमय नल गादा अवजार स	काटा था? काटन क बाद
s हााँ ()	नहीं ()	पता नही ()
7. क््या लोग इस म्बम ार ी क <i>ो</i>	ा धन _{ुवामद} बोलत े ह	ò		
s हााँ ()	नहीं ()	पता नही ()
8. क्या मरीज को धनुवामद के	ं स्टकें नह ी ं गमले	ो थेे?		
हााँ ()	E नहीं ()	पता नही ()
9. क््या मर ी ज क ो मनगलन े में	ं तकलीफ ह ोत ी थ	ी?		
s हााँ ()	नहीं ()	पता नही ()
10. क्या मरीज का पेट बहुत कड़क थ	Π?			
s हा ाँ ()	नहीं ()	पता नही ()
11. क्या मरीज का चेहरा पुतले के ज	से हुआ था?			
S हााँ ()	नहीं ()	पता नही ()

12. क्या मरीज का चेहरा, म्सना, कमर, पेट कड़क थे और हाथ पैर सही थे?

s हााँ ()	नहीं ()		पता नही ()
13. क्या मरीज का शरीर का कोई भाग जल गया ददन के अांदर) जचक ी ह <i>ु</i> ई थी? मनजमतुक				_	
s हााँ ()	नहीं ()		पता नही ()
धनर् ु ाँ द संभव = 1E + 1C अथवा 1C + 2S	20125				
VIII) मपलीया/कावीळ					
1. क्या मरीज की आाँखे मपली थी?					
E हााँ ()	नहीं ()		पता नही ()
2. क्या मरीज की पेशाब मपली थी? E हााँ ()	नहीं ()		पता नही ()
3. क्या मरीज को उलटी जैसे लगता था? S हााँ ()	नहीं ()	पता नही	()
4. क्या मरीज को बार –बार उलटीया होती थी?					
s हा ाँ ()	नहीं ()	पता नही	()
5. क्या मरीज को बुखार था?S हााँ ()	नहीं ()	पता नही	()
6. क्या मर ी ज के पेट दद थ ा? s हा ाँ ()	नहीं ()	पता नही	()

7. क्या मरीज को भुख नहीं लगती

थ ी?				
s हा ँ ()	नहीं ()	पता नही ()
8. क्या मरीज का वजन कम हो	रहा था?			
s हााँ ()	नहीं (()	पता नही ()
9. क्या मरीज को मरने के एक र	माह पहले खुन लगाया?			
s हााँ ()	नहीं (पता नही ()
10. क्या मरीज को बह	ोत द ारू मपने क ी	आदत थ ी ?		
s हााँ ()	नहीं (()	पता नही ()
11. क्या पेट में गोला था?				
s हााँ ()	नहीं (पता नही ()
म पलीय ा संभव = 1E बह _ु ते = 2E + 3S/ 1E + IX) मदू (ममस्तष्क) पर सजन	5S			
1. क्या मरीज बहोत सुस्त या बेह	होश था?			
नहीं ()	s हााँ()	पता नही ()
2. क्या मरीज को झटके /दाटी अ	ाई थी?			
C हा ँ ()	नहीं (()	पता नही ()
3. क्या मरीज को बहोत (मनरथः	मक) करता था? या पाग	ल जैसे बाते करता था	?	
नहीं ()	C हााँ()	पता नही ()
4. क्या मरीज को बुखार था?				
E हााँ ()	नहीं (पता नही (
5. क्या बुखार के साथ पसीना उ	भाता था?			

