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## Second-generation microarrays

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Current microarray analysis uses 'chips' containing either 25-residue **oligonucleotides** synthesized by photolithography or **cDNAs** placed by robotic spotting. In the April **Nature Biotechnology**, Hughes *et al.* describe a microarray technique that exploits an ink-jet printing method and standard phosphoramidite chemistry (*Nature Biotechnology* 2001, **19**:342-347). The ink-jet synthesizer can deliver 25,000 phosphoramidite-containing microdroplets to a 25 x 75 mm glass slide. Hughes *et al.* examined a large range of parameters to define conditions for optimized specificity and sensitivity. They found that 60-mer oligonucleotides hybridized at 30-32% formamide gave the best results. The absolute detection limit was approximately 0.1 copies per cell equivalent. The ink-jet arrays were as effective as spotted cDNA microarrays. Moreover, Hughes *et al.* report that single carefully chosen 60-mer oligonucleotides can be preferable to arrays containing multiple oligonucleotides or cDNAs as they offer maximal specificity. The ink-jet technology provides a very **flexible microarray** system that can be experimentally optimized to detect low abundance mRNAs and spliced variants.

## References

1. Expression monitoring by hybridization to high-density oligonucleotide arrays.
2. Quantitative monitoring of gene expression patterns with a complementary DNA microarray.
3. *Nature Biotechnology*, [<http://biotech.nature.com>]
4. Experimental annotation of the human genome using microarray technology.