# Doubling of world population unlikely

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Most national and international agencies producing population projections avoid addressing explicitly the issue of uncertainty. Typically, they provide either a single projection or a set of low, medium and high variants<sup>1,2</sup>, and only very rarely do they give these projections a probabilistic interpretation. Probabilistic population projections have been developed for specific industrialized countries, mostly the United States, and are based largely on time-series analysis<sup>3</sup>. On a global level, time-series analysis is not applicable because there is a lack of appropriate data, and for conceptual reasons such as the structural discontinuity caused by the demographic transition<sup>4-6</sup>. Here we report on a new probabilistic approach that makes use of expert opinion on trends in fertility, mortality and migration, and on the 90 per cent uncertainty range of those trends in different parts of the world. We have used simulation techniques to derive probability distributions of population sizes and age structures for 13 regions of the world up to the year 2100. Among other things, we find that there is a probability of two-thirds that the world's population will not double in the twenty-first century.

The probabilistic projections are based on distributions for fertility, mortality and migration in all regions, defined in terms of high or low values assumed to cover 90 per cent of all possible future outcomes. For today's high-fertility countries they are based on an assessment of their current standing in the process of demographic transition towards low fertility<sup>7</sup>, together with information about reproductive intentions<sup>8</sup>. These data show that even in sub-Saharan Africa the fertility transition has started, and that it is well advanced in most other developing regions. The high and low assumptions for the years 2030–2035 are total fertility rates (TFR, the number of children per woman) of 4.0 and 2.0 in Africa, central Asia and the Middle East, 3.0 and 1.7 in southern Asia, Pacific Asia and Latin America and 3.0 and 1.5 in Central East Asia (mostly China). For today's industrialized countries the assumptions are based on

a broad survey of possible future societal changes9. The United

Nations and other institutions have assumed that fertility will eventually recover to replacement level (TFR slightly above 2.0), but there is little support for this view<sup>10,11</sup>. Accordingly, TFR values of 2.1 and 1.3 for Europe and the Pacific members of the Organization for Economic Cooperation and Development (OECD countries) and 2.3 and 1.4 for North America have been assumed for 2030–2035.

Assumptions for mortality were set in terms of increase in life expectancy at birth per decade. Contrary to earlier beliefs, there now is a considerable degree of uncertainty about the future course of mortality. In the industrialized countries this stems from the scientific dispute of whether we are already close to a biologically determined limit to life expectancy<sup>12,13</sup>. Accordingly, increases of 3.0 and 1.0 years have been assumed as the 90 per cent range. In the developing countries the uncertainty is more associated with future trends in AIDS<sup>14</sup> and other infectious diseases and the development of health services<sup>15</sup>. For certain regions possible problems with food supply have also been considered<sup>16,17</sup>. Consequently in such cases the assumed range of mortality improvement is wide, for example +4.0 to -2.0 years per decade in sub-Saharan Africa.

Migration is most difficult to handle because of unreliable data and high volatility<sup>18</sup>. For this study a matrix of constant annual interregional migration flows was assumed with the 90 per cent ranges covering two million to zero migration gains in North America, and 1 million to zero in Western Europe.

The projections for 13 regions (see Table 1) show that population growth will probably be most rapid in the middle East, sub-Saharan Africa and North Africa, with a tripling of the population by 2050 and a quadrupling by 2100 likely. Despite this rapid growth, there will also be significant increases in the proportion above 60 years of age. In contrast, in Eastern Europe and the European part of the former Soviet Union, population will probably decrease over the coming decades. By 2050 the Pacific OECD countries and Western Europe are likely to experience little, if any, change in population size. This stagnation or shrinkage in population size in Europe and the Pacific OECD countries will be associated with significant ageing of the population, with the proportion above 60 likely to double from its current values. Even proportions well above 40 per cent are within the 95 per cent confidence interval. These could bring serious consequences for social security systems.

In North America, a younger age distribution, a larger inflow of migrants, and slightly higher fertility than in Europe is likely to result in a roughly 25 per cent increase in population by 2050. Population ageing will occur, but will not be as dramatic as in Europe. Latin America is likely to have a doubling of its population and an increase in its proportion above age 60 to about 20 per cent. By 2050, Central East Asia (mostly China) is likely to grow by 37 per cent and experience an increase in the proportion over 60 from 9 per cent to 25 per cent. Southern Asia (essentially the Indian



Figure 1 Fractiles of the probability distribution of the future size of the world population.



Figure 2 Fractiles of the probability distribution of the proportion of the world population above 60 years of age.