6. क्या बुखार के साथ पसीना आता था?				
s हा ा ँ ()	नहीं ()	पता नही ()
7. क्या लगातार म्सरदद और उत्तरटयााँ ह	ोत ी १	4ٲ?		
s हााँ ()	नहीं ()	पता नही ()
8. क्या प्रकश मरीज को सहन नहीं ह लेता था?	ोत ा थ	ा? आँखोंपर एक	ाश डालने क	े बाद वह तरत आँख बांद कर
s हााँ ()	नहीं ()	पता नही ()
9. क्या मरीज के कान से गपप गनकलत	ा था?			
s हााँ ()	नहीं ()	पता नही ()
10. क्या कड़क या ताि रहत ी गदन थ ी? य ा गदन				
s हााँ ()	नहीं ()	पता नही ()
11. क्या मरीज को बुखार के साथ ख	गा ँ स ी ,	दम, पेेशाब में ज	लन पेट मेें	दद, इत्याद ी ह ोत्त ा था?
s हााँ ()	नहीं ()	पता नही ()
मंपलीया संभव = 1E +1C /1E + 5S बहुते= 1E + 2C या 1E +1C+5S				
12. क्या मरीज का एक हााँथ और एव	क पैर व	ज्मजोर हुआ था? (व	ाह ह ा ँथ,प ै	र महला
नह ी ं सकत ा था?) C हा ाँ ()	नहीं ()	पता
नही ()			
13. क्या मरीज का मुाँह तेढा हुआ था?				
C हााँ ()	नहीं ()	पता नही ()

CVE (ममस्तष्क म**े**ं ख**ु क**ी म**बम**ारी) संभव = 1C

X) ॲनेममया (खुन की कमतरता)

1. क्या मरने के पहले म	ार ीज क ी जबान,ओि,	आँखे क ी पलक े	(अांदरस्े),	हाथ के त	लवे,
नाखुन वैगरे सफ़ेद हुए थब्न	ा गुलाबी नहीं थे?)				(লাল
E हााँ ()	नहीं ()	पता नही ()	

 क्या मरने कु य ददन उलटी,बच्चेदानी) खुन बह र कमल, मचकट तृृी हुई थी? 				•
C हााँ ()	नहीं ()	पता नही ()
3. क्या मरीज को बहोत कमजो	ारी (थकान) महसुस होती १	गी?		
s हााँ ()	नहीं ()	पता नही ()
4. क्या मरीज थोडासा चलने पर	र या काम करने पर सॉस पु	र लती थी?		
s हााँ ()	नहीं ()	पता नही ()
5. क्या मरीज को खुन क	ी कमतरत ा क ी म्बम	ार ी बत ाय ी गः	ई थी?	
s हााँ ()	नहीं ()	पता नही ()
6. क्या मरीज को बार-बार खुन	लगाने की (चढ़ाने की) ज	रुरत पड़ती थी?		
C हाा ँ ()	नहीं ()	पता नही ()
7. क्या मरीज के तृी मे	ं (कृ गम) मगरते थेे?			
8. क्या मरीज को मसकलसेल (ज	नोड़ो के दद) की मबमारी र्थ	† ?		
s हााँ ()	नहीं ()	पता नही ()
संभव = 1E + 2S / अथर बहु ते = 1E = 1C+2S	अॅ गमय ा वा 1C + 2S अथवा 1E + 2C अथवा 20	C + 2S अथवा 1C+3	S	

XI) मह का कसर य**ा ककरोग**

1. क्या मुह में रिक न ह ोन ेव ा ली द	दरमहत जखम (फोड़ा/उज्जु) ब	होत ददन से थी।
E हााँ ()	नहीं ()	पता नही ()

 $2.\ arphi$ ्य**ा मुह में रिक न ह**ोन**ेव**ाली ददरमहत क**ो**ई ग**ि**ान आय**ी** थ**ी**|