Table 1 Result of probabilistic projections for 13 world regions									
Region	Total population (millions)					Population above age 60 (in %)			
	1995	2050				2050			
		Median	2.5%*	97.5%*	1995	Median	2.5%*	97.5%*	
Africa North Africa Sub-Saharan Africa	162 558	439 1,605	309 1,085	583 2,316	5.9 4.7	13.3 9.2	9.4 6.9	19.2 12.8	
East Asia Central East Asia Pacific Asia Pacific OECD countries	1,362 447 147	1,865 796 146	1,351 579 117	2,574 1,047 182	9.2 6.8 19.4	24.9 19.4 39.5	17.8 14.4 31.5	34.1 26.5 48.7	
West Asia Central Asia Middle East Southern Asia	54 151 1,240	137 515 2,368	88 380 1,833	206 692 2,970	7.8 5.4 6.7	15.4 12.5 16.6	10.2 9.1 13.4	24.0 17.3 20.8	
Europe Eastern Europe Former Soviet Union (European part) Western Europe	122 238 447	110 188 471	86 144 370	141 241 584	16.7 16.9 18.6	34.0 34.1 35.0	26.7 26.3 27.5	43.4 44.5 43.9	
Latin America	477	925	707	1,177	-7.6	20.4	15.8	26.4	
North America	297	403	303	534	16.4	30.2	24.0	38.6	

Fertility, mortality and migration are assumed to be independent.

\* Columns labelled 2.5% and 97.5% provide data on the lower and upper bounds, respectively, of the 95 per cent confidence interval.

subcontinent), which still has relatively high fertility and a young population, will probably double its population by 2050 and will be the world's most populous region.

The global results for population size are presented in Fig. 1; those for the proportion above 60 years of age are shown in Fig. 2. The median path of world population growth will increase from 5.8 billion today to 7.9 billion in 2020 and 10.0 billion in 2050. It will reach a peak around 2070-2080, and then begin a slow decline. In 2020 the range of uncertainty of this projection will be rather narrow (with the 95 per cent confidence interval between 7.5 billion and 8.3 billion) because many of the people who will be alive at that date have already been born. By 2050, the 95 per cent confidence interval will widen to between 8.1 billion and 11.9 billion, but the most likely 60 per cent (medium and dark shaded area in Fig. 1) still covers a range of less than one billion. After 2050 the size of the 95 per cent confidence interval will increase substantially, with the 60 per cent confidence interval showing a much smaller rise. The proportion of the world's population above age 60 is likely to increase from 9.5 per cent today to 20 per cent in 2050 to 27 per cent by 2100. A strong increase in the proportion of elderly people is virtually certain, with the low end of the 95 per cent confidence interval showing almost a doubling of today's level.

These probabilistic projections lead us to believe that the focus of public, political and scientific concern will continue to shift from global population growth to population ageing.  $\Box$ 

#### Methods

The probabilistic population projections are based on the multistate cohortcomponent model of population projections, which applies assumed agespecific fertility, mortality and migration rates to the age and sex distribution of the starting population along cohort lines<sup>19,20</sup>. A group of demographers have analysed trends in fertility, mortality and migration in different parts of the world<sup>21</sup>. Their discussions produced a consensus about ranges in 2030–2035 that they thought would cover 90 per cent of all future paths of TFR, life expectancy at birth, and the interregional migration matrix<sup>22</sup>. Because the resulting distributions of assumed values turned out to be symmetric, normal distributions were fitted to those ranges. For each of the three variables, a single draw from a standard normal distribution determined its relative position within its range of future values at selected dates. The values at intermediate dates were determined by piece-wise linear interpolation. This method has been labelled a random scenario approach to population projection<sup>23</sup>. Experiments with less autocorrelated paths for each variable produced very similar means and medians and minor differences in variances<sup>24</sup>.

Beyond 2030–2035, fertility was assumed to reach an average level of between 1.7 and 2.1 children per woman by 2080, with the specific value depending on population density in 2030 (the high the density, the lower the fertility). The 90 per cent range around that value was set at 1.0 children. The range for life-expectancy increases after 2030–2035 was set to 0–2.0 years per decade. Smooth transitions of assumed future life expectancies at birth into age-specific mortality rates were performed by transforming baseline mortality patterns with the help of relational models<sup>25</sup>. Because separate baseline data were needed for males and females in each region, 26 patterns were used. Age-specific fertility rates were derived from the total fertility rates by using a fixed relative age profile of fertility. We used a fixed relative age profile for migrants to determine age-specific migration rates. The projections were performed in five-year steps on populations in five-year age groups.

An unusual feature of this global projection is the explicit consideration of possible correlations in fertility and mortality between and within regions. Four sets of 1,000 simulations were produced, resulting from a cross-classification of perfectly correlated/uncorrelated fertility and mortality trends across regions and within regions. The regional results presented in Table 1 are from the set with uncorrelated trends. The global results are based on the merged distribution of all four sets of simulations, which make up 4,000 projections in total.