	E हााँ ()	नहीं ()	पता नही ()
3.	क्या मुह के गिान या जखम से ख	ान ा मन	गलन े में क ोई तक	लीफ थी।	
	S हााँ ()	नहीं ()	पता नही ()
4.	क्या मुह के गिान या जखम बहोत	त गांद ा व	बास आत ा थ ा?		
	s हााँ ()	नहीं ()	पता नही ()
5.	क्या मरीज के मुह में सफ़ेद दाग () आया	और दफर गिान	फोड़ा जख	म तैयार हर्इ
	C हााँ () सब नहीं होंगा XII पर जाइये एक भी हां) तो सवाल 6 से 9 तव	पता नही (रु पूयीयें)
6.	क्या मरीज का वजन ददन ब ददन कम	ह ो ने ज	ज ा रह ा थ ा ? (क््या मर <i>ी</i> ज	न ददन ब
	ददन सुख रह ा था ?) C हााँ ()	नहीं ()	पता
	नहीं ()			
7.	क्य ा मर ीज क ी भ ुख बह ोत कम ह ुई	ई थ ी ?			
	s हााँ ()	नहीं ()	पता नही ()
8.	क्या मरीज के जखम से खुन बहता था?				
S	ड हा ा ँ ()	नहीं ()	पता नही ()
9.	क्या मरीज को तम्बाखु, बीडी य	ा गचलम	, दारू सेवन कर	न े क ी आदत	थी?
S	ऽ हा ा ँ ()	नहीं ()	पता नही ()
•	यदी हााँ तो कौनसी आदत थी? ()		
	मु का कन्सर				
	संभव = 1E + 1S बहुुते = 1E = 1C+1S अथवा 1E + 4S C	OR 2F +	25		
	400 00 = 15 = 10 110 01401 15 + 40 C	/X ZL T	20		

XII) क्या और दसरी कोई तकलीफ थी?

कौनसी

		·

	अन्य			
वर	जह	पता	नहीं	
ईलाज	ज दकय ा क्या ?			
हााँ	() नहीं (
) पता नहीं ()यदी हााँ तो,			
दकसन	÷			
क्या ईल	गाज दकया			
मबमार	री कौनसी बताई?			
पयमवेक्षव	p/मागमदशमक क ा मनष्कषम			
1)				
2)				
3)				
4)				
5)				
1.	मतृ क के	नाम		सही / अघििा
	माता			

ददनाक: /

/ २०

	मपता	
	पमत	
	पमत	
	लड़का	
	लड़की	
	(दकसी दो का)	
<i>सह</i> ी / उ	भ घ िा	
2. सरप	ग्रांच का नाम:	
<i>सह</i> ी / उ	भ र्घ ि ।	
3. पोर्ल	ीस पटेल का नाम:	
सही / अध	ग ि ा	
मागमद	.शाक / पयमवेक्षक क ा न ाम:	

उम्रः()()()

MAHAN TRUST, KARMGRAM UTAVALI, MELGHAT

पहचान क्रमाांक

६) मरने दक तारीख : () ()/ () ()()()() (ददन /ममहना/साल)

()()		
१) मृतक का पूर ा नाम :	मपता का नाम	_जाटो
२) जन्मतारीख (पता है तो): () ()/ (तलग: पुरुष ()स्नी ()) ()/ ()()()() (ददन /ममहना/स	∃ाल) ३)
४) ग ा ंव क ा नाम (हम ेशा रहन ेक ा) नाम :	: ५) जह ा मृत ्य ू हुआ वह 	ह ग ा ँव क ा

 $(\ \)(\ \)(\ \)-(\ \)(\ \)(\ \)-(\ \)(\ \)(\ \)(\ \)(\ \)-$

७)मृत्यू

समय

	Inference on causes of death	Possible	Most	Disease
			probable	code
1)	Pneumonia			
2)	Tuberculosis			
3)	COAD			
4)	IHD			
5)	Dysentry			
6)	Chronic diarrhoea			

7)	Acute diarrhoea			
8)	Rabies			
9)	AIDS			
10)	Tetanus			
11)	Jaundice			
12)	CNS infection			
13)	CVE			
14)	Fever cause unknown			
15)	Anemia			
16)	Oral cancer/malignancy			
17)	Other			
18	Cause not Known			
Underlying Cause()				
			Singnature:	