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## Attentional requirements in a 'preattentive' feature search task

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It is commonly assumed that certain features are so elementary to the visual system that they require no attentional resources to be perceived. Such 'preattentive' features are traditionally identified by visual search performance<sup>1–3</sup>, in which the reaction time for detecting a feature difference against a set of distractor items does not increase with the number of distractors. This suggests an unlimited capacity for the perception of such features. We provide evidence to the contrary, demonstrating that detection of differences in a simple feature such as orientation is severely impaired by additionally imposing an attentionally demanding rapid serial visual presentation task involving letter identification. The same visual stimuli exhibit non-increasing reaction time versus set-size functions. These results demonstrate that attention can be critical even for the detection of so-called 'preattentive' features.

One basic tenet of modern vision research is that certain attributes of visual stimuli can be processed and detected in parallel across the visual field<sup>4–6</sup>. Visual attributes such as orientation<sup>1,5,6</sup>, colour, or size differences<sup>3</sup> have been put forth as 'preattentively' perceived stimulus properties, a concept introduced by Neisser<sup>4</sup>. Perhaps owing to the emphasis on a dichotomy between 'preattentive' and 'attentive' processing, it is commonly assumed that attentional resources are not necessary for the perception of such image properties. This dichotomy stems from a long history of research with the visual search paradigm in which the time to detect a target is measured as a function of the number of display items (reviewed in ref. 3). Stimulus attributes that require focal attention to be perceived exhibit positive slopes: reaction times increase with increasing display size. In contrast, some search tasks show reaction times that remain flat or even decrease slightly as the number of

items increases. Stimulus attributes leading to this behaviour, such as stimulus orientation, are thought to be processed in parallel across space with unlimited capacity; hence they are called 'preattentive' features, and are sometimes thought to be perceived without the use of attention. In the case of stimulus orientation, this fits well with the orientation selectivity of V1 neurons<sup>7,8</sup>, which could conceivably permit the perception of orientation differences regardless of the attentional state.

If the perception of primitive features enjoys a special status in the visual processing stream, avoiding any bottleneck of limited resources, then performance in detection of a feature difference should be unaffected when attention is diverted elsewhere. We investigated the role of attention in the perception of 'preattentive' orientation features with a dual-task procedure as depicted in Fig. 1. We used a competing task, that of reporting the identity of a single white letter appearing in a rapidly changing stream of otherwise black letters at fixation. This rapid serial visual presentation (RSVP) is very demanding when presented at 12 letters s<sup>-1</sup> and has been shown to effectively consume attentional resources for periods up to half a second<sup>9</sup>. A search array of oriented Gabor patches was presented for 150 ms, immediately followed by 150 ms highcontrast white-noise masks covering their locations. The lag between the onset of the white target letter in the RSVP stream and the onset of the orientation array was randomly varied to examine the temporal extent of interference, if any. In the singletask condition, subjects were instructed to ignore the letters and report only whether an orientation 'oddball', a uniquely oriented item, had been present. In the dual-task condition, subjects were instructed to report both the white letter and whether an orientation oddball was present.

Severe impairments in performance in detecting orientation oddballs resulted when the attentionally demanding RSVP letter identification was additionally imposed (Fig. 2). In the condition of performing only the single task of orientation oddball detection, subjects performed well, averaging 94% correct. However, when performing the dual task of letter identification and orientation oddball detection, oddball detection accuracy was only 60  $\pm$  5% for simultaneous letter and orientation onset (lag 0). Note that the chance level of performance is 50% in this task. Significant degradation in performance persists for several hundred milliseconds after the target letter's appearance, as a result of the attentional demands for processing the target letter<sup>9-11</sup>. For the longest lag of 667 ms, dual-task performance recovered to the single-task level; thus the impairment reflects the temporal dynamics of attentional load rather than just a generic difficulty in encoding and retaining two independent responses. The effects of condition and lag, and the interaction between these variables were all significant (P < 0.01).

One might speculate that we observed attentional effects in the detection performance because these stimuli are unusual in some way and do not qualify as 'preattentive'. Performance was quite high when only the orientation oddball task had to be performed, but do these stimuli exhibit the standard experimental signature of socalled preattentive perception, specifically reaction times that do not increase with the number of items? There was no reason to expect otherwise, because many studies<sup>1,3</sup> have found this for orientation differences that are easily detected. Our stimulus display, however, was slightly different from those used in the usual visual search task in that our search array was on for only a short fixed duration and was masked, whereas it is more customarily presented without a mask and for a longer duration up to the time of the observer's response. To remove any residual doubts, we did a visual search reaction time experiment on visual stimuli that were precisely the same as those used in the first experiment, the only difference being that the number of oriented Gabor items was varied from trial to trial. Subjects were instructed to ignore the letters and respond correctly on the presence of a uniquely oriented item as rapidly as possible. The letter stream was presented as well, although Copyright © 2002 EBSCO Publishing